

Pharmaceutical composition and the use thereof, and application regime of said pharmaceutical composition for on-demand contraception

The present invention relates to the subject matter stated in the patent claims, i.e.:

5 (a) the use of a pharmaceutical composition comprising at least one 2H-indazole as novel EP2 receptor antagonist according to general formula I and at least one cyclooxygenase inhibitor (COX inhibitor) for non-hormonal on-demand contraception and contraception,

(b) a pharmaceutical composition comprising at least one 2H-indazole as novel EP2 receptor antagonist according to general formula Ia and at least one (COX inhibitor)

10 and

(c) the use of at least one EP2 receptor antagonist according to general formula Ia in combination with at least one COX inhibitor for preparing a pharmaceutical composition, and also processes for preparing this pharmaceutical composition and its use for non-hormonal on-demand contraception and fertility control.

15 **Description of the invention**

Hormonal contraception with daily orally administered low dosages of synthetic gestagens and oestrogens is, in addition to administration of a gestagen via an intrauterine system (IUS) such as Mirena®, currently the method showing best pharmacological efficacy in the prevention of pregnancies in clinical studies. Oral contraception requires continuous regular intake of hormone-containing pills, independent of the frequency of sexual intercourse and thus the necessity of contraceptive protection. Women having infrequent sexual intercourse are thus exposed to hormones for an unnecessarily long period of time. Especially for this group of users, there is therefore a need for a preparation which can be taken only when required shortly before or optionally also immediately after intercourse but, at the same time, offers a (contraceptive) protection similar to that of the classic oral contraceptives.

Although there has been such a need for decades (Canzler et al. Zbl. Gynäkol.; (1984); 106:1182-1191), there is hitherto no preparation allowing contraception on demand (alternatively also referred to in the literature as on-demand contraception, fertility control on demand or precoital contraception) (Aitken et al. Contraception; (2008); 78: p.28-p.35). In a study carried out in Ghana, it has been reported that woman use Primolut-N tablets comprising 5 mg of norethisterone, which are approved for the treatment of various gynaecological disorders as on-demand contraceptive in 94% of cases and for the treatment of menstrual disorders in only

6% of cases (Opare-Addo et al. Afr. J. Reprod. Health.; (2011); 15(1):65-7). Anecdotal reports and data collected by Family Health International confirm that women in many emerging countries and South American and Africa, but also in countries such as Great Britain, deliberately use preparations developed for emergency contraception also for on-demand
5 contraception.

Products approved for emergency contraception are Postinor®, Plan B® and Levonelle® (single administration of 1.5 mg of levonorgestrel (LNG) within 72 hours) and ellaOne® (single administration of 30 mg of ulipristal acetate (UPA) within 120 h after sexual intercourse). These preparations are described, for example, in the application EP 1448207 (LNG) and in the
10 applications WO2011/069871 and WO2011/091892 (UPA). Levonorgestrel is a progesterone derivative, and ulipristal acetate is a selective progesterone receptor modulator. Thus, both are hormonal methods employed in very high doses compared to a hormone-comprising oral contraceptive taken regularly. Thus, for example, administration of a single dose of 1.5 mg of LNG corresponds to 10 times the amount that is administered daily when an LNG-containing
15 pill (such as Microgynon®) is taken regularly. In Europe, the mini pill Microlut®, which contains 0.03 mg of LNG and is taken for a period of 28 days, which corresponds to a dose of only 0.84 mg of LNG per cycle, has been approved. Here, the single dose of 1.5 mg of LNG for emergency contraception corresponds to more than two monthly doses of Microlut®. If a pill developed for emergency contraception is used several times a month, the dose is,
20 correspondingly, even higher. Raymond et al. have shown that even single administration may have effects on the cycle (Raymond et al. Obstet Gynecol.; (2011); 117(3):673-81). In addition, these preparations for emergency contraception do not have the same contraceptive reliability as a hormone-containing oral contraceptive taken regularly; however, this would be desirable for on-demand contraception. Evaluation of different studies for pre- and postcoital use of 0.75 mg
25 of LNG showed a Pearl Index of 5.1 [pregnancies per 100 woman years] (95% confidence interval [CI] 3.8-6.7), which represents moderate efficacy (Raymond et al. Obstet Gynecol.; (2011); 117(3):673-81).

Hitherto, for on-demand contraception, only hormonal methods have been described; however, none of these has yet been approved:

30 WO 2010/119030 describes an on-demand preparation where, as gestagen, levonorgestrel or norgestrel is used in a dose range of from 0.5 to 1.5 mg, with the preparation to be used 24 hours prior to intercourse. However, the dose of LNG corresponds to the dose used for emergency contraception.

WO2007/045513 describes a method for on-demand contraception where a gestagen is, if required, administered transdermally using a patch.

WO96/12494 uses competitive progesterone antagonists for on-demand contraception.

WO2010/119029 describes progesterone or progesterone receptor modulators for on-demand
5 contraception which are administered within 72 hours prior to expected sexual intercourse. Here, 30 mg of UPA are employed once prior to sexual intercourse, which corresponds to the dose used for emergency contraception.

Non-hormonal methods for on-demand contraception have not yet been described. However, there is a need for non-hormonal pharmacological methods for women who, for medicinal or
10 personal reasons, do not want to use or cannot use hormonal contraceptives. A further advantage of a non-hormonal approach is the fact that the menstrual cycle is not affected.

It has already long been known that prostaglandins are key molecules in the processes of female reproductive biology, such as for example the regulation of ovulation and fertilization. The effects of the prostaglandins are mediated through their G protein coupled receptors, which are
15 located in the cell membrane. Of particular interest is prostaglandin E₂ (PGE₂), which effects a great variety of cellular actions by binding to functionally different receptor subtypes, namely the EP₁, EP₂, EP₃ and EP₄ receptors. The receptor subtypes to which prostaglandin E₂ binds are of particular importance for the receptor-mediated actions which are involved in the regulation of fertility. Thus it could be shown that the reproductive functions are impaired in EP2 knockout
20 mice (EP2^{-/-}), e.g. in mice which no longer have a functional PGE₂ receptor subtype EP2, and that these animals have a smaller number of offspring per pregnancy (Matsumoto *et al.* Biology of Reproduction; (2001); 64: 1557-1565). Likewise it could be shown that these EP2 knockout mice (Hizaki *et al.* Proc Natl Acad Sci U. S. A.; (1999); 96(18):10501-10506) display markedly decreased cumulus expansion and severe subfertility, which is to be regarded as being causally
25 linked with decreased reproductive processes such as ovulation and fertilization. Accordingly, the EP2 receptor is an important target for the development of drugs for the regulation of female fertility.

The application DE 10 2009 049 662 A1 describes 2-5-disubstituted 2H-indazoles which, with high binding affinity, selectively antagonize the EP2 receptor. These EP2 receptor antagonists
30 are disclosed *inter alia* for use as contraceptive, but not for on-demand contraception.

In a pairing study with cynomolgus monkey, it was shown that an EP2 receptor antagonist administered continuously has a subfertile effect (Peluffo *et al.* 2012, Annual Meeting of the

American Society for Reproductive Medicine). However, continuous administration does not correspond to use for on-demand contraception.

The applications of Summit Corporation Plc. (WO2007/091107), CM Sun et al. (WO2008/070599) and Janssen Pharmaceutica NV (WO2010/124102) describe 2H-indazoles.

5 The compounds disclosed in WO2007/091107 interact with the utropine A promoter and are claimed for the treatment of Duchenne muscular dystrophy, Becker muscular dystrophy or cachexia. WO2008/070599 describes compounds binding to the VGEFR-3 receptor. They are claimed for the treatment of cancer. The compounds described in the application WO2010/124102 inhibit monoacylglycerol lipase (MGL) and are claimed for the treatment of
10 inflammatory pain. None of the three applications mentioned discloses whether the compounds bind to the EP2 receptor and are suitable for fertility control.

The effect of COX inhibitors on female ovulation has been investigated by several research groups. Thus, Pall et al. examined the effect of rofecoxib on ovulation (Pall et al.; Human Reproduction; (2001); 16(7):1323-1328). In this study, 25 mg of the active compound were used
15 on 9 successive days. In 4 of 6 patients, ovulation was delayed by more than 48 hours.

Edelmann et al. 2013 examined the effect of the daily oral administration of 400 mg of celecoxib on ovulation. Depending on time and duration of administration, an increase in ovulatory dysfunction of 25 - 30% was observed. The ovulatory dysfunction was characterized by a flattened LH (luteinizing hormone) peak and/or by anovulation (Edelmann et al. Contraception.
20 (2013); 87(3):352-357). In 2010, Edelmann et al. had already shown that daily administration of 400 mg of celecoxib over the entire cycle increased the cycle length from 27.2 +/- 2.4 days to 28.5 +/- 2.5 days and changed time and height of the LH peak (Edelmann et al. Contraception; (2010); 81(6):496-500). Both results showed that the benefit of celecoxib for emergency contraception is limited.

25 In a pilot study with 6 women treated with 3 x 50 mg of indomethacin per day for at least 3 days, it was shown that administration of indomethacin delayed ovulation by 2 – 12 days in 5 cases. The treatment was initiated when the dominant follicle had reached a maximum size of > 16 mm or LH was detected in the urine (Athanasious et al. Fertil Steril.; (1996); 65(3): 556-60). This study also confirmed the results by Killick S and Elstein M. 1987 where indomethacin
30 administered in a dose of 2 x 100 mg per day over 5 days led to inhibition of ovulation (Killick S and Elstein M. Fertil Steril.; (1987); 47(5):773-7).

A clinical study by Bata et al. 2006 showed that administration of meloxicam reversibly delayed ovulation by 5 days (Bata et al., J Clin Pharmacol (2006); 46:925-932). This result was

confirmed by Jesam et al. 2010 (Jesam et al. Hum Reprod.; (2010); 25(2):368-73). In this clinical study, 30 mg or 15 mg of meloxicam were administered daily over a period of 5 days during the late follicular phase. In 90.9% (30 mg meloxicam group) or 50.0% (15 mg meloxicam group), ovulation was abnormal, which shows that meloxicam may be suitable for emergency contraception.

In principle, COX inhibitors are known to be relatively well tolerated; however, Castellsague et al. 2012 and Varas-Lorenzo et al. 2013 indicated that there is an increased risk of cardiovascular and gastrointestinal events for this class of substances. It is therefore recommended to use a relatively low dosage (Castellsague et al. Drug Saf.; (2012); 35(12):1127-46; Varas-Lorenzo et al. Pharmacoevidenciol Drug Saf.; (2013); doi: 10.1002/pds.3437. [Epub ahead of print]).

In a pilot study, Massai et al. 2007 examined the use of meloxicam in combination with levonorgestrel (LNG) on the time of ovulation in connection with emergency contraception (Massai et al. Human Reproduction; (2007) 22(2):434-439). In this study, 2 tablets of in each case 0.75 mg of LNG (known *inter alia* by the trade name Postinor-2) were administered on their own or together with 15 mg of meloxicam. In the treatment group which received the combination of meloxicam and LNG, there was a trend to reduced frequency of ovulation compared to the group treated exclusively with LNG. This effect was increasingly pronounced the later the tablets were taken relative to the time of ovulation.

In WO2010/149273 (Bayer Pharma AG), it was shown that a low dose of levonorgestrel which, on its own, does not inhibit ovulation, may achieve virtually complete suppression of ovulation in the rat when combined with piroxicam. However, this approach comprises, in addition to the non-hormonally acting COX inhibitor, a hormonal component in the form of levonorgestrel and does therefore not represent a non-hormonal alternative to purely hormonal approaches.

Accordingly, it is an object of the present invention to provide a method and a pharmaceutical composition for on-demand contraception based on a pharmacological non-hormonal approach and thus providing an alternative to hormonal methods.

In accordance with the invention, this object is achieved by using a non-hormonal pharmaceutical composition comprising at least one EP2 receptor antagonist according to formula I and at least one COX inhibitor. In addition, this object is achieved by the pharmaceutical composition itself which comprises at least one EP2 receptor antagonist according to formula Ia and at least one COX inhibitor.

The EP2 receptor antagonists according to formula I or Ia according to the invention have already been described in the patent application PCT/EP2012/073556, unpublished at the

priority date of the present application. There, they are claimed *inter alia* for use for contraception and emergency contraception. In addition, a pharmaceutical composition comprising at least one EP2 receptor antagonist of formula I and at least one COX inhibitor and its use for contraception are described.

5 In the patent application PCT/EP2012/073556, it was shown that the EP2 receptor antagonists according to formula I or Ia have an antagonistic action at the EP2 receptor (see biological examples; Table 1) and are thus suitable for fertility control (see biological examples; Tables 2, 3 and 4). In the present application, in accordance with the invention, it was possible, by combining the EP2 receptor antagonists according to formula I or Ia with a COX inhibitor, to
10 achieve the contraceptive action of the EP2 receptor antagonists or the COX inhibitors surprisingly even at a still ineffective dose of the individual active compounds (see biological examples; Tables 6.1 – 6.5 and 7.1). Thus, the combination of EP2 receptor antagonists according to general formula I or Ia with COX inhibitors allows both the dose of the COX inhibitor and the dose of the EP2 receptor antagonists to be reduced, which in turn reduces
15 possible side effects and thus increases acceptance and safety of the method.

In addition, the contraceptive action was, surprisingly, achieved even after single administration of an EP2 receptor antagonist according to general formula I or Ia in combination with a COX inhibitor shortly before sexual intercourse (see biological examples; Tables 6.1 – 6.5 and 7.1). This shows that EP2 receptor antagonists in combination with COX inhibitors constitute an
20 effective pharmaceutical composition for on-demand contraception which can be administered before or shortly after expected sexual intercourse. Since this pharmaceutical composition is non-hormonal, it represents an alternative to hormonal approaches.

In the pre-ovulatory antral follicle, the oocyte is surrounded by cumulus cells which form a dense cell crown around the oocyte. After the LH peak (luteinizing hormone), a series of
25 processes is activated, which results in a marked morphological change in this cell crown of cumulus cells. During this, the cumulus cells form an extracellular matrix, which leads to the so-called cumulus expansion (Vanderhyden *et al.* Dev Biol.; (1990); 140(2):307-317). This cumulus expansion is an important component of the ovulatory process and the subsequent possibility of fertilization.

30 During the cumulus expansion, prostaglandins and here prostaglandin E₂, synthesis whereof is induced by the LH peak, are of decisive importance. Prostanoid EP2 knockout mice (Hizaki *et al.* Proc Natl Acad Sci U S A.; (1999) 96(18):10501-10506.) show a markedly decreased cumulus expansion and severe subfertility, which demonstrates the importance of the prostanoid EP2 receptor for this process.

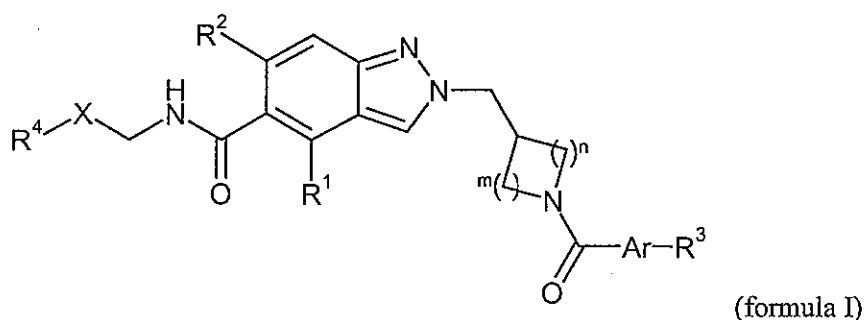
The anti-ovulatory action of COX inhibitors in humans has been demonstrated in clinical studies (as described above). COX inhibitors lead to reduced prostaglandin synthesis. In COX-2 knock-out mice, it was shown that prostaglandins play a role in particular in processes of ovulation, cumulus expansion and fertilization (Cheng JG and Stewart CL. Biol Reprod.; (2003); 68(2):401-4; Lim et al. Cell; (1997) 91(2):197-208).

Since, within the prostaglandin pathway, the EP2 receptor is inhibited directly by the EP2 receptor antagonists, the onset of action by this class of substances is earlier than the onset of action of COX inhibitors, where the action is triggered by lowering of prostaglandin concentrations.

10 *On-demand contraception or contraception on demand*

On-demand contraception, fertility control on demand, precoital contraception or contraception on demand all refer to the on-demand use of one or more substances for preventing unwanted pregnancy. The administration can be before or shortly after expected sexual intercourse. Here, on-demand does not mean optional. Rather, the substance/substances has/have to be administered to prevent unwanted pregnancy.

The invention provides the use of a pharmaceutical composition comprising (a) at least one EP2 receptor antagonist according to formula (I)



wherein

20 R^1 : means H, C_1 - C_2 alkyl or C_1 - C_2 alkyloxy;

R^2 : H or methyl;

subject to the proviso that one of the two residues R^1 or R^2 equals H;

X: $-(CH_2)_l$ -, $-(CH_2)_k-O$ -, $-CH_2-S$ -, $CH_2-S(O)_2$ -, $-CH(CH_3)$ -, $-CH(CH_3)-O$ - or $-C(CH_3)_2-O$ -;

k: 1 or 2;

- l: 0, 1 or 2;
- R⁴: H, C₁-C₄ alkyl, C₃-C₄ cycloalkyl, or CH₂-C₃-C₄ cycloalkyl;
- and in the case that X means -(CH₂)_l- with l meaning 0 or 1 or -CH(CH₃)-
- R⁴: additionally a 4-6-membered heterocyclyl residue;
- 5 and in the case that X means -(CH₂)_l- or -CH(CH₃)-
- R⁴: additionally CN;
- m: 1 or 2;
- n: 1 or 2;
- Ar: a 6-10-membered aryl or 5-10-membered hetaryl residue,
- 10 R³: halogen, CN, SF₅, C₁-C₄ alkyl, C₃-C₆ cycloalkyl, C₄-C₆ heterocyclyl, O-C₁-C₄ alkyl, O-C₃-C₆ cycloalkyl, O-C₄-C₆ heterocyclyl, S-C₁-C₄ alkyl, S(O)₂-C₁-C₄ alkyl, Ar', O-Ar', C(CH₃)₂-CN or C(CH₃)₂-OH;
- Ar': an optionally singly or doubly substituted 6-membered aryl or 5-6-membered heteroaryl residue;
- 15 wherein the substituents are selected from F, Cl, CN, C₁-C₄ alkyl, O-C₁-C₄ alkyl, C(CH₃)₂-CN, C(CH₃)₂-OH and C(O)NH₂;
- and comprising (b) at least one COX inhibitor selected from the list below:
- C1 indomethacin;
- C2 diclofenac;
- 20 C4 naproxen;
- C5 ibuprofen;
- C6 meloxicam;
- C7 piroxicam;
- C8 nimesulide;

- C9 ketoprofen;
- C10 tenoxicam;
- C11 mefenamic acid;
- C12 ketoralac;
- 5 C13 celecoxib(4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl]benzenesulphonamide);
- C14 parecoxib(N-[4-(5-methyl-3-phenyl-4-isoxazolyl)phenyl]sulphonylpropionamide);
- C15 rofecoxib (4-(4-methylphenyl)-3-phenylfuran-2(5H)-one);
- C16 valdecoxib (4-[5-methyl-3-phenyl-4-isoxazolyl]benzenesulphonamide);
- 10 C17 etoricoxib

for on-demand contraception.

The invention furthermore provides the use of a pharmaceutical composition comprising (a) at least one EP2 receptor antagonist according to formula (I), where

R¹: means H, C₁-C₂ alkyl or C₁-C₂ alkyloxy;

15 R²: H or methyl;

subject to the proviso that one of the two residues R¹ or R² equals H;

X: -(CH₂)_l-, -(CH₂)_k-O-, -CH₂-S-, CH₂-S(O)₂-, -CH(CH₃)-, -CH(CH₃)-O- or -C(CH₃)₂-O-;

k: 1 or 2;

l: 0, 1 or 2;

20 R⁴: H, C₁-C₄ alkyl, C₃-C₄ cycloalkyl, or CH₂-C₃-C₄ cycloalkyl;

and in the case that X means -(CH₂)_l- with l meaning 0

R⁴: additionally a 4-6-membered heterocyclyl residue linked via a ring carbon atom;

m: 1 or 2;

n: 1 or 2;

Ar: a 6-10-membered aryl or 5-10-membered hetaryl residue,

R³: halogen, CN, SF₅, C₁-C₄ alkyl, C₃-C₆ cycloalkyl, C₄-C₆ heterocyclyl, O-C₁-C₄ alkyl, O-C₃-C₆ cycloalkyl, O-C₄-C₆ heterocyclyl, S-C₁-C₄ alkyl, S(O)₂-C₁-C₄ alkyl, Ar',
5 O-Ar', C(CH₃)₂-CN or C(CH₃)₂-OH;

Ar': an optionally singly or doubly substituted 6-membered aryl or 5-6-membered heteroaryl residue;

wherein the substituents are selected from F, Cl, CN, C₁-C₄ alkyl, O-C₁-C₄ alkyl, C(CH₃)₂-CN, C(CH₃)₂-OH and C(O)NH₂;

10 and comprising (b) at least one COX inhibitor selected from the list above (COX inhibitors C1-C17) for on-demand contraception.

The 2H-indazoles according to the invention have an antagonistic action on the EP2 receptor.

C₁-C₂ alkyl or C₁-C₄ alkyl should each be understood to mean a linear or branched alkyl residue, such as for example methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl or tert-butyl.

15 The alkyl residues can optionally be singly or multiply substituted with fluorine.

The hydrogen atoms can optionally be replaced by deuterium.

C₁-C₂ alkoxy or C₁-C₄ alkoxy should be understood to mean a linear or branched residue of the formula alkyl-O-, such as for example methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, sec-butoxy, isobutoxy or tert-butyloxy.

20 The alkoxy residues can optionally be singly or multiply substituted with fluorine.

C₃-C₆ cycloalkyl should be understood to mean monocyclic alkyl rings such as cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl.

The cycloalkyl residues can optionally be singly or multiply substituted with fluorine.

4-6-membered heterocyclyl should be understood to mean monocyclic rings which instead of the carbon atoms contain one or more hetero atoms, such as oxygen, sulphur and/or nitrogen or a hetero group such as -S(O)- or -SO₂-. The linking bond can be at any carbon atom or at a nitrogen atom.
25

For example, azetidiny, pyrrolidiny, piperidiny, piperaziny, 1,4-diazepany, morpholiny, thiomorpholiny, oxetany, tetrahydrofurany, tetrahydropyrany, 1,4-dioxany and 2-oxo-oxazolidiny may be mentioned.

The heterocyclyl residues can optionally be singly or multiply substituted with fluorine, hydroxy, methoxy or with oxo.

Halogen should in each case be understood to mean fluorine, chlorine, bromine or iodine.

The C₆-C₁₀-membered aryl residue in each case comprises 6 - 10 carbon atoms and can be benzo-condensed. Phenyl and naphthyl may be mentioned.

The C₆-C₁₀-membered aryl residue can optionally be singly substituted with fluorine, chlorine or a methyl group.

The C₅-C₁₀ heteroaryl residue should be understood to mean mono- or bicyclic ring systems which each contain 5 - 10 ring atoms and which instead of the carbon can contain one or more identical or different hetero atoms, such as oxygen, sulphur or nitrogen. The linking bond can be at any carbon atom or at a nitrogen atom.

For example, thienyl, thiazolyl, furyl, pyrrolyl, oxazolyl, imidazolyl, pyrazolyl, isoxazolyl, isothiazolyl, oxadiazolyl, triazolyl, thiadiazolyl, tetrazolyl, pyridyl, pyridaziny, pyrimidiny, pyraziny, triaziny, phthalidyl, thiophthalidyl, indolyl, isoindolyl, indazolyl, benzothiazolyl, benzofuryl, benzothienyl, benzimidazolyl, benzoxazolyl, azociny, indoliziny, puriny, isoquinoliny, quinoliny, quinoliziny, quinazoliny, quinoxaliny, cinnoliny, phthalaziny, 1,7- or 1,8-naphthyridiny and pteridiny may be mentioned.

The C₅-C₁₀-membered heteroaryl residue can optionally be singly substituted by fluorine, chlorine or a methyl group.

If radicals in the compounds according to the invention are substituted, the radicals can, unless specified otherwise, be mono- or polysubstituted. In the context of the present invention it applies that for all radicals which occur multiple times, their meanings are independent of one another. Substitution with one, two or three identical or different substituents is preferred. Substitution with one substituent is very particularly preferred.

If a basic functional group is contained, the physiologically compatible salts of organic and inorganic acids, such as *inter alia* hydrochloric acid, sulphuric acid, phosphoric acid, citric acid and tartaric acid, are suitable.

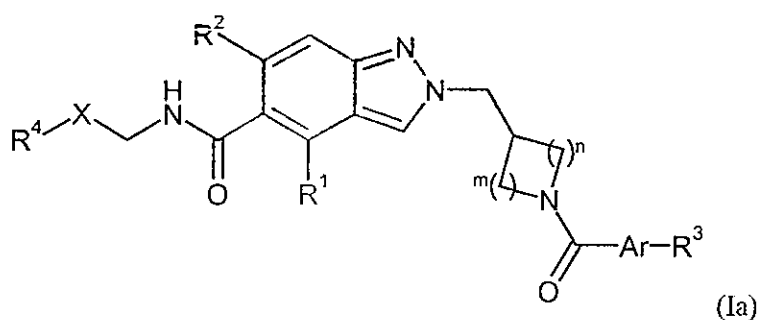
Pharmaceutical composition comprising at least one EP2 receptor antagonist according to general formula Ia and at least one COX inhibitor

The present invention provides a pharmaceutical composition comprising (a) at least one EP2 receptor antagonist according to the formula (Ia) and (b) at least one COX inhibitor.

- 5 The present invention furthermore provides the use of at least one EP2 receptor antagonist of the general formula Ia together with at least one COX inhibitor for preparing a medicament or a pharmaceutical composition comprising suitable formulation auxiliaries and carriers.

EP2 receptor antagonists which, according to the invention, are comprised in the pharmaceutical composition

- 10 EP2 receptor antagonists according to the invention which are comprised in the pharmaceutical composition according to the invention are 2H-indazoles of the general formula Ia, where



R^1, R^2 : mean H or methyl;

subject to the proviso that one of the two residues R^1 or R^2 equals H;

- 15 X: $-(CH_2)_l-$ or $-(CH_2)_k-O-$;

k: 1 or 2;

l: 0, 1 or 2;

R^4 : H, C_1-C_4 alkyl, C_3-C_4 cycloalkyl or $CH_2-C_3-C_4$ cycloalkyl;

m: 1 or 2;

- 20 n: 1 or 2;

Ar: a 6-10-membered aryl or 5-10-membered hetaryl residue,

R³: halogen, CN, SF₅, C₁-C₄ alkyl, C₃-C₆ cycloalkyl, C₂-C₆ heterocyclyl, O-C₁-C₃ alkyl, O-C₃-C₆ cycloalkyl, O-C₄-C₆ heterocyclyl, S-C₁-C₄ alkyl, S(O)₂-C₁-C₄ alkyl, Ar', O-Ar', C(CH₃)₂-CN or C(CH₃)₂-OH; and

Ar': an optionally singly or doubly substituted 6-membered aryl or 5-6-membered heteroaryl
5 residue;

wherein the substituents are selected from F, Cl, CN, C₁-C₄ alkyl, O-C₁-C₄ alkyl, C(CH₃)₂-CN, C(CH₃)₂-OH and C(O)NH₂;

and salts, solvates or solvates of the salts thereof, and isomers, diastereomers, enantiomers and salts or cyclodextrin clathrates thereof, for producing medicaments which overcome the known
10 disadvantages and have improved properties such as good *in-vivo* efficacy, good solubility and stability.

The compounds according to the invention have an antagonistic action at the EP2 receptor.

The radicals and substituents have the same meaning as described above for the EP2 receptor antagonists of the general formula I.

15 Preferred are compounds of the formula Ia, wherein

R¹: means a methyl group;

R²: H;

X: -(CH₂)_l- or -(CH₂)_k-O-;

k: 1;

20 l: 0 or 1;

R⁴: C₁-C₄ alkyl, C₃-C₄ cycloalkyl or CH₂-C₃-C₄ cycloalkyl;

m, n: 2;

Ar: a phenyl residue;

and salts, solvates or solvates of the salts thereof, and isomers, diastereomers, enantiomers and salts or cyclodextrin clathrates thereof, for producing medicaments which overcome the known
25 disadvantages and have improved properties such as good *in-vivo* efficacy, good solubility and stability.

Also preferred are compounds of the formula Ia, wherein

R¹: means a methyl group;

R²: H;

X: -(CH₂)_l- or -(CH₂)_k-O-;

5 k: 1;

l: 0 or 1;

R⁴: C₁-C₄ alkyl, C₃-C₄ cycloalkyl or CH₂-C₃-C₄ cycloalkyl;

m, n: 1;

Ar: a phenyl residue:

10 and salts, solvates or solvates of the salts thereof, and isomers, diastereomers, enantiomers and salts or cyclodextrin clathrates thereof, for producing medicaments which overcome the known disadvantages and have improved properties such as good *in-vivo* efficacy, good solubility and stability.

The following EP2 receptor antagonists according to the invention are particularly preferred:

15 E1 2-([1-(4-cyano-2-fluorobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

E2 2-([1-(4-tert-butoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

20 E3 2-([1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

E4 N-(2-methoxyethyl)-4-methyl-2-([1-(4-morpholinobenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide

E5 2-([1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

25 E6 2-([1-(2-fluoro-4-mesylbenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E7 2-([1-(2-fluoro-4-methoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E8 2-([1-(4-bromo-2-fluorobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E9 2-([1-(2-fluoro-4-methylbenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E10 2-([1-[2-fluoro-4-(trifluoromethyl)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E11 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(pentafluoro- λ^6 -sulphanyl)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E12 N-(2-methoxyethyl)-4-methyl-2-([1-(4-methylbenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E13 2-([1-[4-(4-chlorophenoxy)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E14 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(4-methylphenoxy)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E15 2-([1-[4-(4-tert-butylphenoxy)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E16 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(4-(trifluoromethyl)phenoxy)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E17 N-(2-methoxyethyl)-2-([1-[4-(4-methoxyphenoxy)benzoyl]piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- E18 N-(2-methoxyethyl)-4-methyl-2-([1-(4-phenoxybenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 25 E19 2-([1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E20 2-([1-(4-methoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E21 2-([1-(4-fluorobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E22 N-(2-methoxyethyl)-4-methyl-2-([1-(4-(trifluoromethyl)benzoyl)piperidin-4-yl]-methyl)-2H-indazole-5-carboxamide
- 5 E23 2-([1-(2-methoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E24 N-(2-methoxyethyl)-4-methyl-2-([1-(4-[(trifluoromethyl)sulphonyl]benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E25 N-(2-methoxyethyl)-4-methyl-2-([1-(3-(trifluoromethyl)benzoyl)piperidin-4-yl]-methyl)-2H-indazole-5-carboxamide
- 10 E26 N-(2-methoxyethyl)-4-methyl-2-([1-(3-methylbenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E27 2-([1-(3-chlorobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E28 2-([1-(4-(4-carbamoylphenoxy)benzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E29 2-([1-(4-(cyclopentyloxy)benzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E30 2-([1-(4-(difluoromethyl)benzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E31 2-([1-(4-cyanobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E32 2-([1-(4-(1H-imidazol-1-yl)benzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E33 N-(2-methoxyethyl)-4-methyl-2-([1-(4-(oxazol-2-yl)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E34 N-(2-methoxyethyl)-4-methyl-2-([1-(4-(oxazol-5-yl)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide

- E35 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(isoxazol-5-yl)benzoyl]piperidin-4-yl} methyl)-
2H-indazole-5-carboxamide
- E36 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(1H-pyrazol-1-yl)benzoyl]piperidin-4-yl}-
methyl)-2H-indazole-5-carboxamide
- 5 E37 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(1H-1,2,4-triazol-1-yl)benzoyl]piperidin-4-yl}-
methyl)-2H-indazole-5-carboxamide
- E38 2-({1-[4-(difluoromethoxy)-2-fluorobenzoyl]piperidin-4-yl} methyl)-N-(2-methoxy-
ethyl)-4-methyl-2H-indazole-5-carboxamide
- E39 2-({1-[2-fluoro-4-(pyrrolidin-1-yl)benzoyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-
10 4-methyl-2H-indazole-5-carboxamide
- E40 2-({1-[(3,4'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-
4-methyl-2H-indazole-5-carboxamide
- E41 2-({1-[(3-fluoro-4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-
methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E42 2-({1-[(3-fluoro-4'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-
methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E43 2-[(1-{[3-fluoro-3'-(trifluoromethyl)biphenyl-4-yl]carbonyl} piperidin-4-yl)methyl]-N-
(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E44 2-[(1-{[3-fluoro-2'-(trifluoromethoxy)biphenyl-4-yl]carbonyl} piperidin-4-yl)methyl]-N-
20 (2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E45 2-({1-[(2'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-
methyl-2H-indazole-5-carboxamide
- E46 2-({1-[(2',4'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxy-
ethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E47 2-({1-[(2-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-
methyl-2H-indazole-5-carboxamide
- E48 N-(2-methoxyethyl)-4-methyl-2-({1-[(2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}-
methyl)-2H-indazole-5-carboxamide

- E49 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(4-pyridyloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E50 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(4H-1,2,4-triazol-4-yl)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide
- 5 E51 2-({1-[2-fluoro-4-[morpholin-4-yl]benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E52 2-{{1-[4-(4-bromobenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E53 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide
- 10 E54 2-{{1-[4-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E55 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[(trifluoromethyl)sulphanyl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 15 E56 4-methyl-2-({1-[4-(pentafluoro- λ^6 -sulphanyl)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E57 2-{{1-[4-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E58 4-methyl-N-(2,2,2-trifluoroethyl)-2-({1-[4-(trifluoromethoxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- 20 E59 2-{{1-[4-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide
- E60 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-{{1-[4-(4-methylbenzoyl)piperidin-4-yl]-methyl}-2H-indazole-5-carboxamide
- 25 E61 N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}-methyl)-4-methyl-2H-indazole-5-carboxamide
- E62 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

- E63 N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E64 2-{{1-(4-cyclopropylbenzoyl)piperidin-4-yl}methyl}-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide
- 5 E65 2-{{1-(4-chlorobenzoyl)piperidin-4-yl}methyl}-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E66 4-methyl-2-{{1-(4-methylbenzoyl)piperidin-4-yl}methyl}-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E67 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- 10 E68 4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2-({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E69 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- 15 E70 2-{{1-(4-cyclopropylbenzoyl)piperidin-4-yl}methyl}-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E71 2-{{1-(4-chlorobenzoyl)piperidin-4-yl}methyl}-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E72 N-(2-isopropoxyethyl)-4-methyl-2-{{1-(4-methylbenzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- 20 E73 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E74 N-(2-isopropoxyethyl)-4-methyl-2-({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- 25 E75 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E76 2-{{1-(4-cyclopropylbenzoyl)piperidin-4-yl}methyl}-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E77 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- E78 2-([1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl]methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- 5 E79 2-([1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl]methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- E80 4-methyl-N-[2-(trifluoromethoxy)ethyl]-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E81 N-(2-tert-butoxyethyl)-2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E82 N-(2-tert-butoxyethyl)-2-([1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- E83 N-(2-tert-butoxyethyl)-2-([1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E84 N-(2-tert-butoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E85 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-4-methyl-N-{2-[(²H₃)methyloxy]-(²H₄)ethyl}-2H-indazole-5-carboxamide
- E86 2-([1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl]methyl)-4-methyl-N-{2-[(²H₃)methyloxy](²H₄)ethyl}-2H-indazole-5-carboxamide
- 20 E87 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E88 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[(trifluoromethyl)sulphonyl]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- 25 E89 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl]methyl)-2H-indazole-5-carboxamide
- E90 2-([1-[2-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E91 2-([1-(4-chloro-2-fluorobenzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E92 2-([1-(3-fluoro-4-(trifluoromethyl)benzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E93 2-([1-(4-chloro-3-(trifluoromethyl)benzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E94 2-([1-(4-cyclopropylbenzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E95 N-(2-methoxyethyl)-4-methyl-2-([1-(4-(trifluoromethoxy)phenoxy)benzoyl]azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- 10 E96 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E97 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-(trifluoromethoxy)benzoyl)azetidin-3-yl]methyl)-2H-indazole-5-carboxamide
- 15 E98 2-([1-(2-fluoro-4-(trifluoromethyl)benzoyl)azetidin-3-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E99 4-methyl-2-([1-(4-(pentafluoro- λ^6 -sulphanyl)benzoyl)azetidin-3-yl]methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E100 N-[2-(cyclopropyloxy)ethyl]-4-methyl-2-([1-(4-(trifluoromethoxy)benzoyl)azetidin-3-yl]methyl)-2H-indazole-5-carboxamide
- 20 E101 N-[2-(cyclobutyloxy)ethyl]-4-methyl-2-([1-(4-(trifluoromethoxy)benzoyl)azetidin-3-yl]methyl)-2H-indazole-5-carboxamide
- E102 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide
- 25 E103 N-[2-(cyclopropylmethoxy)ethyl]-2-([1-(4-(4-fluorophenoxy)benzoyl)azetidin-3-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- E104 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-([1-(4-methylbenzoyl)azetidin-3-yl]methyl)-2H-indazole-5-carboxamide

- E105 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- E106 N-(2-tert-butoxyethyl)-2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E107 N-[2-(cyclopropylmethoxy)ethyl]-2-([1-(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl)methyl)-4-methyl-2H-indazole-5-carboxamide
- E108 2-([1-(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E109 2-([1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E110 2-([1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E111 2-([1-(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl)methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 15 E112 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E113 4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2-([1-[4-(trifluoromethyl)phenoxy]benzoyl]azetidin-3-yl)methyl)-2H-indazole-5-carboxamide
- E114 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-[4-(trifluoromethyl)phenoxy]benzoyl]azetidin-3-yl)methyl)-2H-indazole-5-carboxamide
- 20 E115 2-([1-[4-(4-chlorophenoxy)benzoyl]azetidin-3-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E116 2-([1-[4-(4-chlorophenoxy)benzoyl]azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E117 N-(2-methoxyethyl)-4-methyl-2-([1-(4-[5-(trifluoromethyl)pyridin-2-yl]oxy]benzoyl]azetidin-3-yl)methyl)-2H-indazole-5-carboxamide
- E118 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-[5-(trifluoromethyl)pyridin-2-yl]oxy]benzoyl]azetidin-3-yl)methyl)-2H-indazole-5-carboxamide

- E119 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-N-ethyl-4-methyl-2H-indazole-5-carboxamide
- E120 2-([1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 5 E121 2-([1-[4-(4-chlorophenoxy)benzoyl]piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E122 4-methyl-2-([1-[4-(4-methylphenoxy)benzoyl]piperidin-4-yl]methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E123 2-([1-[4'-(4-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 10 E124 4-methyl-2-([1-(4-morpholinobenzoyl)piperidin-4-yl]methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E125 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-[4-[(trifluoromethyl)sulphonyl]benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 15 E126 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-[4-[4-(trifluoromethyl)phenoxy]benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E127 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-([5-(trifluoromethyl)pyridin-2-yl]oxy)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E128 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-([6-(trifluoromethyl)pyridin-3-yl]oxy)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 20 E129 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-([6-(trifluoromethyl)pyridin-2-yl]oxy)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E130 2-([1-[4-(4-cyanophenoxy)benzoyl]piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 25 E131 2-([1-[4-(3-fluorophenoxy)benzoyl]piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E132 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-[4-[3-(trifluoromethyl)phenoxy]benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide

- E133 2-[(1-{4-[(5-cyanopyridin-2-yl)oxy]benzoyl}piperidin-4-yl)methyl]-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E134 2-[(1-{4-[(5-chloropyridin-2-yl)oxy]benzoyl}piperidin-4-yl)methyl]-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 5 E135 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[5-(trifluoromethyl)pyridin-2-yl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E136 2-({1-[4-(2,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E137 2-({1-[4-(3,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 10 E138 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-(trifluoromethyl)biphenyl-4-yl}carbonyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E139 2-({1-[4-(4-bromobenzoyl)piperidin-4-yl]methyl}-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 15 E140 2-({1-[4-(5-chloropyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E141 2-({1-[(4'-methoxy-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E142 4-methyl-2-({1-[4-(6-methylpyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 20 E143 2-({1-[(4'-fluoro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E144 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[6-(trifluoromethyl)pyridin-3-yl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 25 E145 2-({1-[4-(6-methoxypyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E146 2-({1-[4-(6-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide

- E147 4-methyl-2-({1-[4-(5-methylpyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E148 2-({1-[4-(5-fluoropyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 5 E149 2-({1-[4-(5-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E150 4-methyl-2-({1-[4-(2-methylpyrimidin-5-yl)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E151 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[2-(trifluoromethyl)pyrimidin-5-yl]benzoyl}-
10 piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E152 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[6-(trifluoromethyl)pyridin-2-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E153 2-({1-[4-(4-cyanophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E154 2-[[1-(4-bromo-3-methylbenzoyl)piperidin-4-yl]methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E155 2-[[1-(4-tert-butylbenzoyl)piperidin-4-yl]methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E156 2-({1-[4-(1-hydroxy-1-methylethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxy-
20 ethyl)-4-methyl-2H-indazole-5-carboxamide
- E157 2-[[1-(4-cyclohexylbenzoyl)piperidin-4-yl]methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E158 2-({1-[4-(1-cyano-1-methylethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E159 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyrimidin-2-yloxy)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide
- E160 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyridin-3-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

- E161 N-(2-methoxyethyl)-4-methyl-2-{{1-(4-{{5-(trifluoromethyl)pyridin-2-yl}oxy}benzoyl)-piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- E162 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyridin-2-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- 5 E163 N-(2-methoxyethyl)-4-methyl-2-{{1-(4-{{4-(trifluoromethyl)pyrimidin-2-yl}oxy}benzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- E164 N-(2-methoxyethyl)-4-methyl-2-{{1-(4-{{6-(trifluoromethyl)pyridin-3-yl}oxy}benzoyl)-piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- E165 N-(2-methoxyethyl)-4-methyl-2-{{1-(4-{{6-(trifluoromethyl)pyridin-2-yl}oxy}benzoyl)-piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- 10 E166 2-({1-{{4'-(4-methoxybiphenyl-4-yl)carbonyl}piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E167 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(3,4,5,6-tetrahydro-2H-pyran-4-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- 15 E168 N-(2-methoxyethyl)-4-methyl-2-{{1-[4-[3-(trifluoromethyl)phenoxy]benzoyl]piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- E169 2-{{1-[4-{{5-cyanopyridin-2-yl}oxy}benzoyl]piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E170 2-{{1-[4-{{5-chloropyridin-2-yl}oxy}benzoyl]piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E171 2-({1-[4-(2,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E172 2-({1-[4-(3,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E173 N-(2-methoxyethyl)-4-methyl-2-{{1-{{4'-(trifluoromethyl)biphenyl-4-yl}carbonyl}piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- E174 2-({1-[4-(3-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E175 2-([1-(2-fluoro-4-isopropoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E176 2-({1-[(3-fluoro-3',4'-dimethylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E177 2-({1-[(2',3-difluoro-4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E178 2-({1-[4-(difluoromethoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E179 2-({1-[4-(2-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E180 2-({1-[(4'-cyano-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E181 2-({1-[4-(5-chloropyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E182 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[6-(trifluoromethyl)pyridin-2-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E183 N-(2-methoxyethyl)-2-({1-[(4'-methoxy-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E184 2-({1-[(4'-chloro-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E185 2-[(1-{4'-(2-cyanopropan-2-yl)biphenyl-4-yl}carbonyl)piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E186 N-(2-methoxyethyl)-2-({1-[4-(5-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E187 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[6-(trifluoromethyl)pyridin-3-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E188 N-(2-methoxyethyl)-2-({1-[4-(6-methoxypyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide

- E189 2-({1-[(4'-fluoro-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E190 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(6-methylpyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- 5 E191 N-(2-methoxyethyl)-2-({1-[4-(6-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E192 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(2-methylpyrimidin-5-yl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E193 2-({1-[(4'-fluoro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E194 2-({1-[(4'-chloro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E195 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[2-(trifluoromethyl)pyrimidin-5-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 15 E196 2-({1-[(4'-chloro-2'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E197 2-({1-[(2'-chloro-4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E198 2-({1-[4-(5-chloropyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E199 2-({1-[4-(5-fluoropyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E200 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[5-(trifluoromethyl)pyridin-3-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 25 E201 2-[(1-{[4'-(2-hydroxypropan-2-yl)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E202 2-({1-[(3',5'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E203 2-({1-[(4'-fluoro-2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E204 2-({1-[(3',5'-difluoro-2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E205 N-(2-methoxyethyl)-4-methyl-2-({1-[3-methyl-4-(3-pyridyl)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide
- E206 N-(2-methoxyethyl)-4-methyl-2-({1-[3-methyl-4-(4-pyridyl)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide
- E207 N-(2-methoxyethyl)-4-methyl-2-({1-[(2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide
- 10

From the 207 compounds, particular preference is given to the compounds having an EP2 receptor antagonism of > 50 nM:

- E2 2-({1-(4-tert-butoxybenzoyl)piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E3 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E5 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E7 2-({1-(2-fluoro-4-methoxybenzoyl)piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E8 2-({1-(4-bromo-2-fluorobenzoyl)piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E9 2-({1-(2-fluoro-4-methylbenzoyl)piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E10 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E11 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pentafluoro- λ^6 -sulphanyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

- E12 N-(2-methoxyethyl)-4-methyl-2-([1-(4-methylbenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E13 2-([1-[4-(4-chlorophenoxy)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E14 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(4-methylphenoxy)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E15 2-([1-[4-(4-tert-butylphenoxy)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E16 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(4-(trifluoromethyl)phenoxy)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E17 N-(2-methoxyethyl)-2-([1-[4-(4-methoxyphenoxy)benzoyl]piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- E18 N-(2-methoxyethyl)-4-methyl-2-([1-(4-phenoxybenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 15 E19 2-([1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E20 2-([1-(4-methoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E21 2-([1-(4-fluorobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E22 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 25 E24 N-(2-methoxyethyl)-4-methyl-2-([1-[4-((trifluoromethyl)sulphonyl]benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E25 N-(2-methoxyethyl)-4-methyl-2-([1-[3-(trifluoromethyl)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide

- E26 N-(2-methoxyethyl)-4-methyl-2-{{1-(3-methylbenzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- E27 2-{{1-(3-chlorobenzoyl)piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E29 2-({1-[4-(cyclopentyloxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E30 2-({1-[4-(difluoromethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E31 2-{{1-(4-cyanobenzoyl)piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E36 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(1H-pyrazol-1-yl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E38 2-({1-[4-(difluoromethoxy)-2-fluorobenzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E39 2-({1-[2-fluoro-4-(pyrrolidin-1-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E40 2-({1-[(3,4'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E41 2-({1-[(3-fluoro-4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E42 2-({1-[(3-fluoro-4'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E43 2-[(1-{[3-fluoro-3'-(trifluoromethyl)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E44 2-[(1-{[3-fluoro-2'-(trifluoromethoxy)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E45 2-({1-[(2'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E46 2-({1-[(2',4'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E47 2-({1-[(2-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E48 N-(2-methoxyethyl)-4-methyl-2-({1-[(2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E51 2-({1-[2-fluoro-4-(morpholin-4-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E52 2-{{1-(4-bromobenzoyl)piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E53 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E54 2-{{1-(4-chlorobenzoyl)piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E55 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[(trifluoromethyl)sulphonyl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E59 2-{{1-(4-chlorobenzoyl)piperidin-4-yl}methyl}-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide
- 20 E60 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-{{1-(4-methylbenzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- E61 N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E62 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E63 N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide

- E64 2- {[1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl}-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide
- E65 2- {[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- 5 E66 4-methyl-2- {[1-(4-methylbenzoyl)piperidin-4-yl]methyl}-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E67 2- ({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl} methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E68 4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2- ({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl} methyl)-2H-indazole-5-carboxamide
- 10 E69 2- ({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E70 2- {[1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl}-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- 15 E71 2- {[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E72 N-(2-isopropoxyethyl)-4-methyl-2- {[1-(4-methylbenzoyl)piperidin-4-yl]methyl}-2H-indazole-5-carboxamide
- E73 2- ({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl} methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E74 N-(2-isopropoxyethyl)-4-methyl-2- ({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl} methyl)-2H-indazole-5-carboxamide
- E75 2- ({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E76 2- {[1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl}-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E77 2- {[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide

- E78 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- E79 2-({1-[4-(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- 5 E80 4-methyl-N-[2-(trifluoromethoxy)ethyl]-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E81 N-(2-tert-butoxyethyl)-2-({1-[4-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methyl-2H-indazole-5-carboxamide
- 10 E82 N-(2-tert-butoxyethyl)-2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E83 N-(2-tert-butoxyethyl)-2-({1-[4-(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E84 N-(2-tert-butoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 15
- E85 2-({1-[4-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methyl-N-{2-[(2H3)methyloxy](2H4)ethyl}-2H-indazole-5-carboxamide
- E86 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-{2-[(2H3)methyloxy](2H4)ethyl}-2H-indazole-5-carboxamide
- 20
- E87 2-({1-[4-(4-chlorobenzoyl)azetidin-3-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E88 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[(trifluoromethyl)sulphonyl]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- 25
- E89 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide
- E90 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E91 2-([1-(4-chloro-2-fluorobenzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E92 2-([1-[3-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E93 2-([1-[4-chloro-3-(trifluoromethyl)benzoyl]azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E94 2-([1-(4-cyclopropylbenzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E95 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(trifluoromethyl)phenoxy]benzoyl]azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- E100 N-[2-(cyclopropyloxy)ethyl]-4-methyl-2-([1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl]methyl)-2H-indazole-5-carboxamide
- 15 E101 N-[2-(cyclobutyloxy)ethyl]-4-methyl-2-([1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl]methyl)-2H-indazole-5-carboxamide
- E102 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide
- E103 N-[2-(cyclopropylmethoxy)ethyl]-2-([1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E104 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-([1-(4-methylbenzoyl)azetidin-3-yl]methyl)-2H-indazole-5-carboxamide
- E105 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- 25 E107 N-[2-(cyclopropylmethoxy)ethyl]-2-([1-[(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- E108 2-([1-[(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E109 2-({1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E112 2-({1-[4-(4-chlorobenzoyl)azetidin-3-yl]methyl}-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- 5 E113 4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- E114 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- 10 E115 2-({1-[4-(4-chlorophenoxy)benzoyl]azetidin-3-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E116 2-({1-[4-(4-chlorophenoxy)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E117 N-(2-methoxyethyl)-4-methyl-2-({1-[4-({5-(trifluoromethyl)pyridin-2-yl}oxy)benzoyl]azetidin-3-yl)methyl}-2H-indazole-5-carboxamide
- E125 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[(trifluoromethyl)sulphonyl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E126 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 20 E127 4-methyl-N-(2,2,2-trifluoroethyl)-2-({1-[4-({5-(trifluoromethyl)pyridin-2-yl}oxy)benzoyl]piperidin-4-yl)methyl}-2H-indazole-5-carboxamide
- E128 4-methyl-N-(2,2,2-trifluoroethyl)-2-({1-[4-({6-(trifluoromethyl)pyridin-3-yl}oxy)benzoyl]piperidin-4-yl)methyl}-2H-indazole-5-carboxamide
- 25 E131 2-({1-[4-(3-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide

- E132 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[3-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 5 E137 2-({1-[4-(3,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E138 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4'-(trifluoromethyl)biphenyl-4-yl}carbonyl)piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E141 2-({1-[(4'-methoxy-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 10 E143 2-({1-[(4'-fluoro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E153 2-({1-[4-(4-cyanophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E154 2-({1-[4-(4-bromo-3-methylbenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E155 2-({1-[4-(4-tert-butylbenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E157 2-({1-[4-(4-cyclohexylbenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E160 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyridin-3-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E161 N-(2-methoxyethyl)-4-methyl-2-({1-[4-{5-(trifluoromethyl)pyridin-2-yl}oxy]benzoyl}piperidin-4-yl)methyl)-2H-indazole-5-carboxamide
- 25 E162 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyridin-2-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E163 N-(2-methoxyethyl)-4-methyl-2-({1-[4-{4-(trifluoromethyl)pyrimidin-2-yl}oxy]benzoyl}piperidin-4-yl)methyl)-2H-indazole-5-carboxamide

- E164 N-(2-methoxyethyl)-4-methyl-2-[[1-(4-{[6-(trifluoromethyl)pyridin-3-yl]oxy}benzoyl)piperidin-4-yl]methyl]-2H-indazole-5-carboxamide
- E165 N-(2-methoxyethyl)-4-methyl-2-[[1-(4-{[6-(trifluoromethyl)pyridin-2-yl]oxy}benzoyl)piperidin-4-yl]methyl]-2H-indazole-5-carboxamide
- 5 E166 2-({1-[(4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E168 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[3-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 10 E170 2-[(1-{4-[(5-chloropyridin-2-yl)oxy]benzoyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E171 2-({1-[4-(2,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E172 2-({1-[4-(3,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E173 N-(2-methoxyethyl)-4-methyl-2-[(1-{[4'-(trifluoromethyl)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E174 2-({1-[4-(3-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E175 2-[[1-(2-fluoro-4-isopropoxybenzoyl)piperidin-4-yl]methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E176 2-({1-[(3-fluoro-3',4'-dimethylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E177 2-({1-[(2',3-difluoro-4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E178 2-({1-[4-(difluoromethoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E179 2-({1-[4-(2-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E180 2-({1-[(4'-cyano-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E181 2-({1-[4-(5-chloropyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E182 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[6-(trifluoromethyl)pyridin-2-yl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E183 N-(2-methoxyethyl)-2-({1-[(4'-methoxy-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E184 2-({1-[(4'-chloro-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E185 2-[(1-{[4'-(2-cyanopropan-2-yl)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E186 N-(2-methoxyethyl)-2-({1-[4-(5-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E187 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[6-(trifluoromethyl)pyridin-3-yl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E188 N-(2-methoxyethyl)-2-({1-[4-(6-methoxypyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E189 2-({1-[(4'-fluoro-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E190 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(6-methylpyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- 25 E191 N-(2-methoxyethyl)-2-({1-[4-(6-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E193 2-({1-[(4'-fluoro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E194 2-({1-[(4'-chloro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E195 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[2-(trifluoromethyl)pyrimidin-5-yl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 5 E196 2-({1-[(4'-chloro-2'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E197 2-({1-[(2'-chloro-4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E198 2-({1-[4-(5-chloropyridin-3-yl)benzoyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E199 2-({1-[4-(5-fluoropyridin-3-yl)benzoyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E200 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[5-(trifluoromethyl)pyridin-3-yl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 15 E201 2-[(1-{[4'-(2-hydroxypropan-2-yl)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E202 2-({1-[(3',5'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E203 2-({1-[(4'-fluoro-2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E204 2-({1-[(3',5'-difluoro-2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E205 N-(2-methoxyethyl)-4-methyl-2-({1-[3-methyl-4-(3-pyridyl)benzoyl]piperidin-4-yl} methyl)-2H-indazole-5-carboxamide
- 25 E206 N-(2-methoxyethyl)-4-methyl-2-({1-[3-methyl-4-(4-pyridyl)benzoyl]piperidin-4-yl} methyl)-2H-indazole-5-carboxamide
- E207 N-(2-methoxyethyl)-4-methyl-2-({1-[(2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-2H-indazole-5-carboxamide

Particular preference is furthermore given to the following compounds:

E16 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide

E126 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide

E161 N-(2-methoxyethyl)-4-methyl-2-{[1-(4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoyl)piperidin-4-yl]methyl}-2H-indazole-5-carboxamide

Administration of the EP2 receptor antagonist according to general formula I or Ia can take place as a single dose or as two doses. The daily dose is 0.1 – 1000 mg, preferably 50 – 600 mg, where the dose may be administered as a single dose to be administered once or divided into 2 or more partial doses.

It may be necessary to deviate from the amounts mentioned of EP2 receptor antagonist of the general formula I or Ia, depending on body weight, administration route, individual response to the active compound, type of preparation and time or interval at which administration takes place. Thus, in some cases it may be sufficient to use less than the minimum amount mentioned above, whereas in other cases the upper limit mentioned must be exceeded.

COX inhibitors which, according to the invention, are contained in the pharmaceutical composition

In principle, COX inhibitors suitable for use according to the invention are all COX inhibitors available for other indications. This means both selective and non-selective COX inhibitors. According to the invention, those suitable include:

- C1 indomethacin;
- C2 diclofenac;
- C4 naproxen;
- 25 C5 ibuprofen;
- C6 meloxicam;
- C7 piroxicam;
- C8 nimesulide;

- C9 ketoprofen;
- C10 tenoxicam;
- C11 mefanemic acid;
- C12 ketoralac;
- 5 C13 celecoxib (4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl]benzenesulphonamide);
- C14 parecoxib (N-[4-(5-methyl-3-phenyl-4-isoxazolyl)phenyl]sulphonylpropionamide);
- C15 rofecoxib (4-(4-methylphenyl)-3-phenylfuran-2(5H)-one);
- C16 valdecoxib (4-[5-methyl-3-phenyl-4-isoxazolyl]benzenesulphonamide);
- 10 C17 etoricoxib.

Particular preference is given to:

- C1 indomethacin;
- C2 diclofenac;
- C4 naproxen;
- 15 C5 ibuprofen;
- C6 meloxicam;
- C7 piroxicam;
- C8 nimesulide.

Very particular preference is given to:

- 20 C1 indomethacin;
- C2 diclofenac;
- C6 meloxicam;
- C7 piroxicam.

In the case of the COX inhibitors, too, administration of the COX inhibitors may be by administration of single doses or by administration of two doses. In the case of the COX inhibitors, too, different dosages are used, depending on the COX inhibitor employed. The amount of COX inhibitor present in the pharmaceutical composition varies, depending on the chosen inhibitor. Dose ranges for the COX inhibitors used according to the invention result from the therapeutic doses per day for the COX inhibitor in question. These therapeutic doses refer to relatively long-term treatments and continuous therapies, so that for the present indication of on-demand contraception, it is also possible to use four times the recommended therapeutic dose. The lower limit is considered to be a quarter of the recommended therapeutic dose. In the case of the COX inhibitors, too, the dose can be administered as a single dose to be administered once or divided into 2 or more partial doses.

According to the invention, the following dose ranges are suitable for on-demand use:

- C1 indomethacin: 19 – 800 mg/day (daily therapeutic dose: 75-200 mg/day*)
- C2 diclofenac: 37.5 – 800 mg/day (daily therapeutic dose: 150 - 200 mg/day*)
- 15 C4 naproxen: 125 – 4000 mg/day (daily therapeutic dose: 500-1000 mg/day*)
- C5 ibuprofen: 200 – 4800 mg/day (daily therapeutic dose: 800-1200 mg/day*)
- C6 meloxicam: 2 – 60 mg/day (daily therapeutic dose: 7.5-15 mg/day)
- C7 piroxicam: 5 – 80 mg/day (daily therapeutic dose: 20 mg/day*)
- C8 nimesulide: 25 – 800 mg/day (daily therapeutic dose: 100-200 mg/day*)
- 20 C9 ketoprofen: 25 – 1200 mg/day (daily therapeutic dose: 100 -300 mg/day*)
- C10 tenoxicam: 10 – 160 mg/day (daily therapeutic dose: 40 mg/day*)
- C11 mefenamic acid: 125 – 2000 mg/day (daily therapeutic dose: 500 mg/day*)
- C12 ketorolac: 2.5 – 160 mg/day (daily therapeutic dose: 10 - 40 mg/day*)
- C13 celecoxib: 25 – 800 mg/day (daily therapeutic dose: 100 - 200 mg/day*)
- 25 C14 parecoxib: 12.5 – 200 mg/day (daily therapeutic dose: 50 mg/day*)
- C15 rofecoxib: 6 – 200 mg/day (daily therapeutic dose: 25 - 50 mg/day*)

C16 valdecoxib: 10 – 160 mg/day (daily therapeutic dose: 40 mg/day)

C17 etoricoxib: 15 – 360 mg/day (daily therapeutic dose: 60 - 90 mg/day)

*Huntjens et al. Rheumatology; (2005); 44:846-859

In the case of the COX inhibitors, too, it may, if required, be necessary to deviate from the
5 above-mentioned amounts of COX inhibitors, namely depending on body weight, administration
route, individual response to the reactive compound, type of preparation and time or interval at
which the administration takes place. Thus, in some cases it may be sufficient to use less than the
minimum amount mentioned above, whereas in other cases the upper limit mentioned must be
exceeded.

10 The details given for the COX inhibitors refer to their combination with EP2 receptor
antagonists according to the general formula I or Ia.

Combinations which are preferred in accordance with the invention comprise an EP2 receptor
antagonist according to general formula Ia from the list

15 E16 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-
4-yl)methyl]-2H-indazole-5-carboxamide

E126 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-
(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide

E161 N-(2-methoxyethyl)-4-methyl-2-[[1-(4-{[5-(trifluoromethyl)pyridin-2-
yl]oxy}benzoyl)piperidin-4-yl)methyl]-2H-indazole-5-carboxamide

20 and a COX inhibitor from the list

C1 indomethacin;

C2 diclofenac;

C4 naproxen;

C5 ibuprofen;

25 C6 meloxicam;

C7 piroxicam;

C8 nimesulide;

or COX inhibitors which are preferably used in accordance with the invention, from the following list:

C1 indomethacin;

5 C2 diclofenac;

C6 meloxicam;

C7 piroxicam.

Use of the pharmaceutical composition for on-demand contraception and fertility control

10 The invention furthermore provides the use of the pharmaceutical composition comprising at least one EP2 receptor antagonist of formula Ia and at least one COX inhibitor for on-demand contraception and for fertility control.

On-demand contraception, fertility control on demand, precoital contraception or contraception on demand all refer to the on-demand use of one or more substances for preventing unwanted pregnancy. The administration can be before or shortly after expected sexual intercourse. Here,
15 on-demand does not mean optional. Rather, the substance/substances has/have to be administered to prevent unwanted pregnancy.

Contraception, birth control or fertility control all refer likewise to the continuous regular intake of one or more substances at low dosage according to a fixed protocol over a relatively long
20 period of time extending over a plurality of cycles. Intake of the pharmacological contraceptive is independent of the frequency and time of sexual intercourse.

Administration route

The pharmaceutical compositions according to the invention and their use can be systematic and/or local. To this end, they can be administered in a suitable manner, for example orally,
25 pulmonarily, nasally, dermally or transdermally. Further administration routes are vaginal or intrauterine. Suitable for vaginal administration are tampons, gels, suppositories, pessaries or intravaginal rings. Suitable for intrauterine administration are, *inter alia*, intrauterine systems or intrauterine pessaries.

Suitable for oral administration are administration forms described in the prior art as being suitable and comprising the EP2 receptor antagonists according to general formula I or Ia according to the invention and COX inhibitors in crystalline and/or amorphized and/or dissolved form, such as tablets (uncoated tablets or coated tablets, for example with enteric coatings or coatings which dissolve in a delayed manner or insoluble coatings controlling the release of the compound according to the invention), tablets which rapidly disintegrate in the oral cavity or films/wafers, films/lyophilizates, capsules (for example hard or soft gelatine capsules), sugar-coated tablets, wafers, granules, pellets, powders, emulsions, suspensions, aerosols or solutions.

Use is also possible in liquid form, such as for example as syrup, to which a sweetener is optionally added. Likewise, for the oral use of such compounds, the clathrates thereof are also suitable, for example the clathrates with alpha, beta or gamma cyclodextrin or else beta-hydroxypropyl cyclodextrin may be mentioned.

Suitable for the other administration routes are, for example, inhalative medicinal forms (*inter alia* powder inhalers, nebulizers), nasal drops, solutions, sprays; tablets, films/wafers or capsules to be administered lingually, sublingually or buccally, suppositories, aqueous suspensions (lotions, shake mixtures), lipophilic suspensions, ointments, creams, transdermal systems (for example patches), milk, pastes, foams or powders for sprinkling.

The pharmaceutical compositions according to the invention can be converted into the stated administration forms. This can take place in a manner known per se by mixing with inert non-toxic pharmaceutically suitable auxiliaries. These auxiliaries include, *inter alia*, carriers (for example microcrystalline cellulose, lactose, mannitol), solvents (for example liquid polyethylene glycols), emulsifiers and dispersants or wetting agents (for example sodium dodecylsulphate, polyoxysorbitane oleate), binders (for example polyvinylpyrrolidone), synthetic and natural polymers (for example albumin), stabilizers (for example antioxidants such as, for example, ascorbic acid), colorants (for example inorganic pigments such as iron oxides) and taste and/or odour corrigents.

They optionally contain additives, which are for example intended to function as fillers, binders, disintegrants, lubricants, solvents, solution mediators, masking flavours, colorant, or emulsifiers. Types of additive in the sense of the invention are for example saccharides (mono-, di-, tri-, oligo-, and/or polysaccharides), fats, waxes, oils, hydrocarbons, anionic, nonionic, cationic, natural, synthetic or semisynthetic surfactants. In addition, they optionally contain additives such as preservatives, stabilizers, wetting agents or emulsifiers; salts for alteration of the osmotic pressure or buffers.

The present invention furthermore provides medicaments or pharmaceutical compositions comprising at least one EP2 receptor antagonist according to general formula I or Ia according to the invention and at least one COX inhibitor, usually together with one or more inert non-toxic pharmaceutically suitable auxiliaries, and their use for on-demand contraception and fertility control.

The pharmaceutical formulation may comprise at least one EP2 receptor antagonist of the general formula I or Ia and at least one COX inhibitor, either together in one administration form (for example a combination tablet) or in two separate administration forms (for example a tablet), where one administration form comprises at least one EP2 receptor antagonist and the other administration form comprises at least one COX inhibitor.

The person skilled in the art is familiar with how to prepare the pharmaceutical formulation. The preparation of a formulation as combination tablet is described in preparation example 1.

Administration protocol

The invention furthermore provides an administration protocol for on-demand contraception where a pharmaceutical composition comprising at least one COX inhibitor from the list mentioned above and at least one EP2 receptor antagonist according to general formula I or Ia are administered in a single administration dose over a period of up to 24 hours, preferably 6 - 12 hours, prior to expected sexual intercourse or of up to 2 hours after sexual intercourse.

In an alternative embodiment, the single dose preparation may also be taken up to at most 24 hours after sexual intercourse.

Also provided is an administration protocol where two separate pharmaceutical compositions each comprising at least one COX inhibitor from the list mentioned above and at least one EP2 receptor antagonist of the general formula I or Ia are administered in succession, at an interval. The compositions of the two pharmaceutical compositions can be identical or different with respect to selection and dosage of the EP2 receptor antagonist of the general formula I or Ia and the COX inhibitor. Administration of the first pharmaceutical composition is within a period of up to 24 hours, preferably 6 - 12 hours, prior to expected sexual intercourse, and administration of the second pharmaceutical composition is within a period of up to 48 hours after the expected sexual intercourse.

In the cases where the two pharmaceutical compositions are different, the dose of one of the active compounds or of both active compounds of the pharmaceutical composition taken prior to

sexual intercourse may be lower than that for the pharmaceutical composition taken after sexual intercourse.

By an appropriate administration protocol, it is possible to achieve sufficiently high active compound plasma levels for a sufficiently long period of time to counteract unwanted pregnancy. Thus, in the female genital tract sperms are viable for about 5 days. If appropriate, administration after sexual intercourse may also take place as a plurality of doses to achieve active plasma concentrations over a period of up to 5 days.

Furthermore, by administering two pharmaceutical compositions it is possible to achieve an even better efficacy of the preparation. Also, in cases where the expected sexual intercourse has not taken place, the active compound load can be reduced by dispensing with the administration which should have taken place after the expected sexual intercourse.

The invention furthermore provides a kit for on-demand contraception, comprising at least two tablets each comprising at least one COX inhibitor and at least one EP2 receptor antagonist of the general formula I or Ia. A first tablet is taken during a period of up to 24 hours, preferably 6 – 12 hours, prior to the expected sexual intercourse, and at least one further tablet is taken during a period of up to 48 hours after the expected sexual intercourse.

According to the invention, the kit may comprise the two tablets up to 5 times to allow on-demand contraception of 5 expected sexual intercourses at intervals of at least 24 hours.

Process for the production of the compounds according to the invention of the general formula I and Ia:

The following description of the production process is already contained in patent application PCT/EP2012/073556, unpublished at the priority date of the present specification.

- 5 For this, for example a carboxylic acid of the formula VIII is reacted with an amine of the general formula XI by methods known to those skilled in the art to give the compounds according to the invention of the general formula I (Scheme 1).

The reaction takes place in that for example a carboxylic acid of the formula VIII is converted with isobutyl chloroformate in the presence of a tertiary amine, for example triethylamine, into a
10 mixed anhydride, which reacts with an alkali metal salt of the appropriate amine XI in an inert solvent or solvent mixture, for example tetrahydrofuran, N,N-dimethylformamide or dimethoxyethane at temperatures between -30°C and +60°C to give the target compounds of the formula I.

It is also possible to activate a carboxylic acid VIII with reagents such as for example dicyclohexylcarbodiimide (DCC), 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide (EDCI),
15 N-hydroxybenzotriazole (HOBT) or N-[(dimethylamino)-(3H-[1,2,3]triazolo[4,5-b]pyridin-3-ylxy)methyliden]-N-methylmethanaminium hexafluorophosphate (HATU). For example the reaction with HATU takes place in an inert solvent, for example N,N-dimethylformamide or dimethyl sulphoxide in the presence of the appropriate amine XI and a tertiary amine, for example triethylamine or diisopropylamine, at temperatures between -30°C and +60°C.

- 20 It is also possible, to convert a carboxylic acid of the formula VIII into the corresponding carboxylic acid chloride with an inorganic acid chloride, for example phosphorus pentachloride, phosphorus trichloride or thionyl chloride and then into the target compounds of the general formula I in pyridine or an inert solvent, such as for example N,N-dimethylformamide, in the presence of the appropriate amine XI and a tertiary amine, for example triethylamine, at
25 temperatures between -30°C and +60°C.

The compounds according to the invention of the general formula I can also be obtained from bromoindazoles of the general formula VI under palladium(0) catalysis by reaction with an appropriate amine XI and carbon monoxide (CO) or a carbon monoxide source, such as for example molybdenum hexacarbonyl in a suitable solvent or solvent mixture, for example
30 1,4-dioxan/water or tetrahydrofuran, addition of a base such as for example sodium carbonate or 1,8-diazabicyclo(5.4.0)undec-7-ene (DBU) and a catalyst-ligand mixture, for example palladium(II) acetate or trans bis(acetato)bis[o-(di-o-tolylphosphino)benzyl]-dipalladium(II) / tri-tert-butylphosphinotetrafluoroborate at temperatures between 80°C and 160°C (optionally

with microwave irradiation between 80-200 watts), and in case of the use of carbon monoxide at a CO pressure of 5-15 bar (Scheme 1).

The compounds according to the invention of the general formula I can also be obtained from amines of the general formula XV by reaction with carboxylic acids ($Y = OH$), chlorides ($Y = Cl$) or anhydrides (e.g. $Y = O-C(O)-O-CH_2(CH_2)_3CH_3$) of the formula IX in the manner described
5 for the production of the compounds I from the compounds VIII and XI (Scheme 2).

Likewise, the compounds of the general formula I can be obtained from compounds of the general formula XVI, wherein LG' for example means Br or I, by reaction with compounds of the formula XVIII (Scheme 2).

10 Compounds of the formula XVIII are for example (Het)Arylboronic acids ($R^3\text{-Met} = (\text{Het})\text{Ar-B(OH)}_2$) or boronic acid pinacol esters ($R^3\text{-Met} = (\text{Het})\text{Ar-BPin}$), which are converted to biaryl compounds of the formula I by methods known to those skilled in the art in a suitable solvent, for example N,N-dimethylformamide, tetrahydrofuran, dimethoxyethane and optionally water and addition of a base, for example triethylamine, potassium carbonate, caesium carbonate and a
15 catalyst-ligand mixture, for example of palladium(II) acetate / triphenylphosphine, bis(diphenylphosphino)ferrocenedichloropalladium (II) at temperatures between 20°C and 120°C.

Compounds of the formula XVIII can also be (Het)Aryl alcohols ($R^3\text{-Met} = (\text{Het})\text{ArO-H}$), which are converted into biaryl ethers of the formula I by methods known to those skilled in the art in a
20 suitable solvent, for example N,N-dimethylformamide or dimethyl sulphoxide with addition of a base, for example potassium carbonate, caesium carbonate under copper(I) catalysis e.g. with copper(I) bromide at temperatures between 60°C and 120°C.

The carboxylic acids of the general formula VIII can for example be obtained from esters of the formula VII by ester saponification in a suitable solvent or solvent mixture, for example
25 methanol, ethanol or tetrahydrofuran, water with addition of an aqueous solution of an alkali metal hydroxide, for example sodium hydroxide or lithium hydroxide at temperatures between 20°C and 60°C (Scheme 1).

In the same manner, the carboxylic acids XIII can be obtained from the esters XII (Scheme 2, PG: e.g. Boc (tert-butyloxycarbonyl) and the carboxylic acids XXI from the esters XX (Scheme
30 3, LG': e.g. Br or I).

The esters of the general formula VII can be obtained from bromoindazoles of the general formula VI under palladium(0) catalysis by reaction with carbon monoxide or a carbon

monoxide source, such as for example molybdenum hexacarbonyl, and methanol in a suitable solvent, for example dimethyl sulfoxide, N,N-dimethylformamide or tetrahydrofuran and addition of a base, for example triethylamine or 1,8-diazabicyclo(5.4.0)undec-7-ene and a catalyst-ligand mixture, for example of palladium(II) acetate / bis-1,3-diphenylphosphino-
5 propane or trans bis(acetato)bis[o-(di-o-tolylphosphino)benzyl]dipalladium(II) / tri-tert-butylphosphinotetrafluoroborate at temperatures between 20°C and 140°C (optionally with microwave irradiation between 80 - 200 watts), and in case of the use of carbon monoxide at a CO pressure of 5 – 15 bar (Scheme 1).

This method is not restricted to methyl esters, i.e. to the use of methanol, but can also be
10 extended to other esters. Thus for example by use of ethanol instead of methanol in this manner, the corresponding ethyl esters can be synthesized.

In the same manner, the esters XII can be obtained from the bromides IV (Scheme 2, -PG: e.g. -Boc).

The amides of the general formula VII can also be obtained from the amines of the general
15 formula XIX by reaction with compounds of the general formula IX (Scheme 3), analogously to the synthesis of the compounds I from the compounds XV (Scheme 2).

The compounds of the general formula VII can also be obtained from compounds of the formula XX and reaction with compounds of the formula XVIII (Scheme 3), analogously to the described conversion of the compounds XVI to the compounds of the formula I (Scheme 2).

20 The amides of the general formula VI can be obtained from amines of the general formula V by reaction with carboxylic acids (Y = OH), chlorides (Y = Cl) or anhydrides (e.g. Y = O-C(O)-O-CH₂(CH)₃CH₃) of the formula IX (Scheme 1), as described for the conversion of amines XV to amides I (Scheme 2).

The amides of the general formula XVI can be obtained in analogous manner from amines XV
25 and carboxylic acids or carboxylic acid derivatives XVII (Y: e.g. OH, Cl or O-C(O)-O-CH₂(CH)₃CH₃; LG': e.g. Br or I) (Scheme 2).

Analogously, amines XX (LG': e.g. Br or I) can be obtained from amines XIX and carboxylic acids or carboxylic acid derivatives XVII (Y: e.g. OH, Cl or O-C(O)-O-CH₂(CH)₃CH₃; LG': e.g. Br or I) (Scheme 3).

Likewise, carboxylic acids XIII (PG: e.g. Boc) can be converted with amines XI to amides XIV (Scheme 2) and carboxylic acids XXI (LG': e.g. Br, I) with amines XI to amides XVI (Scheme 3) in this manner.

The secondary amines V can be obtained from the corresponding carbamates IV (PG: e.g. Boc) by methods known to those skilled in the art (Scheme 1).

Thus for example tert-butyl carbamates can be converted into the amines V in an acidic medium with the use of e.g. trifluoroacetic acid or hydrochloric acid in a suitable solvent or solvent mixture such as for example dichloromethane, dioxan or acetone/water. In an anhydrous medium the amines V are formed as the corresponding salts.

Analogously, the amines XV can be obtained from the carbamates XIV (Scheme 2) and the amines XIX from the carbamates XII (Scheme 3).

The 2*H*-indazoles of the general formulae IV and VI can be produced in various ways.

For example, 2*H*-indazoles IV can be obtained from 1*H*-indazoles of the general formula II by alkylation with compounds of the general formula III (PG: e.g. Boc, LG: e.g. Br, I, O-Ts (tosyloxy) or O-Ms (mesyloxy)) in a suitable solvent, for example N,N-dimethylformamide, N,N-dimethylacetamide, dimethyl sulphoxide or else THF, 1,4-dioxan and addition of a base such as for example potassium carbonate or caesium carbonate (optionally with addition of tetrabutylammonium iodide) or else sodium bis(trimethylsilyl)amide, at temperatures between 25°C and 100°C or else in case of the use of sodium bis(trimethylsilyl)amide 0°C and 25°C (Scheme 1).

Analogously the 2*H*-indazoles of the general formula VI can be obtained from 1*H*-indazoles of the general formula II and compounds of the general formula X (LG: e.g. Br, I, O-Ts or O-Ms) (Scheme 1).

The 2*H*-indazoles of the general formula IV can also be obtained from ortho-nitrobenzaldehydes of the general formula XXVII by reaction with an appropriate amine XXVIII (PG: e.g. Boc) in a suitable solvent, for example 1,4-dioxan and addition of a reducing agent such as for example triethyl phosphite, possibly with addition of potassium carbonate or caesium carbonate or powdered molecular sieve, at temperatures between 100°C and 160°C (Scheme 4).

Analogously, the 2*H*-indazoles of the general formula VI can be obtained from ortho-nitrobenzaldehydes XXVII by reaction with amines XXIX (Scheme 4).

The compounds of the general formula XIV (PG: e.g. Boc) can also be obtained from bromo-indazoles of the general formula IV (PG: e.g. Boc) under palladium(0) catalysis by reaction with an appropriate amine XI and carbon monoxide or a carbon monoxide source, such as for example molybdenum hexacarbonyl (Scheme 2), analogously to the described process for the
5 conversion of the bromoindazoles VI to the compounds of the general formula I, (Scheme 1).

The ortho-nitrobenzaldehydes of the general formula XXVII can be produced from the ortho-nitrotoluenes of the general formula XXVI by methods known to those skilled in the art in two reaction steps (Scheme 4).

For this, an ortho-nitrotoluene XXVI is dissolved in a suitable solvent such as N,N-dimethyl-
10 formamide and converted with N,N-dimethylformamide dimethyl acetal at temperatures of 100 – 140°C to the corresponding enamine, which is immediately oxidized at temperatures of 0°C – 20°C to the corresponding ortho-nitrobenzaldehyde with a suitable oxidizing agent such as for example NaIO₄ in a suitable aqueous solvent mixture such as water/N,N-dimethylformamide and optionally with addition of a base such as for example triethylamine, N,N-diisopropyl-
15 ethylamine, sodium hydrogen carbonate or sodium carbonate.

The ortho-nitrotoluenes of the general formula XXVI can be produced from the ortho-methylanilines of the general formula XXV by methods known to those skilled in the art (Scheme 4).

For this, an ortho-methylaniline XXV is dissolved in a suitable solvent such as dichloromethane
20 or chloroform, treated with zirconium(IV) tert-butoxide, ground molecular sieve 3Å and tert-butyl hydroperoxide and reacted at temperatures of 20-40°C.

The *1H*-indazoles of the general formula II can be liberated from the acetamides of the general formula XXIV by methods known to those skilled in the art (Scheme 4).

For this, for example an acetamide XXIV is reacted in a suitable solvent such as methanol or
25 ethanol and addition of 37% hydrochloric acid at temperatures of 40 – 80°C.

Analogously thereto, the anilines of the formula XXV can be liberated from the acetanilides of the formula XXIII (Scheme 4).

The acetamides of the general formula XXIV can be produced from the ortho-methyl-acetanilides of the general formula XXIII by methods known to those skilled in the art (Scheme
30 4).

For this, the ortho-methyl-acetanilides XXIII are dissolved in a suitable solvent such as chloroform or toluene, treated with acetic anhydride, isopentyl or tert-butyl nitrite, and optionally potassium acetate and 18-crown-6, and reacted at temperatures of 60 – 100°C.

The ortho-methyl-acetanilides XXIII can be produced from the acetanilides of the general formula XXII by bromination by methods known to those skilled in the art (Scheme 4).

For this, the acetanilides XXII, which can for example be obtained from the corresponding anilines by reaction with acetic anhydride in a suitable solvent (e.g. toluene) at temperatures of 80-110°C, is reacted with bromine in glacial acetic acid at temperatures of 10-25°C.

The compounds of the general formula III wherein LG means -OTs or -OMs can be produced by methods known to those skilled in the art from the corresponding alcohols.

For this, the alcohols are reacted with p-toluenesulphonyl chloride or methanesulphonyl chloride at temperatures between 0°C and 40°C in a suitable solvent, e.g. dichloromethane or tetrahydrofuran or toluene and addition of a suitable base, e.g. pyridine or triethylamine.

The compounds of the general formula X, wherein LG means -OTs or -OMs, can be produced from the corresponding amino alcohols XXX by methods known to those skilled in the art (Scheme 4).

For this, in a first step the amino alcohols XXX are converted to the corresponding amides XXXI in a suitable solvent, e.g. dichloromethane, with arylcarboxylic acid chlorides of the formula Cl-C(O)-Ar-R³ and addition of a suitable base, e.g. triethylamine, at temperatures between 0°C and 25°C. Optionally, corresponding esters formed as by-products can be separated or else saponified to the corresponding alcohols XXXI under standard conditions, e.g. with use of a base, e.g. potassium hydroxide, in a suitable solvent mixture, e.g. ethanol/water, at temperatures between 20°C and 40°C

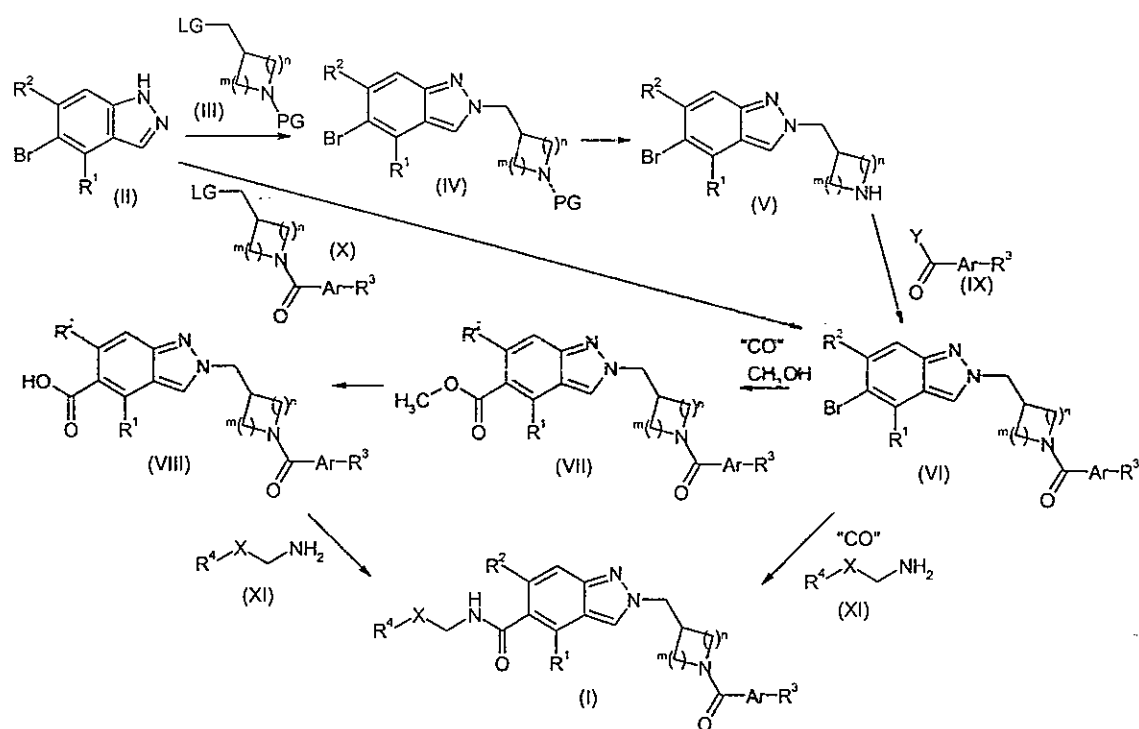
The N-protected alcohols XXXI thus obtained can be converted into the compounds of the formula X with LG: -OTs or -OMs analogously to the process described for the synthesis of the compounds III. Compounds of the formula X with LG: -Br can be obtained from the corresponding alcohols XXXI by methods known to those skilled in the art by reaction with a brominating agent such as for example CBr₄ with addition of PPh₃ as an oxophile in a suitable solvent, e.g. chloroform, at temperatures between 20°C and 40°C.

The compounds of the general formula XXIX, can be produced by methods known to those skilled in the art for example from the amines of the formula XXXII (Scheme 4).

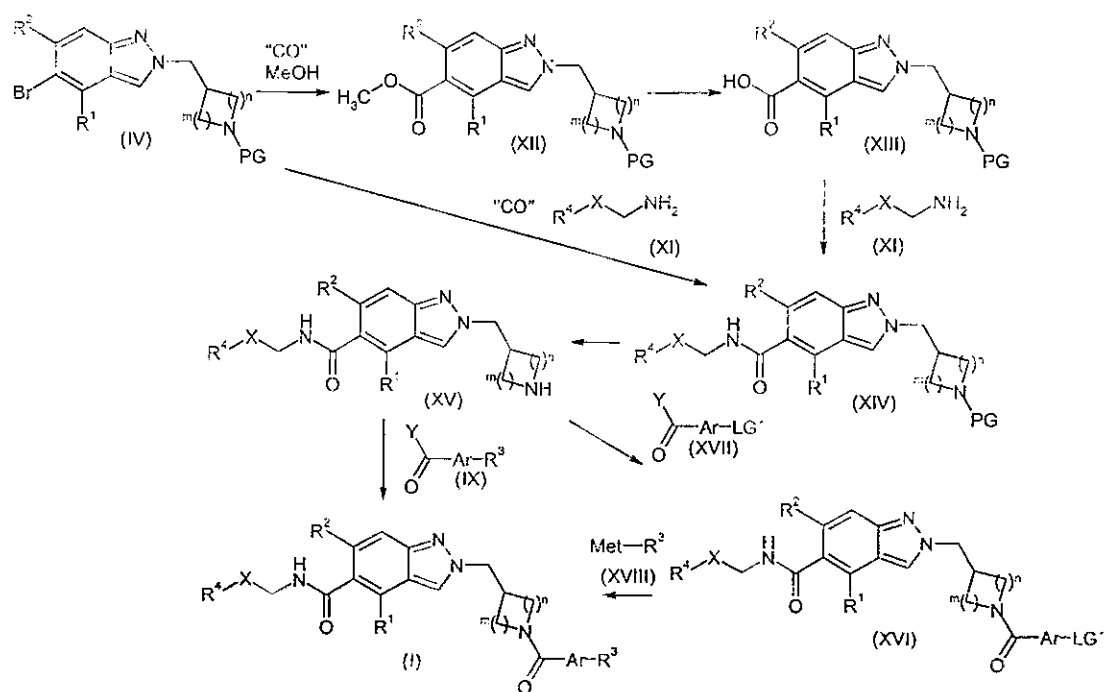
For this, in a first step, the amines XXXII are converted to the corresponding amides XXXIII analogously to the synthesis of the compounds I from the compounds XV.

The tert-butyloxycarbonyl group in compounds of the formula XXXIII is cleaved analogously to the conversion of the compounds IV to the compounds V, hence the compounds of the formula
5 XXIX can be obtained.

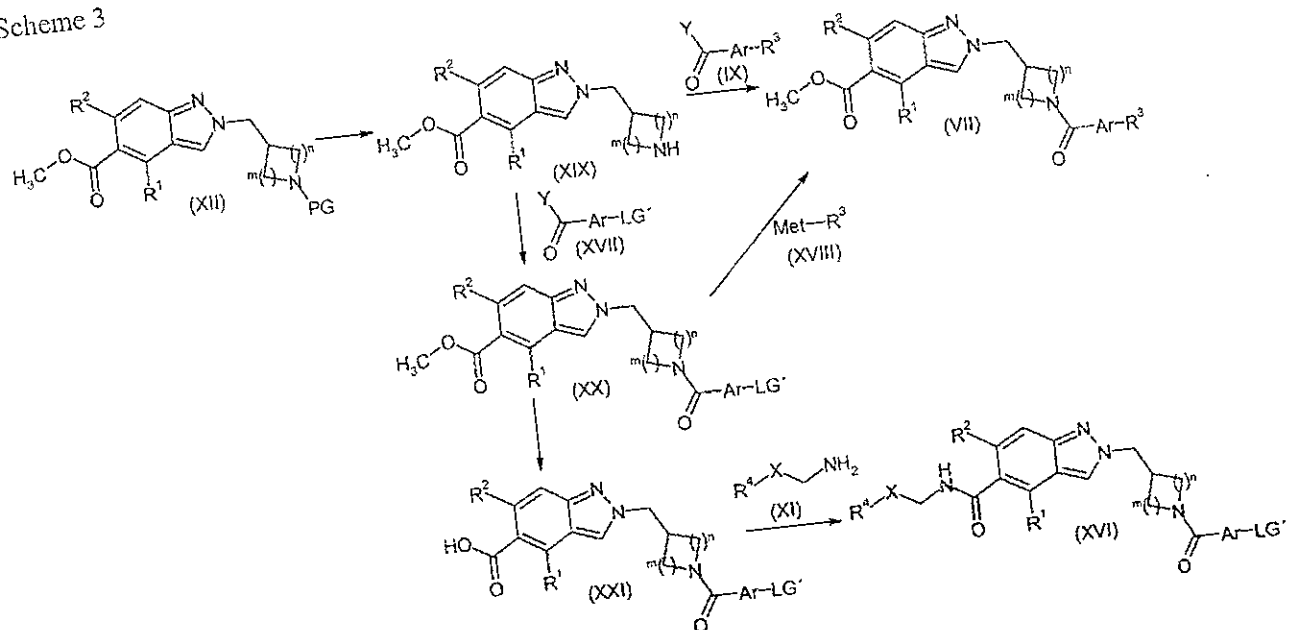
Scheme 1



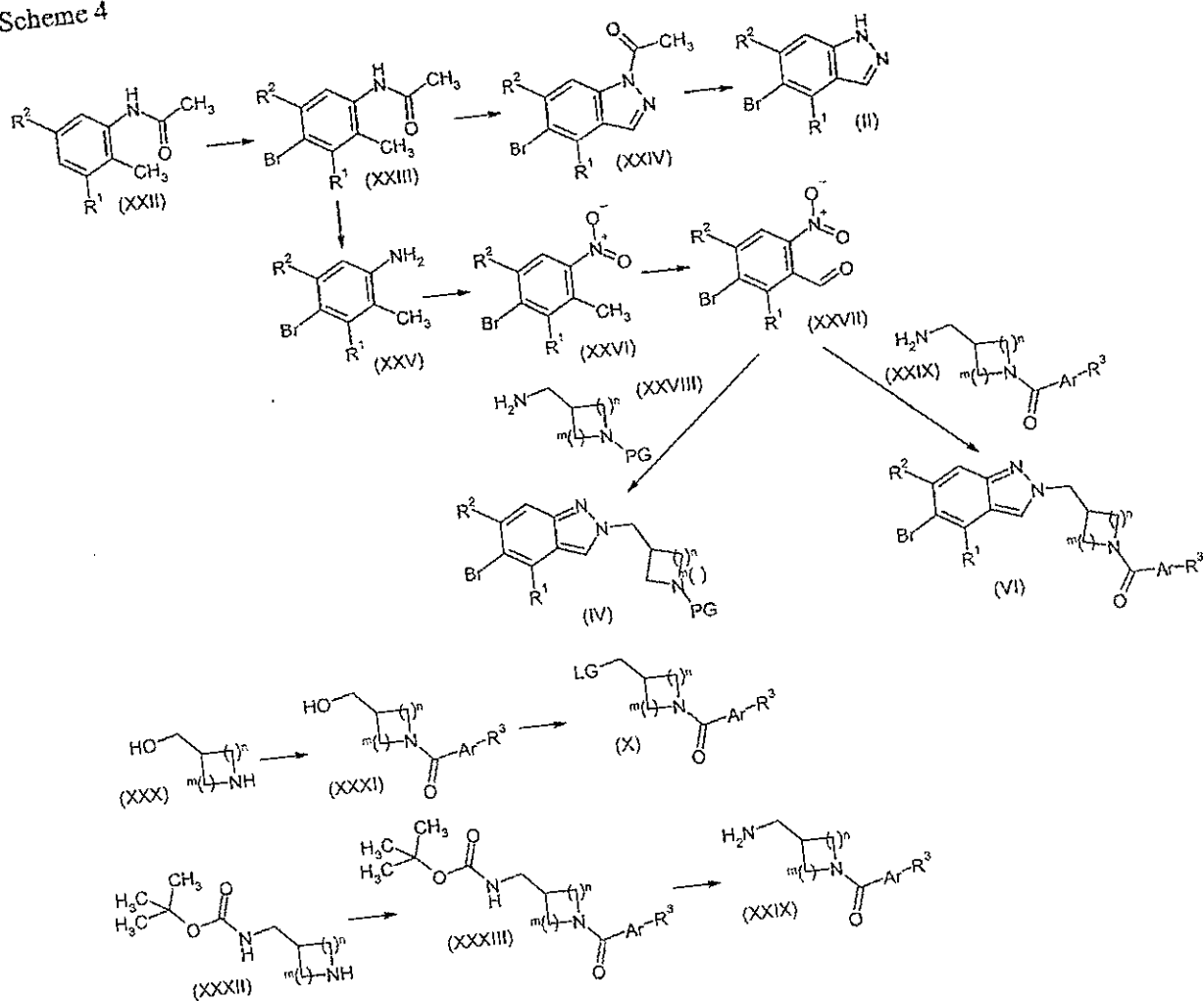
Scheme 2



Scheme 3



Scheme 4



Production of the compounds according to the invention:

The following description of the production of the compounds according to the invention is already contained in patent application PCT/EP2012/073556, unpublished at the priority date of the present specification.

- 5 The following examples illustrate the production of the compounds according to the invention of the general formula (I) and Ia, without limiting the range of the claimed compounds to these examples.

The compounds according to the invention of the general formula (I) can be produced and characterized as described below.

- 10 LC-MS: Waters Acquity HPLC / MS 100-800 Daltons; 20 V (Micromass / Waters ZQ 4000); Column: BEHC 18 (Waters), 2.1 x 50 mm, BEH 1.7 μ m; Mobile phase: A: H₂O/0.05% HCOOH, B: CH₃CN/0.05% HCOOH. Gradient: 10-90% B in 1.7 min, 90% B for 0.2 min, 98-2% B in 0.6 min; Flow rate: 1.3 ml / min, Detection: UV = 200 - 400 nm.

Chiral HPLC separation method A:

- 15 Preparative chiral HPLC: System: Dionex: Pump P 580, Gilson: Liquid Handler 215, Knauer: UV detector K-2501 Column: Chiralpak AD-H 5 μ m 250x20 mm; Solvent: hexane / ethanol 50:50; Flow rate: 15 ml/min; injection volume: 0.5 ml; Detection: UV 254 nm.

Analytical chiral HPLC: System: Waters: Alliance 2695, DAD 996, ESA: Corona; Column: Chiralpak AD-H 5 μ m 150x4.6 mm Solvent: hexane / ethanol 50:50; Flow rate: 1.0 ml/min;

- 20 Temperature: 25°C; injection: 5 μ l; Detection: DAD 254 nm.

Chiral HPLC separation method B:

Preparative chiral HPLC: System: Dionex: Pump P 580, Gilson: Liquid Handler 215, Knauer: UV detector K-2501 Column: Chiralpak IC 5 μ m 250x30 mm; Solvent: ethanol / methanol 50:50; Flow rate: 30 ml/min; injection volume: 0.5 ml; Detection: UV 254 nm.

- 25 Analytical chiral HPLC: System: Waters: Alliance 2695, DAD 996, ESA: Corona; Column: Chiralpak IC 5 μ m 150x4.6 mm Solvent: ethanol / methanol 50:50; Flow rate: 1.0 ml/min; Temperature: 25°C; injection: 5 μ l; Detection: DAD 254 nm.

Chiral HPLC separation method C:

Preparative chiral HPLC: System: Dionex: Pump P 580, Gilson: Liquid Handler 215, Knauer: UV detector K-2501 Column: Chiralpak AD-H 5 μ m 250x20 mm; Solvent: hexane / 2-propanol 50:50; Flow rate: 15 ml/min; injection volume: 0.25 ml; Detection: UV 254 nm.

Analytical chiral HPLC: System: Waters: Alliance 2695, DAD 996, ESA: Corona; Column:
5 Chiralpak AD-H 5 μ m 150x4.6 mm Solvent: hexane / 2-propanol 50:50; Flow rate: 1.0 ml/min; Temperature: 25°C; injection: 5 μ l; Detection: DAD 254 nm.

Chiral HPLC separation method D:

Preparative chiral HPLC: System: Dionex: Pump P 580, Gilson: Liquid Handler 215, Knauer: UV detector K-2501 Column: Chiralpak IB 5 μ m 250x20 mm; Solvent: hexane / ethanol 70:30;
10 Flow rate: 20 ml/min; injection volume: 0.1 ml; Detection: UV 210 nm.

Analytical chiral HPLC: System: Waters: Alliance 2695, DAD 996, ESA: Corona; Column: Chiralpak IB 5 μ m 150x4.6 mm Solvent: hexane / ethanol 70:30; Flow rate: 1.0 ml/min; Temperature: 25°C; injection: 5 μ l; Detection: DAD 230 nm.

Chiral HPLC separation method E:

15 Preparative chiral HPLC: System: Dionex: Pump P 580, Gilson: Liquid Handler 215, Knauer: UV detector K-2501 Column: Chiralpak IC 5 μ m 250x20 mm; Solvent: ethanol / methanol 50:50; Flow rate: 15 ml/min; injection volume: 0.3 ml; Detection: UV 230 nm.

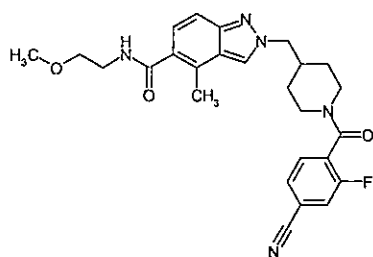
Analytical chiral HPLC: System: Waters: Alliance 2695, DAD 996, ESA: Corona; Column: Chiralpak IC 5 μ m 150x4.6 mm Solvent: ethanol / methanol 50:50; Flow rate: 1.0 ml/min;
20 Temperature: 25°C; injection: 5 μ l; Detection: DAD 230 nm.

Abbreviations

Boc	tert-butoxycarbonyl
CO	carbon monoxide
DBU	1,8-diazabicyclo[5.4.0]undec-7-ene
DCM	dichloromethane
DMF	N,N-dimethylformamide
DMSO	dimethyl sulphoxide

ESI; ES ⁺	<i>electrospray ionization (in MS); positive charged ion trace</i>
hr(s)	<i>hour(s)</i>
HATU	<i>N-[(dimethylamino)-1H-1,2,3-triazolo[4,5-b]pyridin-1-ylmethylene]-N-methylmethanaminium hexafluorophosphate N-oxide</i>
HPLC	<i>high pressure, high performance liquid chromatography</i>
conc.	concentrated
LC-MS	liquid chromatography-coupled mass spectroscopy
M	molar
min(s)	minute(s)
MS	mass spectroscopy
N	normal
NMR	nuclear resonance spectroscopy
R _t	retention time (in HPLC and LC)
RT	room temperature
tert	tertiary
THF	tetrahydrofuran

Example 1: 2-[[1-(4-cyano-2-fluorobenzoyl)piperidin-4-yl]methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

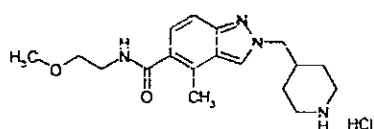


To a solution of 33.7 mg of 4-cyano-2-fluorobenzoic acid in 1.0 ml dimethyl sulphoxide were added 85.5 mg of HATU, 75 mg of the amine prepared in Example 1a and 0.071 ml of N,N-diisopropylethylamine and this was stirred for 1 hour at 25°C. The mixture was concentrated in vacuo and the residue thus obtained purified by HPLC. Yield: 43.3 mg of the title compound.

- 5 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.07 - 1.28 (2H), 1.30 - 1.44 (1H), 1.47 - 1.61 (1H), 2.18 - 2.34 (1H), 2.49 (3H), 2.71 - 2.83 (1H), 2.92 - 3.08 (1H), 3.32 (4H), 3.36 (2H), 3.42 (2H), 4.31 (2H), 4.39 - 4.51 (1H), 7.14 (1H), 7.38 (1H), 7.50 - 7.63 (1H), 7.71 - 7.79 (1H), 7.95 (1H), 8.09 - 8.15 (1H), 8.47 (1H).

The starting material for the above title compound was prepared as follows:

- 10 **Example 1a:** N-(2-methoxyethyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide hydrochloride

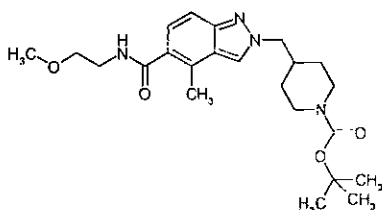


- To 855.1 mg of 1b were added 4.5 ml of 4M hydrochloric acid in dioxan and 1 ml dioxan. An oily mass was formed, which dissolved on vigorous stirring and gentle warming. The reaction mixture was stirred for 1 hr at 30°C. The reaction mixture was concentrated. Yield: 764.7 mg of the title compound, which was further reacted without further purification.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.33 - 1.50 (2H), 1.52 - 1.62 (2H), 2.17 - 2.31 (1H), 2.32 - 2.38 (1H), 2.49 (3H), 2.67 - 2.86 (2H), 3.13 - 3.28 (5H), 3.31 - 3.46 (4H), 4.33 (2H), 7.16 (1H), 7.38 (1H), 8.13 (1H), 8.53 (1H), 8.71 - 8.88 (1H), 8.99 - 9.11 (1H).

20

Example 1b: Tert-butyl 4-({5-[N-(2-methoxyethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}-methyl)piperidin-1-carboxylate



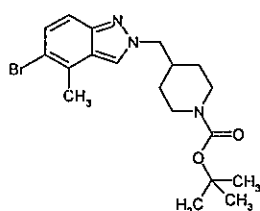
- Version A: 2820 mg of 1c, 1556 mg of 2-methoxyethylamine, 1823 mg of molybdenum hexacarbonyl, 200.4 mg of tri-tert-butylphosphine tetrafluoroborate and 647.5 mg of trans-
- 25

bis(acetato)-bis[o-(di-o-tolylphosphino)benzyl]dipalladium(II) were placed in portions in three microwave tubes and suspended with 56 ml THF. Then 3.1 ml of DBU were added and the mixtures stirred for 20 mins at 125°C and 200 watts in the microwave. The reaction mixtures were combined, filtered and diluted with some ethyl acetate and the organic phase washed twice
5 with water and once with saturated sodium chloride solution. This was dried over sodium sulphate, filtered and concentrated. The residue was chromatographed on the Biotage SP4. Gradient: ethyl acetate/methanol 0-10%. Yield: 885.1 mg of the title compound.

Version B: 780 mg of 1d and 1747 mg of HATU were first dissolved in 10 ml DMSO. Next, 314 mg of 2-methoxyethylamine and 1080 mg of N,N-diisopropylethylamine were added. The
10 mixture was stirred for 1 hr at RT. The reaction mixture was poured into water and extracted three times with ethyl acetate. The combined organic phases were washed with saturated sodium chloride solution, dried over sodium sulphate, filtered and concentrated. The residue was purified chromatographically on the Biotage SP4. Gradient: ethyl acetate/methanol 0-10%. Yield: 740 mg of the title compound.

15 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.01 - 1.13 (2H), 1.34 (9H), 1.36 - 1.43 (2H), 2.06 - 2.19 (1H), 2.49 (3H), 2.59 - 2.70 (2H), 3.29 (3H), 3.33 - 3.39 (2H), 3.40 - 3.45 (2H), 3.82 - 3.93 (2H), 4.28 (2H), 7.14 (1H), 7.38 (1H), 8.09 - 8.14 (1H), 8.46 (1H).

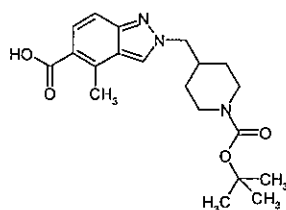
Example 1c: Tert-butyl 4-[(5-bromo-4-methyl-2H-indazol-2-yl)methyl]piperidin-1-carboxylate



20 5 g of 5-bromo-4-methyl-1H-indazole was dissolved in 110 ml DMF and treated with 11.5 g of caesium carbonate and 7.9 g of N-Boc-4-(bromomethyl)piperidine. The mixture was stirred for 3 hrs at 60°C and overnight at RT. The reaction mixture was next diluted with ethyl acetate, and the organic phase was washed twice with water, dried over sodium sulphate, filtered and concentrated. The residue was purified chromatographically on the Biotage SP4 via a 65i-Si
25 column. Gradient: hexane/ethyl acetate 0-100%. Yield: 3.53 g of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.99 - 1.12 (2H), 1.34 (9H), 1.36 - 1.44 (2H), 2.04 - 2.19 (1H), 2.47 (3H), 2.54 - 2.72 (2H), 3.82 - 3.93 (2H), 4.27 (2H), 7.29 (1H), 7.34 (1H), 8.46 (1H).

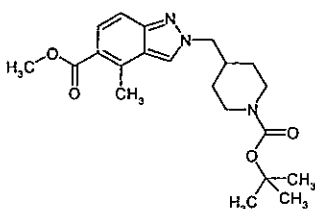
Example 1d: 2-{{1-(tert-butoxycarbonyl)piperidin-4-yl}methyl}-4-methyl-2H-indazol-5-carboxylic acid



1080 mg of 1e were dissolved in 8 ml methanol and treated with 1235 mg of lithium hydroxide in 10 ml water. A further 2 ml THF were added as solubilizer. The mixture was stirred for 24 hrs at RT. Next the methanol and THF were distilled off. The aqueous residue was diluted with water and washed once with ethyl acetate. The aqueous phase was acidified with 10% sulphuric acid and extracted twice with ethyl acetate. The combined organic phases were dried over sodium sulphate, filtered and concentrated. Yield: 880 mg of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.99 - 1.18 (2H), 1.37 - 1.48 (2H), 1.34 (9H), 2.05 - 2.21 (1H), 2.53 - 2.70 (2H), 2.73 (3H), 3.81 - 3.94 (2H), 4.29 (2H), 7.39 (1H), 7.62 (1H), 8.62 (1H), 11.92 - 12.31 (1H).

Example 1e: Methyl 2-{{1-(tert-butoxycarbonyl)piperidin-4-yl}methyl}-4-methyl-2H-indazol-5-carboxylate

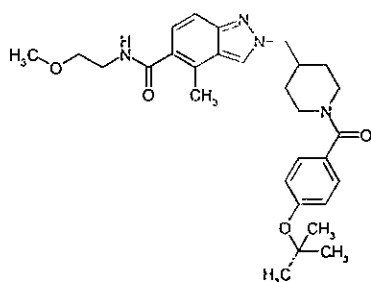


666 mg of 1c, 156.8 mg of methanol, 645.8 mg of molybdenum hexacarbonyl, 47.3 mg of tri-tert-butylphosphine tetrafluoroborate and 123.5 mg of trans-bis(acetato)-bis[o-(di-o-tolylphosphino)benzyl]dipalladium(II) were placed in a microwave tube and suspended in 15 ml THF. Then 0.7 ml of DBU were added and the mixture was stirred for 20 mins at 125°C and 150 watts in the microwave. Next, it was concentrated, taken up in ethyl acetate and the organic phase washed twice with water and once with saturated sodium chloride solution. It was dried over sodium sulphate, filtered and concentrated. The residue was chromatographed on the Biotage SP4. Gradient: hexane/ethyl acetate 0-100%. Yield: 363 mg of the title compound.

- 64 -

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.01 - 1.12 (2H), 1.34 (9H), 1.37 - 1.45 (2H), 2.08 - 2.19 (1H), 2.53 - 2.69 (2H), 2.73 (3H), 3.79 (3H), 3.83 - 3.93 (2H), 4.30 (2H), 7.42 (1H), 7.62 (1H), 8.67 (1H).

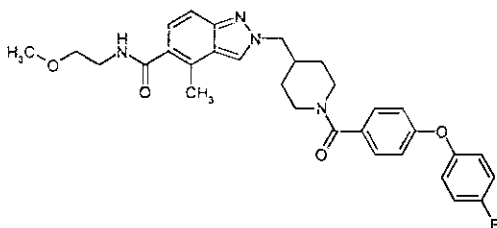
Example 2: 2-({1-[4-tert-(butoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 35.9 mg of the title compound was obtained from 75 mg of the amine prepared in Example 1a and 39.7 mg of 4-(tert-butoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.09 - 1.26 (2H), 1.29 (9H), 1.35 - 1.58 (2H), 2.15 - 2.35 (1H), 2.49 (3H), 2.69 - 2.97 (2H), 3.24 (3H), 3.32 - 3.47 (4H), 4.23 - 4.40 (2H), 6.98 (2H), 7.14 (1H), 7.23 (2H), 7.38 (1H), 8.10 (1H), 8.47 (1H).

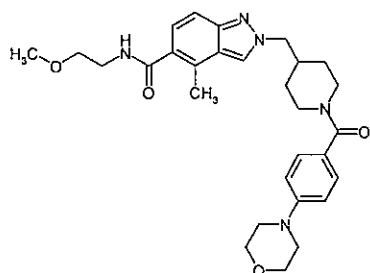
Example 3: 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 51.6 mg of the title compound was obtained from 75 mg of the amine prepared in Example 1a and 47.5 mg of 4-(4-fluorophenoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.10 - 1.30 (2H), 1.32 - 1.58 (2H), 2.19 - 2.35 (1H), 2.49 (3H), 2.71 - 3.02 (2H), 3.25 (3H), 3.32 - 3.39 (2H), 3.40 - 3.47 (2H), 4.32 (2H), 6.95 (2H), 7.06 - 7.14 (3H), 7.15 - 7.27 (3H), 7.31 - 7.42 (3H), 8.07 - 8.15 (1H), 8.47 (1H).

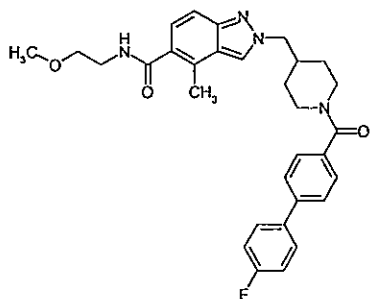
Example 4: N-(2-methoxyethyl)-4-methyl-2--{[1-(4-morpholinobenzoyl)piperidin-4-yl]methyl}-2H-indazole-5-carboxamide



Analogously to Example 1, 48.1 mg of the title compound was obtained from 75 mg of the amine prepared in Example 1a and 42.3 mg of 4-morpholinobenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.09 - 1.29 (2H), 1.36 - 1.53 (2H), 2.16 - 2.34 (1H), 2.49 (3H), 2.72 - 2.94 (2H), 3.13 (4H), 3.24 (3H), 3.42 (4H), 3.64 - 3.75 (4H), 3.92 - 4.15 (1H), 4.32 (2H), 6.92 (2H), 7.14 (1H), 7.20 (2H), 7.38 (1H), 8.07 - 8.15 (1H), 8.47 (1H).

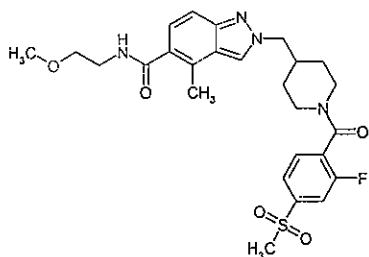
Example 5: 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 50.1 mg of the title compound was obtained from 75 mg of the amine prepared in Example 1a and 44.2 mg of 4-(4-fluorophenyl)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.11 - 1.32 (2H), 1.32 - 1.64 (2H), 2.20 - 2.36 (1H), 2.49 (3H), 2.63 - 3.08 (2H), 3.24 (3H), 3.42 (4H), 3.50 - 3.73 (1H), 4.24 - 4.55 (3H), 7.10 - 7.20 (1H), 7.27 (2H), 7.41 (3H), 7.67 (4H), 8.03 - 8.16 (1H), 8.48 (1H).

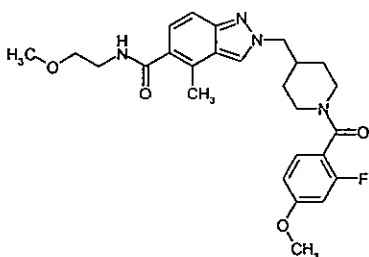
Example 6: 2-({1-(2-fluoro-4-mesylbenzoyl)piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 40.1 mg of the title compound was obtained from 75 mg of the amine prepared in Example 1a and 44.6 mg of 2-fluoro-4-mesyloxybenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.06 - 1.30 (2H), 1.31 - 1.45 (1H), 1.51 - 1.64 (1H),
 5 2.20 - 2.37 (1H), 2.49 (3H), 2.71 - 2.85 (1H), 2.91 - 3.10 (1H), 3.24 (3H), 3.26 (3H), 3.33 - 3.39
 (2H), 3.42 (2H), 4.24 - 4.39 (2H), 4.40 - 4.51 (1H), 7.14 (1H), 7.38 (1H), 7.59 - 7.69 (1H), 7.76 -
 7.89 (2H), 8.07 - 8.14 (1H), 8.47 (1H).

Example 7: 2-({[1-(2-fluoro-4-methoxybenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

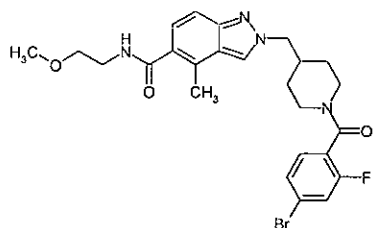


10

Analogously to Example 1, 43.1 mg of the title compound was obtained from 75 mg of the amine prepared in Example 1a and 34.8 mg of 2-fluoro-4-methoxybenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.05 - 1.25 (2H), 1.32 - 1.44 (1H), 1.48 - 1.61 (1H),
 2.18 - 2.33 (1H), 2.49 (3H), 2.63 - 2.79 (1H), 2.88 - 3.05 (1H), 3.24 (3H), 3.42 (d, 5H), 3.75
 15 (3H), 4.27 - 4.35 (2H), 4.37 - 4.49 (1H), 6.77 - 6.89 (2H), 7.14 (1H), 7.24 (1H), 7.38 (1H), 8.07 -
 8.15 (1H), 8.47 (1H).

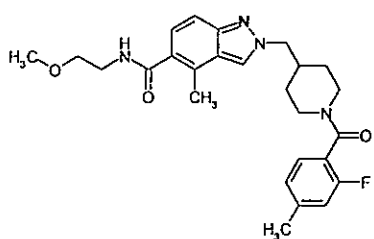
Example 8: 2-({[1-(4-bromo-2-fluorobenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 51.8 mg of the title compound was obtained from 75 mg of the amine prepared in Example 1a and 44.7 mg of 4-bromo-2-fluorobenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.04 - 1.27 (2H), 1.32 - 1.43 (1H), 1.49 - 1.61 (1H),
 5 2.20 - 2.33 (1H), 2.48 (3H), 2.67 - 2.80 (1H), 2.91 - 3.05 (1H), 3.24 (3H), 3.33 - 3.46 (5H), 4.24
 - 4.36 (2H), 4.38 - 4.49 (1H), 7.15 (1H), 7.27 - 7.34 (1H), 7.35 - 7.41 (1H), 7.44 - 7.50 (1H),
 7.60 - 7.66 (1H), 8.07 - 8.15 (1H), 8.47 (1H).

Example 9: 2-({[1-(2-fluoro-4-methylbenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

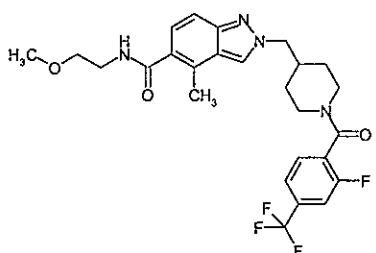


10

Analogously to Example 1, 38.3 mg of the title compound was obtained from 75 mg of the amine prepared in Example 1a and 31.5 mg of 2-fluoro-4-methylbenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.04 - 1.26 (2H), 1.31 - 1.43 (1H), 1.48 - 1.60 (1H),
 2.18 - 2.35 (4H), 2.48 (3H), 2.64 - 2.79 (1H), 2.89 - 3.04 (1H), 3.24 (3H), 3.33 - 3.39 (3H), 3.39
 15 - 3.45 (2H), 4.23 - 4.36 (2H), 4.39 - 4.49 (1H), 7.01 - 7.10 (2H), 7.13 (1H), 7.16 - 7.24 (1H),
 7.38 (1H), 8.07 - 8.14 (1H), 8.47 (1H).

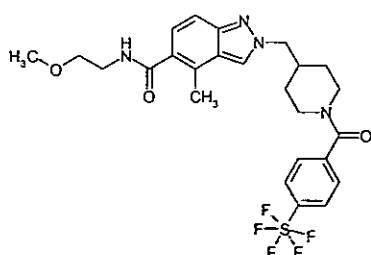
Example 10: 2-({[1-[2-fluoro-4-(trifluoromethyl)benzoyl]piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 57.8 mg of the title compound was obtained from 75 mg of the amine prepared in Example 1a and 42.5 mg of 2-fluoro-4-(trifluoromethyl)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.07 - 1.29 (2H), 1.32 - 1.44 (1H), 1.52 - 1.62 (1H), 2.20 - 2.36 (1H), 2.49 (3H), 2.67 - 2.84 (1H), 2.93 - 3.07 (1H), 3.24 (3H), 3.32 - 3.39 (3H), 3.39 - 3.46 (2H), 4.24 - 4.39 (2H), 4.40 - 4.51 (1H), 7.14 (1H), 7.38 (1H), 7.56 - 7.68 (2H), 7.76 (1H), 8.07 - 8.14 (1H), 8.47 (1H).

Example 11: N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pentafluoro-λ⁶-sulphonyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 135.1 mg of the title compound was obtained from 150 mg of the amine prepared in Example 1a and 101.5 mg of 4-(pentafluoro-λ⁶-sulphonyl)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.10 - 1.30 (2H), 1.31 - 1.42 (1H), 1.50 - 1.62 (1H), 2.20 - 2.33 (1H), 2.49 (3H), 2.67 - 2.83 (1H), 2.91 - 3.06 (1H), 3.24 (3H), 3.32 - 3.39 (2H), 3.41 (3H), 4.32 (2H), 4.37 - 4.48 (1H), 7.14 (1H), 7.38 (1H), 7.55 (2H), 7.94 (2H), 8.12 (1H), 8.47 (1H).

The following compounds were prepared analogously:

Ex.	Structure	IUPAC name	Analysis
12		N-(2-methoxyethyl)-4-methyl-2-{{1-[4-methylbenzoyl]piperidin-4-yl}methyl}-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.20 (2H), 1.42 (2H), 2.29 (4H), 2.49 (3H), 2.73 (1H), 2.93 (1H), 3.25 (3H), 3.37 (2H), 3.43 (2H), 3.57 (1H), 4.31 (2H), 4.40 (1H), 7.14 (1H), 7.21 (4H), 7.38 (1H), 8.09 (1H), 8.46 (1H).

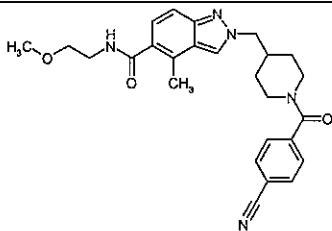
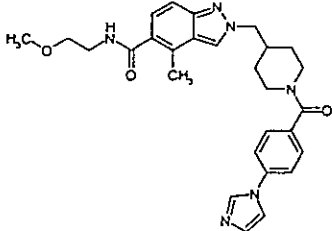
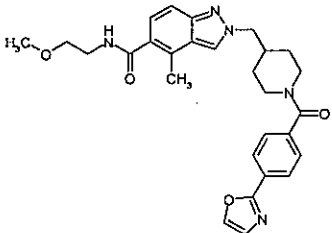
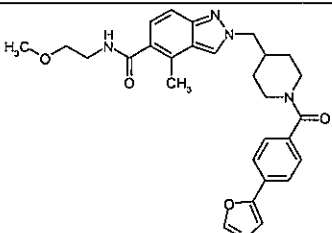
Ex.	Structure	IUPAC name	Analysis
13		N-(2-methoxyethyl)-4-methyl-2-((1-((3-phenylisoxazol-5-yl)carbonyl)piperidin-4-yl)methyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.25 (2H), 1.55 (2H), 2.34 (1H), 2.49 (3H), 2.83 (1H), 3.15 (1H), 3.24 (3H), 3.40 (4H), 3.87 (1H), 4.35 (3H), 7.15 (1H), 7.40 (1H), 7.46 (1H), 7.50 (3H), 7.90 (2H), 8.13 (1H), 8.49 (1H).
14		2-((1-[4-(4-chlorophenoxy)benzoyl]piperidin-4-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.21 (2H), 1.45 (2H), 2.26 (1H), 2.49 (3H), 2.76 (1H), 2.94 (1H), 3.24 (3H), 3.40 (4H), 3.62 (1H), 4.33 (3H), 7.04 (4H), 7.15 (1H), 7.40 (5H), 8.12 (1H), 8.47 (1H).
15		N-(2-methoxyethyl)-4-methyl-2-((1-[4-(4-methylphenoxy)benzoyl]piperidin-4-yl)methyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.20 (2H), 1.43 (2H), 2.26 (4H), 2.48 (3H), 2.76 (1H), 2.90 (1H), 3.24 (3H), 3.38 (4H), 3.62 (1H), 4.32 (3H), 6.92 (4H), 7.71 (3H), 7.34 (3H), 8.12 (1H), 8.47 (1H).
16		2-((1-[4-(4-tert-butylphenoxy)benzoyl]piperidin-4-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.21 (2H), 1.25 (9H), 1.44 (2H), 2.26 (1H), 2.48 (3H), 2.77 (1H), 2.291 (1H), 3.24 (3H), 3.39 (4H), 3.64 (1H), 4.32 (3H), 6.95 (4H), 7.14 (1H), 7.36 (5H), 8.12 (1H), 8.47 (1H).

Ex.	Structure	IUPAC name	Analysis
17		N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)-phenoxy]benzoyl}-piperidin-4-yl)-methyl]-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.22 (2H), 1.45 (2H), 2.27 (1H), 2.49 (3H), 2.77 (1H), 2.97 (1H), 3.24 (3H), 3.39 (4H), 3.61 (1H), 4.33 (3H), 7.15 (5H), 7.39 (3H), 7.73 (2H), 8.12 (1H), 8.48 (1H).
18		N-(2-methoxyethyl)-2-({1-[4-(4-methoxyphenoxy)benzoyl]-piperidin-4-yl}-methyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.19 (2H), 1.44 (2H), 2.25 (1H), 2.48 (3H), 2.86 (2H), 3.24 (3H), 3.40 (4H), 3.64 (1H), 3.72 (3H), 4.32 (3H), 6.88 (2H), 6.95 (2H), 7.02 (2H), 7.14 (1H), 7.31 (2H), 7.38 (1H), 8.12 (1H), 8.47 (1H).
19		N-(2-methoxyethyl)-4-methyl-2-[[1-(4-phenoxybenzoyl)-piperidin-4-yl]-methyl]-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.21 (2H), 1.44 (2H), 2.26 (1H), 2.49 (3H), 2.77 (1H), 2.92 (1H), 3.24 (3H), 3.38 (4H), 3.64 (1H), 4.32 (3H), 6.97 (2H), 7.04 (2H), 7.16 (2H), 7.26 (5H), 8.12 (1H), 8.47 (1H).
20		2-[[1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 0.65 (2H), 0.93 (2H), 1.19 (2H), 1.43 (2H), 1.90 (1H), 2.25 (1H), 2.48 (3H), 2.73 (1H), 2.92 (1H), 3.24 (3H), 3.39 (4H), 3.59 (1H), 4.32 (3H), 7.14 (5H), 7.38 (1H), 8.12 (1H), 8.47 (1H).

Ex.	Structure	IUPAC name	Analysis
21		2-([1-(4-methoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	$^1\text{H-NMR}$ (400 MHz, DMSO- d_6): δ [ppm]= 1.19 (2H), 1.44 (2H), 2.26 (1H), 2.49 (3H), 2.82 (2H), 3.24 (3H), 3.36 (2H), 3.42 (2H), 3.50 (1H), 3.74 (3H), 4.31 (3H), 6.93 (2H), 7.14 (1H), 7.29 (2H), 7.38 (1H), 8.12 (1H), 8.47 (1H).
22		2-([1-(4-fluorobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	$^1\text{H-NMR}$ (400 MHz, DMSO- d_6): δ [ppm]= 1.20 (2H), 1.37 (1H), 1.51 (1H), 2.27 (1H), 2.49 (3H), 2.72 (1H), 2.97 (1H), 3.24 (3H), 3.36 (2H), 3.42 (2H), 3.52 (1H), 4.31 (2H), 4.40 (1H), 7.14 (1H), 7.23 (2H), 7.39 (3H), 8.12 (1H), 8.47 (1H).
23		N-(2-methoxyethyl)-4-methyl-2-([1-(4-(trifluoromethyl)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide	$^1\text{H-NMR}$ (400 MHz, DMSO- d_6): δ [ppm]= 1.22 (2H), 1.34 (1H), 1.56 (1H), 2.28 (1H), 2.49 (3H), 2.75 (1H), 2.99 (1H), 3.24 (3H), 3.35 (2H), 3.42 (3H), 4.32 (2H), 4.44 (1H), 7.14 (1H), 7.38 (1H), 7.55 (2H), 7.77 (2H), 8.12 (1H), 8.47 (1H).
24		2-([1-(2-methoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	$^1\text{H-NMR}$ (400 MHz, DMSO- d_6): δ [ppm]= 1.11 (2H), 1.35 (1H), 1.52 (1H), 2.23 (1H), 2.49 (3H), 2.66 (1H), 2.87 (1H), 3.22 (1H), 3.24 (3H), 3.35 (2H), 3.42 (2H), 3.71 (3H), 4.29 (1H), 4.33 (1H), 4.45 (1H), 6.94 (1H), 7.07 (3H), 7.33 (1H), 7.39 (1H), 8.12 (1H), 8.48 (1H).

Ex.	Structure	IUPAC name	Analysis
25		N-(2-methoxyethyl)-4-methyl-2-((1-(4-((trifluoromethyl)sulphonyl)benzoyl)piperidin-4-yl)methyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.23 (2H), 1.36 (1H), 1.57 (1H), 2.29 (1H), 2.49 (3H), 2.77 (1H), 3.01 (1H), 3.24 (3H), 3.35 (3H), 3.42 (2H), 4.32 (2H), 4.44 (1H), 7.14 (1H), 7.38 (1H), 7.77 (2H), 8.15 (3H), 8.47 (1H).
26		N-(2-methoxyethyl)-4-methyl-2-((1-(3-(trifluoromethyl)benzoyl)piperidin-4-yl)methyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.24 (2H), 1.37 (1H), 1.55 (1H), 2.28 (1H), 2.49 (3H), 2.75 (1H), 3.01 (1H), 3.24 (3H), 3.36 (2H), 3.42 (3H), 4.32 (2H), 4.43 (1H), 7.14 (1H), 7.38 (1H), 7.65 (3H), 7.78 (1H), 8.12 (1H), 8.47 (1H).
27		N-(2-methoxyethyl)-4-methyl-2-((1-(3-methylbenzoyl)piperidin-4-yl)methyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.20 (2H), 1.36 (1H), 1.52 (1H), 2.29 (4H), 2.49 (3H), 2.69 (1H), 2.95 (1H), 3.24 (3H), 3.36 (2H), 3.42 (2H), 3.53 (1H), 4.31 (2H), 4.41 (1H), 7.12 (3H), 7.21 (1H), 7.27 (1H), 7.38 (1H), 8.12 (1H), 8.47 (1H).
28		2-((1-(3-chlorobenzoyl)piperidin-4-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.21 (2H), 1.36 (1H), 1.53 (1H), 2.27 (1H), 2.49 (3H), 2.72 (1H), 2.98 (1H), 3.24 (3H), 3.35 (2H), 3.42 (3H), 4.31 (2H), 4.40 (1H), 7.14 (1H), 7.28 (1H), 7.43 (4H), 8.12 (1H), 8.47 (1H).

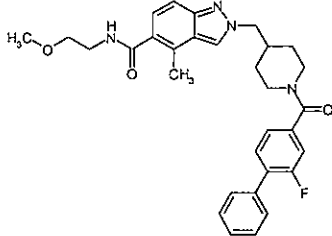
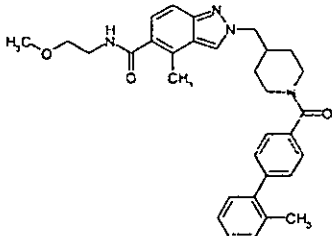
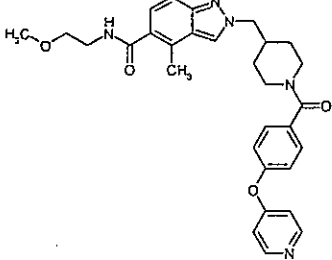
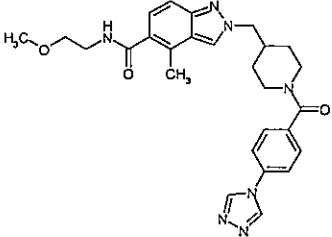
Ex.	Structure	IUPAC name	Analysis
29		N-(2-methoxyethyl)-4-methyl-2-((1-[(1-methyl-1H-indol-2-yl)carbonyl]piperidin-4-yl)methyl)-2H-indazole-5-carboxamide	$^1\text{H-NMR}$ (400 MHz, DMSO- d_6): δ [ppm]= 1.25 (2H), 1.50 (2H), 2.31 (1H), 2.49 (3H), 2.82 (1H), 3.05 (1H), 3.24 (3H), 3.36 (2H), 3.42 (2H), 3.69 (3H), 3.98 (1H), 4.34 (2H), 4.44 (1H), 6.57 (1H), 7.05 (1H), 7.15 (1H), 7.20 (1H), 7.39 (1H), 7.46 (1H), 7.56 (1H), 8.12 (1H), 8.49 (1H).
30		2-((1-[4-(4-carbamoylphenoxy)benzoyl]piperidin-4-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	$^1\text{H-NMR}$ (400 MHz, DMSO- d_6): δ [ppm]= 1.22 (2H), 1.49 (2H), 2.31 (1H), 2.52 (3H), 2.82 (1H), 2.98 (1H), 3.27 (3H), 3.39 (2H), 3.45 (2H), 3.65 (1H), 4.35 (2H), 4.43 (1H), 7.08 (4H), 7.17 (1H), 7.29 (1H), 7.41 (3H), 7.91 (3H), 8.12 (1H), 8.50 (1H).
31		2-((1-[4-(cyclopentyloxy)benzoyl]piperidin-4-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	$^1\text{H-NMR}$ (400 MHz, DMSO- d_6): δ [ppm]= 1.23 (2H), 1.47 (2H), 1.57 (2H), 1.68 (4H), 1.90 (2H), 2.28 (1H), 2.52 (3H), 2.86 (2H), 3.27 (3H), 3.39 (2H), 3.45 (2H), 3.85 (2H), 4.34 (2H), 4.83 (1H), 6.91 (2H), 7.17 (1H), 7.28 (2H), 7.41 (1H), 8.12 (1H), 8.49 (1H).
32		2-((1-[4-(difluoromethyl)benzoyl]piperidin-4-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	$^1\text{H-NMR}$ (400 MHz, DMSO- d_6): δ [ppm]= 1.24 (2H), 1.39 (1H), 1.57 (1H), 2.30 (1H), 2.52 (3H), 2.76 (1H), 3.01 (1H), 3.27 (3H), 3.39 (2H), 3.45 (3H), 4.35 (2H), 4.45 (1H), 7.18 (2H), 7.41 (1H), 7.49 (2H), 7.62 (2H), 8.12 (1H),

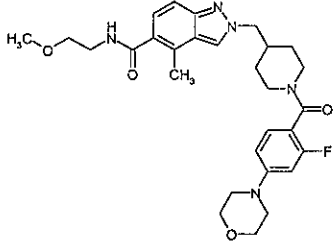
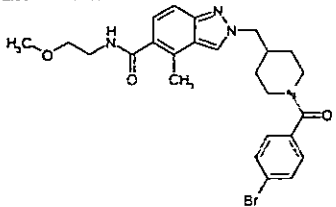
Ex.	Structure	IUPAC name	Analysis
			8.49 (1H).
33		2-([1-(4-cyano-benzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.24 (2H), 1.39 (1H), 1.58 (1H), 2.30 (1H), 2.52 (3H), 2.77 (1H), 3.01 (1H), 3.27 (3H), 3.39 (3H), 3.45 (2H), 4.34 (2H), 4.44 (1H), 7.17 (1H), 7.41 (1H), 7.54 (2H), 7.90 (2H), 8.12 (1H), 8.49 (1H).
34		2-([1-[4-(1H-imidazol-1-yl)-benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.24 (2H), 1.39 (1H), 1.56 (1H), 2.29 (1H), 2.49 (3H), 2.76 (1H), 3.04 (1H), 3.24 (3H), 3.37 (3H), 3.42 (3H), 4.33 (2H), 4.44 (1H), 7.15 (1H), 7.39 (1H), 7.61 (2H), 7.84 (3H), 8.11 (1H), 8.26 (1H), 8.48 (1H).
35		N-(2-methoxyethyl)-4-methyl-2-([1-[4-(oxazol-2-yl)-benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.24 (2H), 1.37 (1H), 1.54 (1H), 2.27 (1H), 2.49 (3H), 2.75 (1H), 3.00 (1H), 3.24 (3H), 3.36 (2H), 3.42 (2H), 3.53 (1H), 4.32 (2H), 4.43 (1H), 7.14 (1H), 7.38 (2H), 7.48 (2H), 8.00 (2H), 8.12 (1H), 8.23 (1H), 8.48 (1H).
36		N-(2-methoxyethyl)-4-methyl-2-([1-[4-(oxazol-5-yl)-benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.22 (2H), 1.38 (1H), 1.53 (1H), 2.28 (1H), 2.49 (3H), 2.74 (1H), 2.98 (1H), 3.24 (3H), 3.36 (2H), 3.41 (2H), 3.55 (1H), 4.32 (2H), 4.41 (1H), 7.15 (1H), 7.42 (3H), 7.74 (3H), 8.12 (1H),

Ex.	Structure	IUPAC name	Analysis
		carboxamide	8.47 (2H).
37		N-(2-methoxyethyl)-4-methyl-2-({1-[4-(isoxazol-5-yl)-benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.22 (2H), 1.36 (1H), 1.55 (1H), 2.28 (1H), 2.49 (3H), 2.74 (1H), 2.99 (1H), 3.24 (3H), 3.39 (5H), 4.31 (2H), 4.43 (1H), 4.74 (2H), 7.14 (1H), 7.38 (1H), 7.50 (2H), 7.95 (2H), 8.12 (1H), 8.47 (1H).
38		N-(2-methoxyethyl)-4-methyl-2-({1-[4-(1H-pyrazol-1-yl)-benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.22 (2H), 1.46 (2H), 2.28 (1H), 2.49 (3H), 2.76 (1H), 2.99 (1H), 3.24 (3H), 3.39 (4H), 3.61 (1H), 4.32 (2H), 4.41 (1H), 6.54 (1H), 7.15 (1H), 7.39 (1H), 7.46 (2H), 7.74 (1H), 7.87 (2H), 8.12 (1H), 8.50 (2H).
39		N-(2-methoxyethyl)-4-methyl-2-({1-[4-(1H-1,2,4-triazol-1-yl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.22 (2H), 1.39 (1H), 1.53 (1H), 2.27 (1H), 2.49 (3H), 2.76 (1H), 3.00 (1H), 3.24 (3H), 3.39 (4H), 3.54 (1H), 4.32 (2H), 4.43 (1H), 7.15 (1H), 7.39 (1H), 7.53 (2H), 7.90 (2H), 8.12 (1H), 8.24 (1H), 8.48 (1H), 9.32 (1H).
40		2-({1-[4-(difluoromethoxy)-2-fluorobenzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.16 (2H), 1.38 (1H), 1.55 (1H), 2.26 (1H), 2.48 (3H), 2.73 (1H), 2.98 (1H), 3.24 (3H), 3.38 (5H), 4.31 (2H), 4.44 (1H), 7.12 (3H), 7.31 (1H), 7.40 (2H),

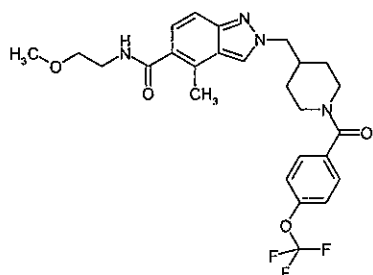
Ex.	Structure	IUPAC name	Analysis
		5-carboxamide	8.12 (1H), 8.47 (1H).
41		2-({1-[2-fluoro-4-methyl-2H-indazole-5-carboxamide]piperidin-4-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.14 (2H), 1.43 (2H), 1.90 (4H), 2.23 (1H), 2.48 (3H), 2.68 (1H), 2.94 (1H), 3.19 (4H), 3.24 (3H), 3.36 (2H), 3.42 (3H), 4.31 (2H), 4.39 (1H), 6.29 (2H), 7.10 (2H), 7.38 (1H), 8.12 (1H), 8.47 (1H).
42		2-({1-[(3,4'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.19 (2H), 1.39 (1H), 1.57 (1H), 2.28 (1H), 2.49 (3H), 2.76 (1H), 3.01 (1H), 3.24 (3H), 3.38 (5H), 4.32 (2H), 4.47 (1H), 7.15 (1H), 7.28 (2H), 7.40 (2H), 7.56 (2H), 7.76 (2H), 8.12 (1H), 8.48 (1H).
43		2-({1-[(3-fluoro-4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.19 (2H), 1.39 (1H), 1.57 (1H), 2.28 (1H), 2.49 (3H), 2.76 (1H), 3.01 (1H), 3.24 (3H), 3.38 (5H), 4.32 (2H), 4.47 (1H), 7.15 (1H), 7.28 (2H), 7.40 (2H), 7.56 (2H), 7.76 (2H), 8.12 (1H), 8.48 (1H).
44		2-({1-[(3-fluoro-4'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.19 (2H), 1.39 (1H), 1.57 (1H), 2.31 (4H), 2.49 (3H), 2.75 (1H), 3.01 (1H), 3.24 (3H), 3.36 (2H), 3.42 (3H), 4.32 (2H), 4.48 (1H), 7.14 (1H), 7.26 (2H), 7.39 (2H), 7.56 (4H), 8.12 (1H),

Ex.	Structure	IUPAC name	Analysis
		5-carboxamide	8.48 (1H).
45		2-[(1-{[3-fluoro-3'-(trifluoromethyl)-biphenyl-4-yl]-carbonyl}-piperidin-4-yl)-methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.20 (2H), 1.40 (1H), 1.57 (1H), 2.29 (1H), 2.49 (3H), 2.76 (1H), 3.03 (1H), 3.24 (3H), 3.36 (2H), 3.41 (3H), 4.33 (2H), 4.48 (1H), 7.15 (1H), 7.39 (1H), 7.45 (1H), 7.69 (4H), 8.03 (2H), 8.12 (1H), 8.49 (1H).
46		2-[(1-{[3-fluoro-2'-(trifluoromethoxy)-biphenyl-4-yl]-carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.20 (2H), 1.42 (1H), 1.57 (1H), 2.29 (1H), 2.49 (3H), 2.76 (1H), 3.04 (1H), 3.24 (3H), 3.36 (5H), 4.32 (2H), 4.48 (1H), 7.15 (1H), 7.44 (8H), 8.12 (1H), 8.49 (1H).
47		2-[(1-{[2'-fluoro-biphenyl-4-yl]-carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.23 (2H), 1.41 (1H), 1.53 (1H), 2.29 (1H), 2.49 (3H), 2.74 (1H), 3.01 (1H), 3.24 (3H), 3.36 (4H), 3.60 (1H), 4.33 (2H), 4.44 (1H), 7.15 (1H), 7.29 (2H), 7.40 (4H), 7.55 (3H), 8.12 (1H), 8.48 (1H).
48		2-[(1-{[2',4'-difluoro-biphenyl-4-yl]-carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.23 (2H), 1.39 (1H), 1.53 (1H), 2.28 (1H), 2.49 (3H), 2.75 (1H), 3.01 (1H), 3.24 (3H), 3.39 (4H), 3.60 (1H), 4.33 (2H), 4.43 (1H), 7.16 (2H), 7.40 (4H),

Ex.	Structure	IUPAC name	Analysis
		5-carboxamide	7.56 (3H), 8.12 (1H), 8.48 (1H).
49		2-({1-[(2-fluoro-biphenyl-4-yl)-carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.24 (2H), 1.39 (1H), 1.56 (1H), 2.29 (1H), 2.49 (3H), 2.74 (1H), 3.02 (1H), 3.25 (3H), 3.37 (2H), 3.42 (2H), 3.59 (1H), 4.33 (2H), 4.43 (1H), 7.15 (1H), 7.28 (2H), 7.40 (2H), 7.46 (2H), 7.54 (3H), 8.12 (1H), 8.48 (1H).
50		N-(2-methoxyethyl)-4-methyl-2-({1-[(2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.24 (2H), 1.42 (1H), 1.55 (1H), 2.20 (3H), 2.29 (1H), 2.49 (3H), 2.75 (1H), 3.02 (1H), 3.25 (3H), 3.37 (2H), 3.42 (2H), 3.64 (1H), 4.33 (2H), 4.42 (1H), 7.21 (5H), 7.38 (5H), 8.12 (1H), 8.49 (1H).
51		N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyridyloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.24 (2H), 1.42 (1H), 1.55 (1H), 2.29 (1H), 2.49 (3H), 2.76 (1H), 3.02 (1H), 3.25 (3H), 3.37 (2H), 3.43 (2H), 3.60 (1H), 4.33 (2H), 4.44 (1H), 7.15 (1H), 7.37 (3H), 7.48 (2H), 7.53 (2H), 8.12 (1H), 8.48 (1H), 8.75 (2H).
52		N-(2-methoxyethyl)-4-methyl-2-({1-[4-(4H-1,2,4-triazol-4-yl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.23 (2H), 1.38 (1H), 1.54 (1H), 2.28 (1H), 2.49 (3H), 2.74 (1H), 3.02 (1H), 3.24 (3H), 3.36 (2H), 3.42 (2H), 3.53 (1H), 4.32 (2H), 4.42 (1H), 7.14 (1H), 7.38 (1H), 7.55 (2H), 7.75 (2H),

Ex.	Structure	IUPAC name	Analysis
		carboxamide	8.14 (1H), 8.48 (1H), 9.15 (2H).
53		2-({1-[2-fluoro-4-(morpholin-4-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (600 MHz, DMSO-d ₆): δ [ppm]= 1.01 (1H), 1.21 (1H), 1.36 (1H), 1.48 (1H), 1.57 (1H), 2.08 (1H), 2.53 (3H), 2.96 (7H), 3.29 (3H), 3.41 (2H), 3.47 (2H), 3.67 (4H), 4.34 (2H), 4.50 (1H), 6.86 (2H), 7.14 (2H), 7.41 (1H), 8.11 (1H), 8.47 (1H), 8.56 (1H).
54		2-({1-[4-bromo-benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]=1.22 (2H), 1.36 (1H), 1.52 (1H), 2.26 (1H), 2.49 (3H), 2.72 (1H), 2.97 (1H), 3.24 (3H), 3.36 (2H), 3.42 (2H), 3.49 (1H), 4.31 (2H), 4.40 (1H), 7.14 (1H), 7.29 (2H), 7.38 (1H), 7.60 (2H), 8.12 (1H), 8.47 (1H).

Example 55: N-(2-methoxyethyl)-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

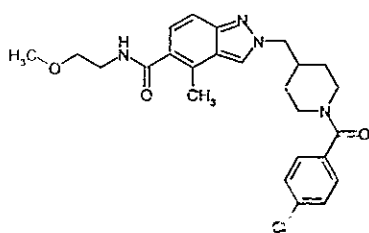


- 5 150 mg of the amine prepared in Example 1a were first dissolved in 1.5 ml pyridine. 101 mg of 4-(trifluoromethoxy)benzoyl chloride were then added and the mixture stirred for 30 min at RT. The reaction mixture was then treated with some toluene and concentrated. The residue was taken up in ethyl acetate and washed twice with water, twice with saturated sodium hydrogen carbonate solution (pH 9) and once with saturated sodium chloride solution. The organic phase

was dried over sodium sulphate, filtered and concentrated. The residue was purified chromatographically on the Biotage SP4. Gradient: ethyl acetate/methanol 0-10%. Yield: 118.2 mg of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.30 (2H), 1.30 - 1.45 (1H), 1.45 - 1.62 (1H),
 5 2.19 - 2.35 (1H), 2.49 (3H), 2.64 - 2.86 (1H), 2.86 - 3.07 (1H), 3.24 (3H), 3.31 - 3.56 (5H), 4.31
 (2H), 4.36 - 4.49 (1H), 7.14 (1H), 7.35 - 7.42 (3H), 7.43 - 7.50 (2H), 8.11 (1H), 8.47 (1H).

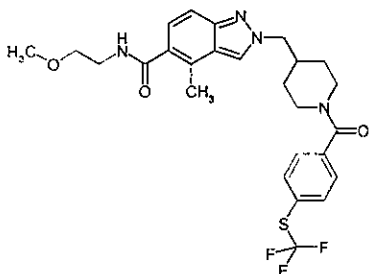
Example 56: 2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



10 Analogously to Example 55, 183 mg of the title compound was obtained from 300 mg of the amine prepared in Example 1a and 157.4 mg of 4-chlorobenzoyl chloride.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.10 - 1.29 (2H), 1.29 - 1.43 (1H), 1.44 - 1.61 (1H),
 2.20 - 2.34 (1H), 2.49 (3H), 2.66 - 2.83 (1H), 2.84 - 3.06 (1H), 3.25 (3H), 3.32 - 3.39 (2H), 3.40
 - 3.45 (2H), 3.45 - 3.55 (1H), 4.26 - 4.35 (2H), 4.35 - 4.46 (1H), 7.14 (1H), 7.32 - 7.41 (3H),
 15 7.47 (2H), 8.09 - 8.15 (1H), 8.47 (1H).

Example 57: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[(trifluoromethyl)sulphanyl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide

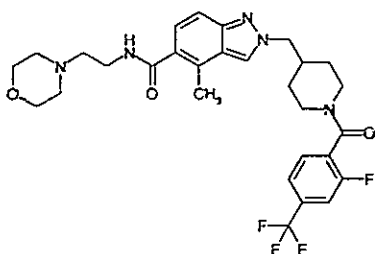


60 mg of the amine prepared in Example 1a were first dissolved in 2 ml DCM at 0°C. 0.034 ml
 20 of triethylamine and 43.2 mg of 4-[(trifluoromethyl)sulphanyl]benzoyl chloride were then added
 and the mixture stirred for 2 hrs at 0°C. The reaction mixture was then diluted with DCM and
 washed with 1-molar hydrochloric acid, saturated sodium hydrogen carbonate solution (pH 9)
 and saturated sodium chloride solution. The organic phase was dried over sodium sulphate,

filtered and concentrated. The residue was purified chromatographically on the Biotage SP4. Gradient: DCM/Methanol 0-10%. The product fraction was taken up in ethyl acetate and extracted twice with saturated sodium hydrogen carbonate solution. The organic phase was concentrated and the residue again purified by HPLC. Yield: 26.5 mg of the title compound.

- 5 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.14 - 1.30 (2H), 1.31 - 1.42 (1H), 1.49 - 1.61 (1H), 2.21 - 2.32 (1H), 2.49 (3H), 2.67 - 2.80 (1H), 2.93 - 3.05 (1H), 3.25 (3H), 3.36 (2H), 3.42 (3H), 4.27 - 4.36 (2H), 4.38 - 4.48 (1H), 7.14 (1H), 7.38 (1H), 7.48 (2H), 7.75 (2H), 8.09 - 8.14 (1H), 8.47 (1H).

- 10 **Example 58:** 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide

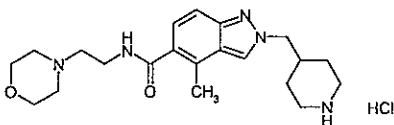


Analogously to Example 1, 34.6 mg of the title compound was obtained from 75 mg of the amine prepared in Example 58a and 37.0 mg of 2-fluoro-4-(trifluoromethyl)benzoic acid.

- 15 ¹H-NMR (400 MHz, methanol-d₄): δ [ppm]= 1.24 - 1.43 (2H), 1.50 - 1.59 (1H), 1.65 - 1.73 (1H), 2.33 - 2.45 (1H), 2.65 (3H), 2.80 - 2.91 (1H), 3.05 - 3.18 (1H), 3.19 - 3.26 (2H), 3.39 - 3.44 (2H), 3.45 - 3.52 (1H), 3.63 - 3.70 (2H), 3.71 - 3.82 (4H), 4.06 - 4.14 (2H), 4.39 (2H), 4.62 - 4.70 (1H), 7.39 (1H), 7.47 (1H), 7.54 - 7.62 (3H), 8.45 (1H).

The starting material was prepared as follows:

- 20 **Example 58a:** 4-methyl-N-(2-morpholinoethyl)-2-(4-piperidylmethyl)-2H-indazol-5-carboxamide hydrochloride

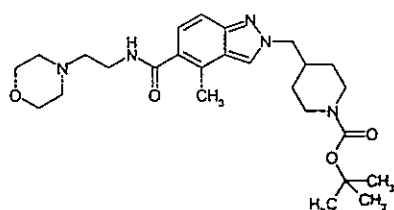


To 624.4 mg of 58b were added 2.9 ml of 4M hydrochloric acid in dioxan and 0.5 ml dioxan. An oily mass was formed, which dissolved on vigorous stirring and gentle warming. The mixture

was stirred for 1 hr at ca. 30°C. The reaction mixture was concentrated. Yield: 980 mg of the title compound, which was further reacted without further purification.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.33 - 1.51 (2H), 1.58 - 1.64 (1H), 2.17 - 2.31 (1H), 2.55 (3H), 2.70 - 2.86 (2H), 3.02 - 3.31 (7H), 3.43 - 3.51 (2H), 3.61 - 3.71 (2H), 3.72 - 3.88 (3H), 3.89 - 3.99 (2H), 4.30 - 4.36 (2H), 7.29 (1H), 7.40 (1H), 8.46 - 8.52 (1H), 8.54 - 8.58 (1H), 8.69 - 8.85 (1H), 8.96 - 9.10 (1H).

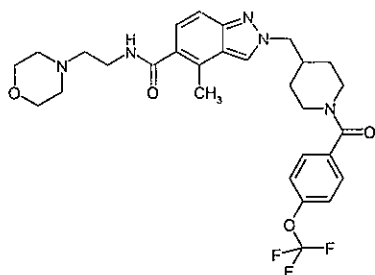
Example 58b: Tert-butyl 4-({4-methyl-5-[N-(2-morpholinoethyl)carbamoyl]-1H-indazol-2-yl}-methyl)piperidin-1-carboxylate



896 mg of 1c, 857 mg of 2-morpholinoethylamine. 579.3 mg of molybdenum hexacarbonyl, 63.6 mg of tri-tert-butylphosphine tetrafluoroborate and 205.8 mg of trans-bis(acetato)-bis[o-(di-o-tolylphosphino)benzyl]dipalladium(II) were placed in a microwave tube and suspended in 18 ml THF. 1.0 ml of DBU was then added and the mixture was stirred for 20 mins at 125°C and 200 watts in the microwave. The reaction mixture was diluted with some ethyl acetate and firstly filtered through Celite. The filtrate was diluted with ethyl acetate and the organic phase washed twice with water and once with saturated sodium chloride solution. It was then dried over sodium sulphate, filtered and concentrated. The residue was chromatographed on the Biotage SP4. Gradient: ethyl acetate/methanol 0-10%. Yield: 624.4 mg of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.98 - 1.13 (2H), 1.34 (11H), 2.05 - 2.27 (2H), 2.33 - 2.41 (5H), 2.51 (3H), 2.56 - 2.73 (2H), 3.25 - 3.37 (2H), 3.51 - 3.57 (4H), 3.81 - 3.93 (2H), 4.28 (2H), 7.14 (1H), 7.36 - 7.41 (1H), 7.95 - 8.02 (1H), 8.46 (1H).

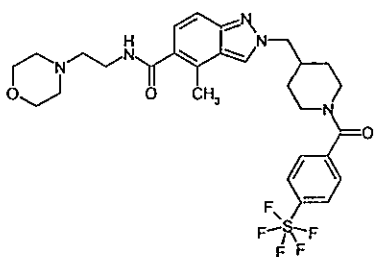
Example 59: 4-methyl-N-(2-morpholinoethyl)-2-({1-[4-(trifluoromethoxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 10.1 mg of the title compound was obtained from 75 mg of the amine prepared in Example 58a and 43.9 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (400 MHz, methanol-d₄): δ [ppm]= 1.24 - 1.44 (2H), 1.44 - 1.58 (1H), 1.59 - 1.74 (1H), 2.32 - 2.44 (1H), 2.51 - 2.58 (4H), 2.59 - 2.65 (5H), 2.77 - 2.90 (1H), 3.02 - 3.17 (1H), 3.53 (2H), 3.62 - 3.72 (5H), 4.35 - 4.41 (2H), 4.55 - 4.67 (1H), 7.29 (1H), 7.34 (2H), 7.43 (1H), 7.49 (2H), 8.39 (1H).

Example 60: 4-methyl-N-(2-morpholinoethyl)-2-({1-[4-(pentafluoro-λ⁶-sulphanyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



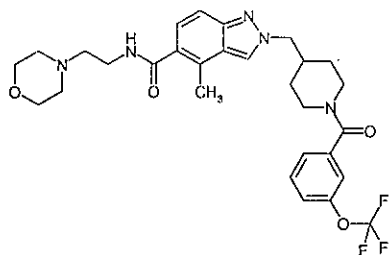
10

Analogously to Example 1, 14.0 mg of the title compound was obtained from 75 mg of the amine prepared in Example 58a and 44.1 mg of 4-(pentafluoro-λ⁶-sulphanyl)benzoic acid.

¹H-NMR (400 MHz, chloroform-d): δ [ppm]= 1.16 - 1.32 (1H), 1.33 - 1.48 (1H), 1.50 - 1.63 (1H), 1.71 - 1.83 (1H), 1.84 - 2.08 (2H), 2.44 (1H), 2.55 (4H), 2.60 - 2.71 (5H), 2.72 - 2.85 (1H), 2.96 - 3.10 (1H), 3.60 (2H), 3.67 - 3.78 (4H), 4.33 (2H), 4.71 - 4.81 (1H), 6.45 (1H), 7.34 (1H), 7.47 (2H), 7.54 (1H), 7.79 (2H), 7.96 (1H).

15

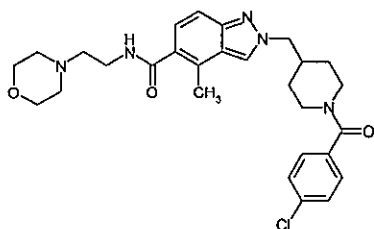
Example 61: 4-methyl-N-(2-morpholinoethyl)-2-({1-[3-(trifluoromethoxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 18.1 mg of the title compound was obtained from 75 mg of the amine prepared in Example 58a and 43.9 mg of 3-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (400 MHz, chloroform-d): δ [ppm]= 1.18 - 1.46 (2H), 1.49 - 1.63 (1H), 1.66 - 1.82 (1H), 2.37 - 2.49 (1H), 2.62 (3H), 2.67 (3H), 2.70 - 2.78 (4H), 2.79 - 2.86 (2H), 2.93 - 3.09 (1H), 3.66 - 3.74 (2H), 3.83 (4H), 4.32 (2H), 4.66 - 4.82 (1H), 6.83 - 6.96 (1H), 7.23 - 7.29 (2H), 7.29 - 7.34 (1H), 7.39 (1H), 7.43 (1H), 7.54 (1H), 7.96 (1H).

Example 62: 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-4-methyl-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide



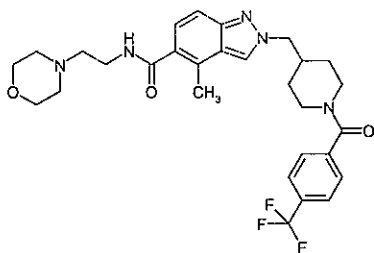
10

Analogously to Example 55, 14.9 mg of the title compound was obtained from 75 mg of the amine prepared in Example 58a and 34.2 mg of 4-chlorobenzoyl chloride.

¹H-NMR (400 MHz, methanol-d₄): δ [ppm]= 1.24 - 1.42 (2H), 1.46 - 1.58 (1H), 1.58 - 1.71 (1H), 2.31 - 2.43 (1H), 2.63 (5H), 2.65 (3H), 2.73 - 2.91 (1H), 3.00 - 3.15 (1H), 3.25 - 3.33 (2H), 3.63 - 3.77 (3H), 3.79 - 4.03 (3H), 4.38 (2H), 4.54 - 4.67 (1H), 7.38 (3H), 7.42 - 7.49 (3H), 8.44 (1H).

15

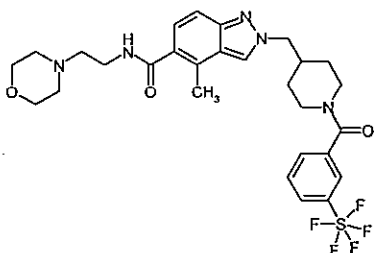
Example 63: 4-methyl-N-(2-morpholinoethyl)-2-([1-(4-(trifluoromethyl)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 31.7 mg of the title compound was obtained from 75 mg of the amine prepared in Example 58a and 33.8 mg of 4-(trifluoromethyl)benzoic acid.

¹H-NMR (400 MHz, methanol-d₄): δ [ppm]= 1.24 - 1.43 (2H), 1.45 - 1.57 (1H), 1.63 - 1.73 (1H), 2.33 - 2.45 (1H), 2.66 (3H), 2.79 - 2.90 (1H), 3.04 - 3.17 (1H), 3.18 - 3.27 (2H), 3.39 - 3.45 (2H), 3.56 - 3.70 (3H), 3.76 (4H), 4.06 - 4.14 (2H), 4.36 - 4.42 (2H), 4.59 - 4.68 (1H), 7.39 (1H), 7.47 (1H), 7.57 (2H), 7.73 (2H), 8.45 (1H).

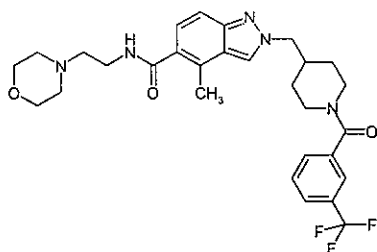
Example 64: 4-methyl-N-(2-morpholinoethyl)-2-({1-[4-(pentafluoro-λ⁶-sulphanyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 24.7 mg of the title compound was obtained from 75 mg of the amine prepared in Example 58a and 44.1 mg of 3-(pentafluoro-λ⁶-sulphanyl)benzoic acid.

¹H-NMR (400 MHz, methanol-d₄): δ [ppm]= 1.27 - 1.47 (2H), 1.48 - 1.60 (1H), 1.60 - 1.74 (1H), 2.31 - 2.46 (1H), 2.66 (3H), 2.78 - 2.93 (1H), 3.04 - 3.18 (1H), 3.18 - 3.26 (2H), 3.39 - 3.43 (2H), 3.57 - 3.70 (3H), 3.72 - 3.82 (4H), 4.06 - 4.14 (2H), 4.39 (2H), 4.55 - 4.67 (1H), 7.38 (1H), 7.47 (1H), 7.60 - 7.67 (2H), 7.82 - 7.86 (1H), 7.89 - 7.94 (1H), 8.45 (1H).

Example 65: 4-methyl-N-(2-morpholinoethyl)-2-({1-[3-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



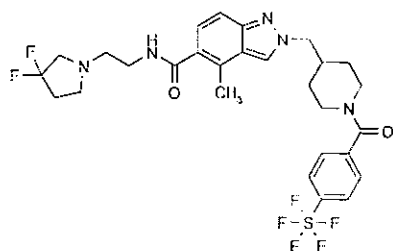
Analogously to Example 1, 14.0 mg of the title compound was obtained from 75 mg of the amine prepared in Example 58a and 33.8 mg of 3-(trifluoromethyl)benzoic acid.

¹H-NMR (400 MHz, methanol-d₄): δ [ppm]= 1.28 - 1.45 (2H), 1.47 - 1.59 (1H), 1.59 - 1.73 (1H), 2.32 - 2.45 (1H), 2.66 (3H), 2.79 - 2.93 (1H), 3.05 - 3.18 (1H), 3.19 - 3.27 (2H), 3.39 - 3.45 (2H), 3.58 - 3.70 (3H), 3.72 - 3.82 (4H), 4.06 - 4.14 (2H), 4.39 (2H), 4.57 - 4.68 (1H), 7.39 (1H), 7.47 (1H), 7.64 (2H), 7.67 - 7.70 (1H), 7.73 - 7.78 (1H), 8.45 (1H).

The following compounds were prepared analogously:

Ex.	Structure	IUPAC name	Analysis
66		2-({1-[(4'-fluorobiphenyl-4-yl)-carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, methanol-d ₄): δ [ppm]= 1.35 (2H), 1.53 (1H), 1.66 (1H), 2.40 (1H), 2.66 (3H), 2.85 (1H), 3.11 (1H), 3.24 (2H), 3.42 (2H), 3.65 (2H), 3.76 (5H), 4.08 (2H), 4.40 (2H), 4.63 (1H), 7.17 (2H), 7.38 (1H), 7.47 (3H), 7.65 (4H), 8.45 (1H).
67		2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, methanol-d ₄): δ [ppm]= 1.33 (2H), 1.68 (2H), 2.38 (1H), 2.66 (3H), 2.84 (1H), 3.11 (1H), 3.25 (2H), 3.42 (2H), 3.65 (2H), 3.76 (5H), 4.11 (2H), 4.38 (2H), 4.59 (1H), 6.98 (2H), 7.04 (2H), 7.11 (2H), 7.38 (3H), 7.47 (1H), 8.44 (1H).

Example 68: N-[2-(3,3-difluoropyrrolidin-1-yl)ethyl]-4-methyl-2-({1-[4-(pentafluoro- λ^6 -sulphanyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

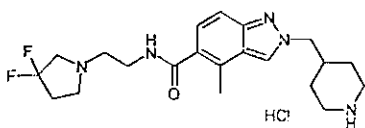


Analogously to Example 57, 61.7 mg of the title compound was obtained from 266.5 mg of the
5 amine prepared in Example 68a and 150.3 mg of 4-(pentafluoro- λ^6 -sulphanyl)benzoyl chloride.

$^1\text{H-NMR}$ (300 MHz, chloroform- d): δ [ppm]= 1.13 - 1.46 (2H), 1.48 - 1.65 (1H), 1.66 - 1.87 (1H), 2.26 - 2.50 (3H), 2.66 (3H), 2.71 - 3.23 (8H), 3.57 - 3.72 (3H), 4.25 - 4.42 (2H), 4.67 - 4.84 (1H), 6.40 - 6.61 (1H), 7.38 (1H), 7.47 (2H), 7.53 (1H), 7.79 (2H), 7.96 (1H).

The starting material was prepared as follows:

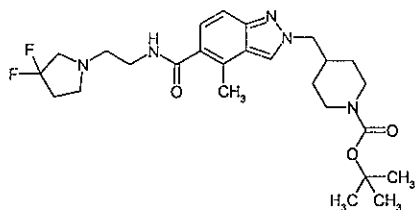
10 **Example 68a:** N-[2-(3,3-difluoropyrrolidin-1-yl)ethyl]-4-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide hydrochloride



To 416.6 mg of 68b were added 1.85 ml of 4M hydrochloric acid in dioxan and 0.5 ml dioxan.
An oily mass was formed, which dissolved on vigorous stirring and gentle warming. The
15 mixture was stirred for 1 hr at ca. 30°C. The reaction mixture was concentrated. Yield: 439.2 mg of the title compound, which was further reacted without further purification.

$^1\text{H-NMR}$ (400 MHz, DMSO- d_6): δ [ppm]= 1.36 - 1.48 (2H), 1.54 - 1.63 (2H), 2.18 - 2.28 (1H), 2.40 - 2.45 (2H), 2.55 (3H), 2.72 - 2.83 (2H), 3.14 - 3.22 (2H), 3.38 - 3.44 (2H), 3.56 - 3.63 (2H), 3.65 - 3.95 (2H), 4.11 - 4.21 (2H), 7.32 (1H), 7.40 (1H), 8.43 - 8.48 (1H), 8.57 (1H), 8.68 -
20 8.82 (1H), 8.96 - 9.07 (1H).

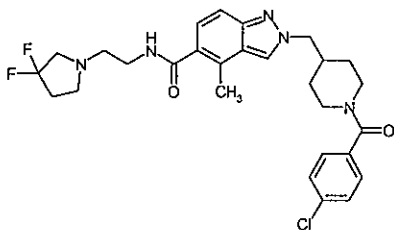
Example 68b: Tert-butyl 4-[(5-{N-[2-(3,3-difluoropyrrolidin-1-yl)ethyl]carbamoyl}-4-methyl-2H-indazol-2-yl)methyl]-piperidin-1-carboxylate



220 mg of 1c, 242.7 mg of 2-(3,3-difluoro-pyrrolidin-1-yl)ethylamine, 142.2 mg of molybdenum hexacarbonyl, 15.6 mg of tri-tert-butylphosphine tetrafluoroborate and 50.5 mg of trans-bis(acetato)-bis-[o-(di-o-tolylphosphino)benzyl]dipalladium(II) were placed in a microwave tube
 5 and suspended in 4.4 ml THF. 0.24 ml of DBU were then added and the mixture stirred for 20 mins at 125°C and 200 watts in the microwave. The reaction mixture was diluted with some ethyl acetate and firstly filtered through Celite. The filtrate was diluted with ethyl acetate and the organic phase washed twice with water and once with saturated sodium chloride solution. It was dried over sodium sulphate, filtered and concentrated. The residue (209.3 mg) was chromatographed on the Biotage SP4. Gradient: DCM/methanol 0-10%. Yield: 53.4 mg of the title compound.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.00 - 1.12 (2H), 1.32 - 1.42 (2H), 1.34 (9H), 2.07 - 2.25 (3H), 2.50 (3H), 2.57 (2H), 2.60 - 2.68 (2H), 2.72 (2H), 2.91 (2H), 3.30 - 3.35 (2H), 3.83 - 3.92 (2H), 4.28 (2H), 7.15 (1H), 7.39 (1H), 8.04 - 8.09 (1H), 8.47 (1H).

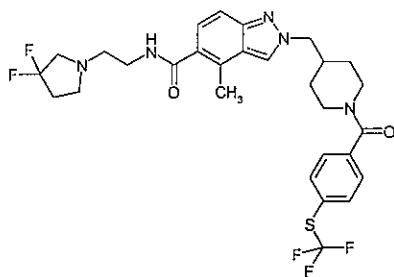
15 **Example 69:** 2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-N-[2-(3,3-difluoropyrrolidin-1-yl)ethyl]-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 57, 79.9 mg of the title compound was obtained from 125.0 mg of the amine prepared in Example 68a and 54.5 mg of 4-chlorobenzoyl chloride.

20 ¹H-NMR (300 MHz, chloroform-d): δ [ppm]= 1.13 - 1.35 (2H), 1.46 - 1.84 (2H), 2.27 - 2.50 (3H), 2.66 (3H), 2.80 - 3.26 (8H), 3.59 - 3.83 (3H), 4.25 - 4.39 (2H), 4.62 - 4.85 (1H), 6.41 - 6.69 (1H), 7.30 - 7.40 (6H), 7.55 (1H), 7.96 (1H).

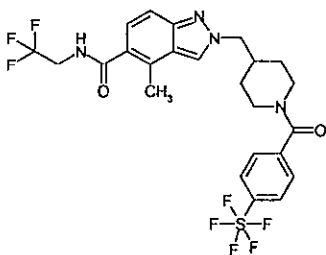
Example 70: N-[2-(3,3-difluoropyrrolidin-1-yl)ethyl]-4-methyl-2-[(1-{4-[(trifluoromethyl)sulphonyl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 55, 15.2 mg of the title compound was obtained from 23.4 mg of the amine prepared in Example 68a and 14.0 mg of 4-[(trifluoromethyl)sulphonyl]benzoyl chloride.

¹H-NMR (400 MHz, chloroform-d): δ [ppm]= 1.23 – 1.26 (1H), 1.33 – 1.46 (1H), 1.48 – 1.61 (1H), 1.77 (1H), 2.20 – 2.35 (2H), 2.36 – 2.49 (1H), 2.64 (3H), 2.73 – 2.85 (5H), 2.90 – 3.10 (3H), 3.56 (2H), 3.63 – 3.76 (1H), 4.26 – 4.39 (2H), 4.70 – 4.82 (1H), 6.28 (1H), 7.33 (1H), 7.42 (2H), 7.53 (1H), 7.68 (2H), 7.96 (1H).

Example 71: 4-methyl-2-({1-[4-(pentafluoro- λ^6 -sulphonyl)benzoyl]piperidin-4-yl} methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



10

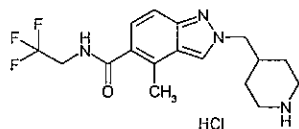
Analogously to Example 1, 124.8 mg of the title compound was obtained from 146.9 mg of the amine prepared in Example 71a and 93.3 mg of 4-(pentafluoro- λ^6 -sulphonyl)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.12 – 1.31 (2H), 1.31 – 1.45 (1H), 1.47 – 1.63 (1H), 2.20 – 2.36 (1H), 2.50 (3H), 2.67 – 2.83 (1H), 2.91 – 3.07 (1H), 3.36 – 3.47 (1H), 3.96 – 4.10 (2H), 4.29 – 4.37 (2H), 4.37 – 4.48 (1H), 7.17 (1H), 7.43 (1H), 7.56 (2H), 7.94 (2H), 8.50 – 8.54 (1H), 8.75 – 8.83 (1H).

15

The starting material was prepared as follows:

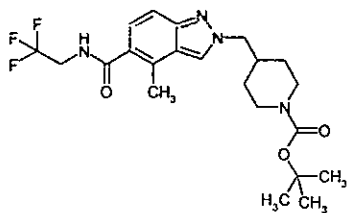
Example 71a: 4-methyl-2-(4-piperidylmethyl)-N-(2,2,2-trifluoroethyl)-2H-indazol-5-carboxamide hydrochloride



To 425.2 mg of 71b were added 2.35 ml of 4M hydrochloric acid in dioxan and 0.5 ml dioxan. An oily mass was formed, which dissolved on vigorous stirring and gentle warming. The mixture was stirred for 1 hr at ca. 30°C. The reaction mixture was concentrated. Yield: 440.6 mg of the title compound, which was further reacted without further purification.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.35 - 1.49 (2H), 1.53 - 1.62 (2H), 2.20 - 2.32 (1H), 2.51 (3H), 2.72 - 2.85 (2H), 3.14 - 3.23 (2H), 3.97 - 4.09 (2H), 4.34 (2H), 7.18 (1H), 7.43 (1H), 8.58 (1H), 8.64 - 8.77 (1H), 8.78 - 8.85 (1H), 8.93 - 9.08 (1H).

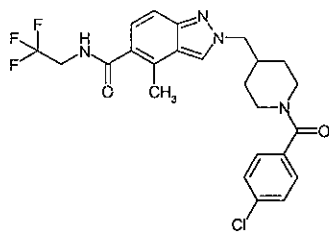
Example 71b: Tert-butyl 4-({4-methyl-5-[N-(2,2,2-trifluoroethyl)carbamoyl]-2H-indazol-2-yl}-methyl)piperidin-1-carboxylate



2820 mg of 1c, 2052 mg of 2,2,2-trifluoroethylamine, 1823.2 mg of molybdenum hexacarbonyl, 200.4 mg of tri-tert-butylphosphine tetrafluoroborate and 647.5 mg of trans-bis(acetato)-bis[o-(di-o-tolylphosphino)-benzyl]dipalladium(II) were placed in a microwave tube and suspended in 56 ml THF. 3.1 ml of DBU were then added and the mixture was stirred for 20 mins at 125°C and 200 watts in the microwave. The reaction mixture was diluted with some ethyl acetate, washed twice with water and once with saturated sodium chloride solution. It was dried over sodium sulphate, filtered and concentrated. The residue was chromatographed on the Biotage SP4. Gradient: hexane/ethyl acetate 0-100%. Yield: 475.2 mg of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.99 - 1.17 (2H), 1.30 - 1.44 (11H), 2.05 - 2.21 (1H), 2.50 (3H), 2.54 - 2.73 (2H), 3.81 - 3.94 (2H), 3.95 - 4.10 (2H), 4.29 (2H), 7.18 (1H), 7.43 (1H), 8.51 (1H), 8.74 - 8.83 (1H).

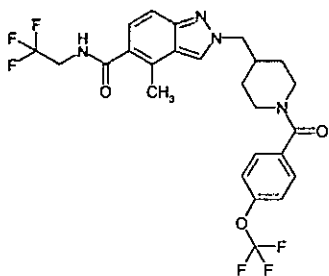
Example 72: 2-{{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 97.8 mg of the title compound was obtained from 146.9 mg of the amine prepared in Example 71a and 58.8 mg of 4-chlorobenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.11 - 1.29 (2H), 1.31 - 1.45 (1H), 1.45 - 1.61 (1H),
 5 2.20 - 2.35 (1H), 2.50 (3H), 2.63 - 2.84 (1H), 2.85 - 3.08 (1H), 3.42 - 3.59 (1H), 3.96 - 4.10
 (2H), 4.29 - 4.47 (3H), 7.17 (1H), 7.36 (2H), 7.40 - 7.50 (3H), 8.52 (1H), 8.75 - 8.82 (1H).

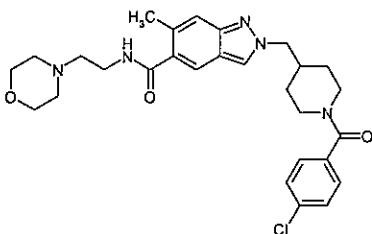
Example 73: 4-methyl-N-(2,2,2-trifluoroethyl)-2-({1-[4-(trifluoromethoxy)benzoyl]piperidin-4-yl} methyl)-2H-indazole-5-carboxamide



10 Analogously to Example 1, 105.4 mg of the title compound was obtained from 146.9 mg of the amine prepared in Example 71a and 77.5 mg of 4-(trifluoromethoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.12 - 1.30 (2H), 1.31 - 1.46 (1H), 1.47 - 1.64 (1H),
 2.20 - 2.36 (1H), 2.50 (3H), 2.66 - 2.85 (1H), 2.86 - 3.09 (1H), 3.40 - 3.55 (1H), 3.96 - 4.10
 (2H), 4.29 - 4.48 (3H), 7.17 (1H), 7.36 - 7.50 (5H), 8.52 (1H), 8.75 - 8.82 (1H).

15 **Example 74:** 2-({1-[4-(4-chlorobenzoyl)piperidin-4-yl]methyl}-6-methyl-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide

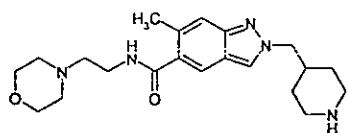


Analogously to Example 55, 30 mg of the title compound was obtained from 79 mg of the amine prepared in Example 74a and 26 mg of 4-chlorobenzoyl chloride.

¹H-NMR (400 MHz, methanol-d₄): δ [ppm]= 1.31 (2H), 1.50 (1H), 1.65 (1H), 2.35 (1H), 2.47 (3H), 2.63 (6H), 2.81 (1H), 3.07 (1H), 3.53 (2H), 3.71 (5H), 4.34 (2H), 4.59 (1H), 7.36 (2H), 7.43 (3H), 7.75 (1H), 8.24 (1H).

The starting material was prepared as follows:

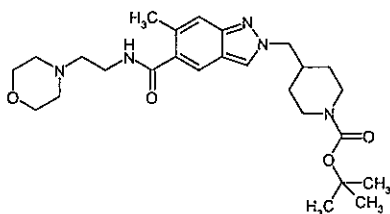
Example 74a: 6-methyl-N-(2-morpholinoethyl)-2-(4-piperidylmethyl)-2H-indazol-5-carboxamide



768 mg of (1.58 mmol) of tert-butyl carbamate 74b were dissolved in 4.0 ml 4M hydrochloric acid in dioxan and stirred under nitrogen at room temperature. A further 0.8 ml 4M hydrochloric acid in dioxan were added and the mixture stirred at room temperature. The reaction solution was concentrated and yielded 760 mg of the title compound, which was used in the next reaction without further purification.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.34 - 1.50 (2H), 1.53 - 1.67 (2H), 2.21 - 2.30 (1H), 2.41 (3H), 2.74 - 2.89 (2H), 3.07 - 3.33 (8H), 3.62 - 3.71 (2H), 3.77 - 3.90 (2H), 3.90 - 4.03 (2H), 4.35 (2H), 7.39 (1H), 7.88 (1H), 8.44 (1H), 8.57 (1H), 8.62 - 8.79 (1H), 8.87 - 9.02 (1H), 11.17 (1H).

Example 74b: Tert-butyl 4-({6-methyl-5-[N-(2-morpholinoethyl)carbamoyl]-2H-indazol-2-yl}-methyl)piperidin-1-carboxylate

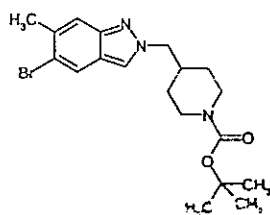


3.00 g (7.35 mmol) of 5-bromoindazole 74c and 2.87 g (22.04 mmol) of 2-morpholinoethylamine were dissolved in 134 ml dioxan and 1.94 g (7.35 mmol) of molybdenum hexacarbonyl, 167 mg (1.47 mmol) of palladium acetate, 213 mg (0.74 mmol) of tributylphosphonium tetrafluoroborate, 2.34 g (22.0 mmol) of sodium carbonate and 2 drops of

water were added. The reaction mixture was heated to 125°C under nitrogen and stirred at this temperature for 2.5 hrs. For the work-up, the suspension was filtered over Celite, rewashed with dioxan, concentrated in vacuo and dried. The crude product was separated by column chromatography. Yield: 518 mg of the title compound.

- 5 ¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.93 - 1.16 (2H), 1.26-1.37 (2H), 1.34 (9H), 1.99 - 2.17 (1H), 2.32 - 2.43 (10H), 2.54 - 2.72 (2H), 3.34 (1H), 3.48 - 3.59 (4H), 3.86 (2H), 4.26 (2H), 7.35 (1H), 7.63 (1H), 8.09 (1H), 8.32 (1H).

Example 74c: Tert-butyl 4-[(5-bromo-6-methyl-2H-indazol-2-yl)methyl]piperidin-1-carboxylate

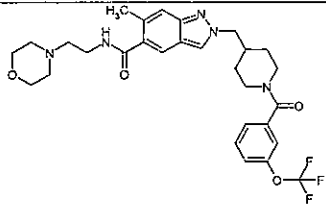


- 10 10.0 g (47.4 mmol) of 5-bromo-4-methyl-1H-indazole and 21.0 g (56.9 mmol) of tert-butyl 4-[(tosyloxy)methyl]piperidin-1-carboxylate were dissolved in 726 ml DMF. To this were added 21.0 g (56.9 mmol) of tetrabutylammonium iodide and 18.5 g (56.9 mmol) of caesium carbonate and the reaction mixture was heated under nitrogen at 80°C for 1.5 hrs. For the work-up, it was treated with water and ethyl acetate, the organic phase separated and the aqueous phase extracted
- 15 several times with ethyl acetate. The organic phases were washed with saturated sodium chloride solution, dried over sodium sulphate and concentrated in vacuo. After purification by column chromatography, this yielded the title compound in 5 g yield.

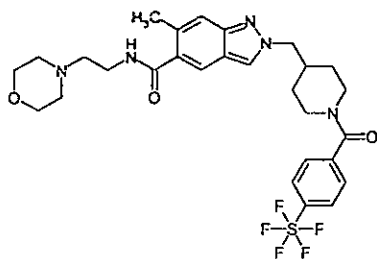
¹H-NMR (600 MHz, chloroform-d): δ [ppm]= 1.16 - 1.30 (2H), 1.45 (9H), 1.54 (2H), 2.22 - 2.33 (1H), 2.51 (3H), 2.67 (2H), 4.03 - 4.17 (2H), 4.27 - 4.32 (2H), 7.60 (1H), 7.81 (1H), 7.90 (1H).

- 20 The following compounds were prepared analogously:

Ex.	Structure	IUPAC name	Analysis
75		6-methyl-N-(2-morpholinoethyl)-2-((1-[4-(trifluoromethoxy)benzoyl]piperidin-4-yl)methyl)-2H-indazole-5-	¹ H-NMR (400 MHz, methanol-d ₄): δ [ppm]= 1.32 (2H), 1.51 (1H), 1.65 (1H), 2.36 (1H), 2.47 (3H), 2.60 (6H), 2.83 (1H), 3.10 (1H), 3.52 (2H), 3.70 (5H), 4.35 (2H), 4.61 (1H), 7.37 (3H), 7.49

		carboxamide	(2H), 7.74 (1H), 8.24 (1H).
76		6-methyl-N-(2-morpholinoethyl)-2-((1-[3-(trifluoromethoxy)benzoyl]piperidin-4-yl)methyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, methanol-d ₄): δ [ppm]= 1.32 (2H), 1.49 (1H), 1.65 (1H), 2.36 (1H), 2.46 (3H), 2.59 (6H), 2.83 (1H), 3.10 (1H), 3.51 (2H), 3.69 (5H), 4.35 (2H), 4.60 (1H), 7.33 (2H), 7.40 (2H), 7.54 (1H), 7.74 (1H), 8.24 (1H).

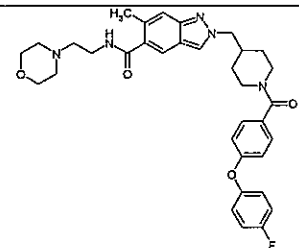
Example 77: 6-methyl-N-(2-morpholinoethyl)-2-((1-[4-(pentafluoro-λ⁶-sulphanyl)benzoyl]piperidin-4-yl)methyl)-2H-indazole-5-carboxamide

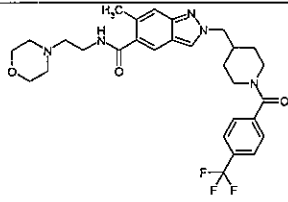
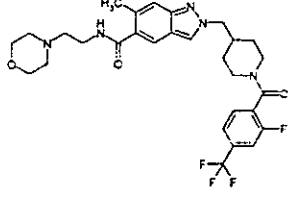
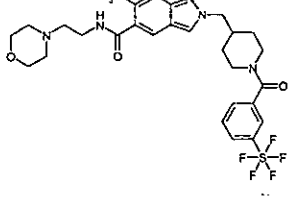
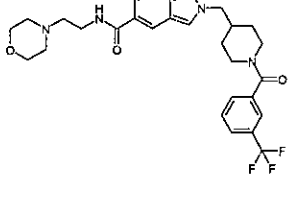


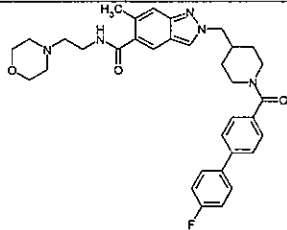
- 5 Analogously to Example 1, 23 mg of the title compound was obtained from 79 mg of the amine prepared in Example 74a and 47 mg of the acid.

¹H-NMR (400 MHz, methanol-d₄): δ [ppm]= 1.34 (2H), 1.52 (1H), 1.66 (1H), 2.37 (1H), 2.49 (3H), 2.84 (1H), 3.10 (1H), 3.25 (2H), 3.42 (2H), 3.63 (3H), 3.75 (4H), 4.10 (2H), 4.36 (2H), 4.62 (1H), 7.43 (1H), 7.56 (2H), 7.89 (3H), 8.28 (1H).

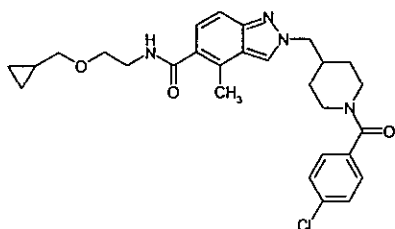
- 10 The following compounds were prepared analogously:

Ex.	Structure	IUPAC name	Analysis
78		2-((1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl)methyl)-6-methyl-N-(2-morpholinoethyl)-2H-indazole-5-	¹ H-NMR (400 MHz, methanol-d ₄): δ [ppm]= 1.31 (2H), 1.58 (2H), 2.35 (1H), 2.49 (3H), 2.40-4.20 (15H), 4.35 (2H), 4.58 (1H), 6.97 (2H), 7.04 (2H), 7.11 (2H), 7.37 (2H), 7.43 (1H), 7.87 (1H),

Ex.	Structure	IUPAC name	Analysis
		carboxamide	8.27 (1H).
79		6-methyl-N-(2-morpholinoethyl)-2-((1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl)methyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, methanol-d ₄): δ [ppm]= 1.34 (2H), 1.51 (1H), 1.67 (1H), 2.38 (1H), 2.50 (3H), 2.85 (1H), 3.10 (1H), 3.25 (2H), 3.43 (2H), 3.64 (3H), 3.76 (4H), 4.10 (2H), 4.38 (2H), 4.64 (1H), 7.44 (1H), 7.57 (2H), 7.76 (2H), 7.89 (1H), 8.31 (1H).
80		2-((1-[2-fluoro-4-(trifluoromethyl)benzoyl]piperidin-4-yl)methyl)-6-methyl-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, methanol-d ₄): δ [ppm]= 1.34 (2H), 1.54 (1H), 1.68 (1H), 2.38 (1H), 2.50 (3H), 2.86 (1H), 3.12 (1H), 3.26 (2H), 3.43 (3H), 3.66 (2H), 3.76 (4H), 4.11 (2H), 4.37 (2H), 4.67 (1H), 7.44 (1H), 7.59 (3H), 7.89 (1H), 8.31 (1H).
81		6-methyl-N-(2-morpholinoethyl)-2-((1-[3-(pentafluoro-λ ⁶ -sulphanyl)benzoyl]piperidin-4-yl)methyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, methanol-d ₄): δ [ppm]= 1.35 (2H), 1.53 (1H), 1.65 (1H), 2.38 (1H), 2.50 (3H), 2.86 (1H), 3.14 (1H), 3.26 (2H), 3.43 (2H), 3.66 (3H), 3.76 (4H), 4.11 (2H), 4.37 (2H), 4.62 (1H), 7.44 (1H), 7.63 (2H), 7.89 (3H), 8.30 (1H).
82		6-methyl-N-(2-morpholinoethyl)-2-((1-[3-(trifluoromethyl)benzoyl]piperidin-4-yl)methyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, methanol-d ₄): δ [ppm]= 1.35 (2H), 1.53 (1H), 1.65 (1H), 2.38 (1H), 2.50 (3H), 2.85 (1H), 3.12 (1H), 3.26 (2H), 3.43 (2H), 3.66 (3H), 3.76 (4H), 4.11 (2H), 4.38 (2H), 4.63 (1H), 7.45 (1H), 7.66 (3H), 7.77

Ex.	Structure	IUPAC name	Analysis
		carboxamide	(1H), 7.90 (1H), 8.33 (1H).
83		2-({1-[(4'-fluoro-biphenyl-4-yl)-carbonyl]piperidin-4-yl}methyl)-6-methyl-N-(2-morpholino-ethyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, methanol-d ₄): δ [ppm]= 1.35 (2H), 1.53 (1H), 1.65 (1H), 2.39 (1H), 2.50 (3H), 2.85 (1H), 3.12 (1H), 3.26 (2H), 3.42 (2H), 3.63 (3H), 3.74 (4H), 4.08 (2H), 4.37 (2H), 4.64 (1H), 7.18 (2H), 7.46 (3H), 7.66 (4H), 7.88 (1H), 8.30 (1H).

Example 84: 2-{{1-(4-chlorobenzoyl)piperidin-4-yl}methyl}-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide

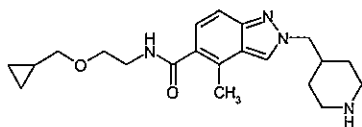


- 5 Analogously to Example 1, 38 mg of the title compound was obtained from 75 mg of the amine prepared in Example 84a and 29 mg of 4-chlorobenzoic acid.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 0.14 (2H), 0.42 (2H), 0.97 (1H), 1.21 (2H), 1.36 (1H), 1.52 (1H), 2.27 (1H), 2.50 (3H), 2.72 (1H), 2.97 (1H), 3.23 (2H), 3.36 (2H), 3.48 (3H), 4.31 (2H), 4.40 (1H), 7.14 (1H), 7.36 (3H), 7.46 (2H), 8.12 (1H), 8.47 (1H).

- 10 The starting material was prepared as follows:

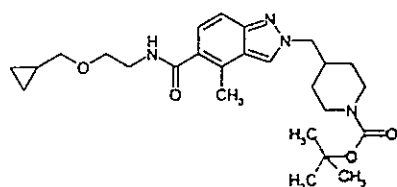
Example 84a: N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide



To 781 mg of 84b were added 3.7 ml of 4M hydrochloric acid in dioxan and 0.5 ml dioxan. The mixture was stirred for 30 mins at ca. 30°C. The reaction mixture was concentrated, taken up in some toluene and again concentrated. Yield: 751 mg of the title compound, which was further reacted without further purification.

- 5 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 0.14 (2H), 0.42 (2H), 0.97 (1H), 1.43 (2H), 1.58 (2H), 2.26 (1H), 2.50 (3H), 2.65 (1H), 2.78(2H), 3.18 (2H), 3.22 (2H), 3.36 (2H), 3.49 (2H), 4.33 (2H), 7.16 (2H), 7.38 (1H), 8.11 (1H), 8.53 (1H), 8.80 (1H), 9.06 (1H).

Example 84b: Tert-butyl 4-[(5-{N-[2-(cyclopropylmethoxy)ethyl]carbamoyl}-4-methyl-2H-indazol-2-yl)methyl]piperidin-1-carboxylate



10

500 mg of 1d was reacted with 203 mg of amine analogously to 1b version B. Yield: 781 mg of the title compound.

- 15 ¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.98 - 1.13 (2H), 1.34 (11H), 2.05 - 2.27 (2H), 2.33 - 2.41 (5H), 2.51 (3H), 2.56 - 2.73 (2H), 3.25 - 3.37 (2H), 3.51 - 3.57 (4H), 3.81 - 3.93 (2H), 4.28 (2H), 7.14 (1H), 7.36 - 7.41 (1H), 7.95 - 8.02 (1H), 8.46 (1H).

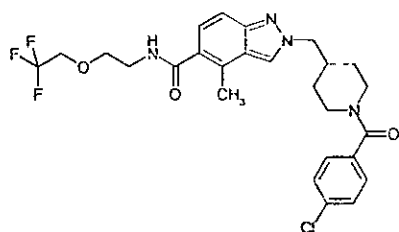
The following compounds were prepared analogously:

Ex.	Structure	IUPAC name	Analysis
85		N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-[[1-(4-methylbenzoyl)-piperidin-4-yl]-methyl]-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 0.14 (2H), 0.42 (2H), 0.97 (1H), 1.19 (2H), 1.37 (1H), 1.50 (1H), 2.26 (1H), 2.29 (3H), 2.50 (3H), 2.72 (1H), 2.94 (1H), 3.23 (2H), 3.36 (2H), 3.48 (2H), 3.57 (1H), 4.30 (2H), 4.39 (1H), 7.14 (1H), 7.21 (4H), 7.38 (1H), 8.12 (1H), 8.47 (1H).

Ex.	Structure	IUPAC name	Analysis
86		N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 0.14 (2H), 0.42 (2H), 0.97 (1H), 1.19 (2H), 1.45 (2H), 2.26 (1H), 2.50 (3H), 2.75 (1H), 2.96 (1H), 3.23 (2H), 3.36 (2H), 3.48 (2H), 3.62 (1H), 4.31 (2H), 4.39 (1H), 6.95 (2H), 7.12 (3H), 7.23 (2H), 7.36 (3H), 8.12 (1H), 8.47 (1H).
87		N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 0.14 (2H), 0.42 (2H), 0.96 (1H), 1.22 (2H), 1.36 (1H), 1.56 (1H), 2.28 (1H), 2.50 (3H), 2.74 (1H), 2.99 (1H), 3.23 (2H), 3.36 (2H), 2.43 (1H), 3.48 (2H), 4.31 (2H), 4.43 (1H), 7.15 (1H), 7.38 (1H), 7.55 (2H), 7.77 (2H), 8.12 (1H), 8.47 (1H).
88		N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[4-(4-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 0.14 (2H), 0.42 (2H), 0.96 (1H), 1.23 (2H), 1.39 (1H), 1.55 (1H), 2.28 (1H), 2.50 (3H), 2.74 (1H), 3.01 (1H), 3.23 (2H), 3.36 (2H), 3.48 (2H), 3.61 (1H), 4.33 (2H), 4.44 (1H), 7.15 (1H), 7.28 (2H), 7.40 (3H), 7.69 (4H), 8.12 (1H), 8.48 (1H).
89		2-({1-[4-(cyclopropylmethoxy)benzoyl]piperidin-4-yl}methyl)-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 0.17 (2H), 0.45 (2H), 0.69 (2H), 0.97 (3H), 1.21 (2H), 1.47 (2H), 1.93 (1H), 2.27 (1H), 2.52 (3H), 2.78 (1H), 2.93 (1H), 3.26

Ex.	Structure	IUPAC name	Analysis
		indazole-5-carboxamide	(2H), 3.37 (2H), 3.51 (2H), 3.62 (1H), 4.34 (2H), 4.41 (1H), 7.10 (2H), 7.17 (1H), 7.22 (2H), 7.41 (1H), 8.11 (1H), 8.49 (1H).

Example 90: 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide

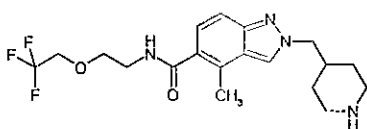


- 5 Analogously to Example 1, 40 mg of the title compound was obtained from 75 mg of the amine prepared in Example 90a and 27 mg of 4-chlorobenzoic acid.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm] = 1.20 (2H), 1.36 (1H), 1.53 (1H), 2.27 (1H), 2.49 (3H), 2.72 (1H), 2.97 (1H), 3.40 (2H), 3.49 (1H), 3.69 (2H), 4.07 (2H), 4.31 (2H), 4.40 (1H), 7.15 (1H), 7.37 (3H), 7.46 (2H), 8.19 (1H), 8.48 (1H).

- 10 The starting material was prepared as follows:

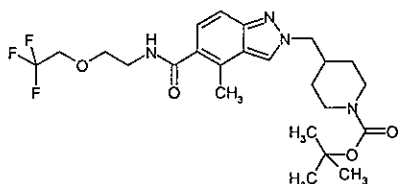
Example 90a: 4-methyl-2-(4-piperidylmethyl)-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide



- 15 To 610 mg of 90b were added 3.0 ml of 4M hydrochloric acid in dioxan and 0.5 ml dioxan. The mixture was stirred for 30 mins at ca. 30°C. The reaction mixture was concentrated, taken up in some toluene and again concentrated. Yield: 619 mg of the title compound, which was further reacted without further purification.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.42 (2H), 1.59 (2H), 2.26 (1H), 2.50 (3H), 2.64 (1H), 2.79 (2H), 3.20 (2H), 3.40 (2H), 3.70 (2H), 4.07 (2H), 4.33 (2H), 5.67(1H), 7.16 (1H), 7.39 (1H), 8.18 (1H), 8.53 (1H), 8.69 (1H), 8.94 (1H).

Example 90b: Tert-butyl 4-[(4-methyl-5-{N-[2-(2,2,2-trifluoroethoxy)ethyl]carbamoyl}-2H-indazol-2-yl)methyl]piperidin-1-carboxylate

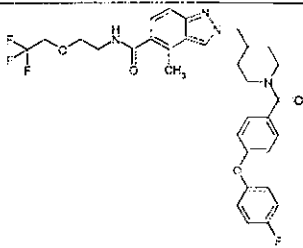
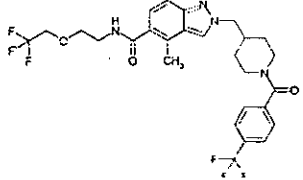
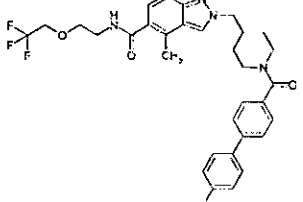
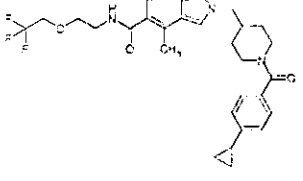


500 mg of 1d was reacted with 240 mg of amine analogously to 1b version B. Yield: 670 mg of the title compound.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.07 (2H), 1.34 (9H), 1.37 (2H), 2.14 (1H), 2.49 (3H), 2.63 (2H), 3.40 (2H), 3.70 (2H), 3.88 (2H), 4.06 (2H), 4.29 (2H), 7.15 (1H), 7.39 (1H), 8.16 (1H), 8.47 (1H).

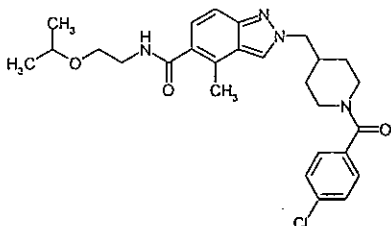
The following compounds were prepared analogously:

Ex.	Structure	IUPAC name	Analysis
91		4-methyl-2-{[1-(4-methylbenzoyl)-piperidin-4-yl]-methyl}-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.19 (2H), 1.36 (1H), 1.49 (1H), 2.25 (1H), 2.29 (3H), 2.49 (3H), 2.71 (1H), 2.93 (1H), 3.40 (2H), 3.56 (1H), 3.69 (2H), 4.07 (2H), 4.31 (2H), 4.40 (1H), 7.15 (1H), 7.21 (4H), 7.39 (1H), 8.19 (1H), 8.48 (1H).

Ex.	Structure	IUPAC name	Analysis
92		2-({1-[4-(4-fluorophenoxy)benzoyl]-piperidin-4-yl}-methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.19 (2H), 1.44 (2H), 2.26 (1H), 2.49 (3H), 2.73 (1H), 2.95 (1H), 3.40 (2H), 3.60 (1H), 3.69 (2H), 4.07 (2H), 4.31 (2H), 4.38 (1H), 6.95 (2H), 7.12 (3H), 7.22 (2H), 7.37 (3H), 8.19 (1H), 8.48 (1H).
93		4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2-({1-[4-(trifluoromethyl)benzoyl]-piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.22 (2H), 1.36 (1H), 1.56 (1H), 2.28 (1H), 2.49 (3H), 2.74 (1H), 2.99 (1H), 3.40 (3H), 3.69 (2H), 4.07 (2H), 4.32 (2H), 4.43 (1H), 7.15 (1H), 7.39 (1H), 7.55 (2H), 7.77 (2H), 8.19 (1H), 8.48 (1H).
94		2-({1-[4-(4'-fluorobiphenyl-4-yl)-carbonyl]piperidin-4-yl}methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.23 (2H), 1.39 (1H), 1.52 (1H), 2.28 (1H), 2.49 (3H), 2.75 (1H), 2.88 (1H), 3.00 (1H), 3.39 (2H), 3.60 (1H), 3.69 (2H), 4.07 (2H), 4.33 (2H), 4.43 (1H), 7.15 (1H), 7.27 (2H), 7.40 (3H), 7.68 (4H), 8.19 (1H), 8.49 (1H).
95		2-({1-[4-(cyclopropylbenzoyl)-piperidin-4-yl]-methyl}-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 0.69 (2H), 0.96 (2H), 1.21 (2H), 1.47 (2H), 1.93 (1H), 2.28 (1H), 2.52 (3H), 2.77 (1H), 2.94 (1H), 3.43 (2H), 3.61 (1H), 3.72 (2H), 4.09 (2H), 4.34 (2H), 4.42 (1H), 7.10 (2H), 7.17 (1H), 7.22

Ex.	Structure	IUPAC name	Analysis
		carboxamide	(2H), 7.41 (1H), 8.18 (1H).

Example 96: 2-{{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide

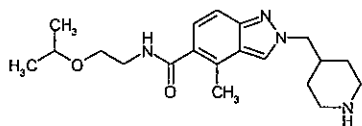


- 5 Analogously to Example 1, 36 mg of the title compound was obtained from 75 mg of the amine prepared in Example 96a and 30 mg of 4-chlorobenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.07 (6H), 1.22 (2H), 1.39 (1H), 1.51 (1H), 2.26 (1H), 2.50 (3H), 2.75 (1H), 2.97 (1H), 3.33 (2H), 3.45 (3H), 3.55 (1H), 4.31 (2H), 4.39 (1H), 7.14 (1H), 7.36 (3H), 7.47 (2H), 8.05 (1H), 8.46 (1H).

- 10 The starting material was prepared as follows:

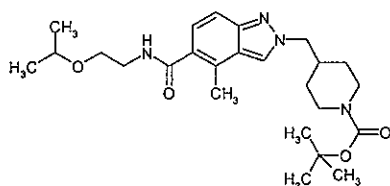
Example 96a: N-(2-isopropoxyethyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazol-5-carboxamide



- 15 To 650 mg of 96b were added 3.2 ml of 4M hydrochloric acid in dioxan and 0.5 ml dioxan. The mixture was stirred for 30 mins at ca. 30°C. The reaction mixture was concentrated, taken up in some toluene and again concentrated. Yield: 529 mg of the title compound, which was further reacted without further purification.

- 20 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.06 (3H), 1.07 (3H), 1.42 (2H), 1.57 (2H), 2.25 (1H), 2.50 (3H), 2.65 (1H), 2.78 (2H), 3.19 (2H), 3.32 (2H), 3.45 (2H), 3.55 (1H), 4.32 (1H), 6.61 (1H), 7.15 (1H), 7.38 (1H), 8.11 (1H), 8.53 (1H), 8.78 (1H), 9.04 (1H).

Example 96b: Tert-butyl 4-({5-[N-(2-isopropoxyethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}-methyl)piperidin-1-carboxylate

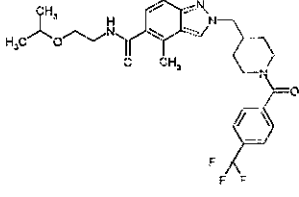
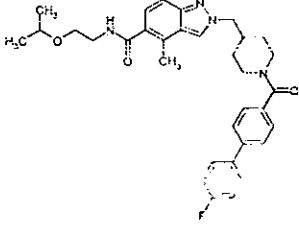
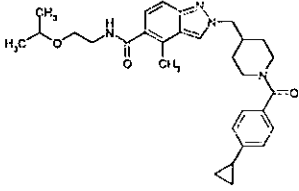


500 mg of 1d was reacted with 138 mg of amine analogously to 1b version B. Yield: 650 mg of the title compound.

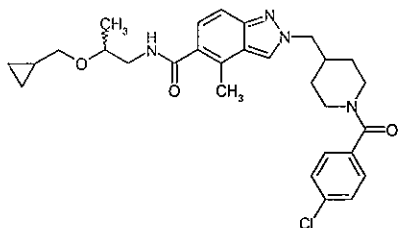
¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.07 (8H), 1.34 (9H), 1.38 (2H), 2.13 (1H), 2.50 (3H), 2.63 (2H), 3.33 (2H), 3.46 (2H), 3.55 (1H), 3.87 (2H), 4.28 (2H), 7.14 (1H), 7.38 (1H), 8.05 (1H), 8.46 (1H).

The following compounds were prepared analogously:

Ex.	Structure	IUPAC name	Analysis
97		N-(2-isopropoxyethyl)-4-methyl-2-({[1-(4-methylbenzoyl)piperidin-4-yl]methyl}-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.07 (6H), 1.19 (2H), 1.45 (2H), 2.29 (4H), 2.50 (3H), 2.75 (1H), 2.92 (1H), 3.33 (2H), 3.45 (2H), 3.55 (2H), 4.31 (2H), 4.40 (1H), 7.14 (1H), 7.20 (4H), 7.38 (1H), 8.05 (1H), 8.46 (1H).
98		2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}-methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.07 (6H), 1.21 (2H), 1.45 (2H), 2.25 (1H), 2.49 (3H), 2.85 (2H), 3.31 (2H), 3.45 (2H), 3.55 (2H), 4.31 (2H), 4.40 (1H), 6.95 (2H), 7.10 (3H), 7.22 (2H), 7.36 (3H), 8.05 (1H), 8.46 (1H).

Ex.	Structure	IUPAC name	Analysis
99		N-(2-isopropoxyethyl)-4-methyl-2-({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.06 (6H), 1.22 (2H), 1.37 (1H), 1.56 (1H), 2.28 (1H), 2.50 (3H), 2.75 (1H), 2.99 (1H), 3.33 (2H), 3.45 (3H), 3.55 (1H), 4.32 (2H), 4.43 (1H), 7.15 (1H), 7.38 (1H), 7.55 (2H), 7.77 (2H), 8.05 (1H), 8.46 (1H).
100		2-({1-[4-(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}-methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.06 (6H), 1.23 (2H), 1.47 (2H), 2.28 (1H), 2.50 (3H), 2.76 (1H), 2.98 (1H), 3.33 (2H), 3.45 (2H), 3.55 (2H), 4.33 (2H), 4.43 (1H), 7.15 (1H), 7.27 (2H), 7.39 (3H), 7.68 (4H), 8.05 (1H), 8.47 (1H).
101		2-({1-[4-(cyclopropylbenzoyl)piperidin-4-yl]-methyl}-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 0.66 (2H), 0.94 (2H), 1.06 (6H), 1.20 (2H), 1.44 (2H), 1.90 (1H), 2.24 (1H), 2.49 (3H), 2.82 (2H), 3.32 (2H), 3.45 (2H), 3.55 (2H), 4.31 (2H), 4.39 (1H), 7.12 (5H), 7.38 (1H), 8.05 (1H), 8.46 (1H).

Example 102: (+/-)-2-({1-[4-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-[2-(cyclopropylmethoxy)propyl]-4-methyl-2H-indazole-5-carboxamide

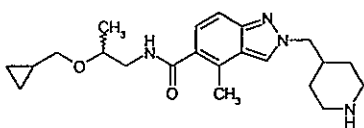


Analogously to Example 1, 41 mg of the title compound was obtained from 100 mg of the amine prepared in Example 102a and 37 mg of 4-chlorobenzoic acid.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 0.16 (2H), 0.43 (2H), 0.97 (1H), 1.10 (3H), 1.23 (2H), 1.40 (1H), 1.55 (1H), 2.29 (1H), 2.53 (3H), 2.75 (1H), 2.99 (1H), 3.23 (2H), 3.20 (2H), 3.51 (1H), 3.59 (1H), 4.34 (2H), 4.43 (1H), 7.17 (1H), 7.39 (3H), 7.49 (2H), 8.09 (1H), 8.49 (1H).

The starting material was prepared as follows:

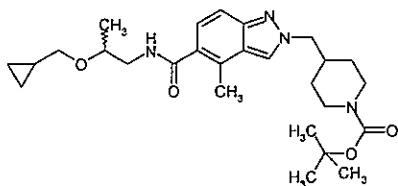
Example 102a: (+/-)-N-[2-(cyclopropylmethoxy)propyl]-4-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide



To 632 mg of 102b were added 2.9 ml of 4M hydrochloric acid in dioxan and 0.5 ml dioxan. The mixture was stirred for 30 mins at ca. 30°C. The reaction mixture was concentrated, taken up in some toluene and again concentrated. Yield: 561 mg of the title compound, which was further reacted without further purification.

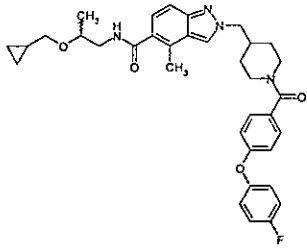
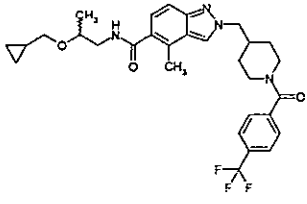
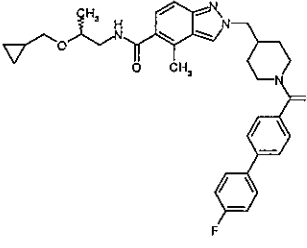
¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.13 (2H), 0.40 (2H), 0.94 (1H), 1.07 (3H), 1.44 (2H), 1.57 (2H), 2.26 (1H), 2.50 (3H), 2.77 (2H), 3.20 (6H), 3.57 (1H), 4.33 (2H), 6.33 (1H), 7.16 (1H), 7.39 (1H), 8.11 (1H), 8.53 (1H), 8.79 (1H), 9.06 (1H).

Example 102b: (+/-)-tert-butyl 4-[(5-{N-[2-(cyclopropylmethoxy)propyl]carbamoyl}-4-methyl-2H-indazol-2-yl)methyl]piperidin-1-carboxylate



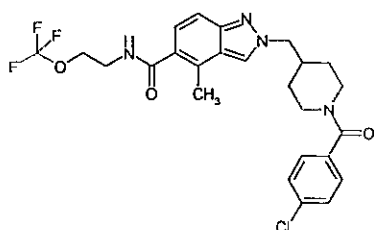
500 mg of 1d was reacted with 222 mg of amine analogously to 1b version B. Yield: 631 mg of the title compound. ¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.13 (2H), 0.40 (2H), 0.94 (1H), 1.07 (3H), 1.14 (2H), 1.34 (9H), 1.38 (2H), 2.13 (1H), 2.50 (3H), 3.24 (6H), 3.57 (1H), 3.87 (2H), 4.28 (2H), 7.15 (1H), 7.39 (1H), 8.09 (1H), 8.46 (1H).

5 The following compounds were prepared analogously:

Ex.	Structure	IUPAC name	Analysis
103		(+/-)-N-[2-(cyclopropylmethoxy)propyl]-2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 0.16 (2H), 0.43 (2H), 0.97 (1H), 1.10 (3H), 1.22 (2H), 1.48 (2H), 2.29 (1H), 2.53 (3H), 2.81 (1H), 2.92 (1H), 3.23 (2H), 3.28 (2H), 3.60 (1H), 3.70 (1H), 4.34 (2H), 4.42 (1H), 6.98 (2H), 7.14 (3H), 7.25 (2H), 7.39 (3H), 8.09 (1H), 8.49 (1H).
104		(+/-)-N-[2-(cyclopropylmethoxy)propyl]-4-methyl-2-({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 0.16 (2H), 0.43 (2H), 0.97 (1H), 1.10 (3H), 1.24 (2H), 1.38 (1H), 1.59 (1H), 2.30 (1H), 2.53 (3H), 2.78 (1H), 3.02 (1H), 3.23 (2H), 3.28 (2H), 3.45 (1H), 3.60 (1H), 4.34 (2H), 4.46 (1H), 7.17 (1H), 7.41 (1H), 7.57 (2H), 7.80 (2H), 8.09 (1H), 8.49 (1H).
105		(+/-)-N-[2-(cyclopropylmethoxy)propyl]-2-({1-[4-(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 0.16 (2H), 0.42 (2H), 0.97 (1H), 1.10 (3H), 1.26 (2H), 1.45 (1H), 1.55 (1H), 2.32 (1H), 2.53 (3H), 2.79 (1H), 3.03 (1H), 3.23 (2H), 3.27 (2H), 3.60 (2H), 4.36 (2H), 4.46 (1H), 7.17 (1H), 7.30 (2H), 7.42 (3H), 7.71 (4H),

Ex.	Structure	IUPAC name	Analysis
		carboxamide	8.09 (1H), 8.50 (1H).
106		(+/-)-2-[[1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl]-N-[2-(cyclopropylmethoxy)propyl]-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 0.16 (2H), 0.43 (2H), 0.69 (2H), 0.96 (3H), 1.10 (3H), 1.20 (2H), 1.47 (2H), 1.93 (1H), 2.28 (1H), 2.53 (3H), 2.77 (1H), 2.93 (1H), 3.23 (2H), 3.28 (2H), 3.59 (2H), 4.34 (2H), 4.40 (1H), 7.10 (2H), 7.17 (1H), 7.22 (2H), 7.41 (1H), 8.09 (1H), 8.49 (1H).

Example 107: 2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide

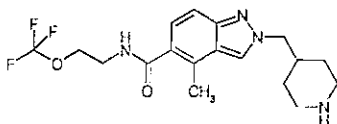


- 5 Analogously to Example 1, 26 mg of the title compound was obtained from 75 mg of the amine prepared in Example 107a and 28 mg of 4-chlorobenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.20 (2H), 1.36 (1H), 1.52 (1H), 2.27 (1H), 2.49 (3H), 2.72 (1H), 2.97 (1H), 3.51 (3H), 4.17 (2H), 4.33 (3H), 7.15 (1H), 7.41 (5H), 8.37 (1H), 8.49 (1H).

- 10 The starting material was prepared as follows:

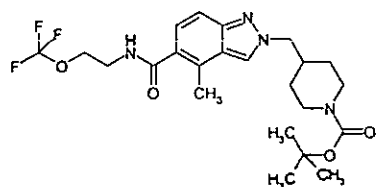
Example 107a: 4-methyl-2-(4-piperidylmethyl)-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide



To 648 mg of 107b were added 3.0 ml of 4M hydrochloric acid in dioxan and 0.5 ml dioxan. The mixture was stirred for 30 mins at ca. 30°C. The reaction mixture was concentrated, taken up in some toluene and again concentrated. Yield: 766 mg of the title compound, which was further reacted without further purification.

- 5 ¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.43 (2H), 1.57 (2H), 2.24 (1H), 2.50 (3H), 2.78 (2H), 3.19 (2H), 3.50 (1H), 4.17 (2H), 4.33 (2H), 6.49 (1H), 7.17 (2H), 7.40 (1H), 8.39 (1H), 8.55 (1H), 8.70 (1H), 8.98 (1H).

Example 107b: Tert-butyl 4-[(4-methyl-5-{N-[2-(trifluoromethoxy)ethyl]carbamoyl}-2H-indazol-2-yl)methyl]piperidin-1-carboxylate



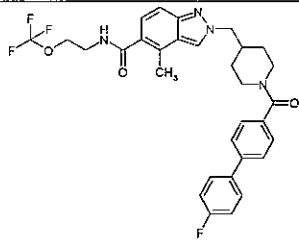
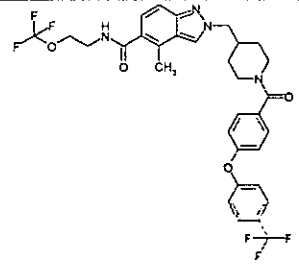
10

500 mg of 1d was reacted with 293 mg of amine analogously to 1b version B. Yield: 748 mg of the title compound.

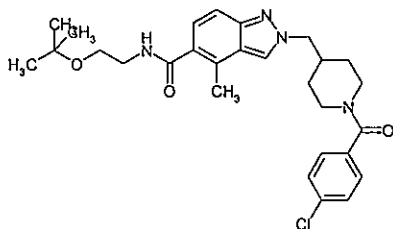
- 15 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.07 (2H), 1.34 (9H), 1.37 (2H), 2.12 (1H), 2.50 (3H), 2.63 (1H), 3.52 (2H), 3.87 (2H), 4.17 (2H), 4.29 (2H), 7.16 (2H), 7.41 (1H), 8.37 (1H), 8.48 (1H).

The following compounds were prepared analogously:

Ex.	Structure	IUPAC name	Analysis
108		2-({1-[4-(4-fluorophenoxy)benzoyl]-piperidin-4-yl}methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.19 (2H), 1.46 (2H), 2.26 (1H), 2.49 (3H), 2.87 (2H), 3.51 (3H), 4.17 (2H), 4.33 (3H), 6.95 (2H), 7.16 (5H), 7.38 (3H), 8.36 (1H), 8.49 (1H).

109		2-({1-[(4'-fluoro-biphenyl-4-yl)-carbonyl]piperidin-4-yl}methyl)-4-methyl-N-[2-(trifluoro-methoxy)ethyl]-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.23 (2H), 1.47 (2H), 2.28 (1H), 2.50 (3H), 2.74 (1H), 2.99 (1H), 3.51 (2H), 3.61 (1H), 4.17 (2H), 4.34 (3H), 7.16 (1H), 7.27 (2H), 7.40 (3H), 7.68 (4H), 8.37 (1H), 8.50 (1H).
110		4-methyl-N-[2-(trifluoromethoxy)ethyl]-2-[(1-{4-[4-(trifluoromethyl)phenoxy]-benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.21 (2H), 1.46 (2H), 2.27 (1H), 2.50 (3H), 2.76 (1H), 2.98 (1H), 3.51 (2H), 3.61 (1H), 4.17 (2H), 4.33 (2H), 4.43 (1H), 7.14 (5H), 7.41 (3H), 7.73 (2H), 8.37 (1H), 8.49 (1H).

Example 111: N-(2-tert-butoxyethyl)-2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-4-methyl-2H-indazole-5-carboxamide

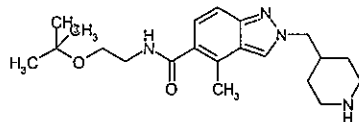


- 5 Analogously to Example 1, 8 mg of the title compound was obtained from 75 mg of the amine prepared in Example 111a and 28 mg of 4-chlorobenzoic acid.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.12 (9H), 1.21 (2H), 1.36 (1H), 1.53 (1H), 2.27 (1H), 2.50 (3H), 2.72 (1H), 2.97 (1H), 3.29 (2H), 3.43 (3H), 4.31 (2H), 4.40 (1H), 7.14 (1H), 7.36 (3H), 7.46 (2H), 8.04 (1H), 8.46 (1H).

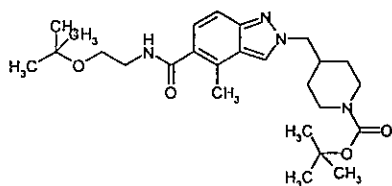
The starting material was prepared as follows:

Example 111a: N-(2-tert-butoxyethyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazol-5-carboxamide



- 5 To 743 mg of 111b were added 3.5 ml of 4M hydrochloric acid in dioxan and 0.5 ml dioxan. The mixture was stirred for 30 mins at ca. 30°C. The reaction mixture was concentrated, taken up in some toluene and again concentrated. Yield: 677 mg of the title compound in a mixture with N-(2-hydroxyethyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide, which was further reacted without further purification.

- 10 **Example 111b:** Tert-butyl 4-({5-[N-(2-tert-butoxyethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}methyl)piperidin-1-carboxylate

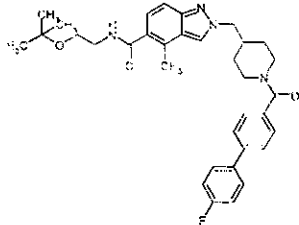
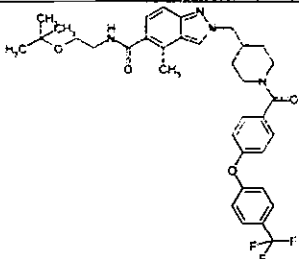


500 mg of 1d was reacted with 206 mg of amine analogously to 1b version B. Yield: 743 mg of the title compound.

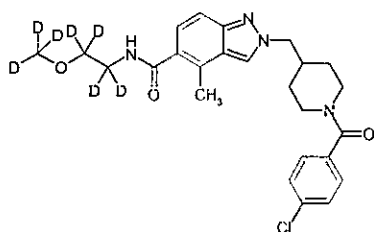
- 15 ¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.06 (2H), 1.11 (9H), 1.34 (9H), 1.39 (2H), 2.13 (1H), 2.50 (3H), 2.65 (2H), 3.27 (2H), 3.40 (2H), 3.87 (2H), 4.28 (2H), 7.14 (1H), 7.38 (1H), 8.05 (1H), 8.46 (1H).

The following compounds were prepared analogously:

Ex.	Structure	IUPAC name	Analysis
112		N-(2-tert-butoxyethyl)-2-({1-[4-(4-fluorophenoxy)benzoyl]-piperidin-4-yl}methyl)-4-methyl-2H-indazole-	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm]= 1.12 (9H), 1.20 (2H), 1.44 (2H), 2.26 (1H), 2.50 (3H), 2.76 (1H), 2.93 (1H), 3.30 (2H), 3.40 (2H), 3.57 (1H), 4.31 (2H), 4.39 (1H), 6.95 (2H), 7.12 (3H),

Ex.	Structure	IUPAC name	Analysis
		5-carboxamide	7.23 (2H), 7.36 (3H), 8.04 (1H), 8.46 (1H).
113		N-(2-tert-butoxyethyl)-2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm] = 1.11 (9H), 1.23 (2H), 1.40 (1H), 1.54 (1H), 2.29 (1H), 2.50 (3H), 2.75 (1H), 3.00 (1H), 3.29 (2H), 3.40 (2H), 3.65 (1H), 4.32 (2H), 4.43 (1H), 7.15 (1H), 7.28 (2H), 7.40 (3H), 7.69 (4H), 8.05 (1H), 8.47 (1H).
114		N-(2-tert-butoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide	¹ H-NMR (400 MHz, DMSO-d ₆): δ [ppm] = 1.12 (9H), 1.21 (2H), 1.40 (1H), 1.52 (1H), 2.27 (1H), 2.50 (3H), 2.75 (1H), 2.99 (1H), 3.28 (2H), 3.40 (2H), 3.60 (1H), 4.32 (2H), 4.40 (1H), 7.15 (5H), 7.40 (3H), 7.73 (2H), 8.04 (1H), 8.47 (1H).

Example 115: 2-{{1-(4-chlorobenzoyl)piperidin-4-yl}methyl}-N-(2-[²H₃]methoxy[²H₄]ethyl)-4-methyl-2H-indazole-5-carboxamide

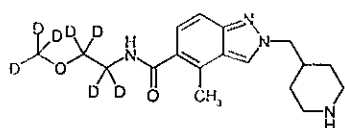


- 5 Analogously to Example 1, 1 mg of the title compound was obtained from 54 mg of the amine prepared in Example 115a and 22 mg of 4-chlorobenzoic acid.

^1H -NMR (300 MHz, DMSO-d_6): δ [ppm]= 1.20 (2H), 1.40 (1H), 1.51 (1H), 2.27 (1H), 2.49 (3H), 2.75 (1H), 2.96 (1H), 3.50 (1H), 4.31 (2H), 4.40 (1H), 7.15 (1H), 7.36 (3H), 7.47 (2H), 8.07 (1H), 8.46 (1H).

The starting material was prepared as follows:

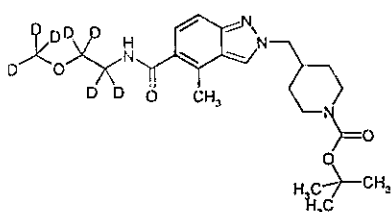
- 5 **Example 115a:** 4-methyl-N-(2-[$^2\text{H}_3$]methyloxy[$^2\text{H}_4$]ethyl)-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide



To 131 mg of 115b were added 0.7 ml of 4M hydrochloric acid in dioxan and 0.5 ml dioxan. The mixture was stirred for 30 mins at ca. 30°C. The reaction mixture was concentrated, taken up in
10 some toluene and again concentrated. Yield: 108 mg of the title compound, which was further reacted without further purification.

^1H -NMR (400 MHz, DMSO-d_6): δ [ppm]= 1.42 (2H), 1.58 (2H), 2.26 (1H), 2.49 (3H), 2.78 (2H), 3.19 (2H), 4.33 (2H), 5.55 (1H), 7.16 (1H), 7.38 (1H), 8.12 (1H), 8.53 (1H), 8.70 (1H), 8.97 (1H).

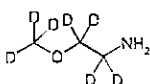
- 15 **Example 115b:** Tert-butyl 4-({4-methyl-5-[N-(2-[$^2\text{H}_3$]methyloxy[$^2\text{H}_4$]ethyl)carbamoyl]-2H-indazol-2-yl}methyl)piperidin-1-carboxylate



500 mg of 1d was reacted analogously to 1b version B with 110 mg of amine prepared in 115c. Yield: 131 mg of the title compound.

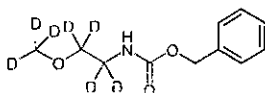
- 20 ^1H -NMR (300 MHz, DMSO-d_6): δ [ppm]= 1.07 (2H), 1.34 (9H), 1.37 (2H), 2.13 (1H), 2.49 (3H), 2.62 (2H), 3.88 (2H), 4.28 (2H), 7.15 (1H), 7.38 (1H), 8.07 (1H), 8.46 (1H).

Example 115c: 2-[$^2\text{H}_3$]methyloxy[$^2\text{H}_4$]ethan-1-amine



110 mg of (0.51 mmol) of 115d was first placed in 2.2 ml toluene. After addition of 11 mg of palladium/charcoal (10%) and hydrogen under normal pressure, the mixture was stirred for 50 mins at room temperature. The reaction mixture was filtered through Celite and rewashed with toluene. The colourless filtrate was used as solution in the subsequent reaction.

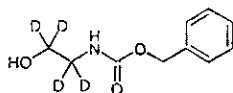
5 **Example 115d:** Benzyl *N*-(2-[²H₃]methyloxy[²H₄]ethyl)carbamate



1.32 g (6.6 mmol) of 115e) was dissolved in 23 ml acetonitrile. 2.3 g (9.9 mmol) of silver oxide and 1.64 ml (3.8 g, 26 mmol) of deuterated iodomethane were then added. The reaction mixture was stirred for 1.5 hrs at 40°C, 7.5 hrs at 72°C and then overnight at room temperature. A further
 10 2.3 g (9.9 mmol) of silver oxide was added and the mixture stirred for 1.5 hrs at 72°C. For the work-up, the black solid was filtered off at the pump and the clear filtrate concentrated. The residue was purified by column chromatography (hexane/ethyl acetate 0-10%). Yield: 1.13 g of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 4.97 (2H), 7.31 (5H).

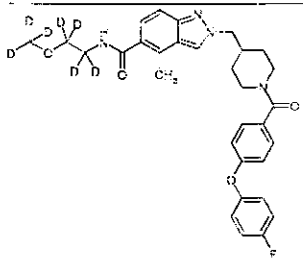
15 **Example 115e:** Benzyl *N*-(2-hydroxy[²H₄]ethyl)carbamate



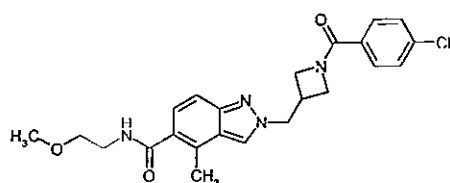
1.0 g (15 mmol) of deuterated aminoethanol was first dissolved in 29 ml dichloromethane. Then 2.5 ml (1.8 g, 18mmol) of triethylamine was added and the reaction mixture cooled to 0°C. At this temperature 2.75 ml (3.34 g, 19.6 mmol) of benzyl chloroformate was now cautiously added
 20 dropwise. After completion of the addition, the mixture was stirred for 1 hr more with no ice-bath and in the process warmed up to R.T. The reaction mixture was diluted once with some dichloromethane and washed once with saturated sodium chloride solution. The organic phase was dried over sodium sulphate, filtered and concentrated. The crude product was purified twice by column chromatography (dichloromethane/methanol 0-10% and ethyl acetate). Yield: 1.32 g
 25 of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 4.54 (1H), 4.97 (2H), 7.12 (1H), 7.81 (5H).

The following compound was prepared analogously:

Ex.	Structure	IUPAC name	Analysis
116		2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2-[² H ₃]-methoxy[² H ₄]ethyl)-2H-indazole-5-carboxamide	¹ H-NMR (300 MHz, DMSO-d ₆): δ [ppm]= 1.19 (2H), 1.46 (2H), 2.25 (1H), 2.49 (3H), 2.85 (2H), 3.66 (1H), 4.31 (2H), 4.38 (1H), 6.95 (2H), 7.16 (5H), 7.36 (3H), 8.07 (1H), 8.46 (1H).

Example 117: 2-{{1-(4-chlorobenzoyl)azetidin-3-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



5

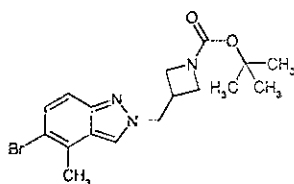
Analogously to Example 55, 471 mg of the title compound was obtained from 500 mg of the compound prepared in Example 117c) and 287 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.22 (1H), 3.25 (3H), 3.32-3.46 (4H), 3.89 (1H), 4.09 (1H), 4.19 (1H), 4.37 (1H), 4.67 (2H), 7.15 (1H), 7.38 (1H), 7.48 (2H), 7.59 (2H), 8.12 (1H), 8.54 (1H).

10

The starting material was prepared as follows:

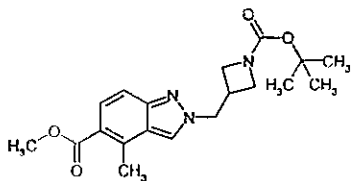
Example 117a: Tert-butyl 3-[(5-bromo-4-methyl-2H-indazol-2-yl)methyl]-azetidin-1-carboxylate



To a solution of 21.1 g of 5-bromo-4-methyl-1H-indazole and 37.6 g of tert-butyl-3-[(tosyloxy)-methyl]azetidin-1-carboxylate in 100 ml dioxan were added 150 ml of a 1M solution of sodium bis(trimethylsilyl)amide in tetrahydrofuran at 25°C and this was stirred for 6 hrs at 90°C. After cooling, the reaction mixture was treated with ethyl acetate and water, the organic phases separated and the aqueous phase extracted twice with 100 ml portions of ethyl acetate. The combined organic phases were washed with saturated sodium chloride solution, dried over sodium sulphate, and concentrated in vacuo after filtration. The residue thus obtained was purified by chromatography on the Flashmaster (hexane/ethyl acetate 1:0-0:1). 15.0 g of the title compound was obtained.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.48 (3H), 3.07 (1H), 3.69 (2H), 3.87 (2H), 4.59 (2H), 7.28 (1H), 7.35 (1H), 8.53 (1H).

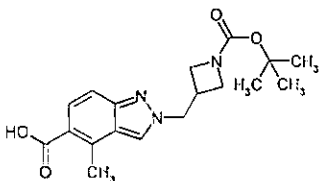
Example 117b: Methyl 2-[[1-(tert-butoxycarbonyl)azetidin-3-yl]methyl]-4-methyl-2H-indazol-5-carboxylate



To a solution of 8.16 g of the bromide prepared in Example 117a in 65.4 ml methanol, 6.5 g of 1,1 bis(diphenylphosphino)ferrocen-palladium(II) dichloride and 15 g of potassium acetate were added at 25°C and the mixture was stirred under CO at 10.15 bar and 120°C for 24 hours in an autoclave. The reaction mixture was then cooled, filtered through Celite at the pump and the filtrate concentrated in vacuo. The residue thus obtained was purified by chromatography on the Flashmaster (hexane/ethyl acetate 1:0-0:1). 6.97g of the title compound was obtained.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.73 (3H), 3.09 (1H), 3.71 (2H), 3.79 (3H), 3.87 (2H), 4.62 (2H), 7.42 (1H), 7.63 (1H), 8.74 (1H).

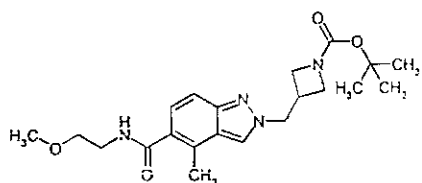
Example 117c: 2-[[1-(tert-butoxycarbonyl)azetidin-3-yl]methyl]-4-methyl-2H-indazol-5-carboxylic acid



Analogously to Example 1d, 5.2 g of the title compound was obtained from 15.1 g of the ester prepared in Example 117b.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.73 (3H), 3.09 (1H), 3.71 (2H), 3.87 (2H), 4.61 (2H), 7.39 (1H), 7.64 (1H), 8.70 (1H), 12.57 (1H).

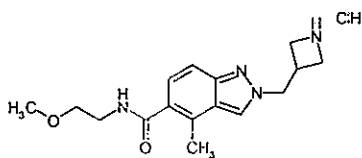
- 5 **Example 117d:** Tert-butyl 3-({5-[N-(2-methoxyethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}-methyl)azetidin-1-carboxylate



Analogously to Example 1, from 2.5 g of the acid prepared in Example 117c and 0.54 g of 2-methoxyethylamine, a crude product was obtained which through purification with a Biotage
10 unit (ethyl acetate : 0 to 30% methanol) yielded 2.69 g of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.49 (3H), 3.08 (1H), 3.25 (3H), 3.32-3.46 (4H), 3.70 (2H), 3.86 (2H), 4.60 (2H), 7.15 (1H), 7.38 (1H), 8.11 (1H), 8.54 (1H).

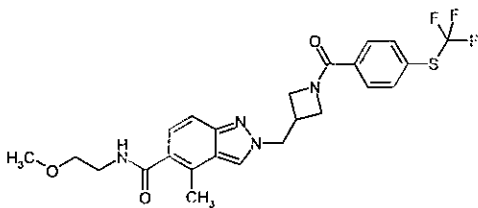
Example 117e: 2-(azetidin-3-ylmethyl)-N-(2-methoxyethyl)-4-methyl-2H-indazol-5-carboxamide hydrochloride



15

Analogously to Example 1a, from 2.69 g of the amide prepared in Example 117d, 2.59 g of the title compound was obtained, which was reacted without further purification.

Example 118: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[(trifluoromethyl)sulphonyl]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide

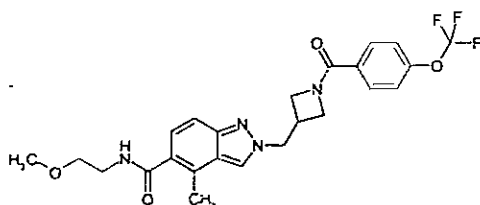


20

Analogously to Example 55, 29 mg of the title compound was obtained from 55 mg of the compound prepared in Example 117e and 43 mg of 4-[(trifluoromethyl)sulphonyl]benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.22 (1H), 3.25 (3H), 3.36 (2H), 3.42 (2H), 3.92 (1H), 4.11 (1H), 4.20 (1H), 4.39 (1H), 4.67 (2H), 7.15 (1H), 7.38 (1H), 7.69 (2H), 7.76 (2H), 8.12 (1H), 8.54 (1H).

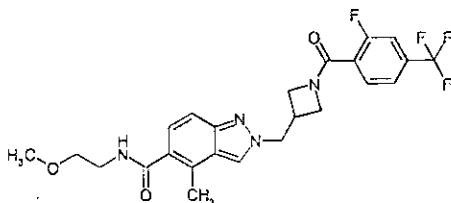
Example 119: N-(2-methoxyethyl)-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 27 mg of the title compound was obtained from 55 mg of the compound prepared in Example 117e and 40 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.20 (1H), 3.24 (3H), 3.32-3.46 (4H), 3.91 (1H), 4.10 (1H), 4.20 (1H), 4.39 (1H), 4.67 (2H), 7.15 (1H), 7.35-7.44 (3H), 7.70 (2H), 8.12 (1H), 8.54 (1H).

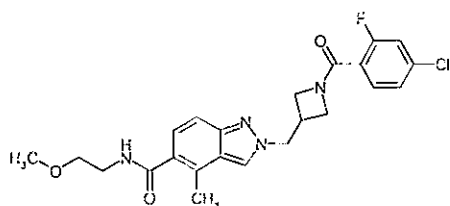
Example 120: 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 17 mg of the title compound was obtained from 55 mg of the compound prepared in Example 117e and 41 mg of 2-fluoro-4-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.48 (3H), 3.23 (1H), 3.25 (3H), 3.36 (2H), 3.42 (2H), 3.94 (2H), 4.11 (2H), 4.66 (2H), 7.15 (1H), 7.38 (1H), 7.64 (2H), 7.79 (1H), 8.12 (1H), 8.54 (1H).

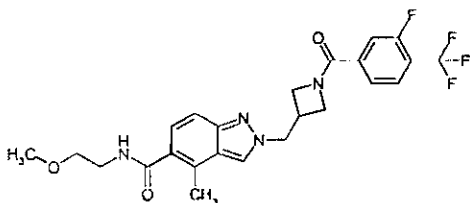
Example 121: 2-([1-(4-chloro-2-fluorobenzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 19 mg of the title compound was obtained from 42 mg of the compound prepared in Example 117e and 26 mg of 4-chloro-2-fluorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.48 (3H), 3.21 (1H), 3.24 (3H), 3.33-3.46 (4H), 3.86-3.97 (2H), 4.08 (2H), 4.65 (2H), 7.15 (1H), 7.34 (1H), 7.37 (1H), 7.44 (1H), 7.52 (1H), 8.12 (1H), 8.53 (1H).

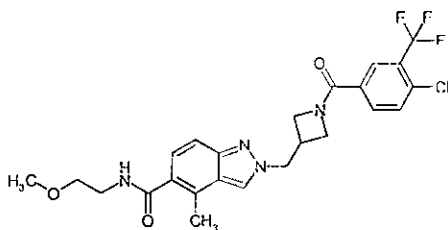
Example 122: 2-([1-[3-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 15 mg of the title compound was obtained from 42 mg of the compound prepared in Example 117c and 31 mg of 3-fluoro-4-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.21 (1H), 3.25 (3H), 3.36 (2H), 3.42 (2H), 3.93 (1H), 4.12 (1H), 4.22 (1H), 4.40 (1H), 4.67 (2H), 7.15 (1H), 7.37 (1H), 7.57 (1H), 7.64 (1H), 7.85 (1H), 8.12 (1H), 8.54 (1H).

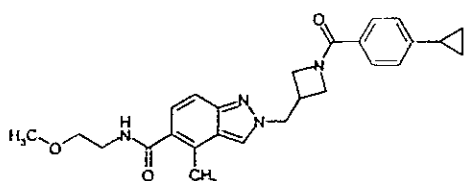
Example 123: 2-([1-[4-chloro-3-(trifluoromethyl)benzoyl]azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 6 mg of the title compound was obtained from 42 mg of the compound prepared in Example 117e and 33 mg of 4-chloro-3-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.21 (1H), 3.25 (3H), 3.31-3.46 (4H), 3.92 (1H), 4.11 (1H), 4.22 (1H), 4.41 (1H), 4.67 (2H), 7.15 (1H), 7.37 (1H), 7.79 (1H), 7.86 (1H), 7.94 (1H), 8.12 (1H), 8.53 (1H).

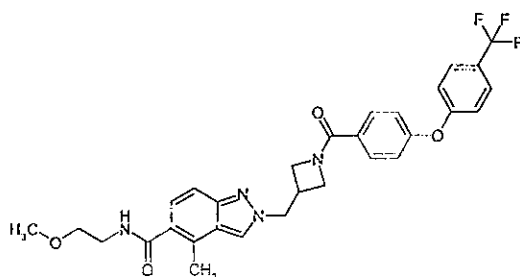
Example 124: 2-[[1-(4-cyclopropylbenzoyl)azetidin-3-yl]methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 76 mg of the title compound was obtained from 163 mg of the compound prepared in Example 117e and 78 mg of 4-cyclopropylbenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.71 (2H), 0.99 (2H), 1.95 (1H), 2.52 (3H), 3.24 (1H), 3.34-3.41 (5H), 3.45 (2H), 3.90 (1H), 4.09 (1H), 4.20 (1H), 4.38 (1H), 4.68 (2H), 7.12 (2H), 7.18 (1H), 7.41 (1H), 7.48 (2H), 8.12 (1H), 8.57 (1H).

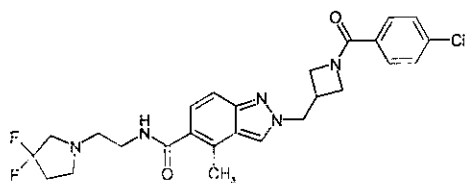
Example 125: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1, 478 mg of the title compound was obtained from 1.20 g of the compound prepared in Example 117e and 850 mg of 4-[4-(trifluoromethyl)phenoxy]benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.52 (3H), 3.25 (1H), 3.27 (3H), 3.39 (2H), 3.45 (2H), 3.94 (1H), 4.12 (1H), 4.24 (1H), 4.42 (1H), 4.70 (2H), 7.14 (2H), 7.18 (1H), 7.22 (2H), 7.41 (1H), 7.69 (2H), 7.77 (2H), 8.12 (1H), 8.57 (1H).

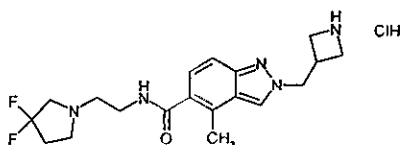
Example 126: 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-N-[2-(3,3-difluoropyrrolidin-1-yl)ethyl]-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 38 mg of the title compound was obtained from 43 mg of the compound prepared in Example 126a and 20 mg of 4-chlorobenzoyl chloride.

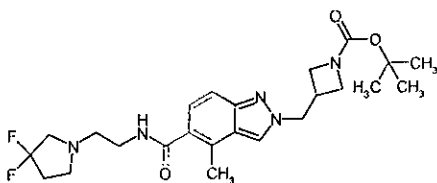
¹H-NMR (300 MHz, DMSO-d₆): δ = 2.10-2.27 (3H), 2.49 (3H), 2.57 (2H), 2.72 (2H), 2.91 (2H), 3.17-3.36 (2H), 3.89 (1H), 4.08 (1H), 4.19 (1H), 4.36 (1H), 4.67 (2H), 7.14 (1H), 7.38 (1H), 7.48 (2H), 7.59 (2H), 8.05 (1H), 8.54 (1H).

Example 126a: 2-(azetidin-3-ylmethyl)-N-[2-(3,3-difluoropyrrolidin-1-yl)ethyl]-4-methyl-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 50 mg of the compound prepared in Example 126b, 43 mg of the title compound was obtained, which was reacted without further purification.

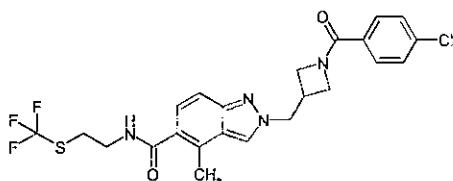
Example 126b: Tert-butyl 3-[(5-{N-[2-(3,3-difluoropyrrolidin-1-yl)ethyl]carbamoyl}-4-methyl-2H-indazol-2-yl)methyl]azetidin-1-carboxylate



Analogously to Example 1b, 71 mg of the title compound was obtained from 70 mg of the compound prepared in Example 117a and 83 mg of 2-(3,3-difluoropyrrolidin-1-yl)ethylamine.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.10-2.28 (2H), 2.50 (3H), 2.57 (2H), 2.66-2.77 (2H), 2.84-2.98 (2H), 3.08 (1H), 3.23-3.36 (2H), 3.69 (2H), 3.86 (2H), 4.60 (2H), 7.14 (1H), 7.39 (1H), 8.05 (1H), 8.55 (1H).

Example 127: 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-4-methyl-N-{2-[(trifluoromethyl)sulphanyl]ethyl}-2H-indazole-5-carboxamide

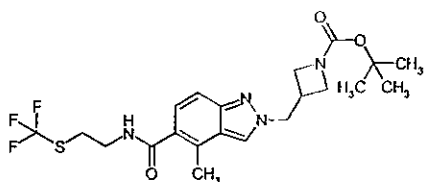


Analogously to Example 55, 143 mg of the title compound was obtained from 130 mg of the compound prepared in Example 127b and 61 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.51 (3H), 3.13-3.29 (3H), 3.50 (2H), 3.90 (1H), 4.09 (1H), 4.19 (1H), 4.37 (1H), 4.67 (2H), 7.19 (1H), 7.40 (1H), 7.48 (2H), 7.59 (2H), 8.38 (1H), 8.57 (1H).

The starting material was prepared as follows:

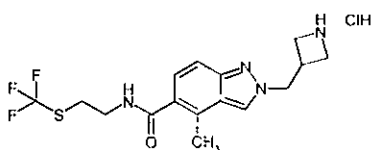
Example 127a: Tert-butyl 3-([4-methyl-5-(N-{2-[(trifluoromethyl)sulphanyl]ethyl}carbamoyl)-indazol-2-yl]methyl)azetidin-1-carboxylate



Analogously to Example 1, 674 mg of the title compound was obtained from 500 mg of the compound prepared in Example 117c.

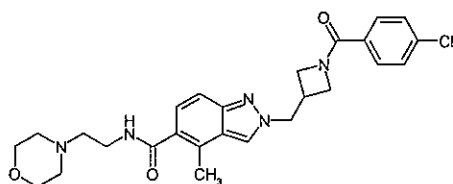
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.52 (3H), 3.08 (1H), 3.17 (2H), 3.50 (2H), 3.69 (2H), 3.86 (2H), 4.61 (2H), 7.19 (1H), 7.41 (1H), 8.38 (1H), 8.57 (1H).

Example 127b: 2-(azetidin-3-ylmethyl)-4-methyl-N-{2-[(trifluoromethyl)sulphanyl]ethyl}-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 150 mg of the compound prepared in Example 127a, 130 mg of the title compound was obtained, which was further reacted without purification.

Example 128: 2-{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-4-methyl-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide

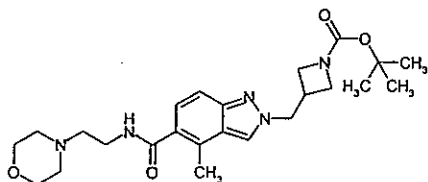


Analogously to Example 55, 30 mg of the title compound was obtained from 50 mg of the compound prepared in Example 128b and 24 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34-2.45 (6H), 2.51 (3H), 3.22 (1H), 3.33 (2H), 3.54 (4H), 3.89 (1H), 4.08 (1H), 4.19 (1H), 4.36 (1H), 4.67 (2H), 7.15 (1H), 7.38 (1H), 7.48 (2H), 7.59 (2H), 7.99 (1H), 8.54 (1H).

The starting material was prepared as follows:

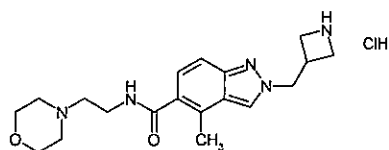
Example 128a: Tert-butyl 3-({4-methyl-5-[N-(2-morpholinoethyl)carbamoyl]-2H-indazol-2-yl}methyl)azetidin-1-carboxylate



Analogously to Example 1b, 59 mg of the title compound was obtained from 200 mg of the compound prepared in Example 117a and 205 mg of 2-morpholinoethylamine.

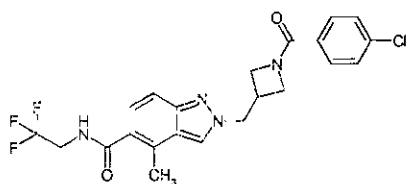
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.34-2.44 (6H), 2.52 (3H), 3.08 (1H), 3.33 (2H), 3.54 (4H), 3.69 (2H), 3.86 (2H), 4.61 (2H), 7.15 (1H), 7.39 (1H), 7.98 (1H), 8.55 (1H).

Example 128b: 2-(azetidin-3-ylmethyl)-4-methyl-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 59 mg of the compound prepared in Example 128a, 63 mg of the title compound was obtained, which was further reacted without purification.

Example 129: 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide

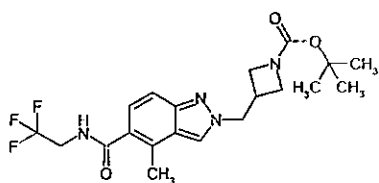


Analogously to Example 55, 127 mg of the title compound was obtained from 128 mg of the compound prepared in Example 129b and 68 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.50 (3H), 3.22 (1H), 3.90 (1H), 3.96-4.13 (3H), 4.19 (1H), 4.37 (1H), 4.68 (2H), 7.18 (1H), 7.42 (1H), 7.48 (2H), 7.59 (2H), 8.59 (1H), 8.79 (1H).

The starting material was prepared as follows:

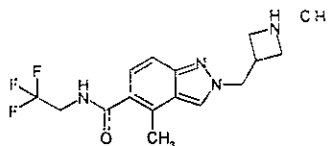
Example 129a: Tert-butyl 3-({4-methyl-5-[N-(2,2,2-trifluoroethyl)carbamoyl]-2H-indazol-2-yl}methyl)azetidin-1-carboxylate



Analogously to Example 1, 1.02 g of the title compound was obtained from 1.0 g of the compound prepared in Example 117c and 287 mg of 2,2,2-trifluoroethylamine.

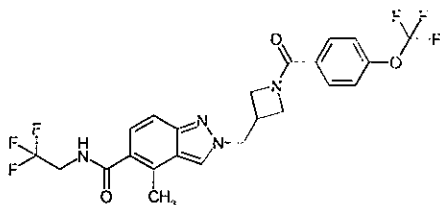
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.51 (3H), 3.09 (1H), 3.70 (2H), 3.86 (2H), 4.03 (2H), 4.62 (2H), 7.17 (1H), 7.43 (1H), 8.60 (1H), 8.79 (1H).

Example 129b: 2-(azetidin-3-ylmethyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 1.0 g of the compound prepared in Example 129a, 852 mg of the title compound was obtained, which was further reacted without purification.

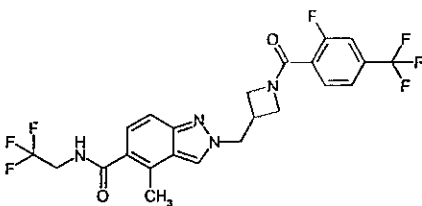
Example 130: 4-methyl-N-(2,2,2-trifluoroethyl)-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 113 mg of the title compound was obtained from 128 mg of the compound prepared in Example 129b and 87 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.50 (3H), 3.23 (1H), 3.92 (1H), 3.96-4.14 (3H), 4.20 (1H), 4.39 (1H), 4.68 (2H), 7.18 (1H), 7.36-7.46 (3H), 7.70 (2H), 8.59 (1H), 8.79 (1H).

Example 131: 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



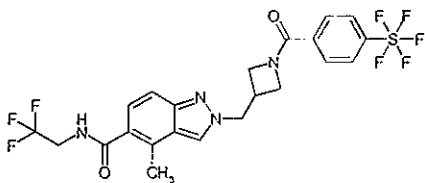
10

Analogously to Example 55, 128 mg of the title compound was obtained from 128 mg of the compound prepared in Example 129b and 88 mg of 2-fluoro-4-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.23 (1H), 3.89-4.16 (6H), 4.67 (2H), 7.17 (1H), 7.42 (1H), 7.60-7.67 (2H), 7.79 (1H), 8.58 (1H), 8.80 (1H).

15

Example 132: 4-methyl-2-({1-[4-(pentafluoro-λ⁶-sulphonyl)benzoyl]azetidin-3-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide

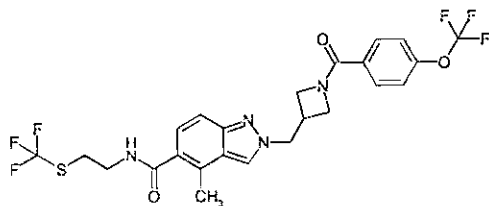


Analogously to Example 1, 121 mg of the title compound was obtained from 128 mg of the compound prepared in Example 129b and 96 mg of 4-(pentafluoro-λ⁶-sulphonyl)benzoic acid.

20

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.50 (3H), 3.24 (1H), 3.94 (1H), 3.98-4.08 (2H), 4.12 (1H), 4.20 (1H), 4.39 (1H), 4.69 (2H), 7.18 (1H), 7.42 (1H), 7.76 (2H), 7.95 (2H), 8.59 (1H), 8.80 (1H).

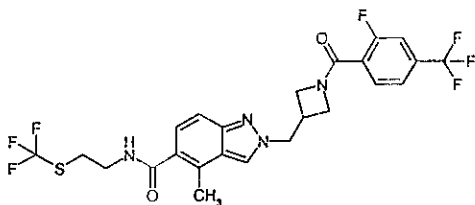
Example 133: 4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-N-{2-[(trifluoromethyl)sulphanyl]ethyl}-2H-indazole-5-carboxamide



Analogously to Example 55, 178 mg of the title compound was obtained from 130 mg of the compound prepared in Example 127b and 78 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.51 (3H), 3.13-3.28 (3H), 3.51 (2H), 3.91 (1H), 4.10 (1H), 4.20 (1H), 4.39 (1H), 4.67 (2H), 7.19 (1H), 7.40 (3H), 7.70 (2H), 8.38 (1H), 8.57 (1H).

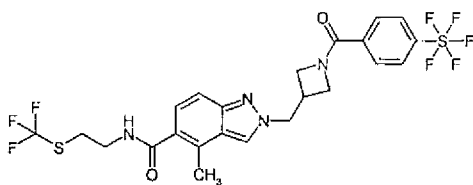
Example 134: 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-4-methyl-N-{2-[(trifluoromethyl)sulphanyl]ethyl}-2H-indazole-5-carboxamide



Analogously to Example 55, 128 mg of the title compound was obtained from 130 mg of the compound prepared in Example 127b and 79 mg of 2-fluoro-4-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.51 (3H), 3.12-3.25 (3H), 3.50 (2H), 3.89-3.98 (2H), 4.06-4.16 (2H), 4.67 (2H), 7.19 (1H), 7.40 (1H), 7.64 (2H), 7.79 (1H), 8.39 (1H), 8.56 (1H).

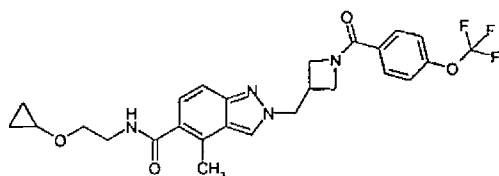
Example 135: 4-methyl-2-({1-[4-(pentafluoro-λ⁶-sulphanyl)benzoyl]azetidin-3-yl}methyl)-N-{2-[(trifluoromethyl)sulphanyl]ethyl}-2H-indazole-5-carboxamide



Analogously to Example 1, 84 mg of the title compound was obtained from 130 mg of the compound prepared in Example 127b and 87 mg of 4-(pentafluoro- λ^6 -sulphonyl)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.51 (3H), 3.18 (2H), 3.23 (1H), 3.51 (2H), 3.94 (1H), 4.12 (1H), 4.20 (1H), 4.39 (1H), 4.68 (2H), 7.19 (1H), 7.41 (1H), 7.76 (2H), 7.95 (2H), 8.39 (1H), 8.56 (1H).

Example 136: N-[2-(cyclopropyloxy)ethyl]-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide

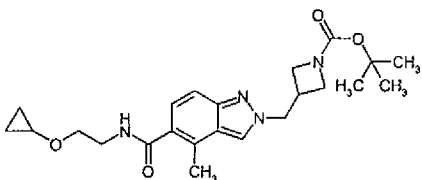


Analogously to Example 55, 172 mg of the title compound was obtained from 161 mg of the compound prepared in Example 136b and 109 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.36-0.48 (4H), 2.48 (3H), 3.22 (1H), 3.27 (1H), 3.35 (2H), 3.53 (2H), 3.91 (1H), 4.10 (1H), 4.20 (1H), 4.39 (1H), 4.67 (2H), 7.14 (1H), 7.35-7.44 (3H), 7.71 (2H), 8.11 (1H), 8.54 (1H).

The starting material was prepared as follows:

Example 136a: Tert-butyl 3-[(5-{N-[2-(cyclopropyloxy)ethyl]carbamoyl}-4-methyl-2H-indazol-2-yl)methyl]azetidin-1-carboxylate

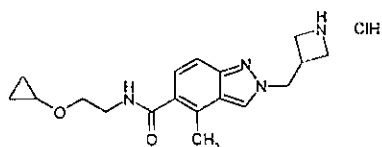


Analogously to Example 1, 189 mg of the title compound was obtained from 150 mg of the compound prepared in Example 118c and 140 mg of 2-cyclopropoxy-ethylammonium trifluoroacetate (preparable according to Example 154a) and 154b) starting from cyclopropanol

and bromoacetamide followed by Boc cleavage with trifluoroacetic acid analogously to Example 1a)).

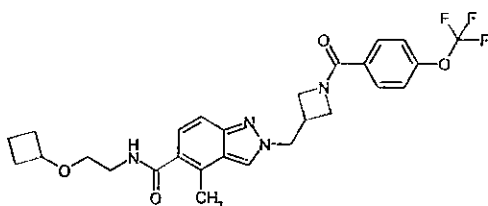
¹H-NMR (300 MHz, DMSO-d₆): δ = 0.36-0.48 (4H), 1.33 (9H), 2.49 (3H), 3.08 (1H), 3.25-3.39 (3H), 3.53 (2H), 3.70 (2H), 3.87 (2H), 4.61 (2H), 7.14 (1H), 7.38 (1H), 8.10 (1H), 8.54 (1H).

- 5 **Example 136b:** 2-(azetidin-3-ylmethyl)-N-[(2-cyclopropyloxy)ethyl]-4-methyl-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 189 mg of the compound prepared in Example 136a, 161 mg of the title compound was obtained, which was further reacted without purification.

- 10 **Example 137:** N-[2-(cyclobutyloxy)ethyl]-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide

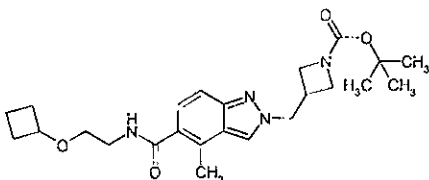


Analogously to Example 55, 207 mg of the title compound was obtained from 254 mg of the compound prepared in Example 137b and 166 mg of 4-(trifluoromethoxy)benzoyl chloride.

- 15 ¹H-NMR (300 MHz, DMSO-d₆): δ = 1.42 (1H), 1.59 (1H), 1.80 (2H), 2.11 (2H), 2.50 (3H), 3.17-3.28 (1H), 3.31-3.41 (4H), 3.85-3.95 (2H), 4.10 (1H), 4.20 (1H), 4.39 (1H), 4.67 (2H), 7.15 (1H), 7.35-7.44 (3H), 7.71 (2H), 8.11 (1H), 8.54 (1H).

The starting material was prepared as follows:

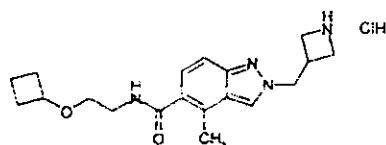
- 20 **Example 137a:** Tert-butyl 3-[(5-{N-[2-(cyclobutyloxy)ethyl]carbamoyl}-4-methyl-2H-indazol-2-yl)methyl]azetidin-1-carboxylate



Analogously to Example 1, 297 mg of the title compound was obtained from 220 mg of the compound prepared in Example 117c and 143 mg of 2-(cyclobutyloxy)ethylammonium trifluoroacetate (preparable according to Example 154a) and 154b) starting from cyclobutanol and bromoacetamide followed by Boc cleavage with trifluoroacetic acid analogously to Example 1a)).

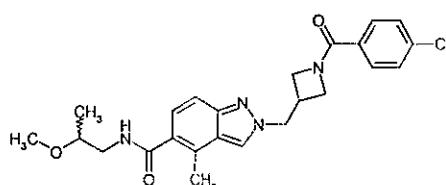
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.33 (9H), 1.35-1.49 (1H), 1.59 (1H), 1.80 (2H), 2.11 (2H), 2.50 (3H), 3.08 (1H), 3.30-3.41 (4H), 3.69 (2H), 3.82-3.95 (3H), 4.61 (2H), 7.15 (1H), 7.38 (1H), 8.10 (1H), 8.54 (1H).

Example 137b: 2-(azetidin-3-ylmethyl)-N-[(2-(cyclobutyloxy)ethyl)-4-methyl-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 297 mg of the compound prepared in Example 137a, 254 mg of the title compound was obtained, which was further reacted without purification.

Example 138: (+/-)-2-[[1-(4-chlorobenzoyl)azetidin-3-yl]methyl]-N-(2-methoxypropyl)-4-methyl-2H-indazole-5-carboxamide

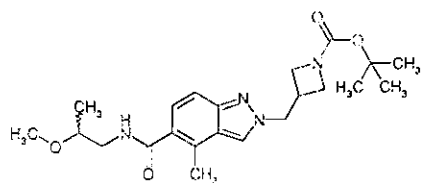


Analogously to Example 55, 166 mg of the title compound was obtained from 237 mg of the compound prepared in Example 138b and 129 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.07 (3H), 2.49 (3H), 3.14-3.28 (3H), 3.43 (1H), 3.89 (1H), 4.08 (1H), 4.19 (1H), 4.36 (1H), 4.67 (2H), 7.15 (1H), 7.38 (1H), 7.48 (2H), 7.59 (2H), 8.11 (1H), 8.54 (1H).

The starting material was prepared as follows:

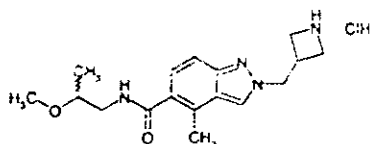
Example 138a: (+/-)-tert-butyl 3-({5-[N-(2-methoxypropyl)carbamoyl]-4-methyl-2H-indazol-2-yl}methyl)azetidin-1-carboxylate



Analogously to Example 1, 331 mg of the title compound was obtained from 323 mg of the compound prepared in Example 117c and 117 mg of 2-methoxypropylamine.

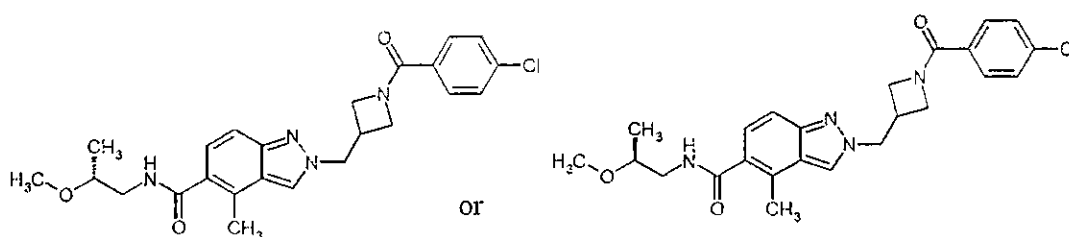
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.08 (3H), 1.33 (9H), 2.50 (3H), 3.09 (1H), 3.16-3.31 (2H), 3.44 (1H), 3.69 (2H), 3.85 (2H), 4.61 (2H), 7.15 (1H), 7.39 (1H), 8.08 (1H), 8.54 (1H).

Example 138b: (+/-)-2-(azetidin-3-ylmethyl)-N-(2-methoxypropyl)-4-methyl-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 280 mg of the compound prepared in Example 138a, 247 mg of the title compound was obtained, which was further reacted without purification.

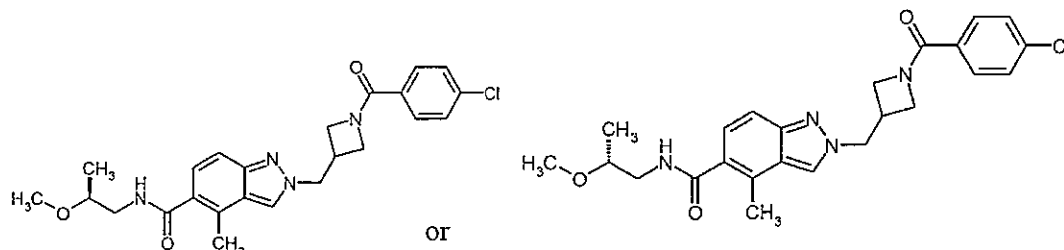
Example 139: (R or S)-2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-N-(2-methoxypropyl)-4-methyl-2H-indazole-5-carboxamide



From 158 mg of the racemate prepared in Example 138, 52 mg of the title compound together with 48 mg of the slower-eluting enantiomer (Example 140) were obtained by racemate separation by means of preparative chiral HPLC (Method A).

Analytical chiral HPLC: 14.5 min.

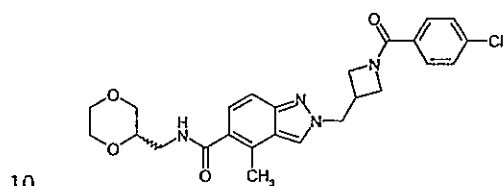
Example 140: (S or R)-2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-N-(2-methoxypropyl)-4-methyl-2H-indazole-5-carboxamide



From 158 mg of the racemate prepared in Example 138, 48 mg of the title compound together
5 with 52 mg of the faster-eluting enantiomer (Example 139) were obtained by racemate separation by means of preparative chiral HPLC (Method A).

Analytical chiral HPLC: 17.1 min.

Example 141: (+/-)-2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-N-(1,4-dioxan-2-ylmethyl)-4-methyl-2H-indazole-5-carboxamide

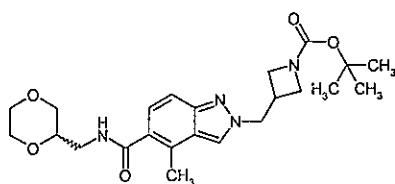


Analogously to Example 55, 177 mg of the title compound was obtained from 269 mg of the
compound prepared in Example 141b and 136 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.13-3.28 (4H), 3.38-3.66 (4H), 3.67-3.77 (2H),
3.89 (1H), 4.08 (1H), 4.19 (1H), 4.37 (1H), 4.67 (2H), 7.15 (1H), 7.38 (1H), 7.48 (2H), 7.59
15 (2H), 8.17 (1H), 8.55 (1H).

The starting material was prepared as follows:

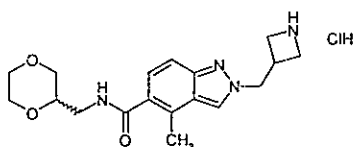
Example 141a: (+/-)-tert-butyl 3-({5-[N-(1,4-dioxan-2-ylmethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}methyl)azetidin-1-carboxylate



Analogously to Example 1, 364 mg of the title compound was obtained from 323 mg of the compound prepared in Example 117c and 110 mg of 1,4-dioxan-2-ylmethylamine.

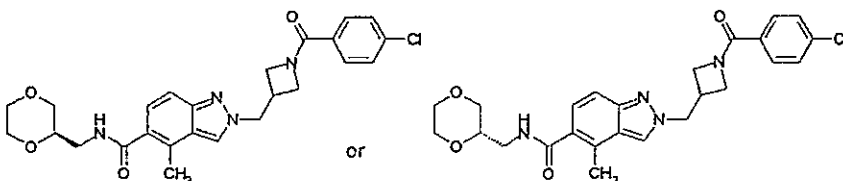
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.33 (9H), 2.50 (3H), 3.09 (1H), 3.16-3.29 (3H), 3.40-3.77 (8H), 3.86 (2H), 4.61 (2H), 7.16 (1H), 7.39 (1H), 8.14 (1H), 8.55 (1H).

- 5 **Example 141b:** (+/-)-2-(azetidin-3-ylmethyl)-N-(1,4-dioxan-2-ylmethyl)-4-methyl-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 314 mg of the compound prepared in Example 141a, 274 mg of the title compound was obtained, which was further reacted without purification.

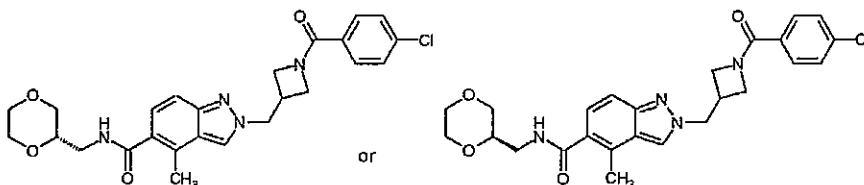
- 10 **Example 142:** (R or S)-2--{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-N-(1,4-dioxan-2-ylmethyl)-4-methyl-2H-indazole-5-carboxamide



- From 172 mg of the racemate prepared in Example 141, 60 mg of the title compound together with 60 mg of the slower-eluting enantiomer (Example 143) were obtained by racemate separation by means of preparative chiral HPLC (Method B).
- 15

Analytical chiral HPLC: 5.84 min.

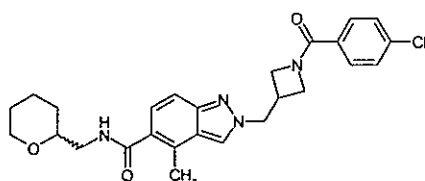
Example 143: (S or R)-2--{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-N-(1,4-dioxan-2-ylmethyl)-4-methyl-2H-indazole-5-carboxamide



From 172 mg of the racemate prepared in Example 141, 60 mg of the title compound together with 60 mg of the faster-eluting enantiomer (Example 142) were obtained by racemate separation by means of preparative chiral HPLC (Method B).

Analytical chiral HPLC: 6.28 min.

- 5 **Example 144:** (+/-)-2-{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-4-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2H-indazole-5-carboxamide

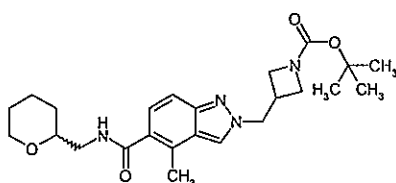


Analogously to Example 55, 144 mg of the title compound was obtained from 255 mg of the compound prepared in Example 144b and 130 mg of 4-chlorobenzoyl chloride.

- 10 $^1\text{H-NMR}$ (300 MHz, DMSO- d_6): δ = 1.10-1.23 (1H), 1.35-1.47 (3H), 1.61 (1H), 1.71-1.81 (1H), 2.48 (3H), 3.15-3.27 (4H), 3.37 (1H), 3.80-3.93 (2H), 4.08 (1H), 4.19 (1H), 4.36 (1H), 4.66 (2H), 7.14 (1H), 7.37 (1H), 7.48 (2H), 7.59 (2H), 8.09 (1H), 8.54 (1H).

The starting material was prepared as follows:

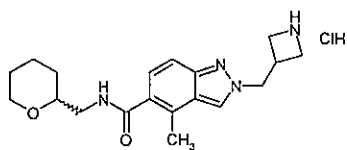
- 15 **Example 144a:** (+/-)-tert-butyl 3-({4-methyl-5-[N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-carbamoyl]-2H-indazol-2-yl}methyl)azetidin-1-carboxylate



Analogously to Example 1, 342 mg of the title compound was obtained from 323 mg of the compound prepared in Example 117c and 141 mg of 3,4,5,6-tetrahydro-2H-pyran-2-ylmethylamine hydrochloride.

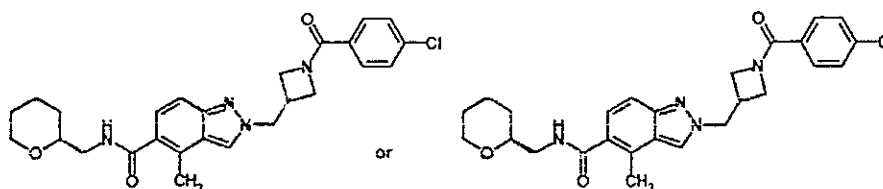
- 20 $^1\text{H-NMR}$ (300 MHz, DMSO- d_6): δ = 1.14-1.21 (1H), 1.33 (9H), 1.37-1.49 (3H), 1.61 (1H), 1.71-1.80 (1H), 2.49 (3H), 3.09 (1H), 3.21 (2H), 3.27-3.43 (2H), 3.69 (2H), 3.81-3.91 (3H), 4.61 (2H), 7.15 (1H), 7.38 (1H), 8.06 (1H), 8.53 (1H).

Example 144b: (+/-)-2-(azetidin-3-ylmethyl)-4-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 298 mg of the compound prepared in Example 144a, 261 mg of the title compound was obtained, which was further reacted without purification.

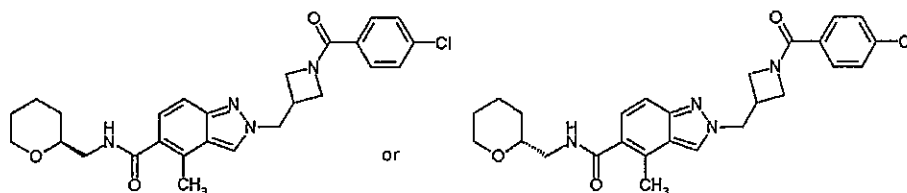
Example 145: (R or S)-2-[[1-(4-chlorobenzoyl)azetidin-3-yl]methyl]-4-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2H-indazole-5-carboxamide



From 140 mg of the racemate prepared in Example 144, 43 mg of the title compound together with 35 mg of the slower-eluting enantiomer (Example 146) were obtained by racemate separation by means of preparative chiral HPLC (Method C).

Analytical chiral HPLC: 4.38 min.

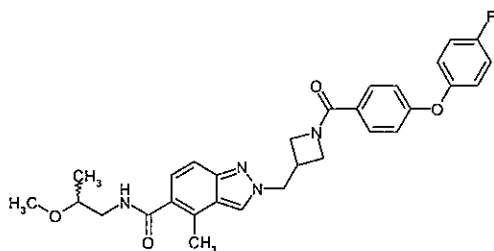
Example 146: (S or R)-2-[[1-(4-chlorobenzoyl)azetidin-3-yl]methyl]-4-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2H-indazole-5-carboxamide



From 140 mg of the racemate prepared in Example 144, 35 mg of the title compound together with 43 mg of the faster-eluting enantiomer (Example 145) were obtained by racemate separation by means of preparative chiral HPLC (Method C).

Analytical chiral HPLC: 4.88 min.

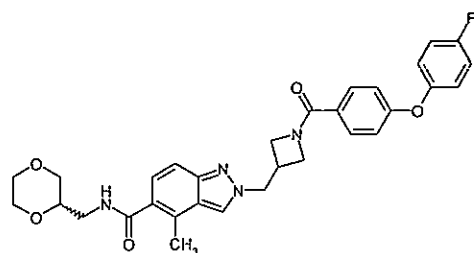
Example 147: (+/-)-2-({1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxypropyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, from 42 mg of the compound prepared in Example 138b and 28 mg of 4-(4-fluorophenoxy)benzoic acid, a material still contaminated after HPLC purification was obtained, which was further purified by an additional preparative thick layer chromatography with ethyl acetate / methanol in the ratio 9:1 as mobile phase. Yield in this manner: 15 mg of the title compound.

¹H-NMR (300 MHz, CDCl₃): δ = 1.22 (3H), 2.64 (3H), 3.28 (1H), 3.36 (3H), 3.40 (1H), 3.57 (1H), 3.76 (1H), 3.97-4.46 (4H), 4.66 (2H), 6.14 (1H), 6.93 (2H), 6.97-7.11 (4H), 7.32 (1H), 7.51 (1H), 7.60 (2H), 8.00 (1H).

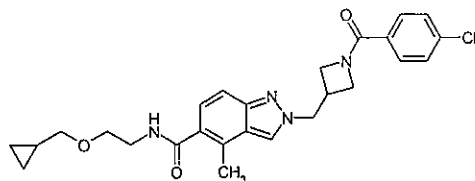
Example 148: (+/-)-N-(1,4-dioxan-2-ylmethyl)-2-({1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl}methyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 18 mg of the title compound was obtained from 43 mg of the compound prepared in Example 141b and 26 mg of 4-(4-fluorophenoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.13-3.28 (4H), 3.38-3.66 (4H), 3.67-3.77 (2H), 3.89 (1H), 4.07 (1H), 4.19 (1H), 4.37 (1H), 4.66 (2H), 6.94 (2H), 7.08-7.18 (3H), 7.24 (2H), 7.38 (1H), 7.60 (2H), 8.17 (1H), 8.55 (1H).

Example 149: 2-({1-[4-(4-chlorobenzoyl)azetidin-3-yl]methyl}-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide

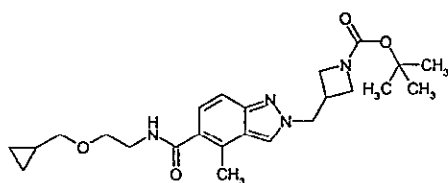


Analogously to Example 55, 72 mg of the title compound was obtained from 127 mg of the compound prepared in Example 149b and 65 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.17 (2H), 0.45 (2H), 0.99 (1H), 2.51-2.54 (3H), 3.20-3.28 (3H), 3.38 (2H), 3.51 (2H), 3.92 (1H), 4.11 (1H), 4.21 (1H), 4.39 (1H), 4.69 (2H), 7.18 (1H), 7.40 (1H), 7.50 (2H), 7.62 (2H), 8.12 (1H), 8.56 (1H).

The starting material was prepared as follows:

Example 149a: Tert-butyl 3-[(5-{N-[2-(cyclopropylmethoxy)ethyl]carbamoyl}-4-methyl-2H-indazol-2-yl)methyl]azetidin-1-carboxylate



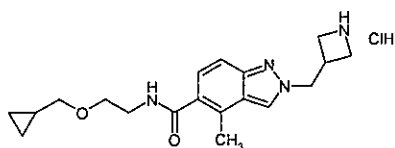
10

Analogously to Example 1, 448 mg of the title compound was obtained from 400 mg of the compound prepared in Example 117c and 133 mg of 2-(cyclopropylmethoxy)ethylamine hydrochloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.14 (2H), 0.42 (2H), 0.97 (1H), 1.33 (9H), 2.50 (3H), 3.09 (1H), 3.24 (2H), 3.36 (2H), 3.49 (2H), 3.69 (2H), 3.86 (2H), 4.61 (2H), 7.15 (1H), 7.38 (1H), 8.08 (1H), 8.54 (1H).

15

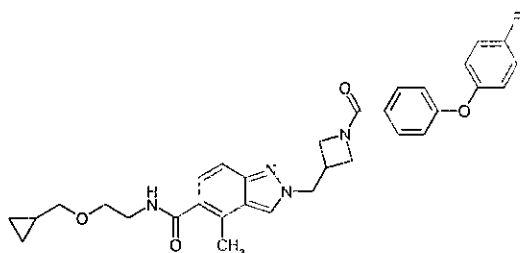
Example 149b: 2-(azetidin-3-ylmethyl)-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide hydrochloride



20

Analogously to Example 1a, from 448 mg of the compound prepared in Example 149a, 390 mg of the title compound was obtained, which was further reacted without purification.

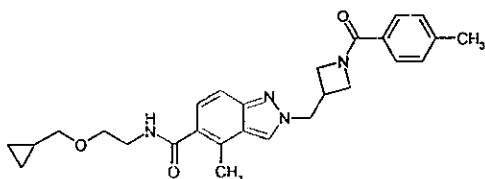
Example 150: N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl}methyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 48 mg of the title compound was obtained from 102 mg of the compound prepared in Example 149b and 63 mg of 4-(4-fluorophenoxy)benzoic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 0.20 (2H), 0.54 (2H), 1.05 (1H), 2.65 (3H), 3.32 (2H), 3.35-3.45 (1H), 3.63-3.75 (4H), 3.99-4.50 (4H), 4.66 (2H), 6.22 (1H), 6.93 (2H), 6.97-7.11 (4H), 7.34 (1H), 7.51 (1H), 7.61 (2H), 8.00 (1H).

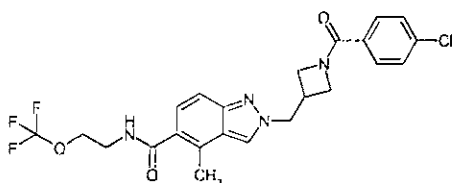
Example 151: N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-{{1-[4-methylbenzoyl]azetidin-3-yl}methyl}-2H-indazole-5-carboxamide



Analogously to Example 1, 55 mg of the title compound was obtained from 127 mg of the compound prepared in Example 149b and 45 mg of 4-methylbenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.17 (2H), 0.45 (2H), 0.99 (1H), 2.33 (3H), 2.53 (3H), 3.19-3.28 (3H), 3.38 (2H), 3.51 (2H), 3.90 (1H), 4.09 (1H), 4.20 (1H), 4.38 (1H), 4.69 (2H), 7.18 (1H), 7.24 (2H), 7.41 (1H), 7.50 (2H), 8.11 (1H), 8.57 (1H).

Example 152: 2-{{1-[4-chlorobenzoyl]azetidin-3-yl}methyl}-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide

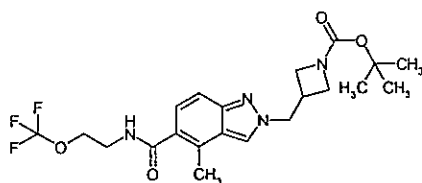


Analogously to Example 55, 122 mg of the title compound was obtained from 234 mg of the compound prepared in Example 152b and 115 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.52 (3H), 3.25 (1H), 3.54 (2H), 3.93 (1H), 4.11 (1H), 4.17-4.25 (3H), 4.40 (1H), 4.70 (2H), 7.19 (1H), 7.43 (1H), 7.50 (2H), 7.61 (2H), 8.38 (1H), 8.58 (1H).

The starting material was prepared as follows:

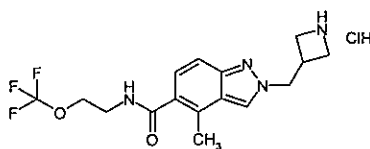
Example 152a: Tert-butyl 3-[(4-methyl-5-{N-[2-(trifluoromethoxy)ethyl]carbamoyl}-2H-indazol-2-yl)methyl]azetidin-1-carboxylate



Analogously to Example 1, 272 mg of the title compound was obtained from 250 mg of the compound prepared in Example 117c and 120 mg of 2-(trifluoromethoxy)ethylamine hydrochloride.

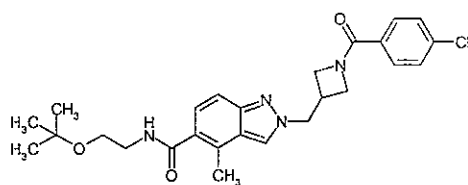
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.50 (3H), 3.08 (1H), 3.51 (2H), 3.70 (2H), 3.86 (2H), 4.17 (2H), 4.61 (2H), 7.16 (1H), 7.41 (1H), 8.36 (1H), 8.56 (1H).

Example 152b: 2-(azetidin-3-ylmethyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 272 mg of the compound prepared in Example 152a, 240 mg of the title compound was obtained, which was further reacted without purification.

Example 153: N-(2-tert-butoxyethyl)-2-[[1-(4-chlorobenzoyl)azetidin-3-yl]methyl]-4-methyl-2H-indazole-5-carboxamide

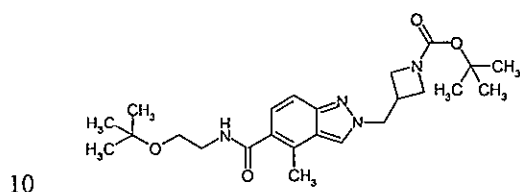


Analogously to Example 55) 12 mg of the title compound was obtained from 160 mg of the compound prepared in Example 153b) and 81 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.14 (9H), 2.53 (3H), 3.19-3.34 (3H), 3.43 (2H), 3.92 (1H), 4.11 (1H), 4.21 (1H), 4.39 (1H), 4.69 (2H), 7.18 (1H), 7.40 (1H), 7.50 (2H), 7.62 (2H), 8.05 (1H), 8.56 (1H).

The starting material was prepared as follows:

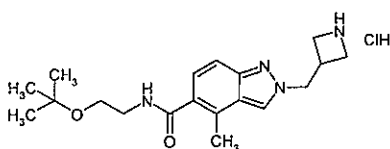
Example 153a: Tert-butyl 3-({5-[N-(2-tert-butoxyethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}-methyl)azetidin-1-carboxylate



Analogously to Example 1, 284 mg of the title compound was obtained from 250 mg of the compound prepared in Example 117c and 111 mg of 2-tert-butoxyethylamine hydrochloride.

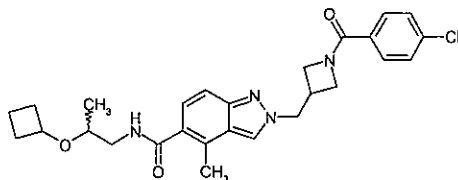
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.11 (9H), 1.32 (9H), 2.50 (3H), 3.08 (1H), 3.27 (2H), 3.40 (2H), 3.69 (2H), 3.86 (2H), 4.60 (2H), 7.15 (1H), 7.38 (1H), 8.04 (1H), 8.54 (1H).

Example 153b: 2-(azetidin-3-ylmethyl)-N-(2-tert-butoxyethyl)-4-methyl-2H-indazol-5-carboxamide hydrochloride



Analogously to Example 1a, from 284 mg of the compound prepared in Example 153a, 240 mg of the title compound was obtained, which was further reacted without purification.

Example 154: (+/-)-2-{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-N-[2-(cyclobutyloxy)propyl]-4-methyl-2H-indazole-5-carboxamide

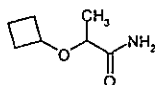


Analogously to Example 55, 135 mg of the title compound was obtained from 222 mg of the compound prepared in Example 154d and 109 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.05 (3H), 1.39 (1H), 1.55 (1H), 1.80 (2H), 2.11 (2H), 2.50 (3H), 3.08-3.23 (3H), 3.55 (1H), 3.89 (1H), 4.03-4.13 (1H), 4.19 (1H), 4.37 (1H), 4.67 (2H), 7.15 (1H), 7.38 (1H), 7.48 (2H), 7.59 (2H), 8.07 (1H), 8.53 (1H).

The starting material was prepared as follows:

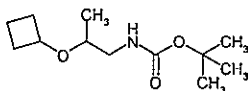
Example 154a: (+/-)-2-(cyclobutyloxy)-propanamide



To a suspension of 732 mg of sodium hydride in 15 ml THF, 2.0 g of cyclobutanol in 0.5 ml THF were added dropwise and stirred for 30 minutes at 25°C. The reaction mixture was then cooled to 0°C and 2.11 g of (+/-) 2-bromopropanamide in 0.5 ml tetrahydrofuran were added dropwise and then stirred for 18 hours at 25°C. The reaction mixture was poured onto ice-water and extracted once with 75 ml dichloromethane. The organic phase was washed with saturated sodium chloride solution, dried over sodium sulphate and concentrated in vacuo after filtration. The crude product thus obtained was purified by column chromatography on silica gel with hexane / 0-50% ethyl acetate. Yield: 0.79 g of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.14 (3H), 1.38 (1H), 1.56 (1H), 1.83 (2H), 2.08 (2H), 3.62 (1H), 3.90 (1H), 7.07 (2H).

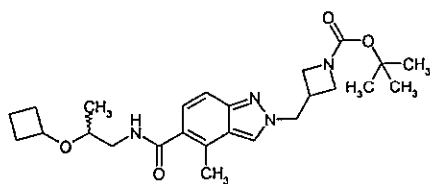
Example 154b: (+/-)-tert-butyl N-[2-(cyclobutyloxy)propyl]carbamate



To a suspension of 6.9 g of lithium aluminium hydride in 250 ml THF, 6.5 g of the amide prepared in Example 154a in 81 ml THF were cautiously added dropwise at 0°C. The mixture was then first stirred for 30 minutes at 0°C and then heated for 5 hours under reflux. After cooling, the mixture was cautiously treated with 20 g of sodium sulphate decahydrate and 20 g of potassium fluoride and solid matter removed by filtration shortly afterwards. The filtrate was concentrated in vacuo. The residue thus obtained (5.8 g) was dissolved without further purification in 174 ml dichloromethane and treated with 11.9 g of di-tert-butyl dicarbonate. After 16 hours' stirring at 25°C, this was concentrated in vacuo and the residue thus obtained purified by two-fold column chromatography on silica gel with hexane / 0-40% ethyl acetate. Yield: 8.54 g of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.94 (3H), 1.28-1.42 (10H), 1.53 (1H), 1.76 (2H), 2.07 (2H), 2.76 (1H), 2.90 (1H), 3.34 (1H), 3.92 (1H), 6.72 (1H).

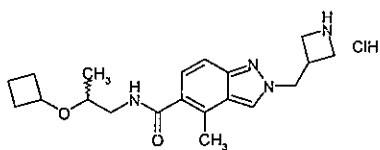
Example 154c: (+/-)-tert-butyl 3-[(5-{N-[2-(cyclobutyloxy)propyl]carbamoyl}-4-methyl-2H-indazol-2-yl)methyl]azetidin-1-carboxylate



Analogously to Example 1a, from 200 mg of the compound prepared in Example 154b), 2-(cyclobutyloxy)propylamine was obtained as the hydrochloride, which without further purification together with 250 mg of the compound prepared in Example 117c analogously to Example 1, yielded 259 mg of the title compound.

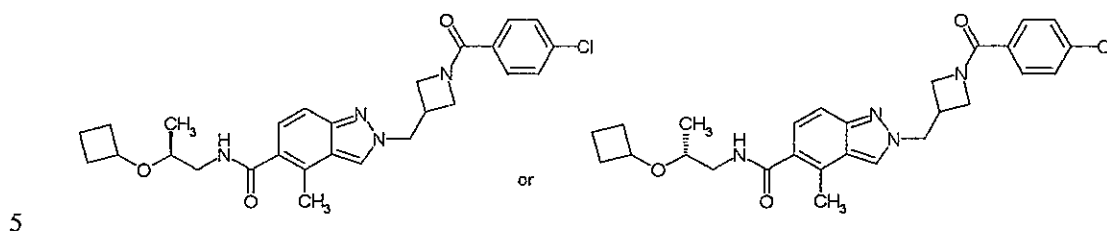
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.05 (3H), 1.32 (9H), 1.39 (1H), 1.55 (1H), 1.80 (2H), 2.11 (2H), 2.50 (3H), 3.02-3.17 (2H), 3.22 (1H), 3.54 (1H), 3.69 (2H), 3.86 (2H), 3.99 (1H), 4.61 (2H), 7.15 (1H), 7.39 (1H), 8.09 (1H), 8.55 (1H).

Example 154d: (+/-)-2-(azetidin-3-ylmethyl)-N-[2-(cyclobutyloxy)propyl]-4-methyl-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 259 mg of the compound prepared in Example 154c, 225 mg of the title compound was obtained, which was further reacted without purification.

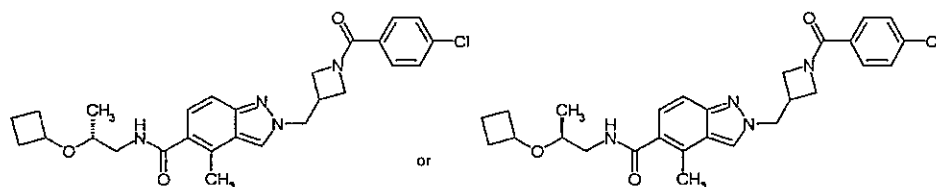
Example 155: (S or R)-2-{{1-(4-chlorobenzoyl)azetidin-3-yl}methyl}-N-[2-(cyclobutyloxy)-propyl]-4-methyl-2H-indazole-5-carboxamide



From 135 mg of the racemate prepared in Example 154, 41 mg of the title compound together with 21 mg of the slower-eluting enantiomer (Example 156) were obtained by racemate separation by means of preparative chiral HPLC (Method A).

Analytical chiral HPLC: 10.05 min.

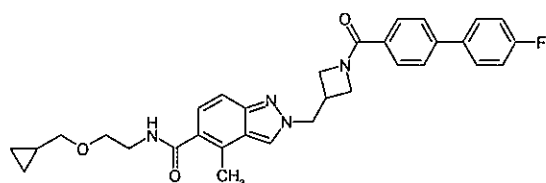
10 **Example 156:** (R or S)-2-{{1-(4-chlorobenzoyl)azetidin-3-yl}methyl}-N-[2-(cyclobutyloxy)-propyl]-4-methyl-2H-indazole-5-carboxamide



15 From 135 mg of the racemate prepared in Example 154, 21 mg of the title compound together with 41 mg of the faster-eluting enantiomer (Example 155) were obtained by racemate separation by means of preparative chiral HPLC (Method A).

Analytical chiral HPLC: 13.14 min.

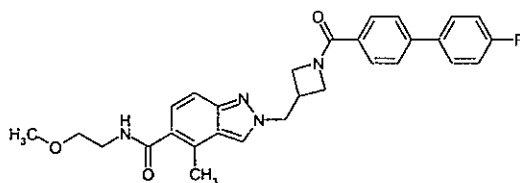
Example 157: N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl}methyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 54 mg of the title compound was obtained from 102 mg of the compound prepared in Example 149b and 58 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 0.20 (2H), 0.54 (2H), 1.05 (1H), 2.66 (3H), 3.32 (2H), 3.42 (1H), 3.61-3.73 (4H), 4.09 (1H), 4.28 (1H), 4.38 (1H), 4.49 (1H), 4.69 (2H), 6.22 (1H), 7.10-7.18 (2H), 7.26 (2H), 7.35 (1H), 7.50-7.59 (3H), 7.70 (2H), 8.01 (1H).

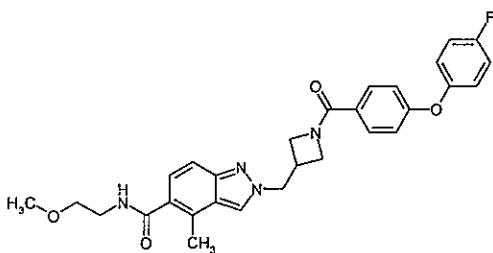
Example 158: 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 54 mg of the title compound was obtained from 108 mg of the compound prepared in Example 117e and 69 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.17-3.29 (4H), 3.36 (2H), 3.43 (2H), 3.92 (1H), 4.11 (1H), 4.24 (1H), 4.42 (1H), 4.68 (2H), 7.15 (1H), 7.29 (2H), 7.39 (1H), 7.63-7.78 (6H), 8.12 (1H), 8.56 (1H).

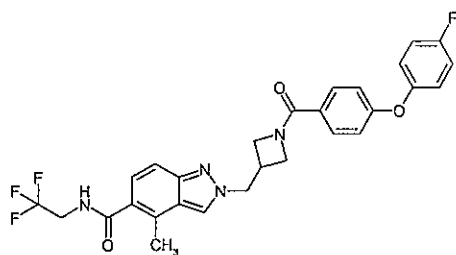
Example 159: 2-({1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 28 mg of the title compound was obtained from 108 mg of the compound prepared in Example 117e and 74 mg of 4-(4-fluorophenoxy)benzoic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 2.65 (3H), 3.31-3.46 (4H), 3.58 (2H), 3.67 (2H), 4.06 (1H), 4.24 (1H), 4.40 (2H), 4.66 (2H), 6.15 (1H), 6.89-7.13 (6H), 7.33 (1H), 7.51 (1H), 7.61 (2H), 8.00 (1H).

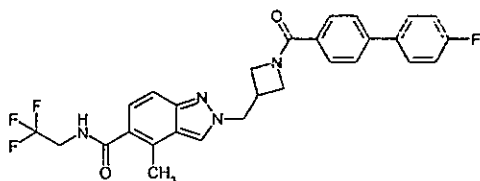
Example 160: 2-({1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 47 mg of the title compound was obtained from 115 mg of the compound prepared in Example 129b and 74 mg of 4-(4-fluorophenoxy)benzoic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 2.65 (3H), 3.39 (1H), 3.97-4.52 (6H), 4.67 (2H), 6.04 (1H),
5 6.89-7.12 (6H), 7.31 (1H), 7.53 (1H), 7.60 (2H), 8.03 (1H).

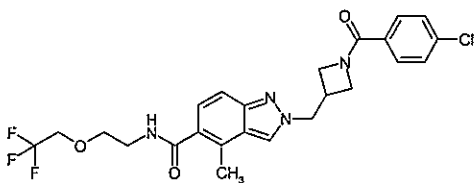
Example 161: 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 28 mg of the title compound was obtained from 115 mg of the
10 compound prepared in Example 129b and 69 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.65 (3H), 3.42 (1H), 4.01-4.54 (6H), 4.69 (2H), 6.02
(1H), 7.15 (2H), 7.32 (1H), 7.51-7.62 (5H), 7.69 (2H), 8.04 (1H).

Example 162: 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide

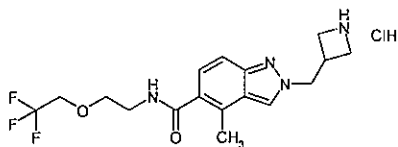


Analogously to Example 55, 197 mg of the title compound was obtained from 252 mg of the
15 compound prepared in Example 162a and 108 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.22 (1H), 3.40 (2H), 3.69 (2H), 3.90 (1H),
4.03-4.13 (3H), 4.19 (1H), 4.37 (1H), 4.67 (2H), 7.15 (1H), 7.38 (1H), 7.48 (2H), 7.59 (2H),
20 8.19 (1H), 8.55 (1H).

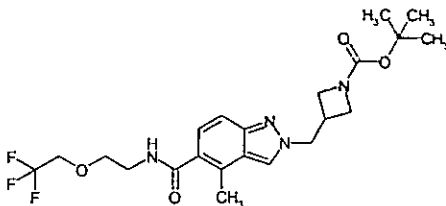
The starting material was prepared as follows:

Example 162a: 2-(azetidin-3-ylmethyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide hydrochloride



- 5 Analogously to Example 1a, from 583 mg of the compound prepared in Example 162b, 505 mg of the title compound was obtained, which was further reacted without purification.

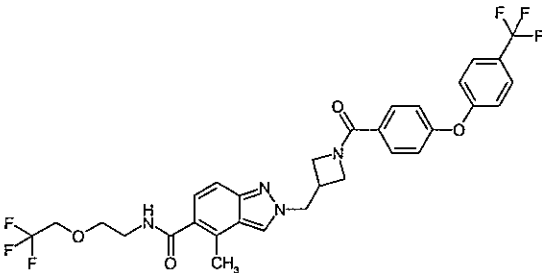
Example 162b: Tert-butyl 3-[(4-methyl-5-{N-[2-(2,2,2-trifluoroethoxy)ethyl]carbamoyl}-2H-indazol-2-yl)methyl]azetidin-1-carboxylate



- 10 Analogously to Example 1, 580 mg of the title compound was obtained from 500 mg of the compound prepared in Example 117c and 260 mg of 2-(2,2,2-trifluoroethoxy)ethylamine hydrochloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.50 (3H), 3.08 (1H), 3.40 (2H), 3.61-3.76 (4H), 3.86 (2H), 4.07 (2H), 4.61 (2H), 7.15 (1H), 7.39 (1H), 8.18 (1H), 8.55 (1H).

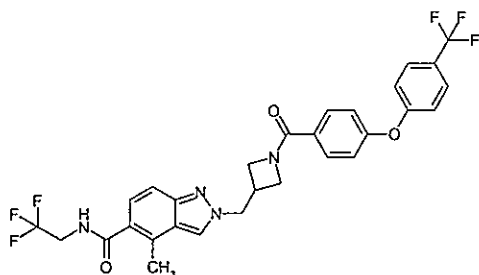
- 15 **Example 163:** 4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide



- Analogously to Example 1, 89 mg of the title compound was obtained from 252 mg of the compound prepared in Example 162a and 175 mg of 4-[4-(trifluoromethyl)phenoxy]benzoic acid.
- 20

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.23 (1H), 3.40 (2H), 3.69 (2H), 3.90 (1H), 4.01-4.14 (3H), 4.21 (1H), 4.40 (1H), 4.68 (2H), 7.11 (2H), 7.15 (1H), 7.20 (2H), 7.39 (1H), 7.66 (2H), 7.75 (2H), 8.19 (1H), 8.56 (1H).

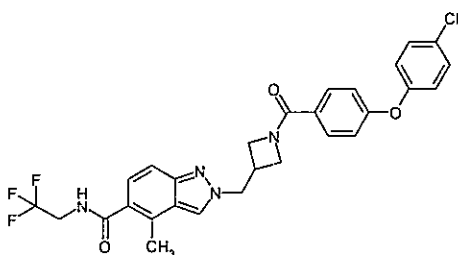
Example 164: 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1, 64 mg of the title compound was obtained from 225 mg of the compound prepared in Example 129b and 149 mg of 4-[4-(trifluoromethyl)phenoxy]benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.50 (3H), 3.24 (1H), 3.91 (1H), 3.95-4.15 (3H), 4.22 (1H), 4.40 (1H), 4.69 (2H), 7.11 (2H), 7.15-7.23 (3H), 7.43 (1H), 7.66 (2H), 7.75 (2H), 8.60 (1H), 8.79 (1H).

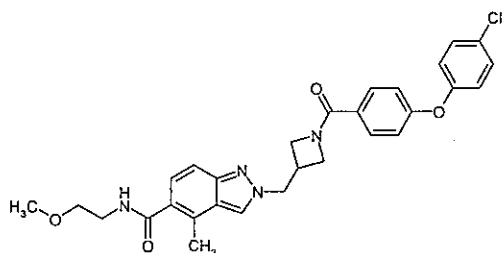
Example 165: 2-({1-[4-(4-chlorophenoxy)benzoyl]azetidin-3-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 97 mg of the title compound was obtained from 225 mg of the compound prepared in Example 129b and 131 mg of 4-(4-chlorophenoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.50 (3H), 3.23 (1H), 3.90 (1H), 3.96-4.12 (3H), 4.20 (1H), 4.38 (1H), 4.68 (2H), 7.00 (2H), 7.09 (2H), 7.18 (1H), 7.40-7.48 (3H), 7.62 (2H), 8.60 (1H), 8.79 (1H)

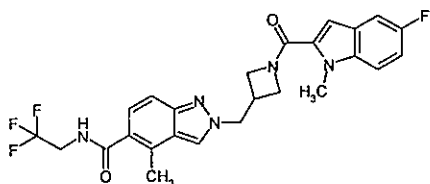
Example 166: 2-({1-[4-(4-chlorophenoxy)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 99 mg of the title compound was obtained from 225 mg of the compound prepared in Example 117e and 140 mg of 4-(4-chlorophenoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.16-3.27 (4H), 3.36 (2H), 3.42 (2H), 3.89 (1H), 4.08 (1H), 4.19 (1H), 4.38 (1H), 4.67 (2H), 7.00 (2H), 7.09 (2H), 7.15 (1H), 7.38 (1H), 7.45 (2H), 7.62 (2H), 8.12 (1H), 8.55 (1H).

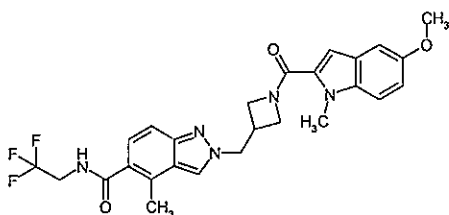
Example 167: 2-({1-[(5-fluoro-1-methyl-1H-indol-2-yl)carbonyl]azetidin-3-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 48 mg of the title compound was obtained from 180 mg of the compound prepared in Example 129b and 82 mg of 5-fluoro-1-methyl-1H-indol-2-carboxylic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 2.66 (3H), 2.80 (3H), 3.36-3.51 (1H), 4.01 (3H), 4.04-4.63 (6H), 4.70 (2H), 6.03 (1H), 6.68 (1H), 7.08 (1H), 7.22-7.36 (3H), 7.55 (1H), 8.05 (1H).

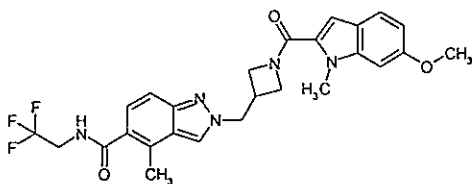
Example 168: 2-({1-[(5-methoxy-1-methyl-1H-indol-2-yl)carbonyl]azetidin-3-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 15 mg of the title compound was obtained from 180 mg of the compound prepared in Example 129b and 87 mg of 5-methoxy-1-methyl-1H-indol-2-carboxylic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 2.66 (3H), 3.43 (1H), 3.84 (3H), 4.00 (3H), 4.03-4.59 (6H),
5 4.70 (2H), 6.02 (1H), 6.66 (1H), 7.02 (1H), 7.27-7.35 (3H), 7.56 (1H), 8.06 (1H).

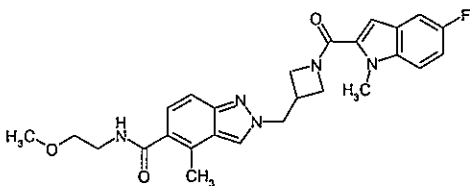
Example 169: 2-({1-[(6-methoxy-1-methyl-1H-indol-2-yl)carbonyl]azetidin-3-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 57 mg of the title compound was obtained from 180 mg of the
10 compound prepared in Example 129b and 87 mg of 6-methoxy-1-methyl-1H-indol-2-carboxylic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 2.66 (3H), 3.43 (1H), 3.89 (3H), 3.99 (3H), 4.05-4.60 (6H),
4.70 (2H), 6.01 (1H), 6.69 (1H), 6.77 (1H), 6.81 (1H), 7.32 (1H), 7.48 (1H), 7.56 (1H), 8.05
(1H).

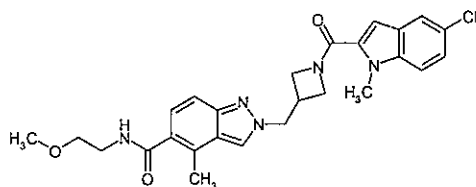
15 **Example 170:** 2-({1-[(5-fluoro-1-methyl-1H-indol-2-yl)carbonyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 95 mg of the title compound was obtained from 180 mg of the
20 compound prepared in Example 117e and 87 mg of 5-fluoro-1-methyl-1H-indol-2-carboxylic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 2.65 (3H), 3.35-3.51 (4H), 3.58 (2H), 3.67 (2H), 4.01 (3H),
4.02-4.62 (4H), 4.69 (2H), 6.15 (1H), 6.68 (1H), 7.08 (1H), 7.21-7.38 (3H), 7.53 (1H), 8.02
(1H).

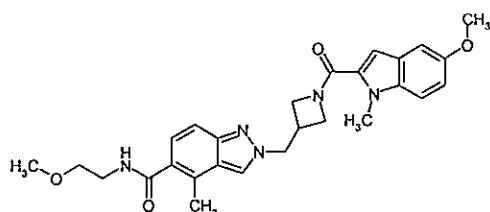
Example 171: 2-(1-[(5-chloro-1-methyl-1H-indol-2-yl)carbonyl]azetidin-3-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 90 mg of the title compound was obtained from 180 mg of the compound prepared in Example 117e and 95 mg of 5-chloro-1-methyl-1H-indol-2-carboxylic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 2.66 (3H), 3.37-3.49 (4H), 3.58 (2H), 3.67 (2H), 4.00 (3H), 4.08 (1H), 4.37 (2H), 4.56 (1H), 4.70 (2H), 6.15 (1H), 6.66 (1H), 7.27-7.38 (3H), 7.53 (1H), 7.58 (1H), 8.02 (1H).

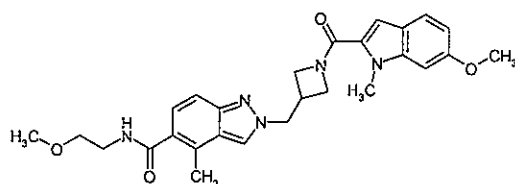
Example 172: N-(2-methoxyethyl)-2-(1-[(5-methoxy-1-methyl-1H-indol-2-yl)carbonyl]azetidin-3-yl)methyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 86 mg of the title compound was obtained from 180 mg of the compound prepared in Example 117e and 93 mg of 5-methoxy-1-methyl-1H-indol-2-carboxylic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 2.65 (3H), 3.36-3.50 (4H), 3.58 (2H), 3.67 (2H), 3.84 (3H), 4.00 (3H), 4.07 (1H), 4.35 (2H), 4.55 (1H), 4.69 (2H), 6.15 (1H), 6.66 (1H), 6.97-7.06 (2H), 7.28 (1H), 7.34 (1H), 7.53 (1H), 8.02 (1H).

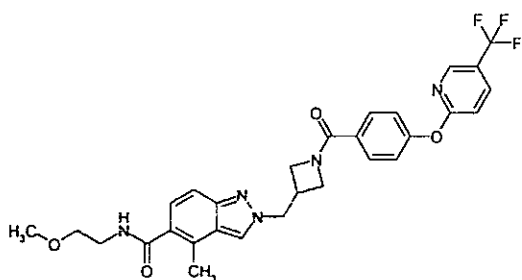
Example 173: N-(2-methoxyethyl)-2-(1-[(6-methoxy-1-methyl-1H-indol-2-yl)carbonyl]azetidin-3-yl)methyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 97 mg of the title compound was obtained from 180 mg of the compound prepared in Example 117e and 93 mg of 6-methoxy-1-methyl-1H-indol-2-carboxylic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 2.66 (3H), 3.37-3.47 (4H), 3.58 (2H), 3.67 (2H), 3.89 (3H),
5 3.99 (3H), 4.02-4.60 (4H), 4.69 (2H), 6.15 (1H), 6.69 (1H), 6.77 (1H), 6.81 (1H), 7.34 (1H),
7.48 (1H), 7.53 (1H), 8.02 (1H).

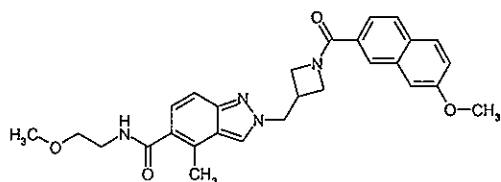
Example 174: N-(2-methoxyethyl)-4-methyl-2-{[1-(4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoyl)azetidin-3-yl]methyl}-2H-indazole-5-carboxamide



10 Analogously to Example 1, 66 mg of the title compound was obtained from 135 mg of the compound prepared in Example 117e and 96 mg of 4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}-benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.50 (3H), 3.18-3.27 (4H), 3.36 (2H), 3.42 (2H), 3.91
(1H), 4.10 (1H), 4.24 (1H), 4.42 (1H), 4.68 (2H), 7.15 (1H), 7.24 (2H), 7.28 (1H), 7.39 (1H),
15 7.67 (2H), 8.12 (1H), 8.24 (1H), 8.56 (2H).

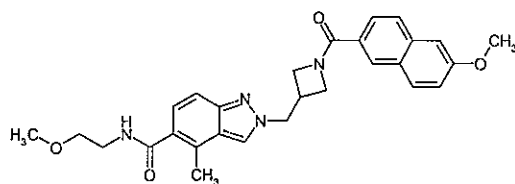
Example 175: N-(2-methoxyethyl)-2-{[1-(7-methoxy-2-naphthoyl)azetidin-3-yl]methyl}-4-methyl-2H-indazole-5-carboxamide



20 Analogously to Example 1, 72 mg of the title compound was obtained from 135 mg of the compound prepared in Example 117e and 69 mg of 7-methoxy-naphthalen-2-carboxylic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.50 (3H), 3.21-3.28 (4H), 3.36 (2H), 3.42 (2H), 3.85
(3H), 3.93-3.99 (1H), 4.14 (1H), 4.26 (1H), 4.47 (1H), 4.69 (2H), 7.15 (1H), 7.22 (1H), 7.38
(1H), 7.43 (1H), 7.50 (1H), 7.81-7.88 (2H), 8.05 (1H), 8.12 (1H), 8.56 (1H).

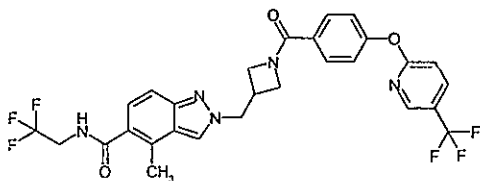
Example 176: N-(2-methoxyethyl)-2-{[1-(6-methoxy-2-naphthoyl)azetidin-3-yl]methyl}-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 72 mg of the title compound was obtained from 135 mg of the compound prepared in Example 117e and 69 mg of 6-methoxynaphthalen-2-carboxylic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.49 (3H), 3.21-3.28 (4H), 3.33-3.39 (2H), 3.42 (2H), 3.86 (3H), 3.91-3.98 (1H), 4.13 (1H), 4.27 (1H), 4.48 (1H), 4.69 (2H), 7.15 (1H), 7.19 (1H), 7.34 (1H), 7.38 (1H), 7.63 (1H), 7.82 (1H), 7.92 (1H), 8.08-8.14 (2H), 8.56 (1H).

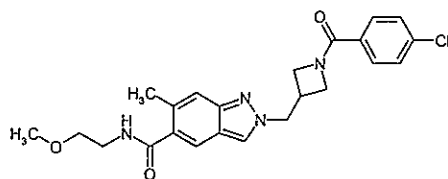
Example 177: 4-methyl-N-(2,2,2-trifluoroethyl)-2-{[1-(4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoyl)azetidin-3-yl]methyl}-2H-indazole-5-carboxamide



Analogously to Example 1, 68 mg of the title compound was obtained from 170 mg of the compound prepared in Example 129b and 113 mg of 4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.35 (3H), 3.21 (1H), 3.89 (1H), 3.96-4.13 (3H), 4.21 (1H), 4.40 (1H), 4.68 (2H), 7.24 (2H), 7.28 (1H), 7.40 (1H), 7.66 (2H), 7.72 (1H), 8.24 (1H), 8.48 (1H), 8.56 (1H), 8.89 (1H).

Example 178: 2-{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide

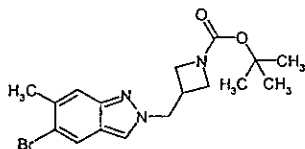


Analogously to Example 55, 154 mg of the title compound was obtained from 210 mg of the compound prepared in Example 178e and 119 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.19 (1H), 3.24 (3H), 3.34 (2H), 3.42 (2H), 3.87 (1H), 4.07 (1H), 4.16 (1H), 4.34 (1H), 4.65 (2H), 7.35 (1H), 7.48 (2H), 7.59 (2H), 7.63 (1H),
5 8.21 (1H), 8.41 (1H).

The starting material was prepared as follows:

Example 178a: Tert-butyl 3-[(5-bromo-6-methyl-2H-indazol-2-yl)methyl]azetidin-1-carboxylate

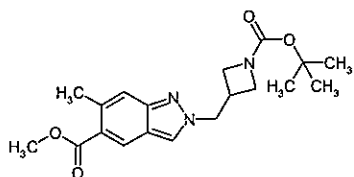


10 To a solution of 6.18 g of 5-bromo-6-methyl-1H-indazole and 15 g of tert-butyl-3-[(tosyloxy)methyl]azetidin-1-carboxylate in 200 ml DMF were added 9.54 g of caesium carbonate and 10.8 g of tetrabutylammonium iodide at 25°C, and this was then heated under reflux for 1.5 hrs. After cooling, the reaction mixture was treated with 1:1 hexane / ether and water, the phases separated and the aqueous phase extracted twice with 250 ml portions of 1:1 hexane / ether. The
15 combined organic phases were washed with saturated sodium chloride solution, dried over sodium sulphate, and concentrated in vacuo after filtration.

The residue thus obtained was purified by twofold column chromatography on silica gel with hexane / 0-20% ethyl acetate. 2.88 g of the title compound was obtained.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.37 (3H), 3.05 (1H), 3.67 (2H), 3.85 (2H), 4.59
20 (2H), 7.56 (1H), 7.97 (1H), 8.33 (1H).

Example 178b: Methyl 2-{[1-(tert-butoxycarbonyl)azetidin-3-yl]methyl}-6-methyl-2H-indazol-5-carboxylate

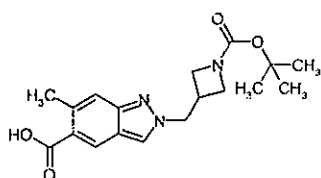


To a solution of 730 mg of the bromide prepared in Example 178a in 17.5 ml tetrahydrofuran
25 were added 0.23 ml of methanol, 145 mg of trans-bis(acetato)-bis[o-(di-o-tolylphosphino)-

benzyl]dipalladium(II), 877 mg of DBU and 760 mg of molybdenum hexacarbonyl. This reaction mixture was then heated in the microwave (120 watts) for 20 minutes at 125°C. Seven further preparations were performed in the same manner and all worked up together. For this, they were concentrated in vacuo and the residue taken up in water and ethyl acetate. After phase separation, the aqueous phase was extracted three times with 75 ml portions of ethyl acetate and the combined organic phases washed with saturated sodium chloride solution, dried over sodium sulphate and concentrated in vacuo after filtration. The residue thus obtained was purified by twofold column chromatography on silica gel with hexane / 0-30% ethyl acetate. Yield: 4.0 g of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.52 (3H), 3.07 (1H), 3.69 (2H), 3.86 (2H), 3.78 (3H), 4.62 (2H), 7.43 (1H), 8.33 (1H), 8.53 (1H).

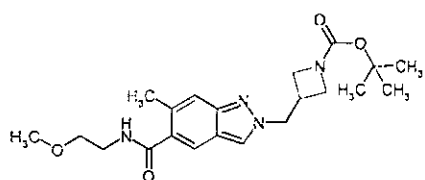
Example 178c: 2-[[1-(tert-butoxycarbonyl)azetidin-3-yl]methyl]-6-methyl-2H-indazol-5-carboxylic acid



Analogously to Example 1d, 3.2 g of the title compound was obtained from 4.0 g of the ester prepared in Example 178b.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.53 (3H), 3.07 (1H), 3.69 (2H), 3.86 (2H), 4.61 (2H), 7.39 (1H), 8.31 (1H), 8.50 (1H), 12.48 (1H).

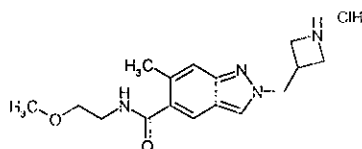
Example 178d: Tert-butyl 3-({5-[N-(2-methoxyethyl)carbamoyl]-6-methyl-2H-indazol-2-yl}-methyl)azetidin-1-carboxylate



Analogously to Example 1, 603 mg of the title compound was obtained from 500 mg of the acid prepared in Example 178c and 109 mg of 2-methoxyethylamine.

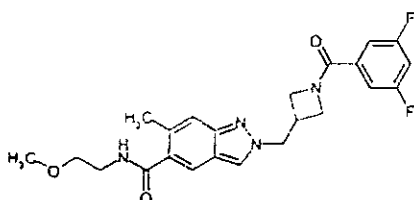
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.34 (3H), 3.06 (1H), 3.25 (3H), 3.32-3.45 (4H), 3.67 (2H), 3.84 (2H), 4.59 (2H), 7.35 (1H), 7.64 (1H), 8.21 (1H), 8.41 (1H).

Example 178e: 2-(azetidin-3-ylmethyl)-N-(2-methoxyethyl)-6-methyl-2H-indazol-5-carboxamide hydrochloride



Analogously to Example 1a, from 250 mg of the amide prepared in Example 178d, 275 mg of the title compound was obtained, which was reacted without further purification.

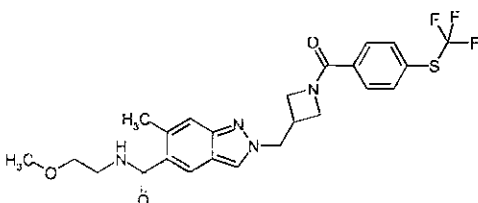
Example 179: 2-{[1-(3,5-Difluorobenzoyl)azetidin-3-yl]methyl}-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 51 mg of the title compound was obtained from 77 mg of the compound prepared in Example 178e and 44 mg of 3,5-difluorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.33 (3H), 3.18 (1H), 3.24 (3H), 3.34 (2H), 3.42 (2H), 3.87 (1H), 4.03-4.11 (1H), 4.20 (1H), 4.38 (1H), 4.65 (2H), 7.20-7.28 (2H), 7.34 (1H), 7.36-7.45 (2H), 7.64 (1H), 8.22 (1H), 8.42 (1H).

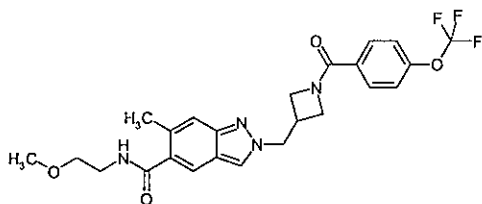
Example 180: N-(2-methoxyethyl)-6-methyl-2-[(1-{4-[(trifluoromethyl)sulphonyl]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 55, 30 mg of the title compound was obtained from 43 mg of the compound prepared in Example 178e and 34 mg of 4-[(trifluoromethyl)sulphonyl]benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.20 (1H), 3.24 (3H), 3.34 (2H), 3.42 (2H), 3.89 (1H), 4.09 (1H), 4.17 (1H), 4.36 (1H), 4.65 (2H), 7.34 (1H), 7.63 (1H), 7.69 (2H), 7.76 (2H), 8.21 (1H), 8.41 (1H).

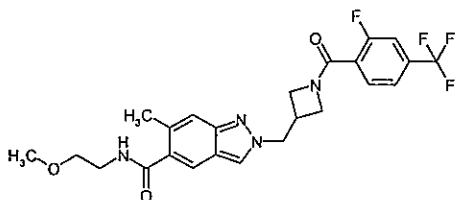
Example 181: N-(2-methoxyethyl)-6-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 221 mg of the title compound was obtained from 294 mg of the compound prepared in Example 178e and 214 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.20 (1H), 3.25 (3H), 3.34 (2H), 3.42 (2H), 3.89 (1H), 4.08 (1H), 4.18 (1H), 4.36 (1H), 4.65 (2H), 7.35 (1H), 7.40 (2H), 7.64 (1H), 7.70 (2H), 8.21 (1H), 8.42 (1H).

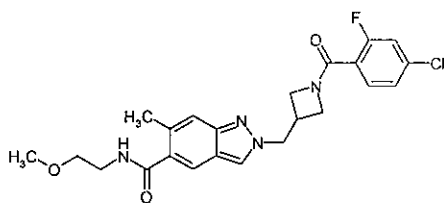
Example 182: 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 11 mg of the title compound was obtained from 43 mg of the compound prepared in Example 178e and 32 mg of 2-fluoro-4-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.33 (3H), 3.20 (1H), 3.24 (3H), 3.35 (2H), 3.42 (2H), 3.86- 3.95 (2H), 4.02-4.16 (2H), 4.65 (2H), 7.34 (1H), 7.61-7.69 (3H), 7.79 (1H), 8.21 (1H), 8.41 (1H).

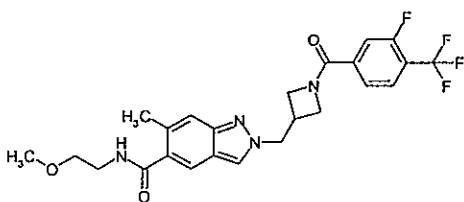
Example 183: 2-([1-(4-chloro-2-fluorobenzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, from 70 mg of the compound prepared in Example 178e and 44 mg of 4-chloro-2-fluorobenzoyl chloride, a material still contaminated after HPLC purification was obtained, which was further purified by additional preparative thick layer chromatography with ethyl acetate / methanol in the ratio 9:1 as mobile phase. This yielded 12 mg of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.19 (1H), 3.25 (3H), 3.34 (2H), 3.42 (2H), 3.84-3.93 (2H), 4.02-4.10 (2H), 4.64 (2H), 7.32-7.36 (2H), 7.46 (1H), 7.53 (1H), 7.63 (1H), 8.22 (1H), 8.41 (1H).

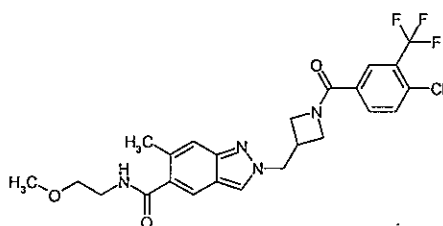
Example 184: 2-({1-[3-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 32 mg of the title compound was obtained from 70 mg of the compound prepared in Example 178e and 52 mg of 3-fluoro-4-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.20 (1H), 3.24 (3H), 3.30-3.38 (2H), 3.42 (2H), 3.90 (1H), 4.10 (1H), 4.19 (1H), 4.38 (1H), 4.66 (2H), 7.34 (1H), 7.57 (1H), 7.60-7.67 (2H), 7.85 (1H), 8.21 (1H), 8.41 (1H).

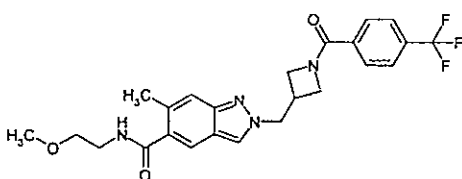
Example 185: 2-({1-[4-chloro-3-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 30 mg of the title compound was obtained from 70 mg of the compound prepared in Example 178e and 55 mg of 4-chloro-3-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.19 (1H), 3.24 (3H), 3.34 (2H), 3.42 (2H), 3.89 (1H), 4.10 (1H), 4.20 (1H), 4.35-4.43 (1H), 4.65 (2H), 7.33 (1H), 7.64 (1H), 7.78 (1H), 7.85 (1H), 7.93 (1H), 8.21 (1H), 8.40 (1H).

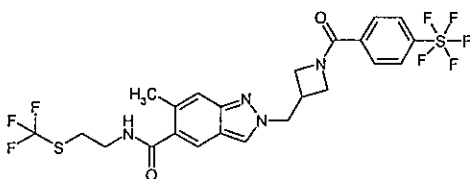
Example 186: N-(2-methoxyethyl)-6-methyl-2-({1-[4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 31 mg of the title compound was obtained from 70 mg of the compound prepared in Example 178e and 47 mg of 4-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.20 (1H), 3.24 (3H), 3.35 (2H), 3.42 (2H), 3.90 (1H), 4.06-4.21 (2H), 4.35 (1H), 4.66 (2H), 7.34 (1H), 7.63 (1H), 7.73-7.82 (4H), 8.21 (1H), 8.41 (1H).

Example 187: 6-methyl-2-({1-[4-(pentafluoro-λ⁶-sulphanyl)benzoyl]azetidin-3-yl}methyl)-N-{2-[(trifluoromethyl)sulphanyl]ethyl}-2H-indazole-5-carboxamide

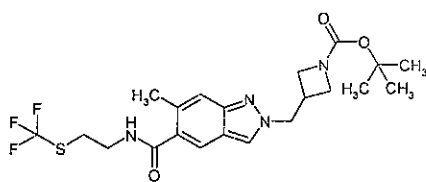


Analogously to Example 1, 35 mg of the title compound was obtained from 170 mg of the compound prepared in Example 187b and 103 mg of 4-(pentafluoro-λ⁶-sulphanyl)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.36 (3H), 3.12-3.23 (3H), 3.49 (2H), 3.91 (1H), 4.06-4.21 (2H), 4.36 (1H), 4.66 (2H), 7.36 (1H), 7.69 (1H), 7.76 (2H), 7.95 (2H), 8.44 (1H), 8.46 (1H).

The starting material was prepared as follows:

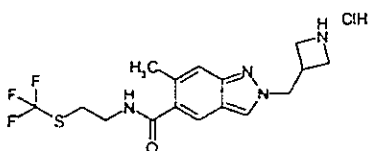
Example 187a: Tert-butyl 3-{{[6-methyl-5-(N-{2-[(trifluoromethyl)sulphanyl]ethyl} carbamoyl)-2H-indazol-2-yl]methyl}azetidin-1-carboxylate



Analogously to Example 1, 998 mg of the title compound was obtained from 800 mg of the acid prepared in Example 178c and 336 mg of 2-[(trifluoromethyl)sulphonyl]ethylamine.

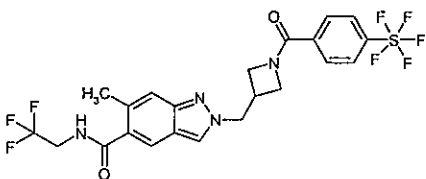
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.36 (3H), 3.06 (1H), 3.13-3.21 (2H), 3.49 (2H), 3.67 (2H), 3.84 (2H), 4.59 (2H), 7.37 (1H), 7.70 (1H), 8.44 (1H), 8.47 (1H).

Example 187b: 2-(azetidin-3-ylmethyl)-6-methyl-N-{2-[(trifluoromethyl)sulphonyl]ethyl}-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a), from 395 mg of the amide prepared in Example 187a), 402 mg of the title compound was obtained, which was reacted without further purification.

Example 188: 6-methyl-2-({1-[4-(pentafluoro-λ⁶-sulphonyl)benzoyl]azetidin-3-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide

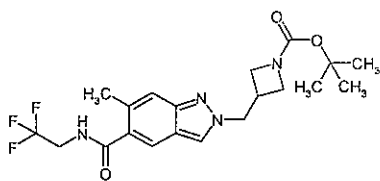


Analogously to Example 1, 25 mg of the title compound was obtained from 136 mg of the compound prepared in Example 188b and 93 mg of 4-(pentafluoro-λ⁶-sulphonyl)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.20 (1H), 3.91 (1H), 3.95-4.21 (4H), 4.36 (1H), 4.67 (2H), 7.38 (1H), 7.71 (1H), 7.76 (2H), 7.95 (2H), 8.45 (1H), 8.86 (1H).

The starting material was prepared as follows:

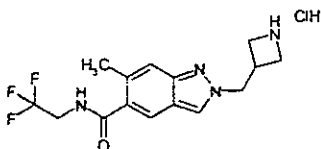
Example 188a: Tert-butyl 3-({6-methyl-5-[N-(2,2,2-trifluoroethyl)carbamoyl]-2H-indazol-2-yl}methyl)azetidin-1-carboxylate



Analogously to Example 1, 550 mg of the title compound was obtained from 560 mg of the acid prepared in Example 178c and 160 mg of 2,2,2-trifluoroethylamine.

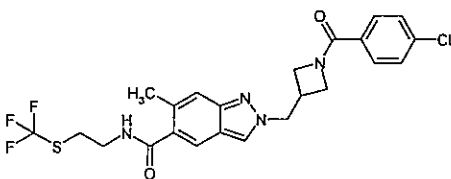
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.34 (3H), 3.06 (1H), 3.67 (2H), 3.84 (2H),
5 3.93-4.10 (2H), 4.60 (2H), 7.40 (1H), 7.71 (1H), 8.46 (1H), 8.88 (1H).

Example 188b: 2-(azetidin-3-ylmethyl)-6-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 550 mg of the amide prepared in Example 188a, 530 mg of
10 the title compound was obtained, which was reacted without further purification.

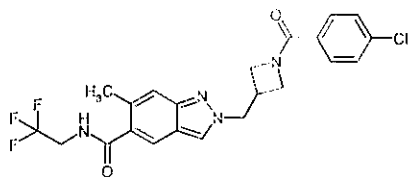
Example 189: 2-[[1-(4-chlorobenzoyl)azetidin-3-yl]methyl]-6-methyl-N-{2-[(trifluoromethyl) sulphanyl]ethyl}-2H-indazole-5-carboxamide



Analogously to Example 55, 45 mg of the title compound was obtained from 170 mg of the
15 compound prepared in Example 187b and 80 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.39 (3H), 3.17-3.27 (3H), 3.52 (2H), 3.90 (1H), 4.10
(1H), 4.19 (1H), 4.37 (1H), 4.68 (2H), 7.39 (1H), 7.50 (2H), 7.61 (2H), 7.72 (1H), 8.47 (1H),
8.49 (1H).

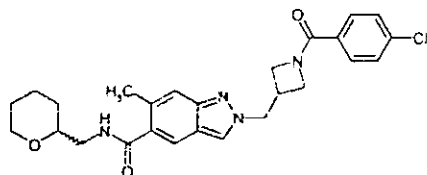
Example 190: 2-[[1-(4-chlorobenzoyl)azetidin-3-yl]methyl]-6-methyl-N-(2,2,2-trifluoroethyl)-
20 2H-indazole-5-carboxamide



Analogously to Example 55, 60 mg of the title compound was obtained from 136 mg of the compound prepared in Example 188b and 72 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.20 (1H), 3.87 (1H), 3.94-4.11 (3H), 4.16 (1H), 4.35 (1H), 4.66 (2H), 7.39 (1H), 7.48 (2H), 7.59 (2H), 7.71 (1H), 8.46 (1H), 8.89 (1H).

Example 191: (+/-)-2-{{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-6-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2H-indazole-5-carboxamide

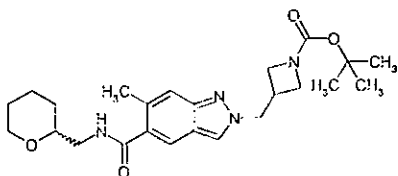


Analogously to Example 55, 49 mg of the title compound was obtained from 120 mg of the compound prepared in Example 191b and 61 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.08-1.23 (1H), 1.36-1.49 (3H), 1.62 (1H), 1.72-1.80 (1H), 2.34 (3H), 3.10-3.43 (5H), 3.80-3.92 (2H), 4.07 (1H), 4.16 (1H), 4.34 (1H), 4.65 (2H), 7.34 (1H), 7.47 (2H), 7.59 (2H), 7.63 (1H), 8.16 (1H), 8.40 (1H).

The starting material was prepared as follows:

Example 191a: (+/-)-tert-butyl 3-({6-methyl-5-[N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-carbamoyl]-2H-indazol-2-yl}methyl)azetidin-1-carboxylate

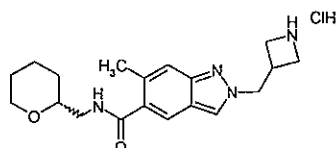


Analogously to Example 1, 140 mg of the title compound was obtained from 300 mg of the acid prepared in Example 178c and 105 mg of 3,4,5,6-tetrahydro-2H-pyran-2-ylmethylamine.

- 160 -

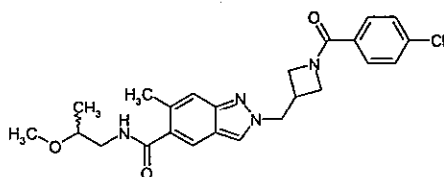
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.07-1.24 (1H), 1.27-1.49 (12H), 1.62 (1H), 1.76 (1H), 2.33 (3H), 3.06 (1H), 3.20 (2H), 3.29-3.45 (2H), 3.67 (2H), 3.78-3.90 (3H), 4.59 (2H), 7.35 (1H), 7.63 (1H), 8.16 (1H), 8.40 (1H).

Example 191b: (+/-)-2-(azetidin-3-ylmethyl)-6-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 140 mg of the amide prepared in Example 191a, 132 mg of the title compound was obtained, which was reacted without further purification.

Example 192: (+/-)-2-[[1-(4-chlorobenzoyl)azetidin-3-yl]methyl]-N-(2-methoxypropyl)-6-methyl-2H-indazole-5-carboxamide

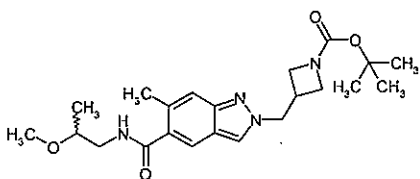


Analogously to Example 55, 89 mg of the title compound was obtained from 151 mg of the compound prepared in Example 192b and 82 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.08 (3H), 2.34 (3H), 3.13-3.32 (6H), 3.43 (1H), 3.87 (1H), 4.07 (1H), 4.16 (1H), 4.34 (1H), 4.65 (2H), 7.35 (1H), 7.48 (2H), 7.59 (2H), 7.63 (1H), 8.18 (1H), 8.41 (1H).

The starting material was prepared as follows:

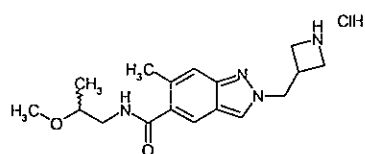
Example 192a: (+/-)-tert-butyl 3-({5-[N-(2-methoxypropyl)carbamoyl]-6-methyl-2H-indazol-2-yl}methyl)azetidin-1-carboxylate



Analogously to Example 1, 358 mg of the title compound was obtained from 300 mg of the acid prepared in Example 178c and 77 mg of 2-methoxypropan-1-amine.

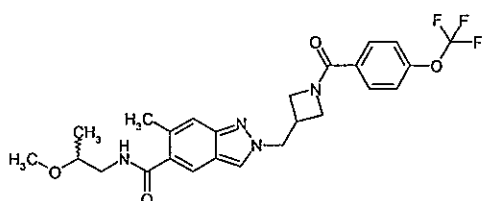
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.08 (3H), 1.32 (9H), 2.34 (3H), 3.06 (1H), 3.13-3.31 (5H), 3.43 (1H), 3.67 (2H), 3.84 (2H), 4.59 (2H), 7.35 (1H), 7.64 (1H), 8.18 (1H), 8.40 (1H).

- 5 **Example 192b:** (+/-)-2-(azetidin-3-ylmethyl)-N-(2-methoxypropyl)-6-methyl-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a), from 358 mg of the amide prepared in Example 192a), 319 mg of the title compound was obtained, which was reacted without further purification.

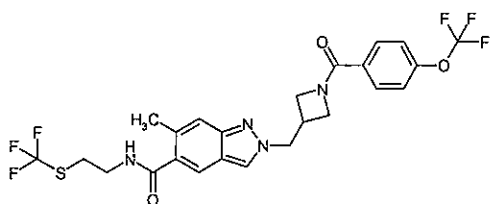
- 10 **Example 193:** (+/-)-N-(2-methoxypropyl)-6-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]-azetidin-3-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 87 mg of the title compound was obtained from 151 mg of the compound prepared in Example 192b and 106 mg of 4-(trifluoromethoxy)benzoyl chloride.

- 15 ¹H-NMR (300 MHz, DMSO-d₆): δ = 1.08 (3H), 2.34 (3H), 3.13-3.31 (6H), 3.43 (1H), 3.88 (1H), 4.08 (1H), 4.17 (1H), 4.36 (1H), 4.65 (2H), 7.35 (1H), 7.40 (2H), 7.64 (1H), 7.70 (2H), 8.18 (1H), 8.41 (1H).

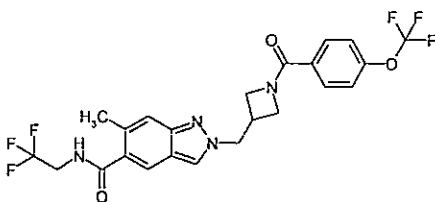
Example 194: 6-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-N-{2-[(trifluoromethyl)sulphanyl]ethyl}-2H-indazole-5-carboxamide



Analogously to Example 55, 87 mg of the title compound was obtained from 173 mg of the compound prepared in Example 187b and 105 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.36 (3H), 3.11-3.24 (3H), 3.49 (2H), 3.89 (1H), 4.09 (1H), 4.17 (1H), 4.36 (1H), 4.66 (2H), 7.36 (1H), 7.40 (2H), 7.66-7.74 (3H), 8.44 (1H), 8.46 (1H).

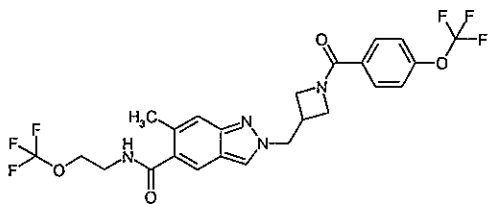
Example 195: 6-methyl-N-(2,2,2-trifluoroethyl)-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 70 mg of the title compound was obtained from 170 mg of the compound prepared in Example 188b and 116 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.35 (3H), 3.21 (1H), 3.89 (1H), 3.95-4.13 (3H), 4.18 (1H), 4.36 (1H), 4.67 (2H), 7.40 (3H), 7.66-7.75 (3H), 8.45 (1H), 8.86 (1H).

Example 196: 6-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide

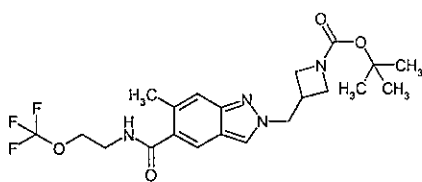


Analogously to Example 55, 22 mg of the title compound was obtained from 179 mg of the compound prepared in Example 196b and 113 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.20 (1H), 3.50 (2H), 3.89 (1H), 4.08 (1H), 4.14-4.21 (3H), 4.36 (1H), 4.66 (2H), 7.36 (1H), 7.40 (2H), 7.66 (1H), 7.70 (2H), 8.44 (1H), 8.46 (1H).

The starting material was prepared as follows:

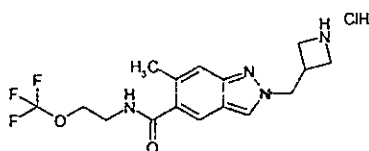
Example 196a: Tert-butyl 3-[(6-methyl-5-{N-[2-(trifluoromethoxy)ethyl]carbamoyl}-2H-indazol-2-yl)methyl]azetidin-1-carboxylate



Analogously to Example 1, 417 mg of the title compound was obtained from 300 mg of the acid prepared in Example 178c and 144 mg of 2-(trifluoromethoxy)ethylamine hydrochloride.

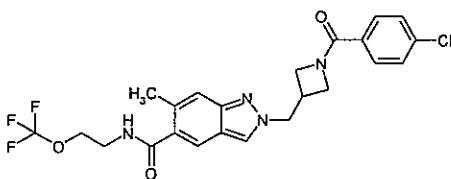
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 2.34 (3H), 3.06 (1H), 3.50 (2H), 3.68 (2H), 3.84 (2H), 4.16 (2H), 4.59 (2H), 7.37 (1H), 7.66 (1H), 8.44 (1H), 8.46 (1H).

Example 196b: 2-(azetidin-3-ylmethyl)-6-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 417 mg of the amide prepared in Example 196a, 369 mg of the title compound was obtained, which was reacted without further purification.

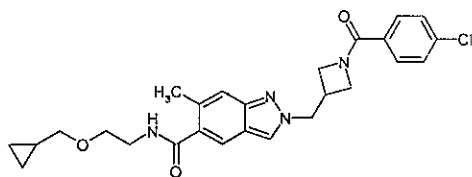
Example 197: 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-6-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide



Analogously to Example 55, 41 mg of the title compound was obtained from 179 mg of the compound prepared in Example 196b and 88 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.19 (1H), 3.50 (2H), 3.87 (1H), 4.07 (1H), 4.13-4.21 (3H), 4.34 (1H), 4.65 (2H), 7.36 (1H), 7.48 (2H), 7.59 (2H), 7.66 (1H), 8.44 (1H), 8.46 (1H).

Example 198: 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-N-[2-(cyclopropylmethoxy)ethyl]-6-methyl-2H-indazole-5-carboxamide

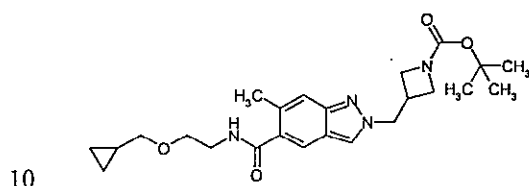


Analogously to Example 55, 107 mg of the title compound was obtained from 214 mg of the compound prepared in Example 198b and 109 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.14 (2H), 0.42 (2H), 0.97 (1H), 2.34 (3H), 3.12-3.26 (3H), 3.34 (2H), 3.47 (2H), 3.87 (1H), 4.07 (1H), 4.16 (1H), 4.34 (1H), 4.65 (2H), 7.35 (1H), 7.48 (2H), 7.59 (2H), 7.63 (1H), 8.21 (1H), 8.42 (1H).

The starting material was prepared as follows:

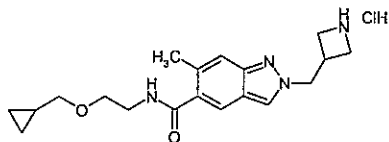
Example 198a: Tert-butyl 3-[(5-{N-[2-(cyclopropylmethoxy)ethyl]carbamoyl}-6-methyl-2H-indazol-2-yl)methyl]azetidin-1-carboxylate



Analogously to Example 1, 247 mg of the title compound was obtained from 245 mg of the acid prepared in Example 178c and 82 mg of 2-(cyclopropylmethoxy)ethylamine hydrochloride.

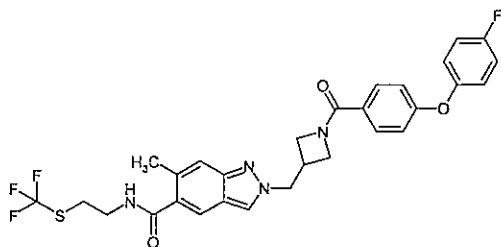
¹H-NMR (300 MHz, DMSO-d₆): δ = 0.14 (2H), 0.43 (2H), 0.97 (1H), 1.32 (9H), 2.35 (3H), 3.06 (1H), 3.23 (2H), 3.34 (2H), 3.48 (2H), 3.67 (2H), 3.85 (2H), 4.59 (2H), 7.35 (1H), 7.64 (1H), 8.18 (1H), 8.41 (1H).

Example 198b: 2-(azetidin-3-ylmethyl)-N-[2-(cyclopropylmethoxy)ethyl]-6-methyl-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 247 mg of the amide prepared in Example 198a, 214 mg of the title compound was obtained, which was reacted without further purification.

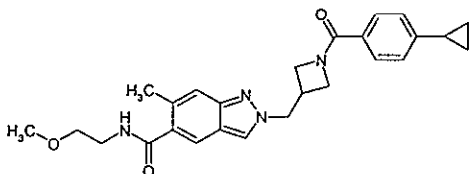
Example 199: 2-({1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl}methyl)-6-methyl-N-{2-[(trifluoromethyl)sulphonyl]ethyl}-2H-indazole-5-carboxamide



Analogously to Example 1, 56 mg of the title compound was obtained from 173 mg of the compound prepared in Example 187b and 98 mg of 4-(4-fluorophenoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.38 (3H), 3.16-3.27 (3H), 3.52 (2H), 3.90 (1H), 4.08 (1H), 4.19 (1H), 4.38 (1H), 4.68 (2H), 6.97 (2H), 7.14 (2H), 7.27 (2H), 7.39 (1H), 7.62 (2H), 7.72 (1H), 8.47 (1H), 8.49 (1H).

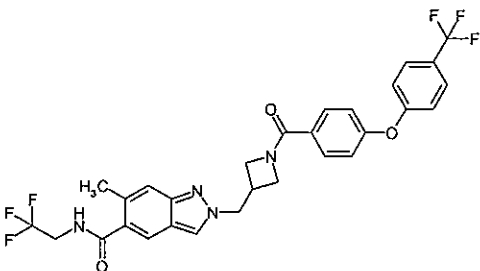
Example 200: 2-{{1-[4-(cyclopropyl)benzoyl]azetidin-3-yl}methyl}-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 86 mg of the title compound was obtained from 168 mg of the compound prepared in Example 178e and 80 mg of 4-cyclopropylbenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.71 (2H), 0.99 (2H), 1.95 (1H), 2.37 (3H), 3.21 (1H), 3.27 (3H), 3.37 (2H), 3.45 (2H), 3.88 (1H), 4.07 (1H), 4.17 (1H), 4.35 (1H), 4.67 (2H), 7.12 (2H), 7.38 (1H), 7.47 (2H), 7.66 (1H), 8.22 (1H), 8.44 (1H).

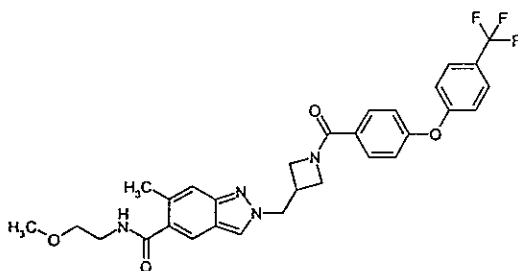
Example 201: 6-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1, 71 mg of the title compound was obtained from 180 mg of the compound prepared in Example 188b and 95 mg of 4-[4-(trifluoromethyl)phenoxy]benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.21 (1H), 3.88 (1H), 3.95-4.13 (3H), 4.19 (1H), 4.37 (1H), 4.67 (2H), 7.11 (2H), 7.19 (2H), 7.39 (1H), 7.65 (2H), 7.69-7.79 (3H), 8.47 (1H), 8.88 (1H).

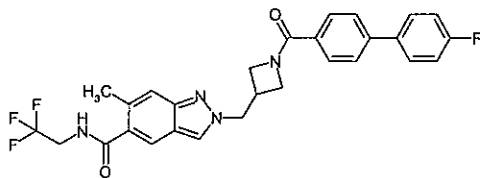
Example 202: N-(2-methoxyethyl)-6-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}-azetidin-3-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1, 108 mg of the title compound was obtained from 180 mg of the compound prepared in Example 178e and 102 mg of 4-[4-(trifluoromethyl)phenoxy]benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.14-3.26 (4H), 3.34 (2H), 3.42 (2H), 3.88 (1H), 4.07 (1H), 4.19 (1H), 4.37 (1H), 4.66 (2H), 7.11 (2H), 7.19 (2H), 7.35 (1H), 7.61-7.68 (3H), 7.75 (2H), 8.21 (1H), 8.42 (1H).

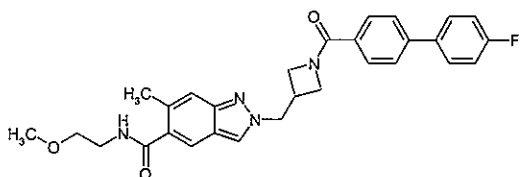
Example 203: 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl}methyl)-6-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 104 mg of the title compound was obtained from 180 mg of the compound prepared in Example 188b and 73 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.22 (1H), 3.90 (1H), 3.95-4.14 (3H), 4.21 (1H), 4.40 (1H), 4.68 (2H), 7.28 (2H), 7.40 (1H), 7.60-7.77 (7H), 8.48 (1H), 8.89 (1H).

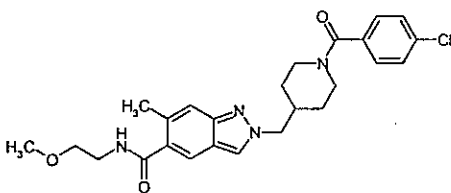
Example 204: 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 73 mg of the title compound was obtained from 180 mg of the compound prepared in Example 178e and 78 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.34 (3H), 3.14-3.26 (4H), 3.34 (2H), 3.41 (2H), 3.89 (1H), 4.09 (1H), 4.21 (1H), 4.39 (1H), 4.66 (2H), 7.28 (2H), 7.35 (1H), 7.61-7.77 (7H), 8.21 (1H), 8.43 (1H).

Example 205: 2-({1-(4-chlorobenzoyl)piperidin-4-yl}methyl)-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide

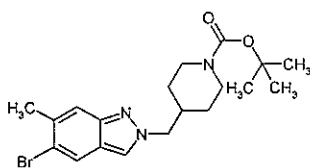


Analogously to Example 55, from 85 mg of the compound prepared in Example 205c and 45 mg of 4-chlorobenzoyl chloride, a material still contaminated after column chromatography was obtained which was further purified by additional preparative thick layer chromatography with dichloromethane / methanol in the ratio 95:5 as mobile phase. Yield: 48 mg of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.09-1.53 (4H), 2.24 (1H), 2.34 (3H), 2.71 (1H), 2.97 (1H), 3.24 (3H), 3.34 (2H), 3.38-3.54 (3H), 7.31-7.37 (3H), 7.46 (2H), 7.63 (1H), 8.21 (1H), 8.33 (H).

The starting material was prepared as follows:

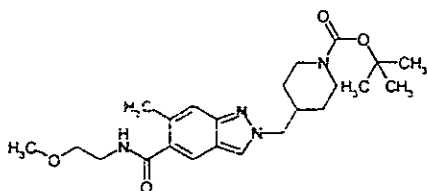
Example 205a: Tert-butyl 4-[(5-bromo-6-methyl-2H-indazol-2-yl)methyl]piperidin-1-carboxylate



Analogously to Example 1c, 2.99 g of the title compound was obtained from 5.7 g of 5-bromo-6-methyl-1H-indazole and 15.0 g of tert-butyl-4-[(tosyloxy)methyl]piperidin-1-carboxylate.

¹H-NMR (300 MHz, CDCl₃): δ = 1.14-1.30 (2H), 1.44 (9H), 1.53 (2H), 2.24 (1H), 2.50 (3H), 2.66 (2H), 4.11 (2H), 4.23 (2H), 7.55 (1H), 7.77 (1H), 7.87 (1H).

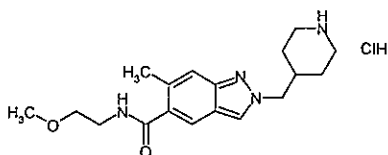
Example 205b: Tert-butyl 4-({5-[N-(2-methoxyethyl)carbamoyl]-6-methyl-2H-indazol-2-yl}-methyl)piperidin-1-carboxylate



Analogously to Example 1b, 3.56 g of the title compound was obtained from 600 mg of the bromide prepared in Example 205a and 331 mg of 2-methoxyethylamine after four runs and twofold purification by chromatography.

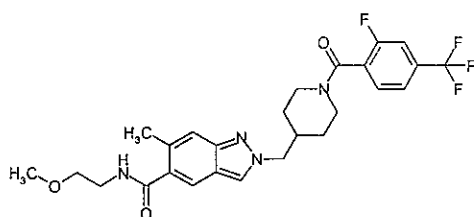
¹H-NMR (300 MHz, DMSO-d₆): δ = 0.96-1.15 (2H), 1.29-1.42 (10H), 2.03-2.18 (1H), 2.34 (3H), 2.51-2.73 (3H), 3.25 (3H), 3.31-3.45 (4H), 3.86 (2H), 4.26 (2H), 7.35 (1H), 7.63 (1H), 8.21 (1H), 8.32 (1H).

Example 205c: N-(2-methoxyethyl)-6-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 100 mg of the amide prepared in Example 205b, 85 mg of the title compound was obtained, which was reacted without further purification.

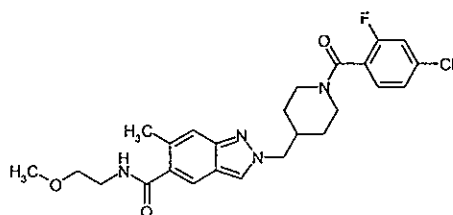
Example 206: 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxy-ethyl)-6-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, from 85 mg of the compound prepared in Example 205c and 58 mg of 2-fluoro-4-(trifluoromethyl)benzoyl chloride, a material still contaminated after column chromatography was obtained which was further purified by additional preparative thick layer chromatography with dichloromethane / methanol in the ratio 95:5 as mobile phase. Yield: 40 mg of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.03-1.28 (2H), 1.37 (1H), 1.53 (1H), 2.23 (1H), 2.34 (3H), 2.77 (1H), 3.00 (1H), 3.26-3.49 (11H), 4.30 (2H), 4.44 (1H), 7.35 (1H), 7.54-7.67 (3H), 7.78 (1H), 8.21 (1H), 8.34 (1H).

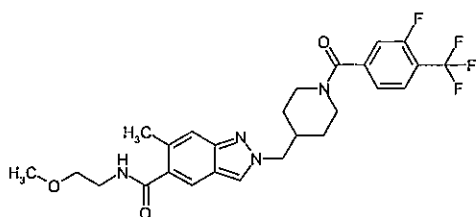
Example 207: 2-({1-[4-chloro-2-fluorobenzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 68 mg of the title compound was obtained from 85 mg of the compound prepared in Example 205c and 49 mg of 4-chloro-2-fluorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.04-1.25 (2H), 1.37 (1H), 1.51 (1H), 2.23 (1H), 2.34 (3H), 2.73 (1H), 2.98 (1H), 3.21-3.48 (3H), 3.30-3.38 (3H), 3.42 (2H), 4.29 (2H), 4.42 (1H), 7.29-7.42 (3H), 7.51 (1H), 7.63 (1H), 8.21 (1H), 8.33 (1H).

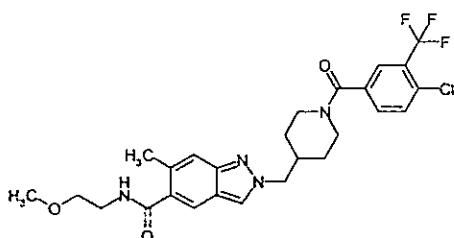
Example 208: 2-({1-[3-fluoro-4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 70 mg of the title compound was obtained from 85 mg of the compound prepared in Example 205c and 58 mg of 3-fluoro-4-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.13-1.27 (2H), 1.35 (1H), 1.51 (1H), 2.23 (1H), 2.34 (3H), 2.74 (1H), 2.98 (1H), 3.22-3.45 (8H), 4.29 (2H), 4.40 (1H), 7.32-7.38 (2H), 7.52 (1H), 7.63 (1H), 7.82 (1H), 8.21 (1H), 8.33 (1H).

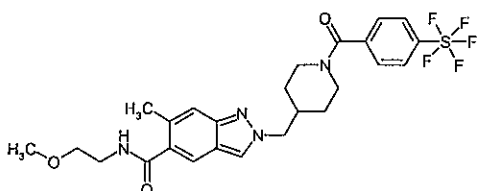
Example 209: 2-({1-[4-chloro-3-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-6-methyl-2H-indazole-5-carboxamide



Analogously to Example 55, 92 mg of the title compound was obtained from 85 mg of the compound prepared in Example 205c and 62 mg of 4-chloro-3-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.14-1.28 (2H), 1.35 (1H), 1.44-1.53 (1H), 2.25 (1H), 2.34 (3H), 2.74 (1H), 3.01 (1H), 3.24 (3H), 3.34 (2H), 3.39-3.50 (3H), 4.29 (2H), 4.39 (1H), 7.35 (1H), 7.61-7.68 (2H), 7.74-7.80 (2H), 8.21 (1H), 8.33 (1H).

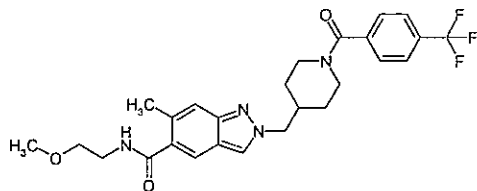
Example 210: N-(2-methoxyethyl)-6-methyl-2-({1-[4-(pentafluoro-λ⁶-sulphanyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 126 mg of the title compound was obtained from 85 mg of the compound prepared in Example 205c and 58 mg of 4-(pentafluoro-λ⁶-sulphanyl)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.12-1.27 (2H), 1.35 (1H), 1.51 (1H), 2.24 (1H), 2.34 (3H), 2.74 (1H), 2.99 (1H), 3.24 (3H), 3.34-3.46 (5H), 4.30 (2H), 4.41 (1H), 7.35 (1H), 7.55 (2H), 7.63 (1H), 7.94 (2H), 8.21 (1H), 8.33 (1H).

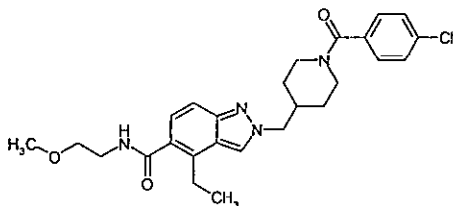
Example 211: N-(2-methoxyethyl)-6-methyl-2-({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 69 mg of the title compound was obtained from 85 mg of the compound prepared in Example 205c and 53 mg of 4-(trifluoromethyl)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.10-1.26 (2H), 1.35 (1H), 1.51 (1H), 2.19-2.30 (1H), 2.34 (3H), 2.74 (1H), 2.98 (1H), 3.21-3.47 (8H), 4.30 (2H), 4.42 (1H), 7.35 (1H), 7.54 (2H), 7.63 (1H), 7.77 (2H), 8.21 (1H), 8.33 (1H).

Example 212: 2-({1-[4-chlorobenzoyl]piperidin-4-yl}methyl)-4-ethyl-N-(2-methoxyethyl)-2H-indazole-5-carboxamide

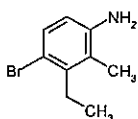


Analogously to Example 55, 61 mg of the title compound was obtained from 91 mg of the compound prepared in Example 212i and 46 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.14-1.28 (5H), 1.31-1.60 (2H), 2.27 (1H), 2.63-3.04 (4H), 3.24 (3H), 3.31-3.56 (5H), 4.31 (2H), 4.39 (1H), 7.10 (1H), 7.32-7.41 (3H), 7.46 (2H), 8.11 (1H), 8.50 (1H).

The starting material was prepared as follows:

Example 212a: 4-bromo-3-ethyl-2-methylaniline

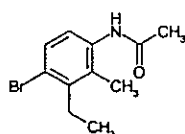


A suspension of 3 g of 3-ethyl-2-methylaniline hydrochloride in 200 ml ethyl acetate was washed twice with 30 ml portions of saturated sodium carbonate solution and twice with 20 ml

portions of saturated sodium chloride solution, dried over sodium sulphate and concentrated in vacuo after filtration. This yielded 2.69 g of ethyl-2-methylaniline, which was further reacted without purification.

To a solution of 2.36 g of the amine prepared above in 29 ml DMF, a solution of 3.1 g of N-bromosuccinimide in 14.5 ml DMF was added dropwise at 0°C and stirred for 30 minutes at 0°C. Then the reaction mixture was diluted with 400 ml ethyl acetate and washed once with 30 ml of a 10% aqueous sodium carbonate solution and once with 30 ml water. After drying over sodium sulphate and filtration, this was concentrated in vacuo. The crude product thus obtained (3.86 g) was used in the next step without purification.

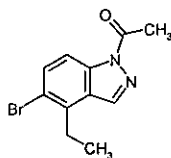
10 **Example 212b:** N-(4-bromo-3-ethyl-2-methylphenyl)acetamide



To a solution of 3.86 g of the bromide prepared in Example 212a in 48 ml pyridine, 2.04 ml of acetic anhydride was added dropwise at 0°C and stirred for 20 hours at 25°C. The reaction mixture was concentrated in vacuo and the crude product thus obtained was purified by column chromatography on silica gel with hexane / 0-100% ethyl acetate. In this manner, 3.73 g of the title compound was obtained.

¹H-NMR (300 MHz, CDCl₃): δ = 1.13 (3H), 2.20 (3H), 2.25 (3H), 2.86 (2H), 6.91 (1H), 7.33 (1H), 7.40 (1H).

Example 212c: 1-(5-bromo-4-ethyl-1H-indazol-1-yl)ethan-1-one



20

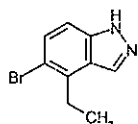
To a solution of 3.72 g of the acetamide prepared in Example 212b in 31.5 ml chloroform were added 4.1 ml of acetic anhydride and 2.85 g of potassium acetate, and 192 mg of 18-crown-6 and 3.4 g of isopentyl nitrite were then added dropwise. The reaction mixture was heated for 20 hours under reflux and after cooling diluted with 200 ml ethyl acetate. The organic phase was washed with 20 ml sodium carbonate solution and once with saturated sodium chloride solution. After drying over sodium sulphate and filtration, this was concentrated in vacuo and the crude

25

product thus obtained purified by column chromatography on silica gel with hexane / 0-50% ethyl acetate. Yield: 3.78 g of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.18 (3H), 2.68 (3H), 3.04 (2H), 7.73 (1H), 8.03 (1H), 8.64 (1H).

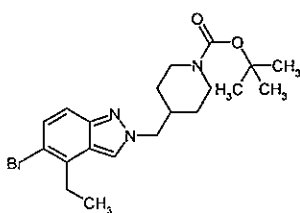
5 **Example 212d:** 5-bromo-4-ethyl-1H-indazole



A mixture of 3.78 g of the compound prepared in Example 212c in 7.3 ml methanol and 26.3 ml of 37% hydrochloric acid was heated under reflux for 2 hours. This was then diluted with 400 ml ethyl acetate. The organic phase was washed three times with 50 ml portions of sodium
10 hydrogen carbonate solution and once with 50 ml saturated sodium chloride solution, dried over sodium sulphate and concentrated in vacuo. The crude product thus obtained was purified by column chromatography on silica gel with hexane / 0-50% ethyl acetate. Yield: 2.97 g of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.17 (3H), 2.98 (2H), 7.30 (1H), 7.41 (1H), 8.16 (1H),
15 13.17 (1H).

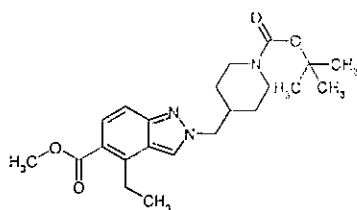
Example 212e: Tert-butyl 4-[(5-bromo-4-ethyl-2H-indazol-2-yl)methyl]piperidin-1-carboxylate



Analogously to Example 1c, 961 mg of the title compound was obtained from 2.97 g of the indazole prepared in Example 212d and 7.3 g of tert-butyl-4-[(tosyloxy)methyl]piperidin-1-
20 carboxylate.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.01-1.13 (2H), 1.16 (3H), 1.34 (9H), 1.41 (2H), 2.12 (1H), 2.63 (2H), 2.90 (2H), 3.87 (2H), 4.27 (2H), 7.27 (1H), 7.35 (1H), 8.49 (1H).

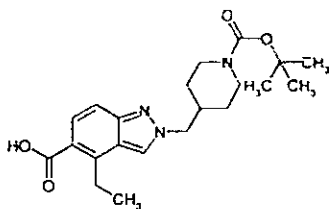
Example 212f: Methyl 2-[[1-(tert-butoxycarbonyl)piperidin-4-yl]methyl]-4-ethyl-2H-indazol-5-carboxylate



Analogously to Example 1e, after two runs 571 mg of the title compound was obtained from 347 mg of the bromide prepared in Example 212e and 0.1 ml of methanol.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.02-1.15 (2H), 1.17-1.24 (3H), 1.34 (9H), 1.42 (2H), 2.14 (1H), 2.63 (2H), 3.17 (2H), 3.79 (3H), 3.88 (2H), 4.30 (2H), 7.43 (1H), 7.60 (1H), 8.68 (1H).

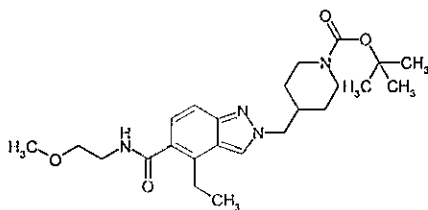
Example 212g: 2-([1-(tert-butoxycarbonyl)piperidin-4-yl]methyl)-4-ethyl-2H-indazol-5-carboxylic acid



Analogously to Example 1d, 427 mg of the title compound was obtained from 371 mg of the ester in Example 212f.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.00-1.15 (2H), 1.20 (3H), 1.34 (9H), 1.42 (2H), 2.14 (1H), 2.64 (2H), 3.19 (2H), 3.88 (2H), 4.29 (2H), 7.39 (1H), 7.61 (1H), 8.63 (1H), 12.33 (1H).

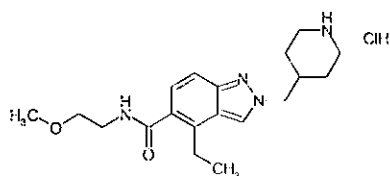
Example 212h: Tert-butyl 4-({4-ethyl-5-[N-(2-methoxyethyl)carbamoyl]-2H-indazol-2-yl}-methyl)piperidin-1-carboxylate



Analogously to Example 1, 215 mg of the title compound was obtained from 427 mg of the acid prepared in Example 212g and 83 mg of 2-methoxy-ethylamine.

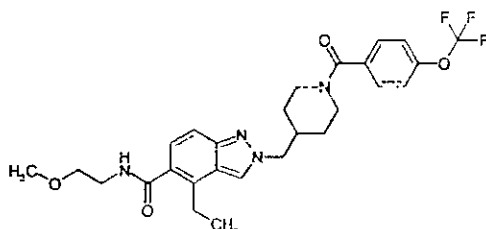
¹H-NMR (300 MHz, DMSO-d₆): δ = 0.98-1.14 (2H), 1.19 (3H), 1.30-1.47 (11H), 2.13 (1H), 2.65 (2H), 2.90 (2H), 3.24 (3H), 3.32-3.45 (4H), 3.87 (2H), 4.28 (2H), 7.10 (1H), 7.38 (1H), 8.12 (1H), 8.50 (1H).

Example 212i: 4-ethyl-N-(2-methoxyethyl)-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 215 mg of the amide prepared in Example 212h, 258 mg of the title compound was obtained, which was reacted without further purification.

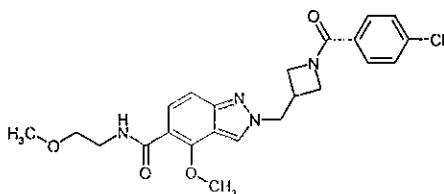
Example 213: 4-Ethyl-N-(2-methoxyethyl)-2-({1-[4-(trifluoromethoxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 32 mg of the title compound was obtained from 91 mg of the compound prepared in Example 212i and 59 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.14-1.29 (5H), 1.31-1.61 (2H), 2.20-2.35 (1H), 2.66-3.06 (4H), 3.24 (3H), 3.30-3.56 (5H), 4.31 (2H), 4.41 (1H), 7.10 (1H), 7.34-7.43 (3H), 7.47 (2H), 8.11 (1H), 8.50 (1H).

Example 214: 2-{{1-[4-chlorobenzoyl]azetidin-3-yl}methyl}-4-methoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide

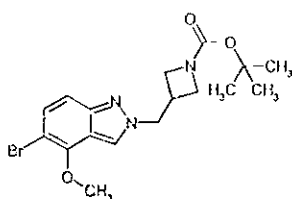


Analogously to Example 55, 691 mg of the title compound was obtained from 68 mg of the compound prepared in Example 214c and 37 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 3.22 (1H), 3.27 (3H), 3.40-3.47 (4H), 3.92 (1H), 4.10 (1H), 4.16 (3H), 4.19 (1H), 4.38 (1H), 4.66 (2H), 7.21 (1H), 7.48 (2H), 7.55-7.67 (3H), 8.18 (1H), 8.87 (1H).

The starting material was prepared as follows:

- 5 **Example 214a:** Tert-butyl 3-[(5-bromo-4-methoxy-2H-indazol-2-yl)methyl]azetidin-1-carboxylate

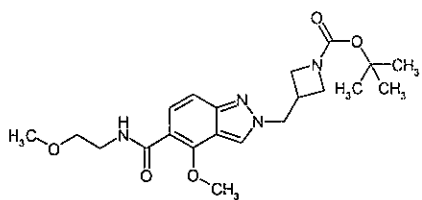


- To a solution of 1.94 g of tert-butyl 3-[(tosyloxy)methyl]azetidin-1-carboxylate in 15.5 ml acetone were added 837 mg of lithium iodide and the reaction mixture was stirred for 16 hours at 35°C. After cooling, it was diluted with 200 ml ethyl acetate and the organic phase washed twice with 30 ml portions of water and once with 20 ml saturated sodium chloride solution. After drying over sodium sulphate and filtration, this was concentrated in vacuo. In this manner, 1.6 g of tert-butyl-3-(iodomethyl)azetidin-1-carboxylate were obtained, which was further reacted without purification.

- 15 To a solution of 620 mg of 5-bromo-7-methoxy-1H-indazole in 24 ml DMF were added 1.11 g of potassium carbonate and the mixture stirred at 25°C for 30 minutes. Then 1.25 g of the iodide prepared above was added and the reaction mixture stirred for 3 hours at 60°C. After cooling, it was diluted with 200 ml 1:1 tert-butyl methyl ether/hexane, washed once each with 20 ml portions of water and saturated sodium chloride solution, dried over sodium sulphate and concentrated in vacuo and the crude product thus obtained purified by column chromatography on silica gel with a hexane / ethyl acetate gradient. Yield: 263 mg of the title compound.

¹H-NMR (300 MHz, CDCl₃): δ = 1.43 (9H), 3.23 (1H), 3.78 (2H), 4.07 (2H), 4.10 (3H), 4.59 (2H), 7.28 (1H), 7.36 (1H), 8.03 (1H).

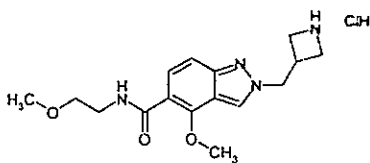
- 25 **Example 214b:** Tert-butyl 3-({4-methoxy-5-[N-(2-methoxyethyl)carbamoyl]-2H-indazol-2-yl)methyl}azetidin-1-carboxylate



Analogously to Example 1b, a total of 193 mg of the title compound was obtained from 110 and 252 mg respectively of the compound prepared in Example 214a and 63 and 143 mg respectively of 2-methoxyethylamine.

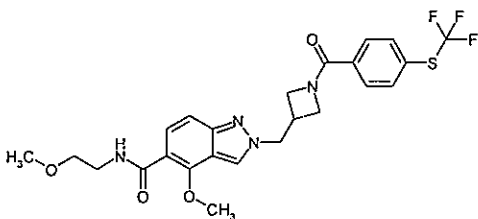
- 5 ¹H-NMR (300 MHz, DMSO-d₆): δ = 1.32 (9H), 3.10 (1H), 3.27 (3H), 3.38-3.48 (4H), 3.72 (2H), 3.82-3.94 (2H), 4.17 (3H), 4.60 (2H), 7.21 (1H), 7.63 (1H), 8.18 (1H), 8.87 (1H).

Example 214c: 2-(azetidin-3-ylmethyl)-4-methoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide hydrochloride



- 10 Analogously to Example 1a, from 50 mg of the compound prepared in Example 214b, 42 mg of the title compound was obtained, which was further reacted without purification.

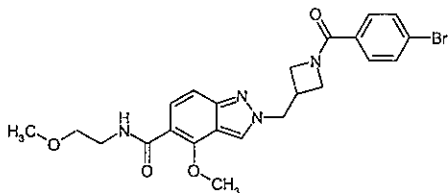
Example 215: 4-methoxy-N-(2-methoxyethyl)-2-[(1-{4-[(trifluoromethyl)sulphonyl]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide



- 15 Analogously to Example 55, 56 mg of the title compound was obtained from 42 mg of the compound prepared in Example 214c and 32 mg of 4-[(trifluoromethyl)sulphonyl]benzoyl chloride.

- ¹H-NMR (300 MHz, DMSO-d₆): δ = 3.21 (1H), 3.27 (3H), 3.38-3.49 (4H), 3.95 (1H), 4.12 (1H), 4.16 (3H), 4.21 (1H), 4.40 (1H), 4.67 (2H), 7.20 (1H), 7.63 (1H), 7.69 (2H), 7.76 (2H),
20 8.18 (1H), 8.87 (1H).

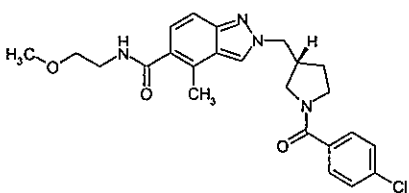
Example 216: 2-[[1-(4-bromobenzoyl)azetidin-3-yl]methyl]-4-methoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 48 mg of the title compound was obtained from 42 mg of the compound prepared in Example 214c and 29 mg of 4-bromobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 3.23 (1H), 3.26 (3H), 3.38-3.48 (4H), 3.92 (1H), 4.09 (1H), 4.16 (3H), 4.19 (1H), 4.38 (1H), 4.66 (2H), 7.20 (1H), 7.51 (2H), 7.58-7.68 (3H), 8.18 (1H), 8.86 (1H).

Example 217: 2-[[[(R)-1-(4-chlorobenzoyl)pyrrolidin-3-yl]methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

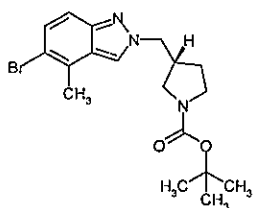


Analogously to Example 55, 50 mg of the title compound was obtained from 106 mg of the compound prepared in Example 217e and 58 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.57-1.73 (1H), 1.81-1.97 (1H), 2.47 / 2.51 (3H), 2.78-2.93 (1H), 3.25 (3H), 3.19-3.61 (8H), 4.39 / 4.51 (2H), 7.11-7.18 (1H), 7.33-7.43 (1H), 7.45 (2H), 7.50 (2H), 8.09-8.16 (1H), 8.47 / 8.55 (1H).

The starting material was prepared as follows:

Example 217a: (R)-tert-butyl-3-[(5-bromo-4-methyl-2H-indazol-2-yl)methyl]pyrrolidine-1-carboxylate

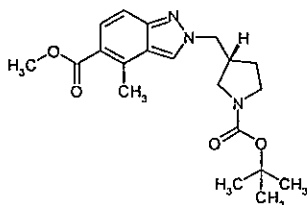


To a solution of 9.0 g of (R)-tert-butyl 3-(hydroxymethyl)pyrrolidin-1-carboxylate in 100 ml pyridine was added 12.8 g of p-toluenesulphonyl chloride at 0°C under nitrogen and stirred for 3 hours at 25°C. The reaction mixture was diluted with ethyl acetate and stirred for 30 minutes with sodium hydrogen carbonate. The phases were then separated and the organic
5 phase washed with saturated sodium chloride solution, dried over sodium sulphate and concentrated in vacuo after filtration. The residue thus obtained was purified by column chromatography on silica gel with hexane / 0-100% ethyl acetate. Yield: 11.4 g of (R)-tert-butyl-3-[(tosyloxy)methyl]pyrrolidin-1-carboxylate.

Analogously to Example 1c, 1.97 g of the title compound was obtained from 4.53 g of 5-bromo-
10 4-methyl-1H-indazole and 11.4 g of the (R)-tert-butyl-3-[(tosyloxy)methyl]pyrrolidin-1-carboxylate prepared above with addition of 7.9 g of tetrabutylammonium iodide.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.34 (9H), 1.50-1.66 (1H), 1.81 (1H), 2.48 (3H), 2.78 (1H), 3.01 (1H), 3.16 (1H), 3.25-3.37 (2H), 4.41 (2H), 7.29 (1H), 7.36 (1H), 8.51 (1H).

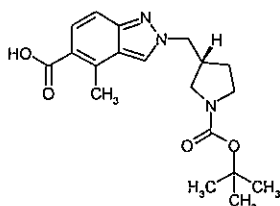
Example 217b: Methyl 2-{[(R)-1-(tert-butoxycarbonyl)pyrrolidin-3-yl]methyl}-4-methyl-2H-
15 indazol-5-carboxylate



Analogously to Example 1e, after two runs 820 mg of the title compound was obtained from 563 mg of the bromide prepared in Example 217a and 0.17 ml of methanol.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.34 (9H), 1.58 (1H), 1.83 (1H), 2.73 (3H), 2.80 (1H), 3.03 (1H), 3.17 (1H), 3.25-3.37 (2H), 3.79 (3H), 4.37-4.49 (2H), 7.44 (1H), 7.64 (1H), 8.72 (1H).
20

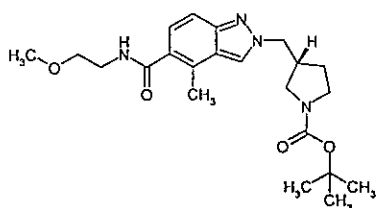
Example 217c: 2-{[(R)-1-(tert-butoxycarbonyl)pyrrolidin-3-yl]methyl}-4-methyl-2H-indazol-5-carboxylic acid



Analogously to Example 1d, 701 mg of the title compound was obtained from 620 mg of the ester prepared in Example 217b.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.34 (9H), 1.50-1.69 (1H), 1.82 (1H), 1.75-1.86 (1H), 2.69-2.88 (4H), 3.03 (1H), 3.17 (1H), 3.24-3.38 (1H), 4.42 (2H), 7.40 (1H), 7.65 (1H), 8.68 (1H), 12.28 (1H).

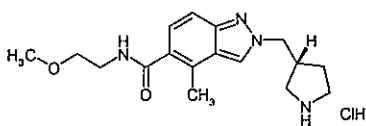
Example 217d: (R)-tert-butyl 3-({5-[N-(2-methoxyethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}methyl)pyrrolidin-1-carboxylate



Analogously to Example 1, 376 mg of the title compound was obtained from 350 mg of the acid prepared in Example 217c and 73 mg of 2-methoxyethylamine.

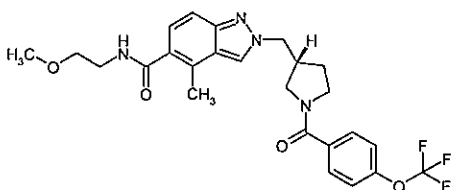
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.35 (9H), 1.57 (1H), 1.80 (1H), 2.50 (3H), 2.73-2.86 (1H), 3.01 (1H), 3.11-3.22 (1H), 3.25 (3H), 3.26-3.39 (3H), 3.39-3.46 (2H), 4.42 (2H), 7.15 (1H), 7.39 (1H), 8.13 (1H), 8.52 (1H).

Example 217e: N-(2-methoxyethyl)-4-methyl-2-[(R)-pyrrolidin-3-ylmethyl]-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 376 mg of the amide prepared in Example 217d, 465 mg of the title compound was obtained, which was reacted without further purification.

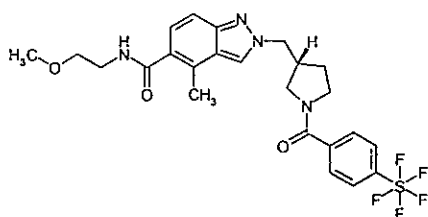
Example 218: N-(2-methoxyethyl)-4-methyl-2-({(R)-1-[4-(trifluoromethoxy)benzoyl]-pyrrolidin-3-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 55, 61 mg of the title compound was obtained from 106 mg of the compound prepared in Example 217e and 74 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.57-1.74 (1H), 1.81-1.97 (1H), 2.47 / 2.51 (3H), 2.77-2.94 (1H), 3.25-3.62 (11H), 4.40 / 4.51 (2H), 7.09-7.19 (1H), 7.31-7.44 (3H), 7.61 (2H), 8.08-8.16 (1H), 8.47 / 8.55 (1H).

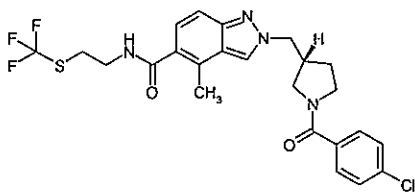
Example 219: N-(2-methoxyethyl)-4-methyl-2-({(3R)-1-[4-(pentafluoro-λ⁶-sulphanyl)benzoyl]-pyrrolidin-3-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 29 mg of the title compound was obtained from 106 mg of the compound prepared in Example 217e and 75 mg of 4-(pentafluoro-λ⁶-sulphanyl)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.57-1.76 (1H), 1.81-1.99 (1H), 2.50 (3H), 2.77-2.94 (1H), 3.16-3.62 (11H), 4.39 / 4.52 (2H), 7.09-7.19 (1H), 7.32-7.44 (1H), 7.68 (2H), 7.90-7.97 (2H), 8.10-8.17 (1H), 8.46 / 8.55 (1H).

Example 220: 2-{{(R)-1-(4-chlorobenzoyl)pyrrolidin-3-yl}methyl}-4-methyl-N-{2-[(trifluoromethyl)sulphanyl]ethyl}-2H-indazole-5-carboxamide

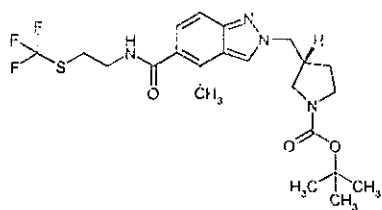


Analogously to Example 55, 29 mg of the title compound was obtained from 106 mg of the compound prepared in Example 220b and 48 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.57-1.73 (1H), 1.81-1.96 (1H), 2.48 / 2.53 (3H), 2.79-2.93 (1H), 3.13-3.60 (8H), 4.40 / 4.51 (2H), 7.14-7.23 (1H), 7.33-7.53 (5H), 8.35-8.42 (1H), 8.49 / 8.58 (1H).

The starting material was prepared as follows:

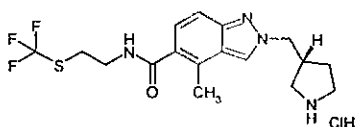
Example 220a: (R)-tert-butyl 3-[[4-methyl-5-(N-{2-[(trifluoromethyl)sulphanyl]ethyl}-carbamoyl)-2H-indazol-2-yl]methyl]pyrrolidin-1-carboxylate



Analogously to Example 1, 245 mg of the title compound was obtained from 350 mg of the acid prepared in Example 217c and 141 mg of 2-[(trifluoromethyl)sulphanyl]ethylamine.

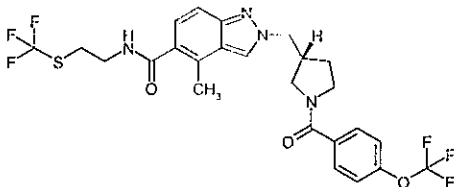
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.35 (9H), 1.52-1.64 (11H), 1.81 (1H), 2.52 (3H), 2.71-2.96 (2H), 2.97-3.05 (12H), 3.18 (2H), 3.23-3.28 (1H), 3.51 (2H), 4.42 (2H), 7.19 (1H), 7.42 (1H), 8.39 (1H), 8.54 (1H).

Example 220b: 4-methyl-2-[(R)-pyrrolidin-3-ylmethyl]-N-{2-[(trifluoromethyl)sulphanyl]ethyl}-ethyl}-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 245 mg of the amide prepared in Example 220a, 410 mg of the title compound was obtained, which was reacted without further purification.

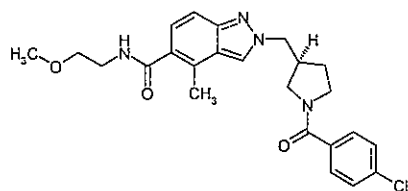
Example 221: 4-methyl-2-[(R)-1-[4-(trifluoromethoxy)benzoyl]pyrrolidin-3-yl]methyl)-N-{2-[(trifluoromethyl)sulphanyl]ethyl}-2H-indazole-5-carboxamide



Analogously to Example 55, 49 mg of the title compound was obtained from 106 mg of the compound prepared in Example 220b and 62 mg of 4-(trifluoromethoxy)benzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.58-1.74 (1H), 1.81-1.98 (1H), 2.48 / 2.53 (3H), 2.78-2.94 (1H), 3.12-3.62 (8H), 4.40 / 4.52 (2H), 7.14-7.23 (1H), 7.32-7.49 (3H), 7.61 (2H), 8.35-8.43 (1H), 8.49 / 8.58 (1H).

Example 222: 2-{[(S)-1-(4-chlorobenzoyl)pyrrolidin-3-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

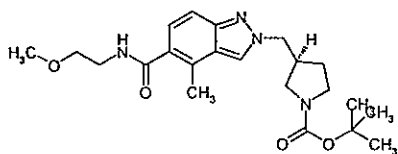


Analogously to Example 55, 67 mg of the title compound was obtained from 134 mg of the compound prepared in Example 222a and 73 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, CDCl₃): δ = 1.72-1.86 (1H), 1.99-2.20 (1H), 2.64 (3H), 2.93-3.13 (1H), 3.29-3.36 (1H), 3.39 (3H), 3.45-3.71 (6H), 3.76-3.86 (1H), 4.31-4.57 (2H), 6.12-6.20 (1H), 7.30-7.42 (3H), 7.43-7.56 (3H), 7.90 / 8.02 (1H).

The starting material was prepared as follows:

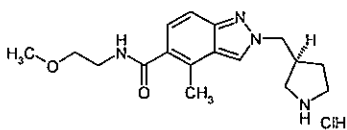
Example 222a: (S)-tert-butyl 3-({5-[N-(2-methoxyethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}methyl)pyrrolidin-1-carboxylate



Analogously to Example 217a to 217d, 760mg of this title compound were prepared starting from (S)-tert-butyl-3-(hydroxymethyl)pyrrolidin-1-carboxylate.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.35 (9H), 1.52-1.64 (1H), 1.77-1.87 (1H), 1.76-1.82 (1H), 2.50 (3H), 2.74-2.86 (1H), 3.02 (1H), 3.12-3.20 (1H), 3.25 (3H), 3.31 (1H), 3.36 (2H), 3.43 (2H), 4.42 (2H), 7.16 (1H), 7.39 (1H), 8.10 (1H), 8.51 (1H).

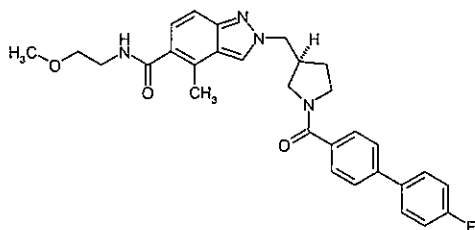
Example 222b: N-(2-methoxyethyl)-4-methyl-2-[(S)-pyrrolidin-3-ylmethyl]-2H-indazole-5-carboxamide hydrochloride



20

Analogously to Example 1a, from 159 mg of the amide prepared in Example 222a, 134 mg of the title compound was obtained, which was reacted without further purification.

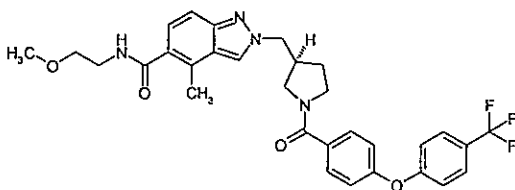
Example 223: 2-({(S)-1-[(4'-fluorobiphenyl-4-yl)carbonyl]pyrrolidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 128 mg of the title compound was obtained from 230 mg of the compound prepared in Example 222b and 141 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 1.73-1.89 (1H), 1.98-2.24 (1H), 2.60 / 2.67 (3H), 2.94-3.18 (1H), 3.37 / 3.39 (3H), 3.49-3.76 (6H), 3.79-3.91 (2H), 4.32-4.61 (2H), 6.08-6.21 (1H), 7.09-7.20 (2H), 7.28-7.38 (1H), 7.46-7.62 (7H), 7.91 / 8.04 (1H).

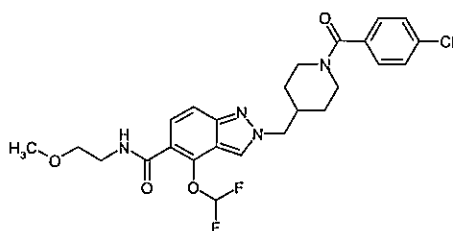
Example 224: N-(2-methoxyethyl)-4-methyl-2-{{(S)-1-[4-(4-(trifluoromethyl)phenoxy)benzoyl]pyrrolidin-3-yl}methyl}-2H-indazole-5-carboxamide



Analogously to Example 1, 102 mg of the title compound was obtained from 230 mg of the compound prepared in Example 222b and 184 mg of 4-[4-(trifluoromethyl)phenoxy]benzoic acid.

¹H-NMR (300 MHz, CDCl₃): δ = 1.70-2.25 (2H), 2.63 / 2.66 (3H), 2.94-3.17 (1H), 3.39 (3H), 3.47-3.92 (8H), 4.33-4.59 (2H), 6.15 (1H), 7.00-7.11 (4H), 7.30-7.40 (1H), 7.47-7.65 (5H), 7.91 / 8.03 (1H).

Example 225: 2-{{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-(difluoromethoxy)-N-(2-methoxyethyl)-2H-indazole-5-carboxamide

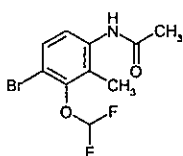


Analogously to Example 55, 90 mg of the title compound was obtained from 106 mg of the compound prepared in Example 225h and 49 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.14-1.58 (4H), 2.19-2.34 (1H), 2.67-2.81 (1H), 2.97 (1H), 3.22-3.53 (9H), 4.37 (2H), 7.13 (1H), 7.31-7.39 (3H), 7.47 (2H), 7.54 (1H), 8.21 (1H), 8.50 (1H).

The starting material was prepared as follows:

Example 225a: N-[4-bromo-3-(difluoromethoxy)-2-methylphenyl]acetamide

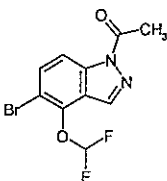


To a solution of 5.0 g of 3-difluoromethoxy-2-methylaniline in 48 ml DMF, a solution of 5.1 g of N-bromosuccinimide in 24 ml DMF was added dropwise at 0°C and stirred for 1 hour at 0°C. Then the reaction mixture was diluted with 400 ml hexane / ethyl acetate, and washed once with 50 ml of a 10% aqueous sodium carbonate solution and three times with 50 ml portions of water. After drying over sodium sulphate and filtration, this was concentrated in vacuo. The crude product thus obtained (6.68 g) was used in the next step without purification.

To a solution of 6.39 g of the bromide prepared above in 70 ml pyridine, 3.0 ml of acetic anhydride were added dropwise at 0°C and stirred for 20 hours at 25°C. The reaction mixture was concentrated in vacuo and the crude product thus obtained purified by column chromatography on silica gel with hexane / 0-100% ethyl acetate. The substance was then recrystallized from hexane. In this manner, 6.39 g of the title compound was obtained.

¹H-NMR (300 MHz, CDCl₃): δ = 2.22 (3H), 2.27 (3H), 6.51 (1H), 6.95 (1H), 7.46 (1H), 7.71 (1H).

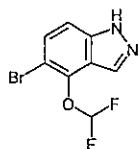
Example 225b: 1-[5-bromo-4-(difluoromethoxy)-1H-indazol-1-yl]ethan-1-one



Analogously to Example 212c, 5.32 g of the title compound was obtained from 6.38 g of the amide prepared in Example 225a.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.70 (3H), 7.34 (1H), 7.90 (1H), 8.16 (1H), 8.46 (1H).

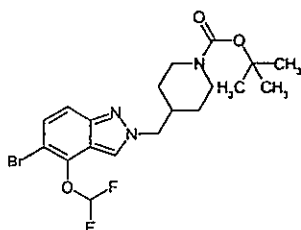
Example 225c: 5-bromo-4-(difluoromethoxy)-1H-indazole



Analogously to Example 212d, 4.12 g of the title compound was obtained from 5.32 g of the
5 indazole prepared in Example 225b.

¹H-NMR (300 MHz, DMSO-d₆): δ = 7.30 (1H), 7.44 (1H), 7.56 (1H), 8.06 (1H), 13.54 (1H).

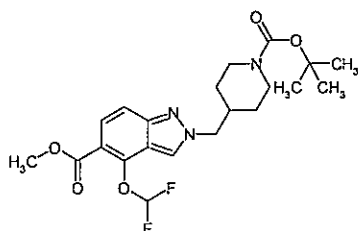
Example 225d: Tert-butyl 4-{{5-bromo-4-(difluoromethoxy)-2H-indazol-2-yl}methyl}-piperidin-1-carboxylate



10 Analogously to Example 212e, 2.06 g of the title compound was obtained from 4.1 g of the indazole prepared in Example 225c and 8.7 g of tert-butyl 4-[(tosyloxy)methyl]piperidin-1-carboxylate.

¹H-NMR (300 MHz, CDCl₃): δ = 1.19-1.28 (2H), 1.45 (9H), 1.51-1.59 (2H), 2.25 (1H), 2.68 (2H), 4.12 (2H), 4.27 (2H), 6.61 (1H), 7.41 (1H), 7.50 (1H), 7.95 (1H).

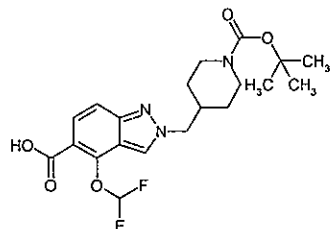
15 **Example 225e:** Methyl 2-{{[1-(tert-butoxycarbonyl)piperidin-4-yl]methyl}-4-(difluoromethoxy)-2H-indazol-5-carboxylate



Analogously to Example 1e, after three runs 1.28 g of the title compound was obtained from 683 mg of the bromide prepared in Example 225d and 0.18 ml of methanol.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.96-1.22 (2H), 1.28-1.47 (11H), 2.14 (1H), 2.54-2.74 (2H), 3.81 (3H), 3.87 (2H), 4.36 (2H), 7.17 (1H), 7.58 (1H), 7.65 (1H), 8.62 (1H).

Example 225f: 2-([1-(tert-butoxycarbonyl)piperidin-4-yl]methyl)-4-(difluoromethoxy)-2H-indazol-5-carboxylic acid

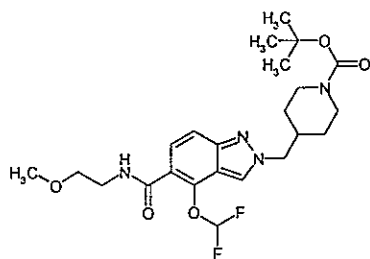


5

Analogously to Example 1d, 1.05 g of the title compound was obtained from 1.28 g of the ester prepared in Example 225e.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.97-1.16 (2H), 1.28-1.47 (11H), 2.13 (1H), 2.63 (2H), 3.87 (2H), 4.35 (2H), 7.14 (1H), 7.54 (1H), 7.65 (1H), 8.58 (1H), 13.05 (1H).

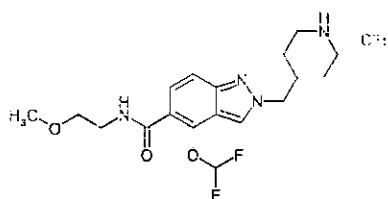
10 **Example 225g:** Tert-butyl 4-({4-(difluoromethoxy)-5-[N-(2-methoxyethyl)carbamoyl]-2H-indazol-2-yl}methyl)piperidin-1-carboxylate



Analogously to Example 1, 503 mg of the title compound was obtained from 520 mg of the compound prepared in Example 225f and 92 mg of 2-methoxyethylamine.

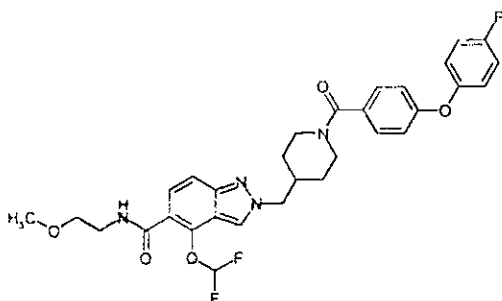
15 ¹H-NMR (300 MHz, DMSO-d₆): δ = 1.01-1.13 (2H), 1.34 (9H), 1.38 (2H), 2.13 (1H), 2.59-2.71 (6H), 3.24 (3H), 3.34-3.44 (2H), 3.87 (2H), 4.34 (2H), 7.14 (1H), 7.33 (1H), 7.54 (1H), 8.21 (1H), 8.50 (1H).

Example 225h: 4-(difluoromethoxy)-N-(2-methoxyethyl)-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 122 mg of the amide prepared in Example 225g, 106 mg of the title compound was obtained, which was reacted without further purification.

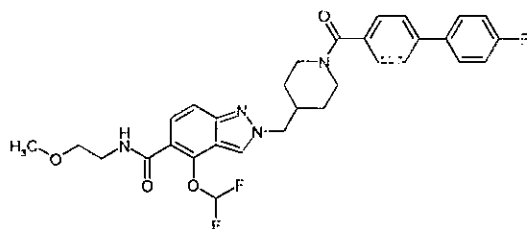
Example 226: 4-(difluoromethoxy)-2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 11 mg of the title compound was obtained from 106 mg of the compound prepared in Example 225h and 59 mg of 4-(4-fluorophenoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.10-1.59 (4H), 2.23-2.33 (1H), 2.75-3.01 (2H), 3.24 (3H), 3.33-3.51 (5H), 3.51-3.72 (1H), 4.37 (2H), 6.95 (2H), 7.07-7.12 (2H), 7.13 (1H), 7.17-7.26 (2H), 7.28-7.39 (3H), 7.54 (1H), 8.21 (1H), 8.50 (1H).

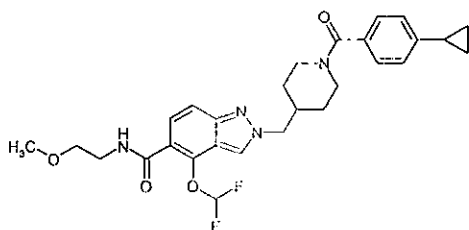
Example 227: 4-(difluoromethoxy)-2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 38 mg of the title compound was obtained from 106 mg of the compound prepared in Example 225h and 55 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.14-1.62 (4H), 2.23-2.35 (1H), 2.68-2.85 (1H), 2.91-3.09 (1H), 3.24 (3H), 3.32-3.48 (5H), 3.50-3.70 (1H), 4.38 (2H), 7.14 (1H), 7.28 (2H), 7.33 (1H), 7.40 (2H), 7.55 (1H), 7.64-7.75 (4H), 8.21 (1H), 8.51 (1H).

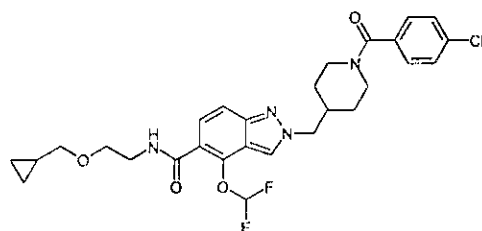
Example 228: 2-{{1-(4-cyclopropylbenzoyl)piperidin-4-yl}methyl}-4-(difluoromethoxy)-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 75 mg of the title compound was obtained from 106 mg of the compound prepared in Example 225h and 41 mg of 4-cyclopropylbenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.66 (2H), 0.94 (2H), 1.10-1.54 (4H), 1.90 (1H), 2.14-2.31 (1H), 2.72-3.02 (2H), 3.03-3.18 (1H), 3.24 (3H), 3.33-3.47 (4H), 3.49-3.68 (1H), 4.37 (2H), 7.07 (2H), 7.13 (1H), 7.19 (2H), 7.33 (1H), 7.54 (1H), 8.21 (1H), 8.50 (1H).

Example 229: 2-{{1-(4-chlorobenzoyl)piperidin-4-yl}methyl}-N-[2-(cyclopropylmethoxy)ethyl]-4-(difluoromethoxy)-2H-indazole-5-carboxamide

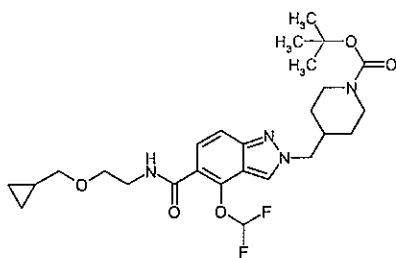


Analogously to Example 55, 170 mg of the title compound was obtained from 127 mg of the compound prepared in Example 229b and 53 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.14 (2H), 0.42 (2H), 0.96 (1H), 1.10-1.27 (2H), 1.37 (1H), 1.51 (1H), 2.26 (1H), 2.71 (1H), 2.97 (1H), 3.20-3.55 (8H), 4.37 (2H), 7.14 (1H), 7.31-7.38 (3H), 7.47 (2H), 7.55 (1H), 8.21 (1H), 8.51 (1H).

The starting material was prepared as follows:

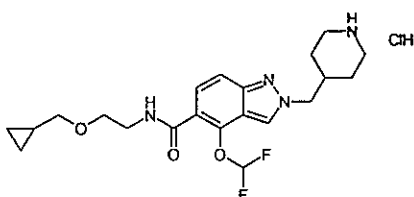
Example 229a: Tert-butyl 4-{{5-{{N-[2-(cyclopropylmethoxy)ethyl]carbonyl}-4-(difluoromethoxy)-2H-indazol-2-yl}methyl}piperidin-1-carboxylate



Analogously to Example 1, 453 mg of the title compound was obtained from 520 mg of the compound prepared in Example 225f and 185 mg of 2-(cyclopropylmethoxy)ethylamine.

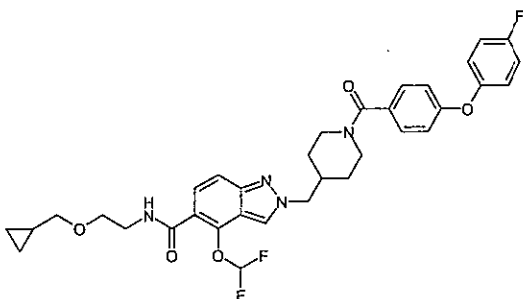
¹H-NMR (300 MHz, DMSO-d₆): δ = 0.14 (2H), 0.42 (2H), 0.97 (1H), 1.07 (2H), 1.34 (9H), 1.39 (2H), 2.13 (1H), 2.56-2.72 (2H), 3.23 (2H), 3.37 (2H), 3.48 (2H), 3.87 (2H), 4.34 (2H), 7.15 (1H), 7.34 (1H), 7.55 (1H), 8.21 (1H), 8.50 (1H).

Example 229b: N-[(2-cyclopropylmethoxy)ethyl]-4-(difluoromethoxy)-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 145 mg of the amide prepared in Example 229a, 127 mg of the title compound was obtained, which was reacted without further purification.

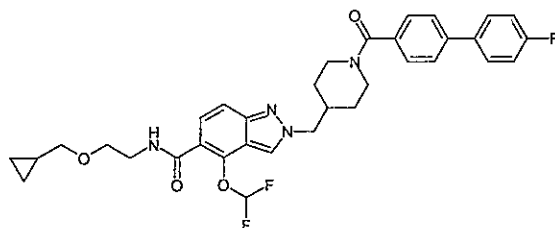
Example 230: N-[2-(cyclopropylmethoxy)ethyl]-4-(difluoromethoxy)-2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 156 mg of the title compound was obtained from 127 mg of the compound prepared in Example 229b and 64 mg of 4-(4-fluorophenoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.14 (2H), 0.42 (2H), 0.96 (1H), 1.10-1.54 (4H), 2.18-2.33 (1H), 2.71-3.02 (2H), 3.03-3.16 (1H), 3.22 (2H), 3.37 (2H), 3.47 (2H), 3.55-3.68 (1H), 4.37 (2H), 6.95 (2H), 7.10 (2H), 7.14 (1H), 7.22 (2H), 7.34 (3H), 7.55 (1H), 8.21 (1H), 8.51 (1H).

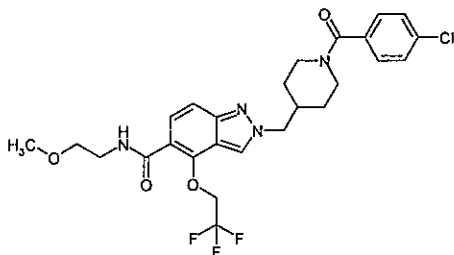
Example 231: N-[2-(cyclopropylmethoxy)ethyl]-4-(difluoromethoxy)-2-({1-[(4'-fluoro-
5 biphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 164 mg of the title compound was obtained from 127 mg of the compound prepared in Example 229b and 60 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.14 (2H), 0.42 (2H), 0.96 (1H), 1.14-1.60 (4H), 2.27 (1H), 2.70-3.04 (1H), 2.87 (1H), 3.03-3.15 (1H), 3.22 (2H), 3.37 (2H), 3.47 (2H), 3.60 (1H), 4.38 (2H), 7.14 (1H), 7.28 (2H), 7.34 (1H), 7.41 (2H), 7.55 (1H), 7.64-7.75 (4H), 8.21 (1H), 8.52 (1H).

Example 232: 2-({1-[(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-(2,2,2-trifluoroethoxy)-2H-indazole-5-carboxamide

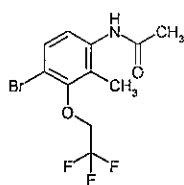


Analogously to Example 55, 113 mg of the title compound was obtained from 109 mg of the compound prepared in Example 232h and 46 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.12-1.61 (3H), 2.19-2.34 (1H), 2.67-3.07 (2H), 3.23 (3H), 3.33-3.56 (4H), 4.28-4.47 (3H), 4.95 (2H), 7.31-7.39 (3H), 7.40-7.50 (3H), 7.99 (1H), 8.65 (1H).

The starting material was prepared as follows:

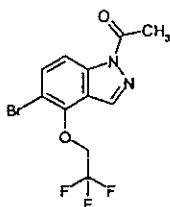
Example 232a: N-[4-bromo-2-methyl-3-(2,2,2-trifluoroethoxy)phenyl]acetamide



Analogously to Example 225a, 6.13 g of the title compound was obtained from 5.0 g of 2-methyl-3-(2,2,2-trifluoroethoxy)aniline.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.07 (3H), 2.18 (3H), 4.55 (2H), 7.29 (1H), 7.45 (1H), 9.42 (1H).

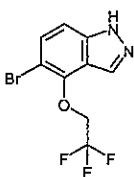
Example 232b: 1-[5-bromo-4-(2,2,2-trifluoroethoxy)-1H-indazol-1-yl]ethan-1-one



Analogously to Example 212c, 5.08 g of the title compound was obtained from 6.13 g of the amide prepared in Example 232a.

¹H-NMR (300 MHz, DMSO-d₆): δ = 2.69 (3H), 5.14 (2H), 7.80 (1H), 7.99 (1H), 8.64 (1H).

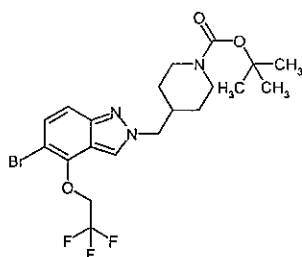
Example 232c: 5-bromo-4-(2,2,2-trifluoroethoxy)-1H-indazole



Analogously to Example 212d, 3.57 g of the title compound was obtained from 5.08 g of the indazole prepared in Example 232b.

¹H-NMR (300 MHz, DMSO-d₆): δ = 5.05 (2H), 7.24 (1H), 7.45 (1H), 8.26 (1H), 13.38 (1H).

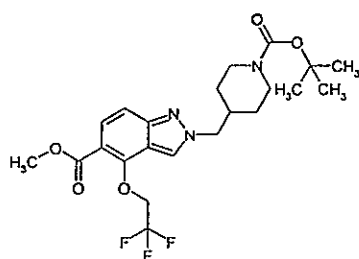
Example 232d: Tert-butyl 4-{{[5-bromo-4-(2,2,2-trifluoroethoxy)-2H-indazol-2-yl]methyl}-piperidin-1-carboxylate



Analogously to Example 212e, 1.77 g of the title compound was obtained from 3.57 g of the indazole prepared in Example 232c and 6.7 g of tert-butyl 4-[(tosyloxy)methyl]piperidin-1-carboxylate.

- 5 $^1\text{H-NMR}$ (300 MHz, CDCl_3): δ = 1.16-1.31 (2H), 1.45 (9H), 1.54 (2H), 2.25 (1H), 2.68 (2H), 4.05-4.21 (2H), 4.27 (2H), 4.54 (2H), 7.35 (1H), 7.41 (1H), 7.91 (1H).

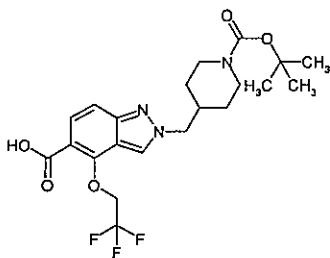
Example 232e: Methyl 2-{[1-(tert-butoxycarbonyl)piperidin-4-yl]methyl}-4-(2,2,2-trifluoroethoxy)-2H-indazole-5-carboxylate



- 10 Analogously to Example 1e, after three runs 857 mg of the title compound was obtained from 590 mg of the bromide prepared in Example 232d and 0.15 ml of methanol.

$^1\text{H-NMR}$ (300 MHz, DMSO-d_6): δ = 1.00-1.17 (2H), 1.34 (9H), 1.37-1.47 (2H), 2.13 (1H), 2.55-2.74 (2H), 3.79 (3H), 3.87 (2H), 4.32 (2H), 4.89 (2H), 7.39 (1H), 7.56 (1H), 8.67 (1H).

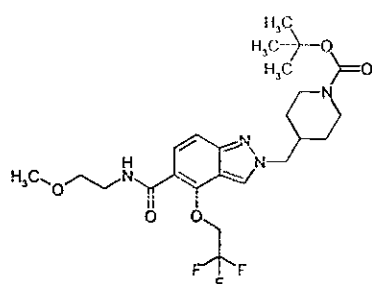
- Example 232f:** 2-{[1-(tert-butoxycarbonyl)piperidin-4-yl]methyl}-4-(2,2,2-trifluoroethoxy)-2H-indazole-5-carboxylic acid
- 15



Analogously to Example 1d, 724 mg of the title compound was obtained from 854 mg of the ester prepared in Example 232e.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.01-1.17 (2H), 1.34 (9H), 1.40 (2H), 2.13 (1H), 2.53-2.75 (2H), 3.87 (2H), 4.32 (2H), 4.84 (2H), 7.37 (1H), 7.58 (1H), 8.60 (1H), 12.74 (1H).

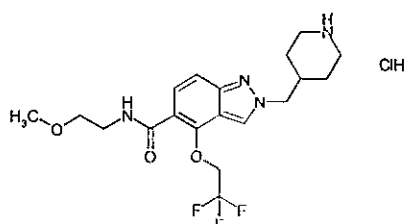
- 5 **Example 232g:** Tert-butyl 4-({5-[N-(2-methoxyethyl)carbamoyl]-4-(2,2,2-trifluoroethoxy)-2H-indazol-2-yl}methyl)piperidin-1-carboxylate



Analogously to Example 1, 386 mg of the title compound was obtained from 360 mg of the compound prepared in Example 232f and 59 mg of 2-methoxyethylamine.

- 10 ¹H-NMR (300 MHz, DMSO-d₆): δ = 0.98-1.18 (2H), 1.34 (9H), 1.43 (2H), 2.14 (1H), 2.65 (2H), 3.23 (3H), 3.35-3.47 (4H), 3.88 (2H), 4.30 (2H), 4.96 (2H), 7.35 (1H), 7.43 (1H), 7.99 (1H), 8.65 (1H).

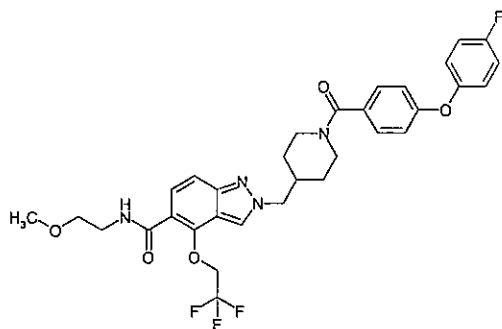
Example 232h: N-(2-methoxyethyl)-2-(4-piperidylmethyl)-4-(2,2,2-trifluoroethoxy)-2H-indazole-5-carboxamide hydrochloride



15

Analogously to Example 1a, from 109 mg of the amide prepared in Example 232g, 113 mg of the title compound was obtained, which was reacted without further purification.

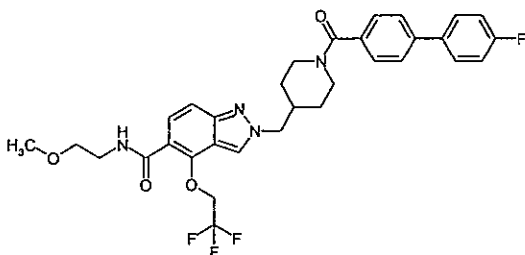
Example 233: 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-(2,2,2-trifluoroethoxy)-2H-indazole-5-carboxamide



Analogously to Example 1, 51 mg of the title compound was obtained from 109 mg of the compound prepared in Example 232h and 56 mg of 4-(4-fluorophenoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.10-1.31 (1H), 1.48 (2H), 2.18-2.37 (1H), 2.69-3.04 (2H),
 5 3.23 (3H), 3.37-3.48 (4H), 3.56-3.71 (1H), 4.33 (2H), 4.95 (2H), 6.95 (2H), 7.06-7.15 (2H),
 7.18-7.27 (2H), 7.31-7.38 (3H), 7.40-7.47 (1H), 7.99 (1H), 8.66 (1H).

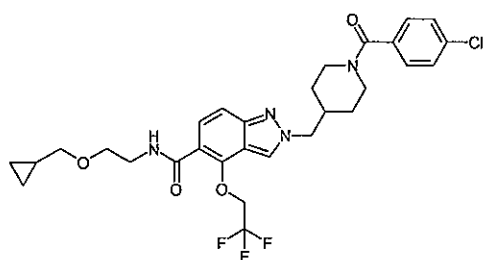
Example 234: 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-(2,2,2-trifluoroethoxy)-2H-indazole-5-carboxamide



10 Analogously to Example 1, 61 mg of the title compound was obtained from 109 mg of the compound prepared in Example 232h and 52 mg of 4-(4-fluorophenyl)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.14-1.63 (3H), 2.19-2.37 (1H), 2.68-3.09 (2H), 3.23 (3H),
 3.34-3.48 (4H), 3.62 (1H), 4.28-4.42 (2H), 4.96 (2H), 7.22-7.47 (6H), 7.64-7.76 (4H), 7.99 (1H),
 8.66 (1H).

15 **Example 235:** 2-{{1-(4-chlorobenzoyl)piperidin-4-yl}methyl}-N-[2-(cyclopropylmethoxy)ethyl]-4-(2,2,2-trifluoroethoxy)-2H-indazole-5-carboxamide

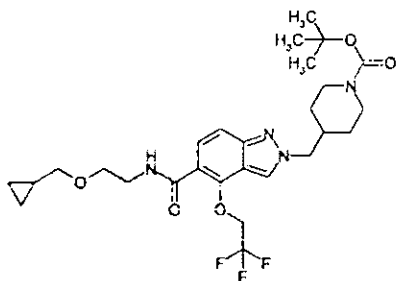


Analogously to Example 55, 122 mg of the title compound was obtained from 122 mg of the compound prepared in Example 235b and 48 mg of 4-chlorobenzoyl chloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.13 (2H), 0.41 (2H), 0.95 (1H), 1.11-1.62 (3H), 2.19-2.34 (1H), 2.73 (1H), 2.98 (1H), 3.21 (2H), 3.33-3.57 (5H), 4.33 (2H), 4.95 (2H), 7.31-7.51 (6H),
 5 8.00 (1H), 8.65 (1H).

The starting material was prepared as follows:

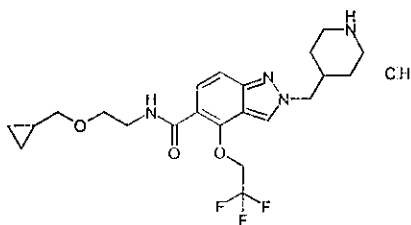
Example 235a: Tert-butyl 4-{{5-{N-[2-(cyclopropylmethoxy)ethyl]carbamoyl}-4-(2,2,2-trifluoroethoxy)-2H-indazol-2-yl]methyl}piperidin-1-carboxylate



10 Analogously to Example 1, 429 mg of the title compound was obtained from 360 mg of the compound prepared in Example 232f and 119 mg of 2-(cyclopropylmethoxy)ethylamine.

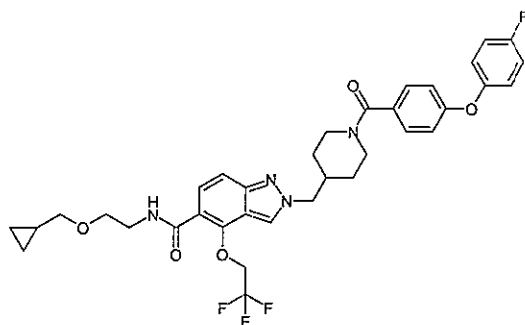
¹H-NMR (300 MHz, DMSO-d₆): δ = 0.13 (2H), 0.42 (2H), 0.95 (1H), 1.08 (2H), 1.34 (9H), 1.41 (2H), 2.14 (1H), 3.22 (2H), 3.41 (2H), 3.48 (2H), 3.88 (2H), 4.30 (2H), 4.95 (2H), 7.35 (1H),
 7.42 (1H), 8.00 (1H), 8.65 (1H).

15 **Example 235b:** N-[2-(cyclopropylmethoxy)ethyl]-2-(4-piperidylmethyl)-4-(2,2,2-trifluoroethoxy)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 138 mg of the amide prepared in Example 235a, 122 mg of the title compound was obtained, which was reacted without further purification.

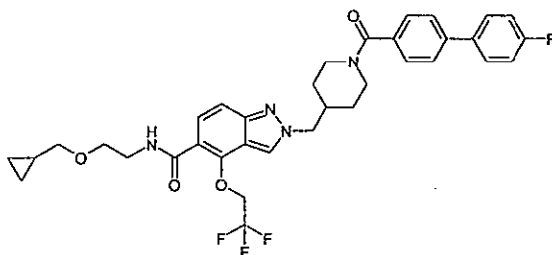
20 **Example 236:** N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-(2,2,2-trifluoroethoxy)-2H-indazole-5-carboxamide



Analogously to Example 1, 95 mg of the title compound was obtained from 122 mg of the compound prepared in Example 235b and 58 mg of 4-(4-fluorophenoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.13 (2H), 0.41 (2H), 0.95 (1H), 1.09-1.60 (3H), 2.17-2.36 (1H), 2.68-3.06 (2H), 3.22 (2H), 3.34-3.76 (5H), 4.33 (2H), 4.95 (2H), 6.95 (2H), 7.06-7.14 (2H), 7.17-7.27 (2H), 7.30-7.38 (3H), 7.42 (1H), 7.99 (1H), 8.65 (1H).

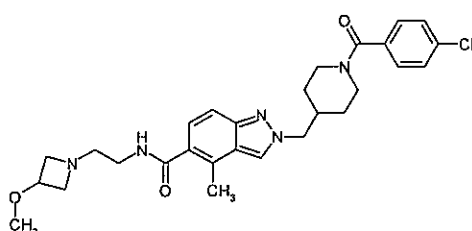
Example 237: N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-(2,2,2-trifluoroethoxy)-2H-indazole-5-carboxamide



Analogously to Example 1, 98 mg of the title compound was obtained from 122 mg of the compound prepared in Example 235b and 54 mg of 4-(4-fluorophenoxy)benzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.12 (2H), 0.41 (2H), 0.95 (1H), 1.15-1.64 (3H), 2.21-2.36 (1H), 2.70-3.10 (2H), 3.21 (2H), 3.35-3.69 (5H), 4.35 (2H), 4.95 (2H), 7.22-7.46 (6H), 7.63-7.77 (4H), 7.99 (1H), 8.66 (1H).

Example 238: 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-N-[2-(3-methoxyazetidin-1-yl)ethyl]-4-methyl-2H-indazole-5-carboxamide

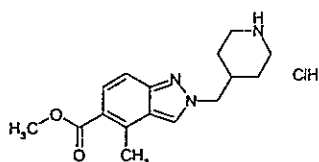


Analogously to Example 1, 33 mg of the title compound was obtained from 120 mg of the compound prepared in Example 238c and 40 mg of 2-(3-methoxyazetidin-1-yl)ethylamine (prepared analogously to WO2006/104406).

¹H-NMR (300 MHz, CDCl₃): δ = 0.83 (1H), 1.20-2.01 (4H), 2.38 (1H), 2.66 (3H), 2.69-3.09 (2H), 3.31 (3H), 3.43 (2H), 3.56-3.90 (5H), 4.18-4.32 (3H), 4.43 (2H), 6.91 (1H), 7.30-7.41 (5H), 7.52 (1H), 7.95 (1H).

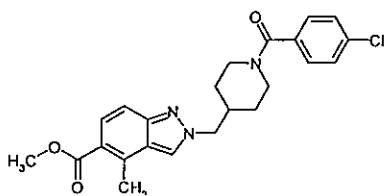
The starting material was prepared as follows:

Example 238a: methyl 4-methyl-2-(4-piperidylmethyl)-2H-indazol-5-carboxylate hydrochloride



Analogously to Example 1a, from 3.0 g of the ester prepared in Example 1e, 2.51 g of the title compound was obtained, which was reacted without further purification.

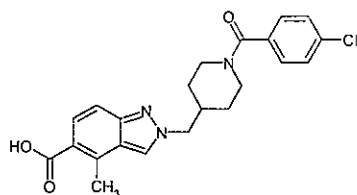
Example 238b: Methyl 2-{{1-(4-chlorobenzoyl)piperidin-4-yl}methyl}-4-methyl-2H-indazol-5-carboxylate



Analogously to Example 55, 3.27 g of the title compound was obtained from 2.5 g of the compound prepared in Example 238a and 1.5 g of 4-chlorobenzoyl chloride.

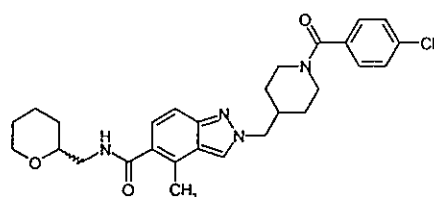
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.16-1.63 (4H), 2.21-2.33 (1H), 2.73 (4H), 2.90-3.05 (1H), 3.44-3.55 (1H), 3.79 (3H), 4.32 (2H), 4.36-4.47 (1H), 7.36 (2H), 7.42 (1H), 7.46 (2H), 7.63 (1H), 8.67 (1H).

Example 238c: 2-{{1-(4-chlorobenzoyl)piperidin-4-yl}methyl}-4-methyl-2H-indazol-5-carboxylic acid



Analogously to Example 1d, 470 mg of the title compound was obtained from 485 mg of the ester prepared in Example 238b.

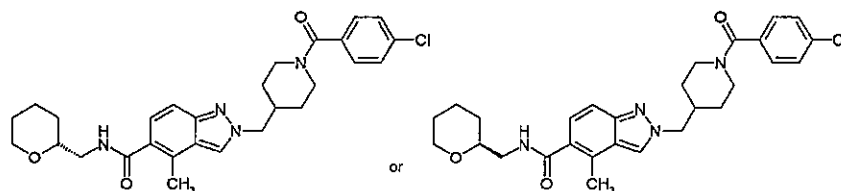
Example 239: (+/-)-2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-4-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 188 mg of the title compound was obtained from 300 mg of the compound prepared in Example 238c and 92 mg of 3,4,5,6-tetrahydro-2H-pyran-2-ylmethylamine hydrochloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.01-1.27 (3H), 1.29-1.64 (7H), 2.19-2.32 (1H), 2.48 (3H), 2.66-2.80 (1H), 2.89-3.00 (1H), 3.00-3.08 (1H), 3.16-3.25 (2H), 3.34-3.42 (1H), 3.43-3.55 (1H), 3.77-3.88 (1H), 4.31 (2H), 4.39 (1H), 7.14 (1H), 7.32-7.40 (2H), 7.46 (2H), 7.83 (1H), 8.09 (1H), 8.46 (1H).

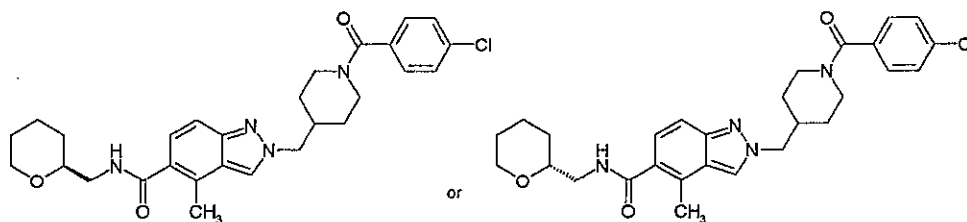
Example 240: (R or S)-2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-4-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2H-indazole-5-carboxamide



From 185 mg of the racemate prepared in Example 239, 51 mg of the title compound together with 53 mg of the slower-eluting enantiomer (Example 241) were obtained by racemate separation by means of preparative chiral HPLC (Method B).

Analytical chiral HPLC: 7.02 min.

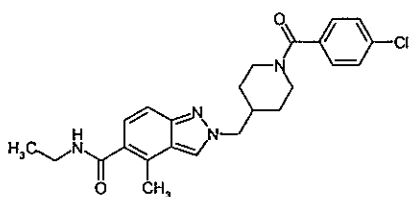
Example 241: (S or R)-2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-4-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2H-indazole-5-carboxamide



From 185 mg of the racemate prepared in Example 239, 53 mg of the title compound together
5 with 51 mg of the faster-eluting enantiomer (Example 240) were obtained by racemate separation by means of preparative chiral HPLC (Method B).

Analytical chiral HPLC: 8.24 min.

Example 242: 2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-N-ethyl-4-methyl-2H-indazole-5-carboxamide

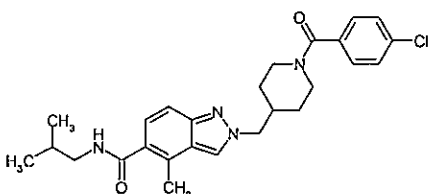


10

Analogously to Example 1, 85 mg of the title compound was obtained from 300 mg of the compound prepared in Example 238c and 43 mg of ethylamine 2M in THF.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.08 (3H), 1.16-1.60 (4H), 2.20-2.33 (1H), 2.48 (3H),
2.63-3.07 (2H), 3.22 (2H), 3.48 (1H), 4.24-4.46 (3H), 7.14 (1H), 7.31-7.42 (3H), 7.46 (2H), 8.08
15 (1H), 8.46 (1H).

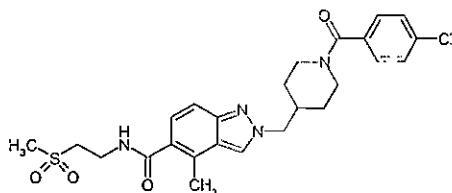
Example 243: 2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-N-isobutyl-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 92 mg of the title compound was obtained from 300 mg of the
20 compound prepared in Example 238c and 69 mg of isobutylamine.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.85-0.90 (6H), 1.13-1.57 (4H), 1.75-1.83 (1H), 2.21-2.34 (1H), 2.49 (3H), 3.03 (4H), 3.48 (1H), 4.27-4.45 (3H), 7.14 (1H), 7.31-7.41 (3H), 7.46 (2H), 8.11 (1H), 8.47 (1H).

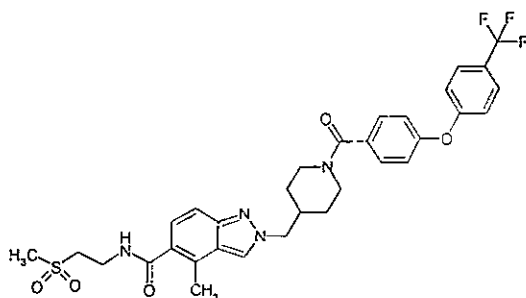
Example 244: 2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-N-(2-mesyloethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 132 mg of the title compound was obtained from 300 mg of the compound prepared in Example 238c and 151 mg of 2-mesyloethylamine hydrochloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.19-1.62 (4H), 2.24-2.36 (1H), 2.54 (3H), 2.70-3.05 (2H), 3.05 (3H), 3.38 (2H), 3.52 (1H), 3.65 (2H), 4.30-4.49 (3H), 7.22 (1H), 7.38 (2H), 7.42 (1H), 7.49 (2H), 8.33 (1H), 8.52 (1H).

Example 245: N-(2-mesyloethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide

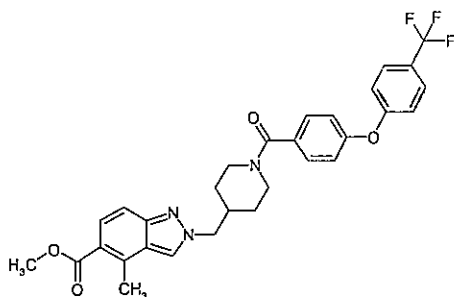


Analogously to Example 1, 67 mg of the title compound was obtained from 137 mg of the compound prepared in Example 245b and 41 mg of 2-mesyloethylamine hydrochloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.15-1.62 (4H), 2.22-2.33 (1H), 2.50 (3H), 2.68-3.01 (2H), 3.02 (2H), 3.35 (1H), 3.52-3.69 (3H), 4.28-4.47 (2H), 7.12 (1H), 7.15-7.22 (2H), 7.37-7.44 (2H), 7.73 (1H), 8.32 (1H), 8.50 (1H).

The starting material was prepared as follows:

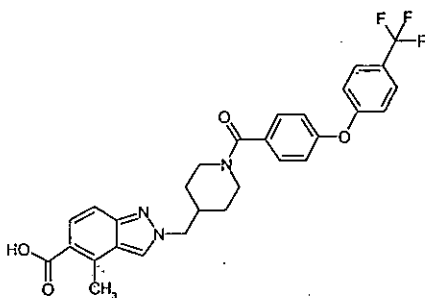
Example 245a: Methyl 4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxylate



Analogously to Example 1, 2.73 g of the title compound was obtained from 3.76 g of the compound prepared in Example 238a and 3.28 g of 4-[4-(trifluoromethyl)phenoxy]benzoic acid.

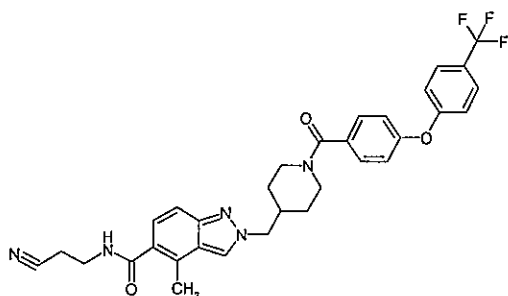
¹H-NMR (300 MHz, DMSO-d₆): δ = 1.25 (2H), 1.35-1.70 (2H), 2.20-2.37 (1H), 2.65-2.70 (3H),
 5 2.75-3.10 (2H), 3.52-3.72 (1H), 3.82 (4H), 4.36 (2H), 7.14 (2H), 7.20 (2H), 7.39-7.48 (3H), 7.66 (1H), 7.75 (2H), 8.70 (1H).

Example 245b: 4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)-methyl]-2H-indazol-5-carboxylic acid



10 Analogously to Example 1d, 1.1 g of the title compound was obtained from 2.73 g of the ester prepared in Example 245a.

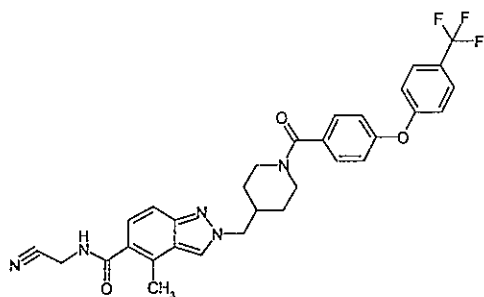
Example 246: N-(2-cyanoethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



15 Analogously to Example 1, 144 mg of the title compound was obtained from 137 mg of the compound prepared in Example 245b and 18 mg of 3-aminopropanenitrile.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.13-1.58 (4H), 2.21-2.33 (1H), 2.52 (3H), 2.75 (2H), 2.80-3.16 (2H), 3.44 (2H), 3.59 (1H), 4.27-4.50 (3H), 7.12 (2H), 7.15-7.21 (3H), 7.37-7.44 (3H), 7.73 (2H), 8.46 (1H), 8.51 (1H)

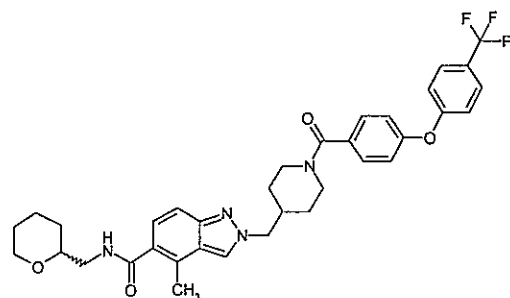
Example 247: N-(cyanomethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl} piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1, 125 mg of the title compound was obtained from 137 mg of the compound prepared in Example 245b and 24 mg of 2-aminoacetonitrile hydrochloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.16-1.60 (4H), 2.23-2.34 (1H), 2.53 (3H), 2.69-3.06 (2H), 3.59 (1H), 4.26 (2H), 4.29-4.50 (3H), 7.08-7.23 (5H), 7.38-7.46 (3H), 7.73 (2H), 8.54 (1H), 8.83 (1H).

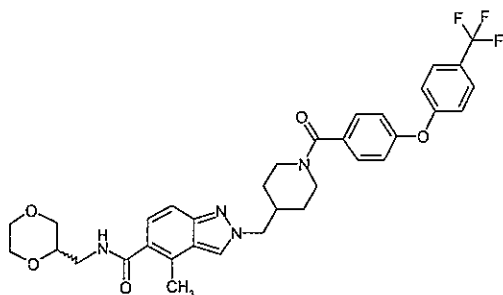
Example 248: (+/-)-4-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl} piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1, 231 mg of the title compound was obtained from 254 mg of the compound prepared in Example 245b and 72 mg of 3,4,5,6-tetrahydro-2H-pyran-2-ylmethylamine.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.01-1.66 (10H), 2.24-2.32 (1H), 2.70-3.15 (4H), 3.21 (3H), 3.34-3.43 (1H), 3.54-3.64 (1H), 3.76-3.89 (2H), 4.32 (3H), 7.09-7.21 (5H), 7.35-7.44 (3H), 7.73 (2H), 8.09 (1H), 8.47 (1H).

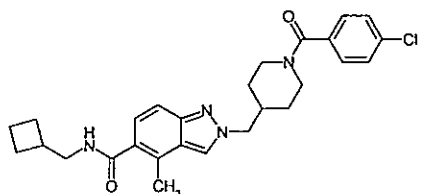
Example 249: (+/-)-N-(1,4-dioxan-2-ylmethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1, 280 mg of the title compound was obtained from 254 mg of the compound prepared in Example 245b and 73 mg of 1,4-dioxan-2-ylmethylamine.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.13-1.61 (4H), 2.23-2.32 (1H), 2.49 (3H), 2.75 (1H), 2.94-3.27 (4H), 3.38-3.77 (8H), 4.32 (2H), 7.09-7.21 (5H), 7.36-7.44 (3H), 7.73 (2H), 8.16 (1H), 8.48 (1H).

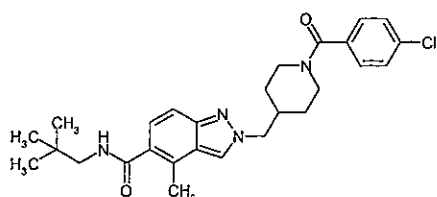
Example 250: 2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(cyclobutylmethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 79 mg of the title compound was obtained from 300 mg of the compound prepared in Example 238c and 115 mg of cyclobutylmethylamine hydrochloride.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.16-1.58 (4H), 1.64-1.86 (4H), 1.90-2.01 (2H), 2.19-2.32 (1H), 2.48 (3H), 2.50 (1H), 2.66-3.05 (2H), 3.19-3.25 (2H), 3.48 (1H), 4.27-4.47 (3H), 7.12 (1H), 7.31-7.40 (3H), 7.46 (2H), 8.09 (1H), 8.46 (1H).

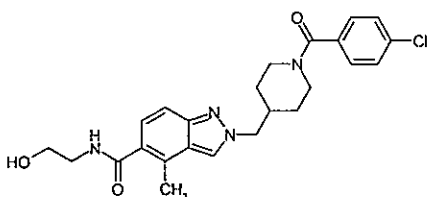
Example 251: 2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(2,2-dimethylpropyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 108 mg of the title compound was obtained from 300 mg of the compound prepared in Example 238c and 83 mg of 2,2-dimethylpropan-1-amine.

¹H-NMR (300 MHz, DMSO-d₆): δ = 0.91 (9H), 1.17-1.63 (4H), 2.24-2.36 (1H), 2.52 (3H), 2.70-3.04 (2H), 3.07 (2H), 3.52 (1H), 4.30-4.50 (3H), 7.17 (1H), 7.38 (2H), 7.42 (1H), 7.49 (2H), 8.07 (1H), 8.49 (1H).

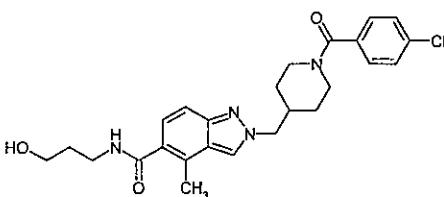
Example 252: 2-{{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(2-hydroxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 84 mg of the title compound was obtained from 370 mg of the compound prepared in Example 238c and 54 mg of 2-aminoethanol.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.14-1.64 (4H), 2.24-2.35 (1H), 2.53 (3H), 2.69-3.09 (2H), 3.30 (2H), 3.43-3.58 (3H), 4.27-4.50 (3H), 4.67 (1H), 7.20 (1H), 7.35-7.43 (3H), 7.49 (2H), 8.02 (1H), 8.49 (1H).

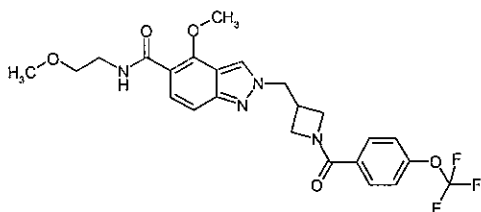
Example 253: 2-{{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(3-hydroxypropyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1, 95 mg of the title compound was obtained from 370 mg of the compound prepared in Example 238c and 66 mg of 3-aminopropan-1-ol.

¹H-NMR (300 MHz, DMSO-d₆): δ = 1.17-1.60 (4H), 1.66 (2H), 2.22-2.35 (1H), 2.53 (3H), 2.70-3.09 (2H), 3.27 (2H), 3.47 (2H), 3.47-3.67 (1H), 4.34 (2H), 4.41 (1H), 4.45 (1H), 7.17 (1H), 7.35-7.43 (3H), 7.49 (2H), 8.07 (1H), 8.49 (1H).

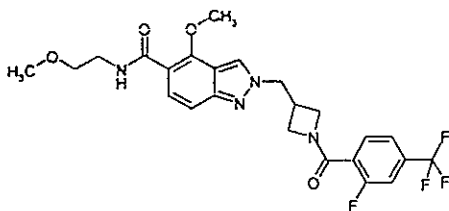
Example 254: 4-methoxy-N-(2-methoxyethyl)-2-({[1-(4-(trifluoromethoxy)benzoyl]azetidin-3-yl]methyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 90 mg of the title compound was obtained from 100 mg of the compound prepared in Example 214c and 97 mg of 4-(trifluoromethoxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 3.24 - 3.29 (1H), 3.29 - 3.32 (3H), 3.43 - 3.51 (4H),
 5 3.91 - 4.04 (1H), 4.10 - 4.18 (1H), 4.20 (3H), 4.22 - 4.29 (1H), 4.44 (1H), 4.70 (2H), 7.24 (1H),
 7.37 - 7.51 (2H), 7.66 (1H), 7.70 - 7.80 (2H), 8.22 (1H), 8.91 (1H).

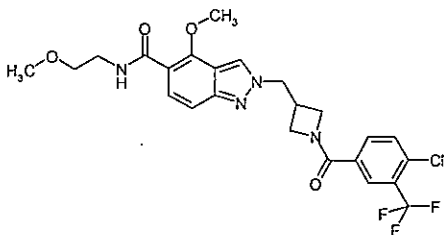
Example 255: 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-4-methoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



10 Analogously to Example 1, 40 mg of the title compound was obtained from 100 mg of the compound prepared in Example 214c and 98 mg of 2-fluoro-4-(trifluoromethyl)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 3.30 (4H), 3.44 - 3.50 (4H), 3.94 - 4.04 (2H), 4.15
 (2H), 4.20 (3H), 4.70 (2H), 7.24 (1H), 7.60 - 7.75 (3H), 7.78 - 7.89 (1H), 8.16 - 8.27 (1H), 8.90
 15 (1H).

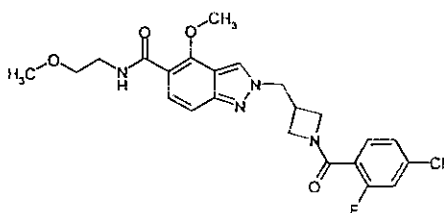
Example 256: 2-({1-[4-chloro-3-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-4-methoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 40 mg of the title compound was obtained from 100 mg of the compound prepared in Example 214c and 106 mg of 4-chloro-3-(trifluoromethyl)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 3.30 (4H), 3.43 - 3.53 (4H), 3.95 - 4.05 (1H), 4.20 (4H), 4.23 - 4.31 (1H), 4.47 (1H), 4.71 (2H), 7.23 (1H), 7.67 (1H), 7.78 - 7.85 (1H), 7.86 - 7.93 (1H), 7.96 (1H), 8.22 (1H), 8.90 (1H).

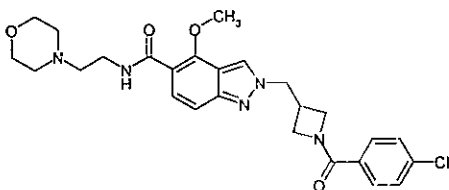
Example 257: 2-{[1-(4-chloro-2-fluorobenzoyl)azetidin-3-yl]methyl}-4-methoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



Analogously to Example 1, 20 mg of the title compound was obtained from 100 mg of the compound prepared in Example 214c and 82 mg of 4-chloro-2-fluorobenzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 3.30 (4H), 3.42 - 3.51 (4H), 3.98 (2H), 4.14 (2H), 4.20 (3H), 4.69 (2H), 7.24 (1H), 7.38 (1H), 7.48 (1H), 7.56 (1H), 7.66 (1H), 8.15 - 8.30 (1H), 8.90 (1H).

Example 258: 2-{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-4-methoxy-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide

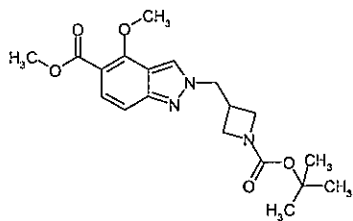


Analogously to Example 1b Version B, 67 mg of the title compound was obtained from 66 mg of the compound prepared in Example 258d and 32 mg of 2-morpholinoethylamine in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 2.44 (4H), 3.21 - 3.36 (3H), 3.42 (2H), 3.61 (4H), 3.88 - 4.04 (1H), 4.07 - 4.18 (1H), 4.23 (4H), 4.35 - 4.50 (1H), 4.70 (2H), 7.25 (1H), 7.41 - 7.56 (2H), 7.59 - 7.66 (2H), 7.70 (1H), 8.32 (1H), 8.91 (1H).

The starting material was prepared as follows:

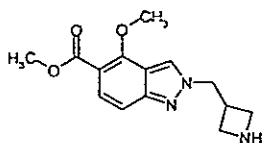
Example 258a: Methyl 2-{[1-(tert-butoxycarbonyl)azetidin-3-yl]methyl}-4-methoxy-2H-indazol-5-carboxylate



Analogously to Example 1e, 448 mg of the title compound was obtained from 750 mg of the compound prepared in Example 214a and 546 mg of methanol.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.27 - 1.44 (9H), 3.01 - 3.22 (1H), 3.78 (5H), 3.85 - 3.99 (2H), 4.14 (3H), 4.64 (2H), 7.24 (1H), 7.51 (1H), 8.94 (1H).

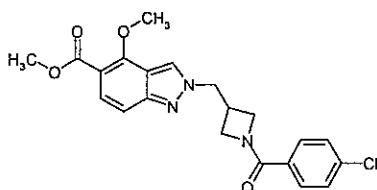
Example 258b: Methyl 2-(azetidin-3-ylmethyl)-4-methoxy-2H-indazol-5-carboxylate



800 mg of 258a were first placed in 40 ml acetone, treated with 40 ml of semiconcentrated hydrochloric acid, stirred until complete conversion and concentrated to dryness. Yield: 414 mg of the title compound.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 3.39 (1H), 3.79 (3H), 3.84 - 3.95 (2H), 3.96 - 4.07 (2H), 4.15 (3H), 4.72 (2H), 7.25 (1H), 7.52 (1H), 8.92 (1H).

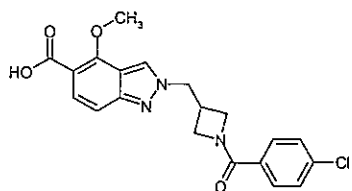
Example 258c: Methyl 2-{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-4-methoxy-2H-indazol-5-carboxylate



404 mg of 258b were first placed in 30 ml DCM at 0°C, treated with 0.21 ml of 4-chlorobenzoyl chloride and 0.75 ml of N-ethyl-diisopropylamine and stirred at room temperature until complete conversion. For the work-up, the reaction solution was treated with saturated sodium hydrogen carbonate solution, extracted with DCM and the combined organic phases dried with

sodium sulphate and concentrated to dryness. This yielded 608 mg of the title compound, which was used in the next step without further purification.

Example 258d: 2-{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-4-methoxy-2H-indazol-5-carboxylic acid



5

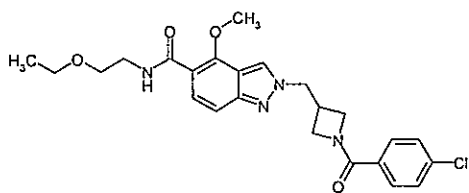
550 mg of 258c were first placed in 20 ml ethanol, treated with 20 ml of 2N sodium hydroxide and stirred until complete conversion. For the work-up, the reaction solution was adjusted to a pH of 3 with 1N hydrochloric acid, extracted with ethyl acetate, and the combined organic phases dried with sodium sulphate and concentrated to dryness. Yield after purification by column chromatography on silica gel with a hexane/ ethyl acetate gradient: 213 mg of the title compound.

10

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 3.27 (1H), 3.97 (1H), 4.07 - 4.17 (4H), 4.24 (1H), 4.42 (1H), 4.70 (2H), 7.22 (1H), 7.46 - 7.57 (3H), 7.59 - 7.68 (2H), 8.88 (1H), 12.61 (m, 1H).

15

Example 259: 2-{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-N-(2-ethoxyethyl)-4-methoxy-2H-indazole-5-carboxamide

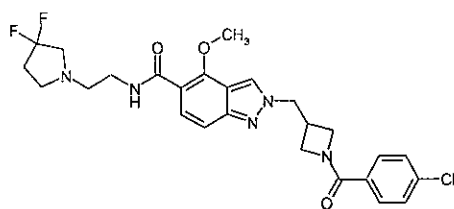


Analogously to Example 1b Version B, 30 mg of the title compound was obtained from 66 mg of the compound prepared in Example 258d and 22 mg of 2-ethoxyethylamine in DMF.

20

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.07 - 1.21 (3H), 3.30 (1H), 3.40 - 3.57 (6H), 3.90 - 4.02 (1H), 4.08 - 4.17 (1H), 4.21 (4H), 4.34 - 4.48 (1H), 4.70 (2H), 7.24 (1H), 7.45 - 7.57 (2H), 7.60 - 7.74 (3H), 8.24 (1H), 8.91 (1H).

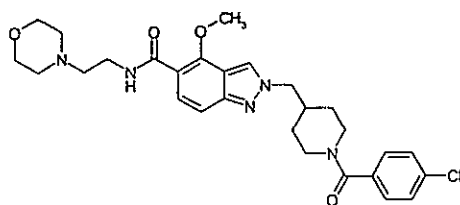
Example 260: 2-{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-N-[2-(3,3-difluoropyrrolidin-1-yl)ethyl]-4-methoxy-2H-indazole-5-carboxamide



Analogously to Example 1b Version B, 58 mg of the title compound was obtained from 66 mg of the compound prepared in Example 258d and 37 mg of 2-(3,3-difluoropyrrolidin-1-yl)ethylamine in DMF.

- 5 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 2.27 (2H), 2.64 (2H), 2.77 (2H), 2.96 (2H), 3.22 - 3.30 (1H), 3.41 (2H), 3.92 - 4.01 (1H), 4.11 (1H), 4.19 (4H), 4.35 - 4.48 (1H), 4.70 (2H), 7.24 (1H), 7.51 (2H), 7.57 - 7.75 (3H), 8.32 (1H), 8.90 (1H).

Example 261: 2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-4-methoxy-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide



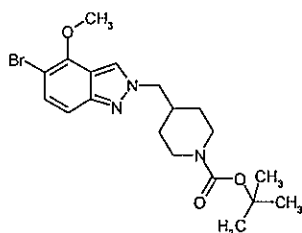
10

Analogously to Example 1b, Version B, 77 mg of the title compound was obtained from 100 mg of the compound prepared in Example 261e and 46 mg of 2-morpholinoethylamine in DMF.

- 15 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.11 - 1.33 (2H), 1.35 - 1.75 (2H), 2.18 - 2.39 (1H), 2.65 - 2.88 (2H), 2.90 - 3.23 (3H), 3.41 - 3.82 (8H), 3.84 - 4.12 (2H), 4.24 (3H), 4.35 (3H), 7.26 (1H), 7.33 - 7.44 (2H), 7.46 - 7.58 (2H), 7.69 (1H), 8.27 - 8.52 (1H), 8.87 (1H).

The starting material was prepared as follows:

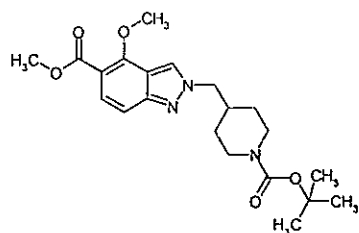
Example 261a: Tert-butyl 4-[(5-bromo-4-methoxy-2H-indazol-2-yl)methyl]piperidin-1-carboxylate



Analogously to Example 1c, 5.57 g of the title compound was obtained from 9.65 g of 5-bromo-4-methoxy-1H-indazole and 23.55 g of tert-butyl-4-[(tosyloxy)methyl]piperidin-1-carboxylate.

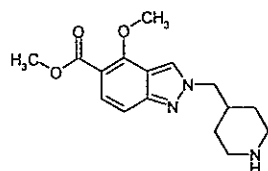
¹H-NMR (400 MHz, chloroform-d): δ [ppm]= 1.16 - 1.31 (2H), 1.39 - 1.49 (9H), 1.52 - 1.64 (2H), 2.16 - 2.36 (1H), 2.56 - 2.80 (2H), 4.11 (5H), 4.26 (2H), 7.28 - 7.33 (1H), 7.34 - 7.39 (1H),
5 7.99 (1H).

Example 261b: Methyl 2-{[1-(tert-butoxycarbonyl)piperidin-4-yl]methyl}-4-methoxy-2H-indazol-5-carboxylate



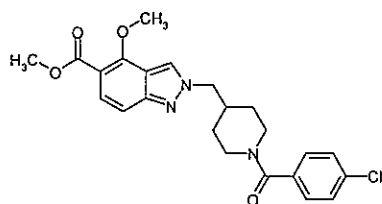
Analogously to Example 1e, 1.1 g of the title compound was obtained from 2.4 g of the
10 compound prepared in Example 261a and 1.63 g of methanol.

Example 261c: Methyl 4-methoxy-2-(4-piperidylmethyl)-2H-indazol-5-carboxylate



Analogously to Example 258b, from 1.1 g of the compound prepared in Example 261b 1.1 g of the title compound was obtained, which was reacted in the next step without further purification.

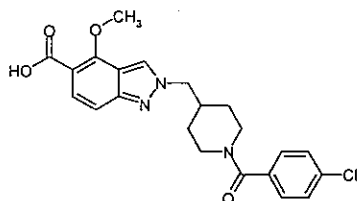
15 **Example 261d:** Methyl 2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methoxy-2H-indazol-5-carboxylate



Analogously to Example 258c, from 1.1 g of the compound prepared in Example 261c and 762 mg of 4-chlorobenzoyl chloride, 980 mg of the title compound was obtained after
20 purification by column chromatography on silica gel with a hexane/ethyl acetate gradient.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.24 (2H), 1.36 - 1.76 (2H), 2.18 - 2.40 (1H), 2.65 - 2.88 (1H), 2.91 - 3.12 (1H), 3.42 - 3.67 (1H), 3.78 (3H), 4.05 - 4.21 (3H), 4.35 (3H), 7.24 (1H), 7.34 - 7.43 (2H), 7.45 - 7.56 (3H), 8.86 (1H).

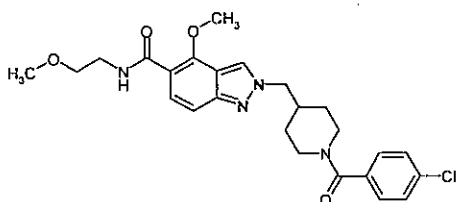
Example 261e: 2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methoxy-2H-indazol-5-carboxylic acid



Analogously to Example 258d, 547 mg of the title compound was obtained from 980 mg of the compound prepared in Example 261d.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.12 - 1.35 (2H), 1.35 - 1.73 (2H), 2.21 - 2.40 (1H), 2.63 - 3.12 (2H), 3.14 - 3.70 (3H), 4.34 (3H), 5.76 (s, 1H), 7.21 (1H), 7.32 - 7.45 (2H), 7.46 - 7.58 (3H), 8.76 (1H).

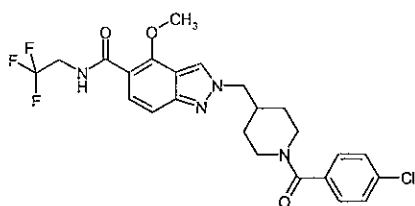
Example 262: 2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 50 mg of the title compound was obtained from 100 mg of the compound prepared in Example 261e and 26 mg of 2-methoxyethylamine in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.11 - 1.35 (2H), 1.35 - 1.72 (2H), 2.21 - 2.39 (1H), 2.69 - 2.88 (1H), 2.92 - 3.11 (1H), 3.30 (s, 3H), 3.38 - 3.64 (5H), 4.20 (3H), 4.34 (3H), 7.25 (1H), 7.34 - 7.43 (2H), 7.45 - 7.57 (2H), 7.66 (1H), 8.22 (1H), 8.83 (1H).

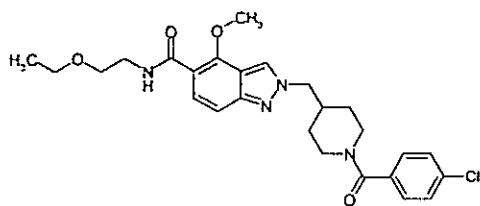
Example 263: 2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methoxy-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 68 mg of the title compound was obtained from 100 mg of the compound prepared in Example 261e and 35 mg of 2,2,2-trifluoroethylamine in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.09 - 1.35 (2H), 1.37 - 1.75 (2H), 2.21 - 2.41 (1H),
 5 2.69 - 2.88 (1H), 2.92 - 3.19 (1H), 3.42 - 3.69 (1H), 4.13 (2H), 4.23 (3H), 4.35 (3H), 7.26 (1H),
 7.34 - 7.45 (2H), 7.46 - 7.55 (2H), 7.61 (1H), 8.59 (1H), 8.88 (1H).

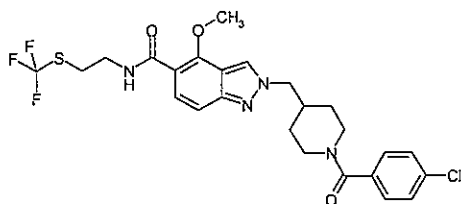
Example 264: 2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-N-(2-ethoxyethyl)-4-methoxy-2H-indazole-5-carboxamide



10 Analogously to Example 1b, Version B, 60 mg of the title compound was obtained from 100 mg of the compound prepared in Example 261e and 31 mg of 2-ethoxyethylamine in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.14 (3H), 1.19 - 1.35 (2H), 1.37 - 1.72 (2H), 2.23 -
 2.43 (1H), 2.68 - 2.88 (1H), 2.91 - 3.15 (1H), 3.41 - 3.56 (m, 7H), 4.20 (3H), 4.34 (3H), 7.25
 (1H), 7.33 - 7.45 (2H), 7.46 - 7.55 (2H), 7.67 (1H), 8.23 (1H), 8.83 (1H).

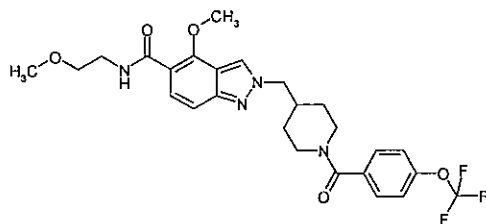
15 **Example 265:** 2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-4-methoxy-N-{2-[(trifluoromethyl)sulphonyl]ethyl}-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 68 mg of the title compound was obtained from 100 mg of the compound prepared in Example 261e and 51 mg of 2-[(trifluoromethyl)sulphonyl]-
 20 ethylamine in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.14 - 1.35 (2H), 1.36 - 1.72 (2H), 2.18 - 2.40 (1H), 2.66 - 2.87 (1H), 2.91 - 3.11 (1H), 3.21 (2H), 3.42 - 3.70 (3H), 4.22 (3H), 4.34 (3H), 7.24 (1H), 7.32 - 7.45 (2H), 7.46 - 7.55 (2H), 7.64 (1H), 8.45 (1H), 8.85 (1H).

Example 266: 4-methoxy-N-(2-methoxyethyl)-2-({1-[4-(trifluoromethoxy)benzoyl]piperidin-4-yl)methyl)-2H-indazole-5-carboxamide

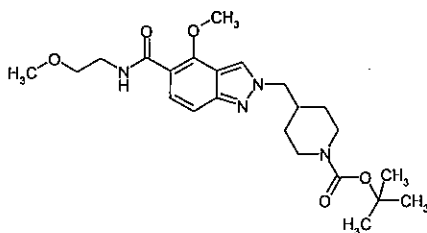


Analogously to Example 1b, Version B, 84 mg of the title compound was obtained from 100 mg of the compound prepared in Example 266b and 89 mg of 4-(trifluoromethoxy)benzoic acid in DMF.

LC-MS: R_t = 1.16 min, MS (ES⁺): m/z = 435 (M+H)⁺.

The starting material was prepared as follows:

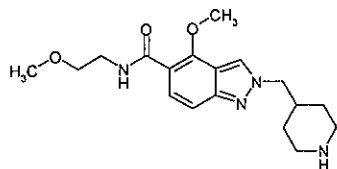
Example 266a: Tert-butyl 4-({4-methoxy-5-[N-(2-methoxyethyl)carbamoyl]-2H-indazol-2-yl)methyl)piperidin-1-carboxylate



3 g of compound 261a, 1.59 g of 2-methoxyethylamine, 1.87 g of molybdenum hexacarbonyl, 204 mg of tri-tert-butylphosphine tetrafluoroborate and 316 mg of palladium(II) acetate were first suspended in 100 ml 1,4-dioxan. Then 2.25 g of sodium carbonate and a few drops of water were added, and the mixture stirred for 25 mins at 140°C and 150 watts in the microwave. The reaction mixture was concentrated and the residue was purified by column chromatography on silica gel using a hexane/ ethyl acetate/ methanol gradient. Yield: 1.3 g of the title compound.

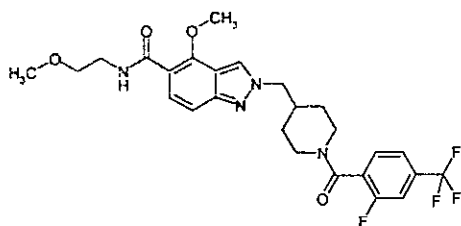
¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.99 - 1.22 (2H), 1.29 - 1.55 (11H), 2.03 - 2.33 (1H), 2.60 - 2.83 (2H), 3.28 - 3.31 (3H), 3.43 - 3.52 (4H), 3.82 - 4.03 (2H), 4.20 (3H), 4.31 (2H), 7.25 (1H), 7.66 (1H), 8.21 (1H), 8.82 (1H).

Example 266b: 4-methoxy-N-(2-methoxyethyl)-2-(4-piperidylmethyl)-2H-indazol-5-carboxamide



Analogously to Example 258b, from 1.3 g of the compound prepared in Example 266a, 1.48 g of the title compound was obtained, which was used in the subsequent reactions without further purification.

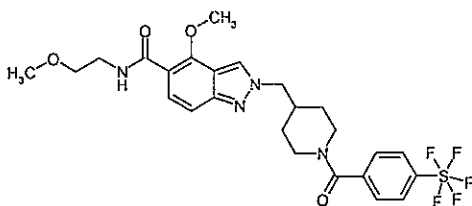
Example 267: 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-4-methoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 20 mg of the title compound was obtained from 100 mg of the compound prepared in Example 266b and 90 mg of 2-fluoro-4-(trifluoromethyl)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.05 - 1.36 (2H), 1.39 - 1.53 (1H), 1.56 - 1.72 (1H), 2.21 - 2.40 (1H), 2.75 - 2.90 (1H), 2.92 - 3.15 (1H), 3.27 - 3.35 (4H), 3.44 - 3.53 (4H), 4.20 (3H), 4.27 - 4.43 (2H), 4.44 - 4.58 (1H), 7.25 (1H), 7.54 - 7.74 (3H), 7.81 (1H), 8.20 (1H), 8.83 (1H).

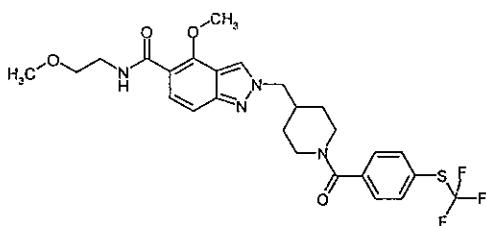
Example 268: 4-methoxy-N-(2-methoxyethyl)-2-({1-[4-(pentafluoro-λ⁶-sulphonyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 39 mg of the title compound was obtained from 100 mg of the compound prepared in Example 266b and 107 mg of 4-(pentafluoro- λ^6 -sulphanyl)benzoic acid in DMF.

LC-MS: R_t = 1.18 min, MS (ES⁺): m/z = 577 (M+H)⁺.

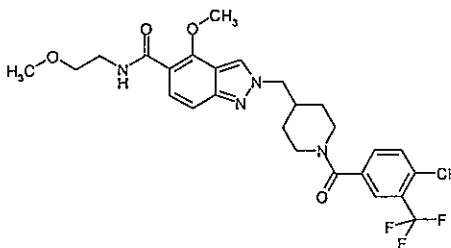
- 5 **Example 269:** 4-methoxy-N-(2-methoxyethyl)-2-[(1-{4-[(trifluoromethyl)sulphanyl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



- 10 Analogously to Example 1b, Version B, 25 mg of the title compound was obtained from 100 mg of the compound prepared in Example 266b and 96 mg of 4-[(trifluoromethyl)sulphanyl]benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm] = 1.09 - 1.37 (2H), 1.38 - 1.72 (2H), 2.23 - 2.39 (1H), 2.70 - 2.93 (1H), 2.95 - 3.16 (1H), 3.30 (3H), 3.40 - 3.55 (5H), 4.20 (3H), 4.35 (d, 3H), 7.25 (1H), 7.46 - 7.58 (2H), 7.66 (1H), 7.78 (2H), 8.20 (1H), 8.82 (1H).

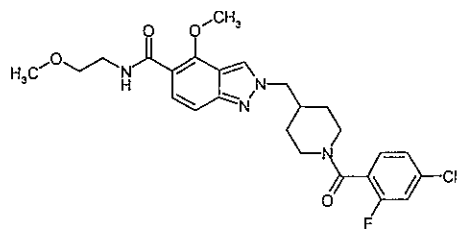
- 15 **Example 270:** 2-({1-[4-chloro-3-(trifluoromethyl)benzoyl]piperidin-4-yl)methyl}-4-methoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 27 mg of the title compound was obtained from 100 mg of the compound prepared in Example 266b and 97 mg of 4-chloro-3-(trifluoromethyl)benzoic acid in DMF.

- 20 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm] = 1.15 - 1.37 (2H), 1.38 - 1.73 (2H), 2.14 - 2.41 (1H), 2.70 - 2.88 (1H), 2.95 - 3.18 (1H), 3.30 (3H), 3.39 - 3.62 (5H), 4.20 (3H), 4.35 (3H), 7.25 (1H), 7.59 - 7.73 (2H), 7.75 - 7.86 (2H), 8.20 (1H), 8.82 (1H).

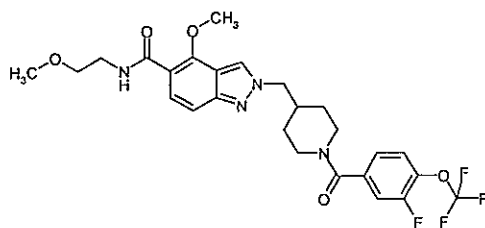
Example 271: 2-([1-(4-chloro-2-fluorobenzoyl)piperidin-4-yl]methyl)-4-methoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 17 mg of the title compound was obtained from 100 mg of the compound prepared in Example 266b and 75 mg of 4-chloro-2-fluorobenzoic acid in DMF.

$^1\text{H-NMR}$ (400 MHz, DMSO-d_6): δ [ppm] = 1.08 - 1.33 (2H), 1.37 - 1.52 (1H), 1.53 - 1.70 (1H), 2.20 - 2.42 (1H), 2.72 - 2.88 (1H), 2.94 - 3.12 (1H), 3.30 (3H), 3.37 (1H), 3.43 - 3.51 (4H), 4.20 (3H), 4.34 (2H), 4.42 - 4.55 (1H), 7.25 (1H), 7.31 - 7.48 (2H), 7.55 (1H), 7.66 (1H), 8.20 (1H), 8.83 (1H).

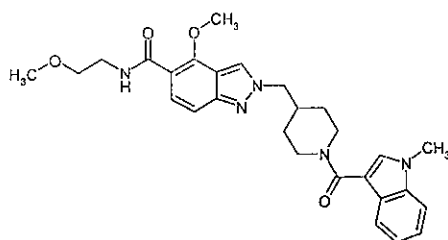
Example 272: 2-([1-[3-fluoro-4-(trifluoromethoxy)benzoyl]piperidin-4-yl]methyl)-4-methoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 12 mg of the title compound was obtained from 100 mg of the compound prepared in Example 266b and 97 mg of 3-fluoro-4-(trifluoromethoxy)benzoic acid in DMF.

LC-MS: R_t = 1.18 min, MS (ES⁺): m/z = 553 ($M+H$)⁺.

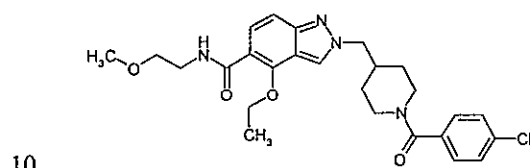
Example 273: 4-methoxy-N-(2-methoxyethyl)-2-([1-[(1-methyl-1H-indol-3-yl)carbonyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 19 mg of the title compound was obtained from 100 mg of the compound prepared in Example 266b and 76 mg of 1-methyl-1H-indol-3-carboxylic acid in DMF.

- 5 $^1\text{H-NMR}$ (300 MHz, DMSO-d_6): δ [ppm]= 1.11 - 1.40 (2H), 1.54 (2H), 2.18 - 2.42 (1H), 2.93 (2H), 3.30 (3H), 3.40 - 3.45 (4H), 3.70 - 3.91 (3H), 4.12 - 4.48 (7H), 7.00 - 7.34 (3H), 7.48 (1H), 7.66 (3H), 8.22 (1H), 8.84 (1H).

Example 274: 2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-4-ethoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



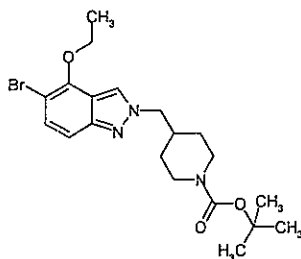
10

Analogously to Example 1b, Version B, 47 mg of the title compound was obtained from 110 mg of the compound prepared in Example 274c and 72 mg of 4-chlorobenzoic acid in DMF.

- 15 $^1\text{H-NMR}$ (400 MHz, DMSO-d_6): δ [ppm]= 1.10 - 1.33 (2H), 1.44 (5H), 2.20 - 2.39 (1H), 2.68 - 2.85 (1H), 2.92 - 3.12 (1H), 3.32 (7H), 3.51 - 3.62 (1H), 4.20 - 4.59 (5H), 7.26 (1H), 7.33 - 7.44 (2H), 7.47 - 7.55 (2H), 7.70 (1H), 8.30 (1H), 8.79 (1H).

The starting material was prepared as follows:

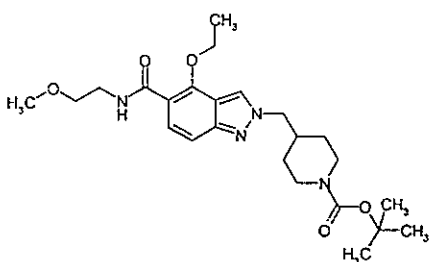
Example 274a: Tert-butyl 4-[(5-bromo-4-ethoxy-2H-indazol-2-yl)methyl]piperidin-1-carboxylate



Analogously to Example 74c, 285 mg of the title compound was obtained from 484 mg of 5-bromo-4-ethoxy-1H-indazole and 1.11 g of tert-butyl-4-[(5-bromo-4-ethoxy-2H-indazol-2-yl)-methyl]piperidin-1-carboxylate.

¹H-NMR (400 MHz, chloroform-d): δ [ppm]= 1.13 - 1.34 (2H), 1.46 (9H), 1.49 (3H), 1.56 - 1.63 (2H), 2.16 - 2.37 (1H), 2.54 - 2.80 (2H), 4.02 - 4.21 (2H), 4.26 (2H), 4.33 (2H), 7.28 - 7.43 (2H), 7.93 (1H).

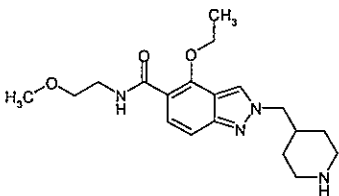
Example 274b: Tert-butyl 4-({4-ethoxy-5-[N-(2-methoxyethyl)carbamoyl]-2H-indazol-2-yl}-methyl)piperidin-1-carboxylate



Analogously to Example 266a, 235 mg of the title compound was obtained from 285 mg of the compound prepared in Example 274a and 146 mg of 2-methoxyethylamine.

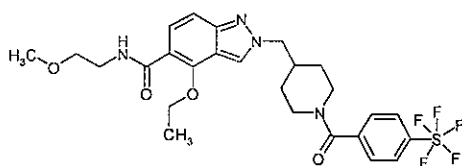
¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.97 - 1.22 (2H), 1.29 - 1.56 (14H), 2.06 - 2.31 (1H), 2.59 - 2.86 (2H), 3.33 (3H), 3.49 (4H), 3.80 - 4.02 (2H), 4.30 (2H), 4.51 (2H), 7.26 (1H), 7.71 (1H), 8.31 (1H), 8.79 (1H).

Example 274c: 4-ethoxy-N-(2-methoxyethyl)-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide



Analogously to Example 258b, from 235 mg of the compound prepared in Example 274b, 237 mg of the title compound was obtained, which was reacted in the next step without further purification.

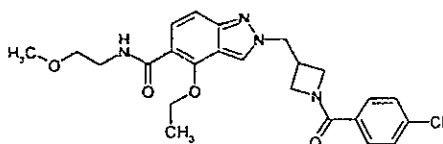
Example 275: 4-ethoxy-N-(2-methoxyethyl)-2-({1-[4-(pentafluoro- λ^6 -sulphonyl)benzoyl]-piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 30 mg of the title compound was obtained from 110 mg of the compound prepared in Example 274c and 113 mg of 4-(pentafluoro- λ^6 -sulphonyl)benzoic acid in DMF.

- 5 $^1\text{H-NMR}$ (400 MHz, DMSO-d_6): δ [ppm]= 1.10 - 1.36 (2H), 1.44 (4H), 1.54 - 1.69 (1H), 2.22 - 2.40 (1H), 2.70 - 2.87 (1H), 2.93 - 3.13 (1H), 3.30 (3H), 3.48 (5H), 4.34 (2H), 4.41 - 4.63 (3H), 7.26 (1H), 7.59 (2H), 7.70 (1H), 7.98 (2H), 8.30 (1H), 8.79 (1H).

Example 276: 2-{[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-4-ethoxy-N-(2-methoxyethyl)-2H-indazole-5-carboxamide



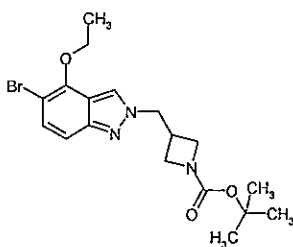
10

Analogously to Example 1b, Version B, 25 mg of the title compound was obtained from 50 mg of the compound prepared in Example 276c and 35 mg of 4-chlorobenzoic acid in DMF.

- 15 $^1\text{H-NMR}$ (400 MHz, DMSO-d_6): δ [ppm]= 1.44 (3H), 3.21 - 3.29 (1H), 3.31 (3H), 3.49 (4H), 3.96 (1H), 4.06 - 4.30 (2H), 4.34 - 4.59 (3H), 4.69 (2H), 7.25 (1H), 7.51 (2H), 7.62 (2H), 7.71 (1H), 8.30 (1H), 8.86 (1H).

The starting material was prepared as follows:

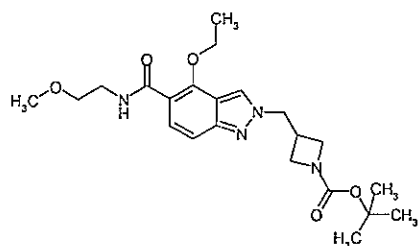
Example 276a: Tert-butyl 3-[(5-bromo-4-ethoxy-2H-indazol-2-yl)methyl]azetidin-1-carboxylate



- 20 Analogously to Example 74c, 204 mg of the title compound was obtained from 500 mg of 5-bromo-4-ethoxy-1H-indazole and 1.06 g of tert-butyl-3-[(tosyloxy)methyl]azetidin-1-carboxylate.

¹H-NMR (600 MHz, chloroform-d): δ [ppm]= 1.44 (9H), 1.49 (3H), 3.14 - 3.30 (1H), 3.72 - 3.83 (2H), 4.07 (2H), 4.33 (2H), 4.59 (2H), 7.29 (1H), 7.37 (1H), 7.98 (1H).

Example 276b: Tert-butyl 3-({4-ethoxy-5-[N-(2-methoxyethyl)carbamoyl]-2H-indazol-2-yl}-methyl)azetidin-1-carboxylate

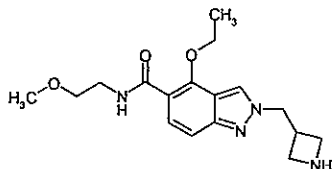


5

Analogously to Example 266a, from 191 mg of the compound prepared in Example 276a and 105 mg of 2-methoxyethylamine, 121 mg of the title compound was obtained, which was reacted in the next step without further purification.

Example 276c: 2-(azetidin-3-ylmethyl)-4-ethoxy-N-(2-methoxyethyl)-2H-indazol-5-carboxamide

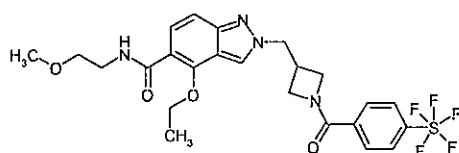
10



Analogously to Example 258b, from 121 mg of the compound prepared in Example 276b, 93 mg of the title compound was obtained, which was reacted in the next step without further purification.

Example 277: 4-ethoxy-N-(2-methoxyethyl)-2-({1-[4-(pentafluoro- λ^6 -sulphonyl)benzoyl]-azetidin-3-yl}methyl)-2H-indazole-5-carboxamide

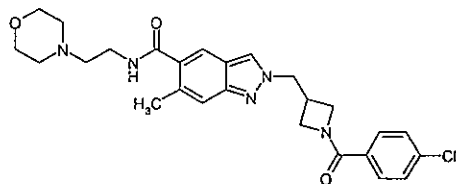
15



Analogously to Example 1b, Version B, 35 mg of the title compound was obtained from 50 mg of 276c and 56 mg of 4-(pentafluoro- λ^6 -sulphonyl)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.37 - 1.50 (3H), 3.26 (1H), 3.31 (3H), 3.43 - 3.57 (4H), 3.92 - 4.05 (1H), 4.10 - 4.31 (2H), 4.38 - 4.57 (3H), 4.70 (2H), 7.25 (1H), 7.66 - 7.87 (3H), 7.98 (2H), 8.30 (1H), 8.86 (1H).

Example 278: 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-6-methyl-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide

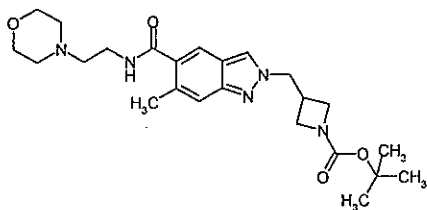


Analogously to Example 1b, Version B, 23 mg of the title compound was obtained from 100 mg of 278b and 66 mg of 4-chlorobenzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 2.42 (s, 3H), 3.09 - 3.27 (m, 3H), 3.29 - 3.33 (2H), 3.49 - 3.73 (m, 6H), 3.84 - 3.96 (m, 1H), 3.97 - 4.15 (m, 3H), 4.16 - 4.27 (m, 1H), 4.31 - 4.45 (m, 1H), 4.62 - 4.80 (m, 2H), 7.36 - 7.47 (1H), 7.48 - 7.57 (2H), 7.58 - 7.70 (2H), 7.77 - 7.89 (1H), 8.41 - 8.58 (2H).

The starting material was prepared as follows:

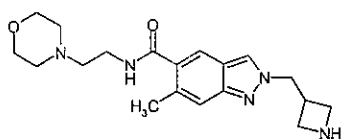
Example 278a: Tert-butyl 3-({6-methyl-5-[N-(2-morpholinoethyl)carbamoyl]-2H-indazol-2-yl}methyl)azetidin-1-carboxylate



Analogously to Example 1b, Version B, 418 mg of the title compound was obtained from 410 mg of 178c and 232 mg of 2-morpholinoethylamine in DMF.

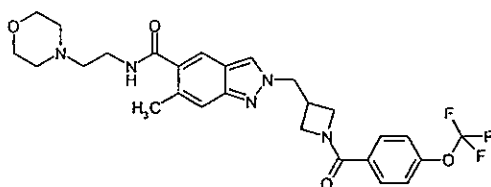
¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.36 (9H), 2.39 (3H), 2.44 (4H), 2.95 - 3.17 (1H), 3.24 - 3.43 (4H), 3.58 (4H), 3.64 - 3.79 (2H), 3.88 (2H), 4.62 (2H), 7.39 (1H), 7.68 (1H), 8.04 - 8.22 (1H), 8.45 (1H).

Example 278b: 2-(azetidin-3-ylmethyl)-6-methyl-N-(2-morpholinoethyl)-2H-indazole-5-carboxamide



Analogously to Example 258b, from 400 mg of the compound prepared in Example 278a, 372 mg of the title compound was obtained, which was reacted in the next step without further purification.

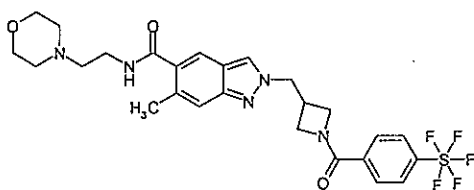
- 5 **Example 279:** 6-methyl-N-(2-morpholinoethyl)-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 44 mg of the title compound was obtained from 100 mg of 278b and 86 mg of 4-(trifluoromethoxy)benzoic acid in DMF.

- 10 ¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 2.42 (3H), 3.07 - 3.27 (5H), 3.48 - 3.74 (6H), 3.85 - 4.17 (4H), 4.18 - 4.29 (1H), 4.33 - 4.48 (1H), 4.61 - 4.82 (2H), 7.30 - 7.54 (3H), 7.68 - 7.78 (2H), 7.79 - 7.88 (1H), 8.43 - 8.60 (2H).

Example 280: 6-methyl-N-(2-morpholinoethyl)-2-({1-[4-(pentafluoro-λ⁶-sulphanyl)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide

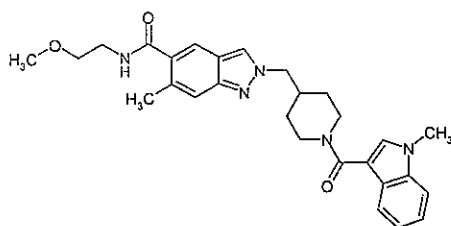


15

Analogously to Example 1b, Version B, 37 mg of the title compound was obtained from 100 mg of 278b and 104 mg of 4-(pentafluoro-λ⁶-sulphanyl)benzoic acid in DMF.

- 20 ¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 2.42 (3H), 3.06 - 3.27 (5H), 3.47 - 3.75 (6H), 3.87 - 4.08 (3H), 4.09 - 4.29 (2H), 4.33 - 4.49 (1H), 4.61 - 4.80 (2H), 7.34 - 7.52 (1H), 7.71 - 7.88 (3H), 7.92 - 8.07 (2H), 8.39 - 8.59 (2H).

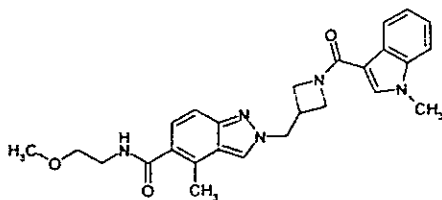
Example 281: N-(2-methoxyethyl)-6-methyl-2-({1-[(1-methyl-1H-indol-3-yl)carbonyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 4.2 mg of the title compound was obtained from 150 mg of 205c and 119 mg of 1-methyl-1H-indol-3-carboxylic acid in DMF.

LC-MS: $R_t = 1.01$ min, MS (ES⁺): $m/z = 489$ (M+H)⁺.

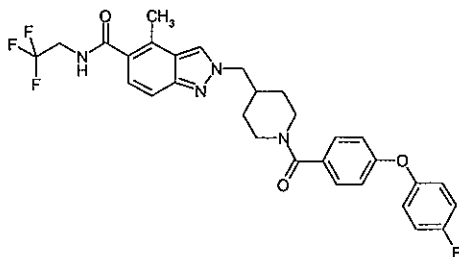
- 5 **Example 282:** N-(2-methoxyethyl)-4-methyl-2-({1-[(1-methyl-1H-indol-3-yl)carbonyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 35 mg of the title compound was obtained from 175 mg of 117e and 122 mg of 1-methyl-1H-indol-3-carboxylic acid in DMF.

- 10 LC-MS: $R_t = 0.95$ min, MS (ES⁺): $m/z = 460$ (M+H)⁺.

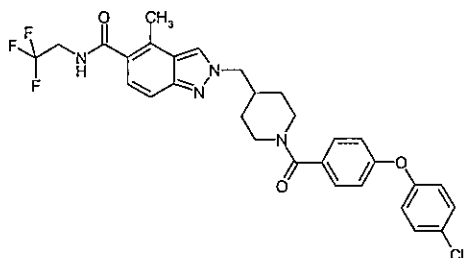
Example 283: 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



- 15 Analogously to Example 1b, Version B, 267 mg of the title compound was obtained from 250 mg of 71a and 246 mg of 4-(4-fluorophenoxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm] = 1.16 - 1.32 (2H), 1.35 - 1.69 (2H), 2.20 - 2.39 (1H), 2.54 (3H), 2.70 - 3.15 (2H), 3.50 - 3.83 (1H), 4.07 (2H), 4.36 (3H), 6.91 - 7.05 (2H), 7.08 - 7.31 (5H), 7.34 - 7.42 (2H), 7.46 (1H), 8.56 (1H), 8.83 (1H).

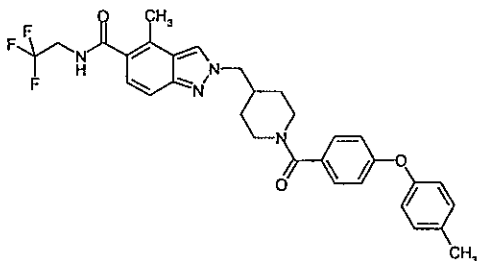
Example 284: 2-({1-[4-(4-chlorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 55 mg of the title compound was obtained from 100 mg of 71a and 105 mg of 4-(4-chlorophenoxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.34 - 1.72 (2H), 2.21 - 2.41 (1H), 2.54 (3H), 2.67 - 3.20 (2H), 3.51 - 3.83 (1H), 4.07 (2H), 4.36 (3H), 6.92 - 7.15 (4H), 7.20 (1H), 7.35 - 7.53 (5H), 8.56 (1H), 8.84 (1H).

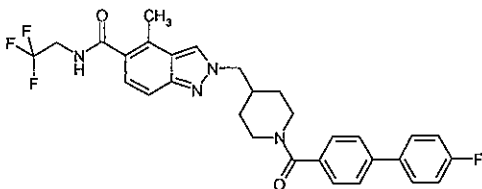
Example 285: 4-methyl-2-({1-[4-(4-methylphenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 60 mg of the title compound was obtained from 100 mg of 71a and 97 mg of 4-(4-methylphenoxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.36 - 1.71 (2H), 2.30 (4H), 2.54 (3H), 2.64 - 3.17 (2H), 3.49 - 3.82 (1H), 4.07 (2H), 4.36 (3H), 6.85 - 7.04 (4H), 7.21 (3H), 7.29 - 7.41 (2H), 7.46 (1H), 8.56 (1H), 8.84 (s1H).

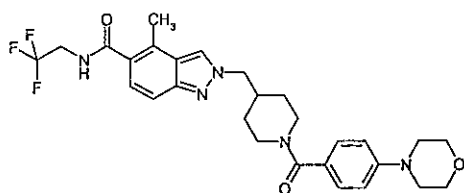
Example 286: 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 39 mg of the title compound was obtained from 100 mg of 71a and 92 mg of 4-(4-fluorophenyl)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.14 - 1.36 (2H), 1.36 - 1.73 (2H), 2.18 - 2.41 (1H), 2.54 (3H), 2.70 - 3.16 (2H), 3.51 - 3.82 (1H), 4.07 (2H), 4.38 (3H), 7.21 (1H), 7.31 (2H), 7.40 - 7.53 (3H), 7.64 - 7.85 (4H), 8.56 (1H), 8.81 (1H).

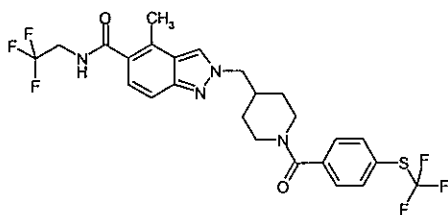
Example 287: 4-methyl-2-{{1-[4-morpholinobenzoyl]piperidin-4-yl)methyl}-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 45 mg of the title compound was obtained from 100 mg of 71a and 88 mg of 4-morpholinobenzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.24 (2H), 1.40 - 1.60 (2H), 2.20 - 2.40 (1H), 2.54 (3H), 2.73 - 3.01 (2H), 3.38 (4H), 3.64 - 3.81 (5H), 4.07 (3H), 4.36 (2H), 6.94 (2H), 7.16 - 7.30 (3H), 7.46 (1H), 8.55 (1H), 8.81 (1H).

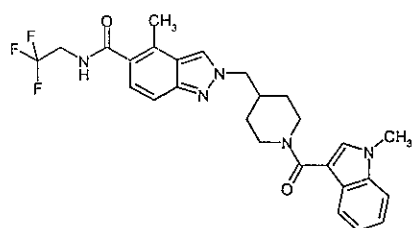
Example 288: 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[(trifluoromethyl)sulphonyl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 67 mg of the title compound was obtained from 100 mg of 71a and 94 mg of 4-[(trifluoromethyl)sulphonyl]benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.12 - 1.35 (2H), 1.35 - 1.70 (2H), 2.21 - 2.40 (1H), 2.54 (3H), 2.70 - 2.88 (1H), 2.94 - 3.12 (1H), 3.38 - 3.61 (1H), 4.07 (2H), 4.36 (3H), 7.21 (1H), 7.37 - 7.60 (3H), 7.78 (2H), 8.55 (1H), 8.81 (1H).

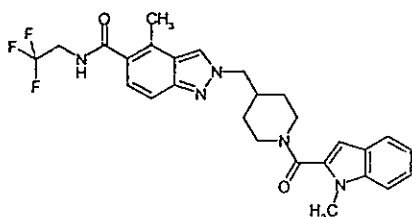
Example 289: 4-methyl-2-({1-[(1-methyl-1H-indol-3-yl)carbonyl]piperidin-4-yl)methyl}-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 43 mg of the title compound was obtained from 100 mg of 71a and 74 mg of 1-methyl-1H-indol-3-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.27 (2H), 1.52 (2H), 2.20 - 2.40 (1H), 2.54 (3H),
 5 2.92 (2H), 3.82 (3H), 4.07 (2H), 4.27 (2H), 4.38 (2H), 6.92 - 7.33 (3H), 7.47 (2H), 7.61 - 7.76 (2H), 8.56 (1H), 8.81 (1H).

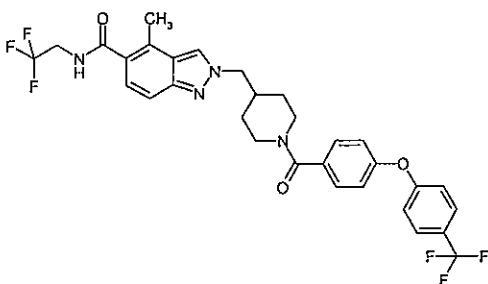
Example 290: 4-methyl-2-({1-[(1-methyl-1H-indol-2-yl)carbonyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



10 Analogously to Example 1b, Version B, 51 mg of the title compound was obtained from 100 mg of 71a and 74 mg of 1-methyl-1H-indol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.41 (2H), 1.43 - 1.73 (2H), 2.19 - 2.43 (1H),
 2.54 (3H), 2.74 - 3.23 (2H), 3.73 (3H), 4.07 (3H), 4.39 (3H), 6.60 (s, 1H), 7.03 - 7.14 (1H), 7.16
 - 7.30 (2H), 7.48 (2H), 7.59 (1H), 8.57 (1H), 8.81 (1H).

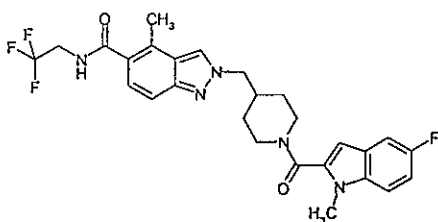
15 **Example 291:** 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 156 mg of the title compound was obtained from 150 mg of 71a and 179 mg of 4-[4-(trifluoromethyl)phenoxy]benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.36 - 1.68 (2H), 2.20 - 2.43 (1H), 2.54 (3H), 2.68 - 3.17 (2H), 3.50 - 3.79 (1H), 4.07 (2H), 4.37 (3H), 7.09 - 7.28 (5H), 7.35 - 7.51 (3H), 7.76 (2H), 8.56 (1H), 8.83 (1H).

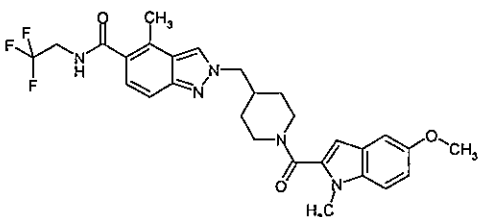
Example 292: 2-({1-[(5-fluoro-1-methyl-1H-indol-2-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 28 mg of the title compound was obtained from 100 mg of 71a and 82 mg of 5-fluoro-1-methyl-1H-indol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.18 - 1.38 (2H), 1.39 - 1.72 (2H), 2.25 - 2.42 (1H), 2.54 (3H), 2.74 - 3.25 (2H), 3.73 (3H), 4.07 (3H), 4.38 (3H), 6.58 (1H), 7.09 (1H), 7.21 (1H), 7.36 (1H), 7.47 (2H), 8.57 (1H), 8.83 (1H).

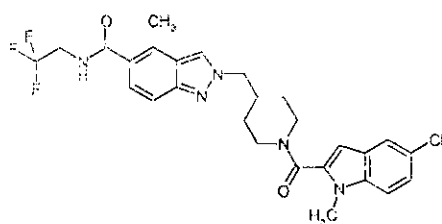
Example 293: 2-({1-[(5-methoxy-1-methyl-1H-indol-2-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 37 mg of the title compound was obtained from 100 mg of 71a and 87 mg of 5-methoxy-1-methyl-1H-indol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.30 (2H), 1.41 - 1.71 (2H), 2.26 - 2.43 (1H), 2.54 (3H), 2.71 - 3.22 (2H), 3.69 (3H), 3.75 (3H), 4.07 (3H), 4.39 (3H), 6.51 (1H), 6.88 (1H), 7.07 (1H), 7.21 (1H), 7.34 - 7.54 (2H), 8.57 (1H), 8.83 (1H).

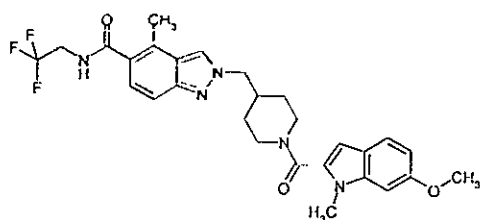
Example 294: 2-({1-[(5-chloro-1-methyl-1H-indol-2-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 43 mg of the title compound was obtained from 100 mg of 71a and 89 mg of 5-chloro-1-methyl-1H-indol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.29 (2H), 1.38 - 1.72 (2H), 2.26 - 2.43 (1H), 2.54 (3H), 2.71 - 3.21 (2H), 3.73 (3H), 3.84 - 4.18 (3H), 4.38 (3H), 6.59 (1H), 7.15 - 7.28 (2H), 7.47 (1H), 7.55 (1H), 7.65 (1H), 8.57 (1H), 8.83 (1H).

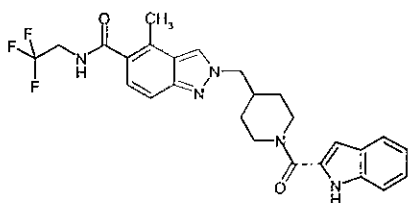
Example 295: 2-({1-[(6-methoxy-1-methyl-1H-indol-2-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 36 mg of the title compound was obtained from 100 mg of 71a and 87 mg of 6-methoxy-1-methyl-1H-indol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.28 (2H), 1.52 (2H), 2.25 - 2.43 (1H), 2.54 (3H), 2.78 - 3.16 (2H), 3.70 (3H), 3.82 (3H), 3.97 - 4.63 (6H), 6.54 (1H), 6.73 (1H), 7.01 (1H), 7.21 (1H), 7.39 - 7.53 (2H), 8.57 (1H), 8.83 (1H).

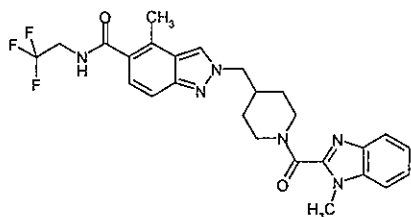
Example 296: 2-({1-(1H-Indol-2-ylcarbonyl)piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 43 mg of the title compound was obtained from 100 mg of 71a and 68 mg of indol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.16 - 1.40 (2H), 1.58 (2H), 2.27 - 2.44 (1H), 2.55 (3H), 2.77 - 3.24 (2H), 4.07 (2H), 4.30 - 4.59 (4H), 6.74 (1H), 6.95 - 7.08 (1H), 7.10 - 7.26 (2H), 7.40 (1H), 7.48 (1H), 7.59 (1H), 8.58 (1H), 8.84 (1H), 11.54 (1H).

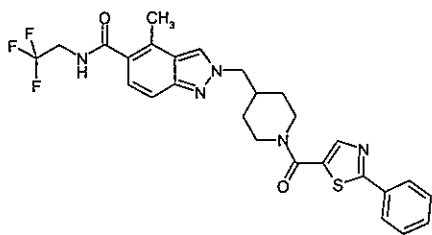
Example 297: 4-methyl-2-({1-[(1-methyl-1H-benzimidazol-2-yl)carbonyl]piperidin-4-yl}-methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 26 mg of the title compound was obtained from 100 mg of 71a and 75 mg of 1-methyl-1H-benzimidazol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.42 (2H), 1.46 (1H), 1.57 - 1.75 (1H), 2.27 - 2.44 (1H), 2.54 (3H), 2.87 (1H), 3.11 (1H), 3.83 (3H), 4.07 (3H), 4.40 (2H), 4.48 - 4.62 (1H), 7.13 - 7.41 (3H), 7.47 (1H), 7.58 - 7.76 (2H), 8.58 (1H), 8.83 (1H).

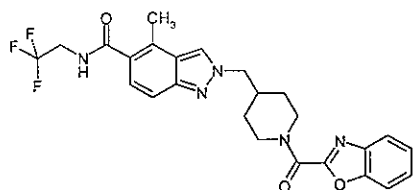
Example 298: 4-methyl-2-({1-[(2-phenylthiazol-5-yl)carbonyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 53 mg of the title compound was obtained from 100 mg of 71a and 87 mg of 2-phenylthiazol-5-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.18 - 1.41 (2H), 1.56 (2H), 2.27 - 2.44 (1H), 2.54 (3H), 2.73 - 3.29 (2H), 3.93 - 4.53 (6H), 7.21 (1H), 7.42 - 7.61 (4H), 7.93 - 8.03 (2H), 8.14 (1H), 8.57 (1H), 8.83 (1H).

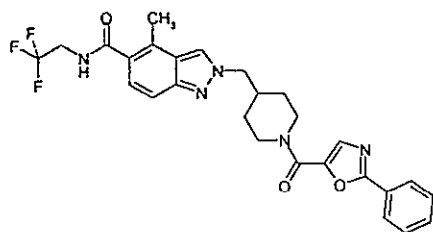
Example 299: 2-([1-(benzoxazol-2-ylcarbonyl)piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 20 mg of the title compound was obtained from 100 mg of 71a and 69 mg of benzoxazol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.44 (2H), 1.48 - 1.72 (2H), 2.29 - 2.46 (1H),
 5 2.54 (3H), 2.82 - 3.01 (1H), 3.12 - 3.31 (1H), 4.07 (2H), 4.40 (4H), 7.21 (1H), 7.42 - 7.61 (3H),
 7.78 - 7.95 (2H), 8.58 (1H), 8.83 (1H).

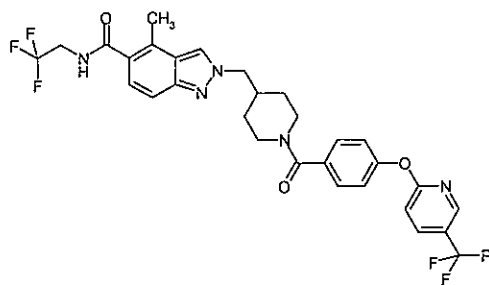
Example 300: 4-methyl-2-({1-[(2-phenyloxazol-5-yl)carbonyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



10 Analogously to Example 1b, Version B, 20 mg of the title compound was obtained from 100 mg of 71a and 80 mg of 2-phenyloxazol-5-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.15 - 1.43 (2H), 1.59 (2H), 2.31 - 2.45 (1H), 2.55 (3H), 2.70 - 3.35 (2H), 4.07 (2H), 4.39 (4H), 7.21 (1H), 7.48 (1H), 7.53 - 7.67 (3H), 7.80 (1H), 7.94 - 8.07 (2H), 8.58 (1H), 8.83 (1H).

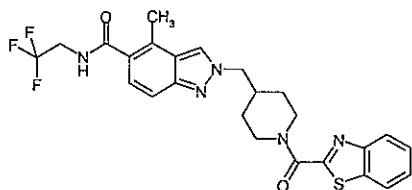
15 **Example 301:** 4-methyl-N-(2,2,2-trifluoroethyl)-2-{{1-(4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 23 mg of the title compound was obtained from 75 mg of 71a and 90 mg of 4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.13 - 1.35 (2H), 1.37 - 1.72 (2H), 2.19 - 2.42 (1H), 2.54 (3H), 2.69 - 3.17 (2H), 3.55 - 3.82 (1H), 3.91 - 4.18 (2H), 4.26 - 4.60 (3H), 7.28 (4H), 7.38 - 7.56 (3H), 8.16 - 8.33 (1H), 8.49 - 8.63 (2H), 8.75 - 8.91 (1H).

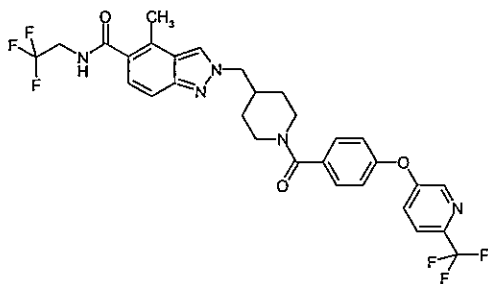
Example 302: 2-{{1-(benzothiazol-2-ylcarbonyl)piperidin-4-yl}methyl}-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 35 mg of the title compound was obtained from 75 mg of 71a and 57 mg of benzothiazol-2-carboxylic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.36 (2H), 1.59 (2H), 2.31 - 2.46 (1H), 2.55 (3H), 2.91 (1H), 3.26 (1H), 4.07 (2H), 4.31 - 4.61 (3H), 4.95 - 5.23 (1H), 7.21 (1H), 7.41 - 7.69 (3H), 8.05 - 8.28 (2H), 8.58 (1H), 8.83 (1H).

Example 303: 4-methyl-N-(2,2,2-trifluoroethyl)-2-{{1-(4-{{6-(trifluoromethyl)pyridin-3-yl}oxy}benzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide

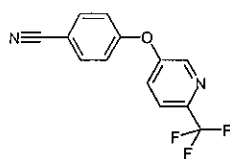


Analogously to Example 1b, Version B, 42 mg of the title compound was obtained from 75 mg of 71a and 90 mg of the carboxylic acid prepared in Example 303b) in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.16 - 1.33 (2H), 1.34 - 1.67 (2H), 2.21 - 2.39 (1H), 2.54 (3H), 2.68 - 3.16 (2H), 3.52 - 3.76 (1H), 3.95 - 4.16 (2H), 4.28 - 4.61 (3H), 7.13 - 7.31 (3H), 7.46 (3H), 7.59 - 7.71 (1H), 7.92 (1H), 8.52 - 8.65 (2H), 8.84 (1H).

The starting material was prepared as follows:

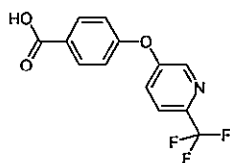
Example 303a: 4-{{6-(trifluoromethyl)pyridin-3-yl}oxy}benzonitrile



2.5 g of 5-bromo-2-(trifluoromethyl)pyridine, 5.8 g of 4-hydroxybenzonitrile, 10.8 g of caesium carbonate and 1.87 g of molybdenum hexacarbonyl in 100 ml 1,4-dioxan were heated under reflux until complete conversion. The reaction mixture was diluted with water, extracted several times with ethyl acetate, and the combined organic phases washed with saturated sodium chloride solution, dried over sodium sulphate and concentrated. The residue was purified by column chromatography on silica gel using a hexane/ ethyl acetate gradient. Yield: 1.35 g of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 7.35 (2H), 7.75 - 7.84 (1H), 7.89 - 8.03 (3H), 8.66 (1H).

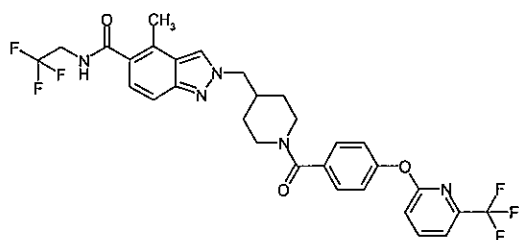
Example 303b: 4-{{6-(trifluoromethyl)pyridin-2-yl}oxy}benzoic acid



1.33 g of the compound prepared in Example 303a in 64 ml ethanol was treated with 32 ml of 40% potassium hydroxide solution and heated under reflux until complete conversion. The reaction mixture was diluted with water and extracted with ethyl acetate. The organic phase was washed with saturated sodium chloride solution, dried over sodium sulphate and concentrated. Yield: 1.32 g of the title compound.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 7.19 - 7.33 (2H), 7.68 - 7.81 (1H), 7.90 - 8.08 (3H), 8.64 (1H), 12.71 - 13.28 (1H).

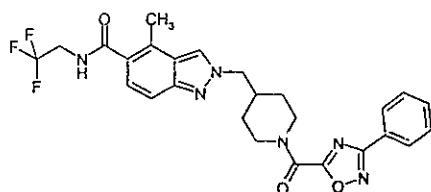
Example 304: 4-methyl-N-(2,2,2-trifluoroethyl)-2-{{1-(4-{{6-(trifluoromethyl)pyridin-2-yl}oxy}benzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 48 mg of the title compound was obtained from 75 mg of 71a and 90 mg of 4-{{6-(trifluoromethyl)pyridin-2-yl}oxy}benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.15 - 1.34 (2H), 1.35 - 1.71 (2H), 2.23 - 2.41 (1H), 2.54 (3H), 2.67 - 3.17 (2H), 3.51 - 3.77 (1H), 3.97 - 4.18 (2H), 4.22 - 4.65 (3H), 7.13 - 7.31
5 (3H), 7.37 (1H), 7.41 - 7.53 (3H), 7.67 (1H), 8.07 - 8.20 (1H), 8.57 (1H), 8.75 - 8.92 (1H).

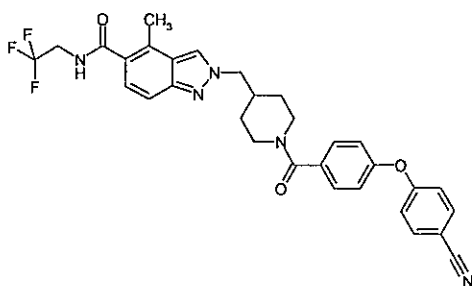
Example 305: 4-methyl-2-({1-[(3-phenyl-1,2,4-oxadiazol-5-yl)carbonyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 40 mg of the title compound was obtained from 90 mg
10 of 71a and 73 mg of 3-phenyl-1,2,4-oxadiazol-5-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.18 - 1.43 (2H), 1.48 - 1.72 (2H), 2.33 - 2.47 (1H), 2.54 (3H), 2.95 (1H), 3.24 (1H), 3.94 - 4.17 (3H), 4.33 - 4.53 (3H), 7.21 (1H), 7.47 (1H), 7.55 - 7.70 (3H), 7.96 - 8.12 (2H), 8.57 (1H), 8.83 (1H).

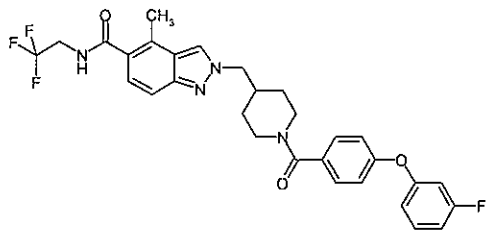
Example 306: 2-({1-[4-(4-cyanophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
15



Analogously to Example 1b, Version B, 59 mg of the title compound was obtained from 110 mg of 71a and 101 mg of 4-(4-cyanophenoxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.34 (2H), 1.35 - 1.70 (2H), 2.18 - 2.41 (1H), 2.54 (3H), 2.70 - 3.18 (2H), 3.54 - 3.79 (1H), 4.07 (2H), 4.37 (3H), 7.10 - 7.28 (5H), 7.41 - 7.52
20 (3H), 7.79 - 7.94 (2H), 8.56 (s1H), 8.81 (s1H).

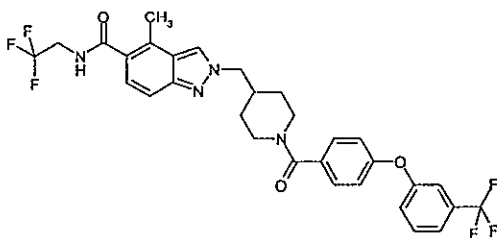
Example 307: 2-({1-[4-(3-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 40 mg of the title compound was obtained from 100 mg
5 of 71a and 89 mg of 4-(3-fluorophenoxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.14 - 1.34 (2H), 1.36 - 1.70 (2H), 2.22 - 2.44 (1H), 2.54 (3H), 2.70 - 3.13 (2H), 3.52 - 3.86 (1H), 4.07 (2H), 4.37 (3H), 6.79 - 7.12 (5H), 7.21 (1H), 7.35 - 7.52 (4H), 8.55 (1H), 8.81 (1H).

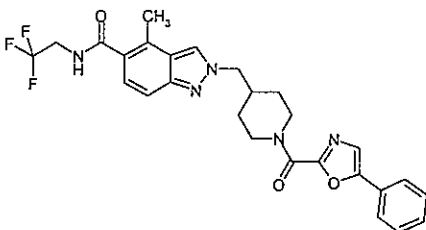
Example 308: 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[3-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
10



Analogously to Example 1b, Version B, 47 mg of the title compound was obtained from 100 mg
of 71a and 119 mg of 4-[3-(trifluoromethyl)phenoxy]benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.16 - 1.34 (2H), 1.37 - 1.66 (2H), 2.22 - 2.38 (1H), 2.54 (3H), 2.69 - 3.14 (2H), 3.50 - 3.83 (1H), 4.07 (2H), 4.37 (3H), 7.10 (2H), 7.21 (1H), 7.36 (1H), 7.39 - 7.50 (4H), 7.54 (1H), 7.61 - 7.75 (1H), 8.55 (1H), 8.81 (1H).
15

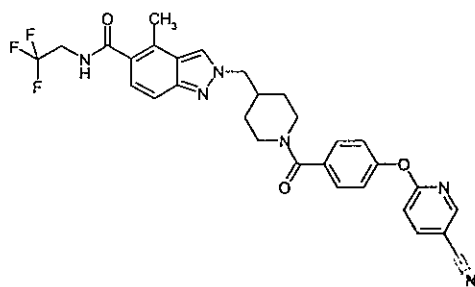
Example 309: 4-methyl-2-({1-[(5-phenyloxazol-2-yl)carbonyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 10 mg of the title compound was obtained from 100 mg of 71a and 73 mg of 5-phenyloxazol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.24 (2H), 1.58 (2H), 2.29 - 2.45 (1H), 2.55 (3H), 2.86 (1H), 3.21 (1H), 4.07 (2H), 4.31 - 4.52 (3H), 4.62 (1H), 7.21 (1H), 7.39 - 7.57 (4H), 7.74 - 7.83 (2H), 7.88 (1H), 8.57 (1H), 8.81 (1H).

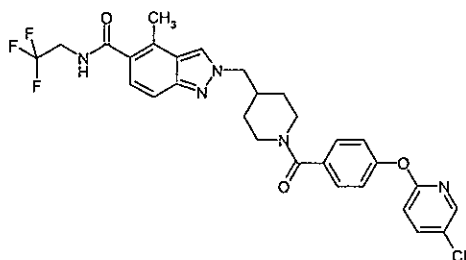
Example 310: 2-[(1-{4-[(5-cyanopyridin-2-yl)oxy]benzoyl}piperidin-4-yl)methyl]-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 47 mg of the title compound was obtained from 100 mg of 71a and 119 mg of 4-[3-(trifluoromethyl)phenoxy]benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.16 - 1.34 (2H), 1.37 - 1.66 (2H), 2.22 - 2.38 (1H), 2.54 (3H), 2.69 - 3.14 (2H), 3.50 - 3.83 (1H), 4.07 (2H), 4.37 (3H), 7.10 (2H), 7.21 (1H), 7.36 (1H), 7.39 - 7.50 (4H), 7.54 (1H), 7.61 - 7.75 (1H), 8.55 (1H), 8.81 (1H).

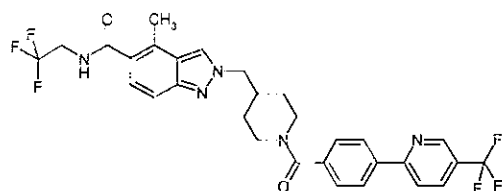
Example 311: 2-[(1-{4-[(5-chloropyridin-2-yl)oxy]benzoyl}piperidin-4-yl)methyl]-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 72 mg of the title compound was obtained from 100 mg of 71a and 96 mg of 4-[(5-chloropyridin-2-yl)oxy]benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.34 (2H), 1.36 - 1.73 (2H), 2.22 - 2.41 (1H), 2.54 (3H), 2.69 - 3.18 (2H), 3.47 - 3.88 (1H), 4.07 (2H), 4.37 (3H), 7.06 - 7.28 (4H), 7.34 - 7.52 (3H), 7.99 (1H), 8.22 (1H), 8.56 (1H), 8.81 (1H).

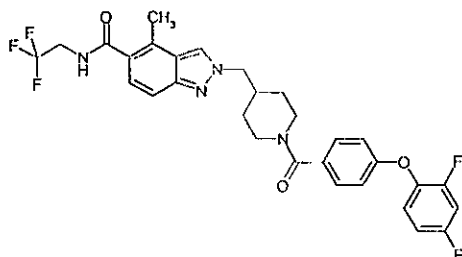
Example 312: 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[5-(trifluoromethyl)pyridin-2-yl]-benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 83 mg of the title compound was obtained from 100 mg of 71a and 103 mg of 4-[5-(trifluoromethyl)pyridin-2-yl]benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.28 (2H), 1.36 - 1.73 (2H), 2.33 (1H), 2.54 (3H), 2.70 - 3.18 (2H), 3.47 - 3.76 (1H), 3.96 - 4.16 (2H), 4.38 (3H), 7.21 (1H), 7.41 - 7.60 (3H), 8.17 - 8.40 (4H), 8.56 (1H), 8.81 (1H), 9.07 (1H).

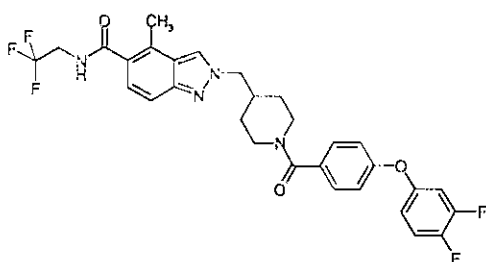
Example 313: 2-({1-[4-(2,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 87 mg of the title compound was obtained from 100 mg of 71a and 96 mg of 4-(2,4-difluorophenoxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.34 - 1.72 (2H), 2.18 - 2.42 (1H), 2.52 - 2.58 (3H), 2.69 - 3.20 (2H), 3.43 - 3.84 (1H), 4.07 (2H), 4.36 (3H), 6.92 - 7.06 (2H), 7.09 - 7.25 (2H), 7.28 - 7.42 (3H), 7.43 - 7.61 (2H), 8.55 (1H), 8.81 (1H).

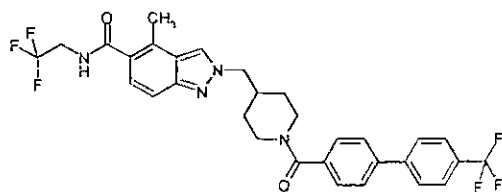
Example 314: 2-({1-[4-(3,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 48 mg of the title compound was obtained from 100 mg of 71a and 96 mg of 4-(3,4-difluorophenoxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.24 (2H), 1.36 - 1.72 (2H), 2.19 - 2.42 (1H), 2.54 (3H), 2.70 - 3.17 (2H), 3.44 - 3.86 (1H), 3.99 - 4.17 (2H), 4.36 (3H), 6.86 - 6.97 (1H), 7.01 - 7.11 (2H), 7.21 (1H), 7.29 (1H), 7.35 - 7.55 (4H), 8.55 (1H), 8.81 (1H).

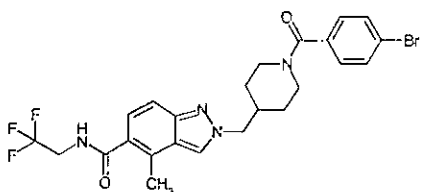
Example 315: 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{[4'-(trifluoromethyl)biphenyl-4-yl]-carbonyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 32 mg of the title compound was obtained from 100 mg of 71a and 102 mg of 4'-(trifluoromethyl)biphenyl-4-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.27 (2H), 1.37 - 1.71 (2H), 2.24 - 2.39 (1H), 2.53 (3H), 2.70 - 3.18 (2H), 3.43 - 3.51 (2H), 3.54 - 3.86 (1H), 4.37 (3H), 7.18 (1H), 7.38 - 7.57 (3H), 7.72 - 7.88 (4H), 7.89 - 7.98 (2H), 8.13 (1H), 8.51 (1H).

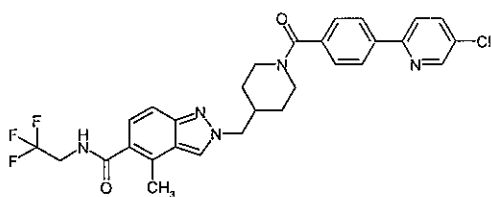
Example 316: 2-{[1-(4-bromobenzoyl)piperidin-4-yl)methyl]-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 6.96 g of the title compound was obtained from 5.61 g 71a and 2.89 g of 4-bromobenzoic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.32 (2H), 1.34 - 1.74 (2H), 2.19 - 2.41 (1H), 2.54 (3H), 2.88 - 3.22 (2H), 3.42 - 3.74 (1H), 3.94 - 4.18 (2H), 4.36 (3H), 7.20 (1H), 7.32 (2H), 7.46 (1H), 7.64 (2H), 8.55 (1H), 8.83 (1H).

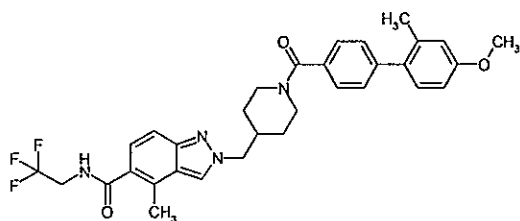
Example 317: 2-({1-[4-(5-chloropyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



80 mg of the aryl bromide prepared in Example 316 together with 32 mg of (5-chloropyridin-2-yl)boronic acid was first placed in 1 ml tetrahydrofuran, treated with 17.3 mg of 1,1'-bis(diphenylphosphino)ferrocenodichloropalladium(II) and 0.17 ml of 1M potassium carbonate solution and heated in the microwave for 10 minutes at 120°C (100 watts). After fresh addition of 11 mg of (5-chloropyridin-2-yl)boronic acid and 17.3 mg of the palladium(II) catalyst, the reaction mixture was again heated in the microwave for 10 minutes at 120°C (100 watts) until complete conversion. The reaction mixture was concentrated. After HPLC purification, this yielded 15 mg of the title compound.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.18 - 1.36 (2H), 1.37 - 1.75 (2H), 2.28 - 2.38 (1H), 2.54 (3H), 2.73 - 3.18 (2H), 3.49 - 3.74 (1H), 3.95 - 4.16 (2H), 4.28 - 4.62 (3H), 7.22 (1H), 7.33 - 7.65 (5H), 7.98 - 8.22 (3H), 8.56 (1H), 8.77 - 8.91 (1H).

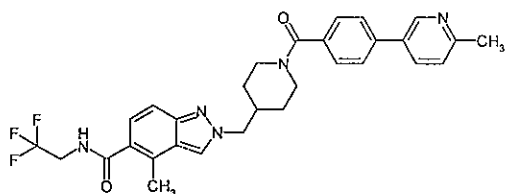
Example 318: 2-({1-[(4'-methoxy-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 317, 50 mg of the title compound was obtained from 125 mg of 316 and 53 mg of (4-methoxy-2-methyl-phenyl)boronic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.16 - 1.36 (2H), 1.37 - 1.74 (2H), 2.23 (3H), 2.29 - 2.38 (1H), 2.54 (3H), 2.72 - 3.12 (2H), 3.77 (4H), 3.97 - 4.17 (2H), 4.38 (3H), 6.75 - 6.94 (2H), 7.07 - 7.26 (2H), 7.29 - 7.56 (5H), 8.56 (1H), 8.81 (1H).

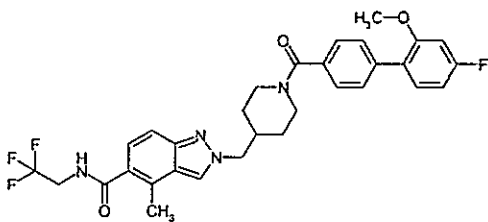
Example 319: 4-methyl-2-({1-[4-(6-methylpyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 317, 37 mg of the title compound was obtained from 125 mg of 316 and 44 mg of (6-methylpyridin-3-yl)boronic acid under reflux.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.15 - 1.35 (2H), 1.37 - 1.71 (2H), 2.20 - 2.40 (1H), 2.52 (3H), 2.54 (3H), 2.74 - 3.15 (2H), 3.52 - 3.82 (1H), 4.07 (2H), 4.38 (3H), 7.20 (1H), 7.37 (1H), 7.47 (3H), 7.77 (2H), 8.01 (1H), 8.57 (1H), 8.72 - 8.92 (2H).

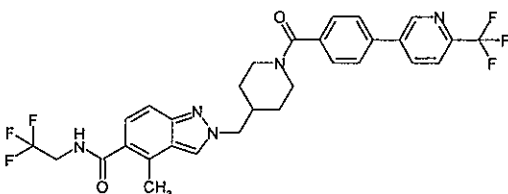
Example 320: 2-({1-[(4'-fluoro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 317, 60 mg of the title compound was obtained from 125 mg of 316 and 75 g of (4-fluoro-2-methoxyphenyl)boronic acid under reflux.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.16 - 1.37 (2H), 1.38 - 1.75 (2H), 2.25 - 2.37 (1H), 2.54 (3H), 2.72 - 3.17 (2H), 3.79 (4H), 4.07 (2H), 4.38 (3H), 6.87 (1H), 7.04 (1H), 7.21 (1H), 7.27 - 7.58 (6H), 8.56 (1H), 8.81 (1H).

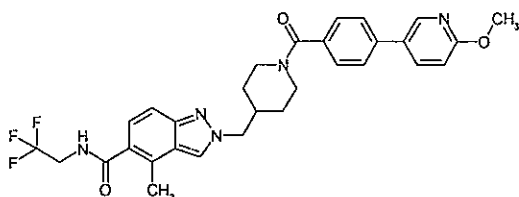
Example 321: 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[6-(trifluoromethyl)pyridin-3-yl]-benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 317, 45 mg of the title compound was obtained from 125 mg of 316 and 84 mg of [6-(trifluoromethyl)pyridin-3-yl]boronic acid under reflux.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.15 - 1.37 (2H), 1.37 - 1.71 (2H), 2.22 - 2.41 (1H), 2.54 (3H), 2.73 - 3.20 (2H), 3.45 - 3.74 (1H), 3.93 - 4.17 (2H), 4.29 - 4.62 (3H), 7.21 (1H), 7.39 - 7.62 (4H), 7.89 (2H), 8.01 (1H), 8.57 (1H), 8.73 - 8.92 (1H), 9.13 (1H).

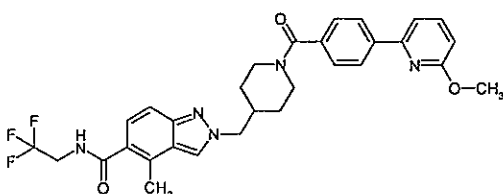
Example 322: 2-({1-[4-(6-methoxypyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 317, 54 mg of the title compound was obtained from 125 mg of 316 and 68 mg of (6-methoxypyridin-3-yl)boronic acid under reflux.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.13 - 1.35 (2H), 1.36 - 1.71 (2H), 2.20 - 2.40 (1H), 2.54 (3H), 2.73 - 3.16 (2H), 3.51 - 3.77 (1H), 3.90 (3H), 3.96 - 4.19 (2H), 4.37 (3H), 6.93 (1H), 7.20 (1H), 7.39 - 7.52 (3H), 7.73 (2H), 8.05 (1H), 8.47 - 8.63 (2H), 8.83 (1H).

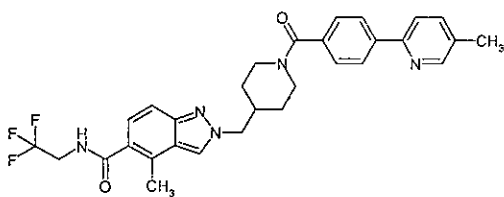
Example 323: 2-({1-[4-(6-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 317, 55 mg of the title compound was obtained from 125 mg of 316 and 49 mg of (6-methoxypyridin-2-yl)boronic acid under reflux.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.16 - 1.36 (2H), 1.37 - 1.69 (2H), 2.20 - 2.41 (1H), 2.54 (3H), 2.74 - 3.16 (2H), 3.47 - 3.81 (1H), 3.96 (3H), 4.06 (2H), 4.38 (3H), 6.81 (1H), 7.21 (1H), 7.47 (3H), 7.60 (1H), 7.74 - 7.90 (1H), 8.15 (2H), 8.56 (1H), 8.81 (1H).

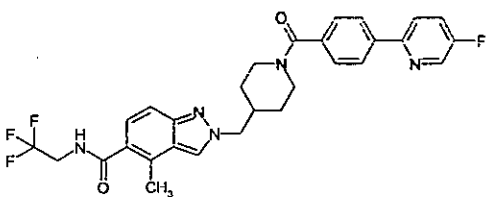
Example 324: 4-methyl-2-({1-[4-(5-methylpyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 317, 9 mg of the title compound was obtained from 125 mg of 316 and 44 mg of (5-methylpyridin-2-yl)boronic acid under reflux.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.16 - 1.36 (2H), 1.37 - 1.71 (2H), 2.24 - 2.30 (1H), 2.34 (3H), 2.54 (3H), 2.74 - 3.16 (2H), 3.52 - 3.79 (1H), 3.93 - 4.17 (2H), 4.38 (3H), 7.20 (1H), 7.37 - 7.54 (3H), 7.66 - 7.78 (1H), 7.89 (1H), 8.11 (2H), 8.45 - 8.61 (2H), 8.74 - 8.88 (1H).

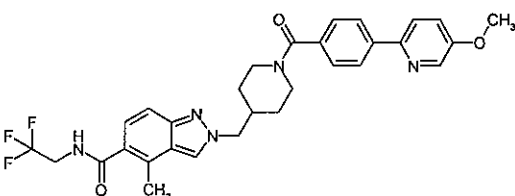
Example 325: 2-({1-[4-(5-fluoropyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 317, 35 mg of the title compound was obtained from 125 mg of 316 and 45 mg of (5-fluoropyridin-2-yl)boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.34 (2H), 1.35 - 1.69 (2H), 2.22 - 2.41 (1H), 2.54 (3H), 2.69 - 3.19 (2H), 3.51 - 3.78 (1H), 4.07 (2H), 4.38 (3H), 7.20 (1H), 7.40 - 7.53 (3H), 7.85 (1H), 8.04 - 8.17 (3H), 8.57 (1H), 8.68 (1H), 8.83 (1H).

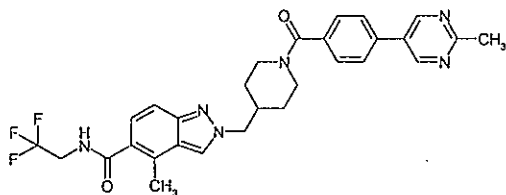
Example 326: 2-({1-[4-(5-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 317, 94 mg of the title compound was obtained from 125 mg of 316 and 49 mg of (5-methoxypyridin-2-yl)boronic acid under reflux.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.36 (2H), 1.37 - 1.69 (2H), 2.28 - 2.40 (1H), 2.54 (3H), 2.73 - 3.19 (2H), 3.48 - 3.79 (1H), 3.88 (3H), 4.07 (2H), 4.38 (3H), 7.20 (1H), 7.38 - 7.55 (4H), 7.96 (1H), 8.07 (2H), 8.39 (1H), 8.57 (1H), 8.82 (1H).

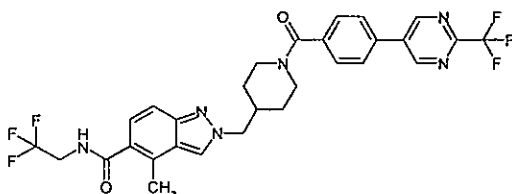
Example 327: 4-methyl-2-({1-[4-(2-methylpyrimidin-5-yl)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide



Analogously to Example 317, 46 mg of the title compound was obtained from 125 mg of 316 and 102 mg of (2-methylpyrimidin-5-yl)boronic acid pinacol ester under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.16 - 1.35 (2H), 1.35 - 1.67 (2H), 2.21 - 2.42 (1H), 2.54 (3H), 2.67 (3H), 2.71 - 2.89 (1H), 2.93 - 3.20 (1H), 3.50 - 3.73 (1H), 4.07 (2H), 4.37 (3H), 7.20 (1H), 7.40 - 7.57 (3H), 7.85 (2H), 8.57 (1H), 8.84 (1H), 9.05 (2H).

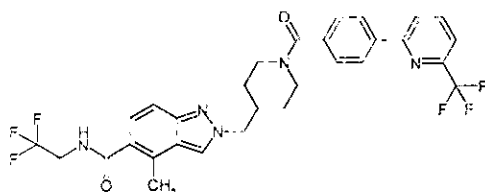
Example 328: 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[2-(trifluoromethyl)pyrimidin-5-yl]benzoyl]piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 317, 34 mg of the title compound was obtained from 125 mg of 316 and 89 mg of [2-(trifluoromethyl)pyrimidin-5-yl]boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.21 - 1.35 (2H), 1.36 - 1.71 (2H), 2.23 - 2.41 (1H), 2.54 (3H), 2.71 - 2.89 (1H), 2.94 - 3.17 (1H), 3.48 - 3.72 (1H), 4.05 (2H), 4.28 - 4.61 (3H), 7.21 (1H), 7.47 (1H), 7.57 (2H), 7.98 (2H), 8.57 (1H), 8.84 (1H), 9.44 (2H).

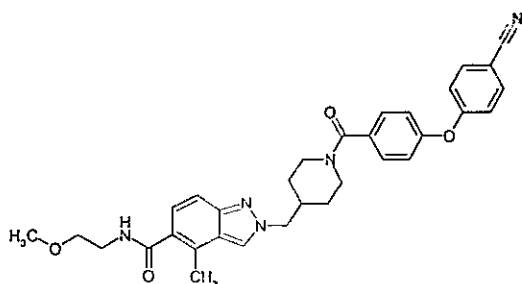
Example 329: 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[6-(trifluoromethyl)pyridin-2-yl]benzoyl]piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 317, 70 mg of the title compound was obtained from 125 mg of 316 and 121 mg of [6-(trifluoromethyl)pyridin-2-yl]boronic acid pinacol ester under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.36 (2H), 1.37 - 1.72 (2H), 2.23 - 2.39 (1H),
 5 2.54 (3H), 2.68 - 2.90 (1H), 2.92 - 3.16 (1H), 3.50 - 3.73 (1H), 4.06 (2H), 4.38 (3H), 7.20 (1H),
 7.41 - 7.60 (3H), 7.89 (1H), 8.11 - 8.27 (3H), 8.33 (1H), 8.56 (1H), 8.81 (1H).

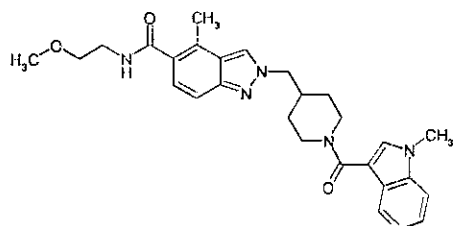
Example 330: 2-({1-[4-(4-cyanophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



10 Analogously to Example 1b, Version B, 151 mg of the title compound was obtained from 213 mg of 71a and 153 mg of 4-(4-cyanophenoxy)benzoic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.22 (2H), 1.31 - 1.66 (2H), 2.15 - 2.36 (1H), 2.49
 (3H), 2.70 - 3.12 (2H), 3.24 (3H), 3.36 (2H), 3.39 - 3.46 (2H), 3.50 - 3.79 (1H), 4.32 (3H), 7.00 -
 7.23 (5H), 7.30 - 7.49 (3H), 7.75 - 7.91 (2H), 8.12 (1H), 8.48 (1H).

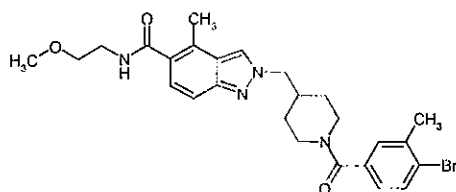
15 **Example 331:** N-(2-methoxyethyl)-4-methyl-2-({1-[(1-methyl-1H-indol-3-yl)carbonyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 26 mg of the title compound was obtained from 38 mg of 71a and 36 mg of 1-methyl-1H-indol-3-carboxylic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.43 - 1.64 (2H), 2.20 - 2.39 (1H), 2.53 (3H), 2.80 - 3.03 (2H), 3.28 (3H), 3.37 - 3.50 (4H), 3.81 (3H), 4.37 (4H), 7.18 (3H), 7.36 - 7.53 (2H), 7.67 (2H), 8.04 - 8.25 (1H), 8.52 (1H).

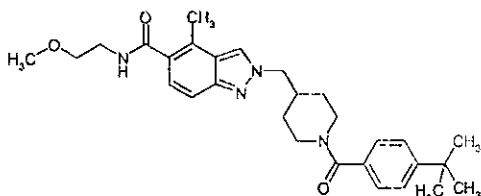
Example 332: 2-([1-(4-bromo-3-methylbenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 498 mg of the title compound was obtained from 807 mg of 71a and 788 mg of 4-bromo-3-methylbenzoic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.20 - 1.32 (2H), 1.33 - 1.67 (2H), 2.24 - 2.33 (1H), 2.36 (3H), 2.52 (3H), 2.73 (1H), 2.91 - 3.13 (1H), 3.28 (3H), 3.35 - 3.63 (5H), 4.34 (3H), 7.03 - 7.23 (2H), 7.30 - 7.49 (2H), 7.63 (1H), 8.15 (1H), 8.51 (1H).

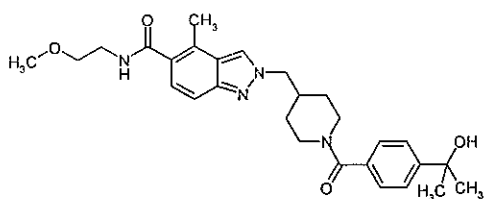
Example 333: 2-([1-(4-tert-butylbenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 50 mg of the title compound was obtained from 100 mg of 71a and 81 mg of 4-tert-butylbenzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.20 - 1.33 (11H), 1.33 - 1.67 (2H), 2.21 - 2.37 (1H), 2.52 (3H), 2.68 - 3.10 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.45 (2H), 3.52 - 3.74 (1H), 4.35 (3H), 7.18 (1H), 7.25 - 7.35 (2H), 7.36 - 7.52 (3H), 8.15 (1H), 8.50 (1H).

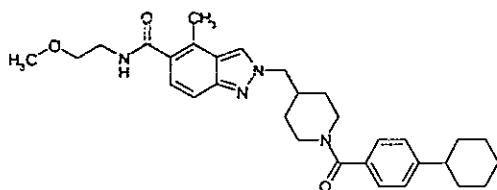
Example 334: 2-([1-[4-(1-hydroxy-1-methylethyl)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 28 mg of the title compound was obtained from 100 mg of 71a and 82 mg of 4-(1-hydroxy-1-methylethyl)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.24 (2H), 1.42 (8H), 2.20 - 2.40 (1H), 2.52 (3H),
 5 2.64 - 3.11 (2H), 3.28 (3H), 3.34 - 3.43 (2H), 3.43 - 3.51 (2H), 3.53 - 3.71 (1H), 4.35 (3H), 4.76
 - 5.43 (1H), 7.18 (1H), 7.28 (2H), 7.42 (1H), 7.46 - 7.56 (2H), 8.15 (1H), 8.51 (1H).

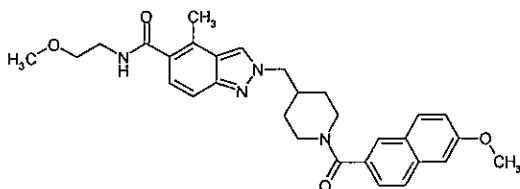
Example 335: 2-([1-(4-cyclohexylbenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



10 Analogously to Example 1b, Version B, 75 mg of the title compound was obtained from 100 mg of 71a and 93 mg of 4-cyclohexylbenzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.10 - 1.63 (9H), 1.64 - 1.89 (5H), 2.17 - 2.40 (1H),
 2.52 (3H), 2.69 - 3.08 (2H), 3.28 (3H), 3.38 - 3.43 (2H), 3.43 - 3.49 (2H), 3.53 - 3.73 (1H), 4.35
 (3H), 7.18 (1H), 7.27 (4H), 7.42 (1H), 8.15 (1H), 8.50 (1H).

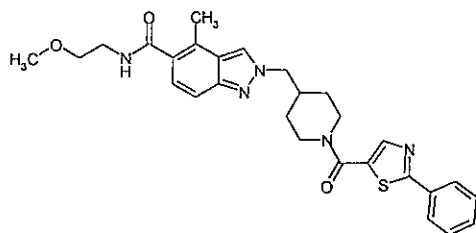
15 **Example 336:** N-(2-methoxyethyl)-2-([1-(6-methoxy-2-naphthoyl)piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 23 mg of the title compound was obtained from 100 mg of 71a) and 92 mg of 6-methoxy-naphthalen-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.37 - 1.72 (2H), 2.23 - 2.39 (1H), 2.52 (3H), 2.72 - 3.12 (2H), 3.28 (3H), 3.36 - 3.42 (2H), 3.43 - 3.49 (2H), 3.59 - 3.81 (1H), 3.89 (3H), 4.36 (3H), 7.09 - 7.27 (2H), 7.33 - 7.49 (3H), 7.79 - 7.95 (3H), 8.15 (1H), 8.51 (1H).

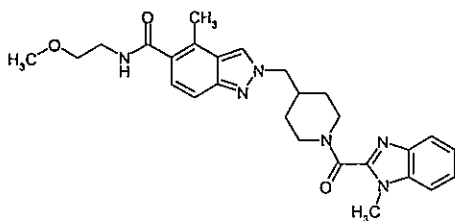
Example 337: N-(2-methoxyethyl)-4-methyl-2-({1-[(2-phenylthiazol-5-yl)carbonyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 36 mg of the title compound was obtained from 100 mg of 71a and 93 mg of 2-phenylthiazol-5-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.55 (2H), 2.19 - 2.45 (1H), 2.52 - 2.56 (3H), 2.70 - 3.24 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.44 - 3.49 (2H), 4.37 (4H), 7.19 (1H), 7.43 (1H), 7.49 - 7.62 (3H), 7.90 - 8.03 (2H), 8.08 - 8.25 (2H), 8.52 (1H).

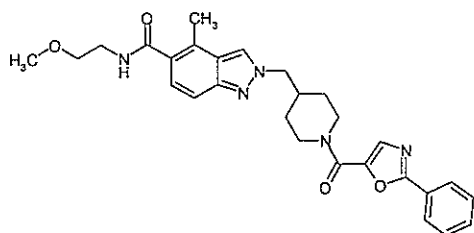
Example 338: N-(2-methoxyethyl)-4-methyl-2-({1-[(1-methyl-1H-benzimidazol-2-yl)carbonyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 39 mg of the title compound was obtained from 100 mg of 71a and 80 mg of 1-methyl-1H-benzimidazol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.22 - 1.43 (2H), 1.43 - 1.53 (1H), 1.59 - 1.70 (1H), 2.27 - 2.44 (1H), 2.53 (3H), 2.87 (1H), 3.11 (1H), 3.28 (3H), 3.36 - 3.42 (2H), 3.43 - 3.48 (2H), 3.83 (3H), 4.00 - 4.14 (1H), 4.39 (2H), 4.48 - 4.61 (1H), 7.18 (1H), 7.23 - 7.46 (3H), 7.58 - 7.74 (2H), 8.10 - 8.20 (1H), 8.53 (1H).

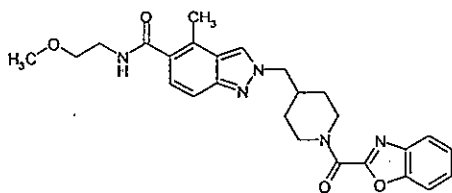
Example 339: N-(2-methoxyethyl)-4-methyl-2-({1-[(2-phenyloxazol-5-yl)carbonyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 30 mg of the title compound was obtained from 100 mg of 71a and 86 mg of 2-phenyloxazol-5-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.14 - 1.48 (2H), 1.59 (2H), 2.24 - 2.46 (1H), 2.52 - 2.56 (3H), 2.71 - 3.01 (1H), 3.11 - 3.26 (1H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 4.38 (4H), 7.19 (1H), 7.43 (1H), 7.52 - 7.67 (3H), 7.80 (1H), 7.97 - 8.05 (2H), 8.16 (1H), 8.53 (1H).

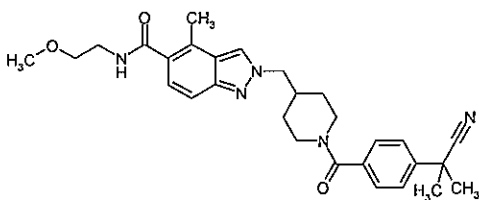
Example 340: 2-({1-(benzoxazol-2-ylcarbonyl)piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 37 mg of the title compound was obtained from 100 mg of 71a and 84 mg of benzoxazol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.43 (2H), 1.48 - 1.73 (2H), 2.28 - 2.45 (1H), 2.53 (3H), 2.91 (1H), 3.22 (1H), 3.33 - 3.57 (m, 7H), 4.25 - 4.60 (4H), 7.18 (1H), 7.36 - 7.62 (3H), 7.78 - 7.95 (2H), 8.16 (1H), 8.53 (1H).

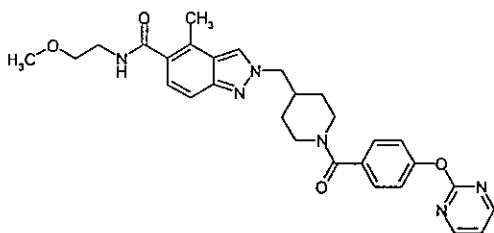
Example 341: 2-({1-[4-(1-cyano-1-methylethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 20 mg of the title compound was obtained from 100 mg of 71a and 86 mg of 4-(1-cyano-1-methylethyl)benzoic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.34 (2H), 1.35 - 1.64 (2H), 1.70 (6H), 2.16 - 2.40 (1H), 2.52 (3H), 2.64 - 3.13 (2H), 3.28 (3H), 3.40 (2H), 3.43 - 3.49 (2H), 3.49 - 3.67 (1H),
5 4.35 (3H), 7.18 (1H), 7.42 (3H), 7.58 (2H), 8.15 (1H), 8.50 (1H).

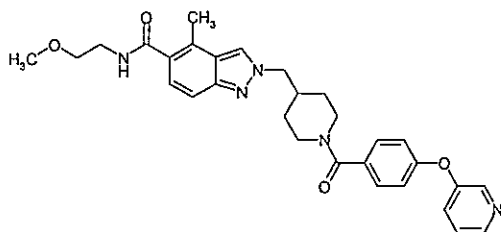
Example 342: N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyrimidin-2-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 37 mg of the title compound was obtained from 100 mg
10 of 71a and 98 mg of 4-(2-pyrimidin-yloxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.26 (2H), 1.38 - 1.66 (2H), 2.20 - 2.40 (1H), 2.52 - 2.56 (3H), 2.70 - 3.16 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.49 (2H), 3.64 - 3.77 (m, 1H),
4.36 (3H), 7.18 (1H), 7.22 - 7.33 (3H), 7.39 - 7.49 (3H), 8.15 (1H), 8.43 - 8.57 (1H), 8.66 (2H).

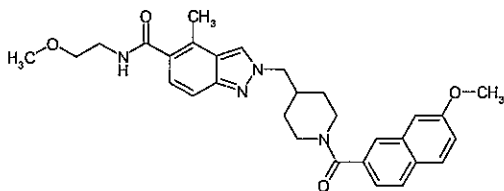
Example 343: N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyridin-3-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
15



Analogously to Example 1b, Version B, 7 mg of the title compound was obtained from 75 mg of
71a and 73 mg of 4-(3-pyridyloxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.32 (2H), 1.35 - 1.69 (2H), 2.19 - 2.38 (1H),
20 2.52 (3H), 2.70 - 3.14 (2H), 3.28 (3H), 3.34 - 3.43 (2H), 3.43 - 3.51 (2H), 3.55 - 3.70 (1H), 4.35
(3H), 7.04 - 7.13 (2H), 7.18 (1H), 7.37 - 7.46 (3H), 7.47 - 7.54 (1H), 7.58 (1H), 8.15 (1H), 8.38 -
8.55 (3H).

Example 344: N-(2-methoxyethyl)-2-[[1-(7-methoxy-2-naphthoyl)piperidin-4-yl]methyl]-4-methyl-2H-indazole-5-carboxamide

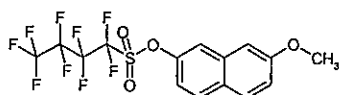


Analogously to Example 1b, Version B, 28 mg of the title compound was obtained from 75 mg of 71a and 69 mg of the carboxylic acid prepared in Example 344c) in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.21 - 1.35 (2H), 1.37 - 1.75 (2H), 2.23 - 2.39 (1H), 2.52 (3H), 2.73 - 3.13 (2H), 3.28 (3H), 3.35 - 3.42 (2H), 3.45 (2H), 3.53 - 3.74 (1H), 3.88 (3H), 4.37 (3H), 7.13 - 7.25 (2H), 7.29 (1H), 7.36 - 7.48 (2H), 7.77 - 7.93 (3H), 8.15 (1H), 8.52 (1H).

The starting material was prepared as follows:

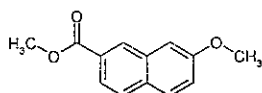
Example 344a: 7-methoxynaphthalen-2-yl-1,1,2,2,3,3,4,4,4-nonafluorobutan-1-sulphonate



5.12 g of 7-methoxy-2-naphthol were dissolved in 50 ml toluene and cooled to 0°C. 30.7 ml of N,N-diisopropylethylamine and 17.6 ml of nonafluorobutanesulphonyl fluoride were successively added dropwise to this and the reaction mixture was then stirred at room temperature until complete conversion. The reaction mixture was poured into 600 ml of saturated ammonium chloride solution, the deposited brown oil separated and the aqueous phase extracted several times with ethyl acetate. The combined organic phases were washed with water and saturated sodium chloride solution, dried and concentrated. The residue was purified by column chromatography on silica gel using a hexane/ ethyl acetate gradient. Yield: 11.45 g of the title compound.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 3.89 (3H), 7.28 (1H), 7.41 (1H), 7.52 (1H), 7.90 - 8.00 (2H), 8.05 (1H).

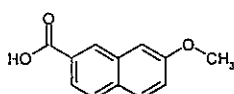
Example 344b: Methyl 7-methoxynaphthalen-2-carboxylate



Analogously to Example 1e, 2.34 g of the title compound was obtained from 11.0 g of 344a and 8.8 ml of methanol.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 3.89 (3H), 3.91 (3H), 7.31 (1H), 7.57 (1H), 7.82 (1H), 7.94 (2H), 8.54 (1H).

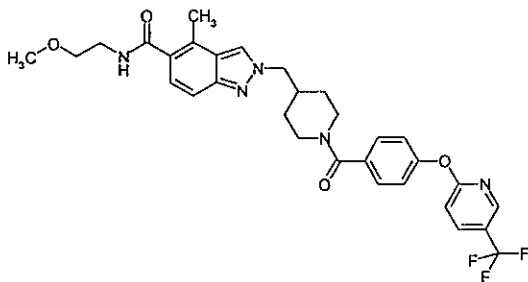
Example 344c: 7-methoxynaphthalen-2-carboxylic acid



Analogously to Example 258d, 1.27 g of the title compound was obtained from 2.0 g of 344b.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 3.89 (3H), 7.30 (1H), 7.53 (1H), 7.78 - 7.86 (1H), 7.87 - 8.01 (2H), 8.50 (1H), 12.99 (1H).

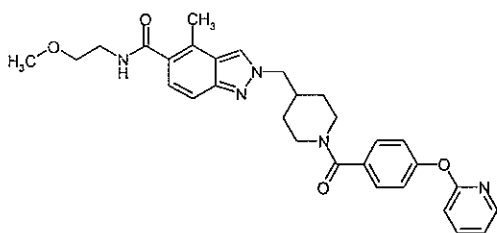
Example 345: N-(2-methoxyethyl)-4-methyl-2-({[1-(4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoyl)piperidin-4-yl]methyl}-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 221 mg of the title compound was obtained from 200 mg of 71a and 232 mg of 4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.21 - 1.34 (2H), 1.36 - 1.72 (2H), 2.23 - 2.39 (1H), 2.53 (3H), 2.89 (2H), 3.31 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.53 - 3.86 (1H), 4.36 (3H), 7.18 (1H), 7.23 - 7.33 (3H), 7.37 - 7.50 (3H), 8.13 (1H), 8.25 (1H), 8.51 (1H), 8.55 - 8.63 (1H).

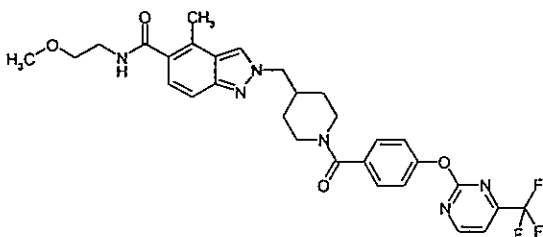
Example 346: N-(2-methoxyethyl)-4-methyl-2-({[1-(4-(pyridin-2-yloxy)benzoyl)piperidin-4-yl]methyl}-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 29 mg of the title compound was obtained from 100 mg of 71a and 98 mg of 4-(2-pyridyloxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.39 - 1.68 (2H), 2.24 - 2.39 (1H), 2.52 - 2.56 (3H), 2.71 - 3.13 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.45 (2H), 3.53 - 3.87 (1H), 4.36 (3H), 7.08 (1H), 7.13 - 7.26 (4H), 7.34 - 7.52 (3H), 7.88 (1H), 8.07 - 8.23 (2H), 8.51 (1H).

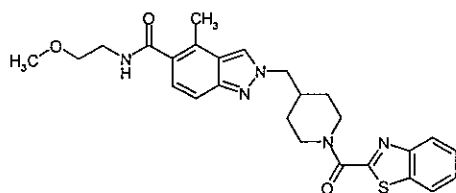
Example 347: N-(2-methoxyethyl)-4-methyl-2-({[1-(4-{[4-(trifluoromethyl)pyrimidin-2-yl]-oxy}benzoyl)piperidin-4-yl]methyl}-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 11 mg of the title compound was obtained from 75 mg of 71a and 97 mg of 4-{[4-(trifluoromethyl)pyrimidin-2-yl]oxy}benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.37 - 1.69 (2H), 2.19 - 2.39 (1H), 2.53 (3H), 2.68 - 3.18 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.45 (2H), 3.51 - 3.79 (1H), 4.36 (3H), 7.18 (1H), 7.31 - 7.39 (2H), 7.39 - 7.53 (3H), 7.81 (1H), 8.15 (1H), 8.51 (1H), 9.00 (1H).

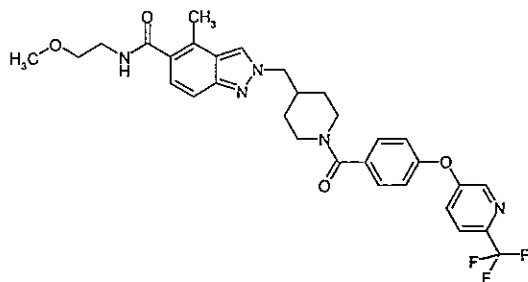
Example 348: 2-({[1-(benzothiazol-2-ylcarbonyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 28 mg of the title compound was obtained from 75 mg of 71a and 61 mg of benzothiazol-2-carboxylic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.34 (2H), 1.62 (2H), 2.32 - 2.45 (1H), 2.53 (3H), 2.82 - 3.00 (1H), 3.26 (1H), 3.28 (3H), 3.40 (2H), 3.43 - 3.50 (2H), 4.38 (3H), 4.98 - 5.23 (1H), 7.19 (1H), 7.43 (1H), 7.59 (2H), 8.01 - 8.30 (3H), 8.53 (1H).

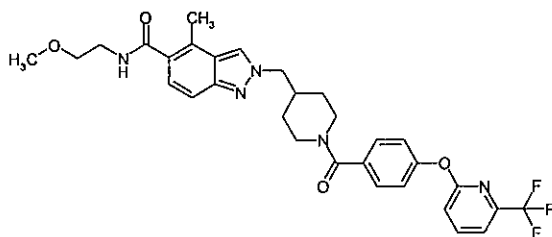
Example 349: N-(2-methoxyethyl)-4-methyl-2-{{1-(4-{{6-(trifluoromethyl)pyridin-3-yl}oxy}-benzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 34 mg of the title compound was obtained from 75 mg of 71a and 96 mg of 303b in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.24 (2H), 1.37 - 1.68 (2H), 2.22 - 2.38 (1H), 2.53 (3H), 2.71 - 3.14 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.44 - 3.49 (2H), 3.52 - 3.84 (1H), 4.36 (3H), 7.09 - 7.29 (3H), 7.35 - 7.53 (3H), 7.65 (1H), 7.92 (1H), 8.13 (1H), 8.51 (1H), 8.60 (1H).

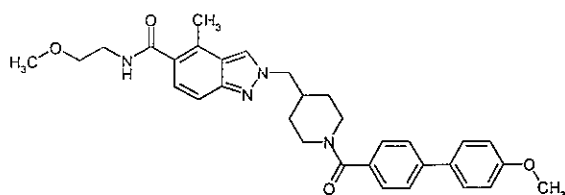
Example 350: N-(2-methoxyethyl)-4-methyl-2-{{1-(4-{{6-(trifluoromethyl)pyridin-2-yl}oxy}-benzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 36 mg of the title compound was obtained from 100 mg of 71a and 129 mg of 4-{{6-(trifluoromethyl)pyridin-2-yl}oxy}benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.20 - 1.34 (2H), 1.36 - 1.66 (2H), 2.24 - 2.39 (1H), 2.53 (3H), 2.70 - 3.16 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.44 - 3.49 (2H), 3.53 - 3.82 (1H), 4.36 (3H), 7.18 (1H), 7.22 - 7.30 (2H), 7.32 - 7.50 (4H), 7.67 (1H), 8.06 - 8.21 (2H), 8.51 (1H).

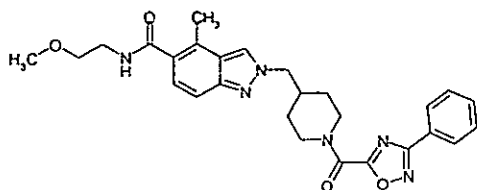
Example 351: 2-({1-[(4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 15 mg of the title compound was obtained from 100 mg of 71a and 104 mg of 4'-methoxy-biphenyl-4-carboxylic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.24 (2H), 1.37 - 1.64 (2H), 2.19 - 2.32 (1H), 2.53 (3H), 2.74 - 3.10 (2H), 3.28 (3H), 3.40 (2H), 3.45 (2H), 3.67 - 3.91 (4H), 4.21 - 4.64 (3H), 7.04 (2H), 7.19 (1H), 7.41 (2H), 7.66 (5H), 8.04 - 8.22 (1H), 8.51 (1H).

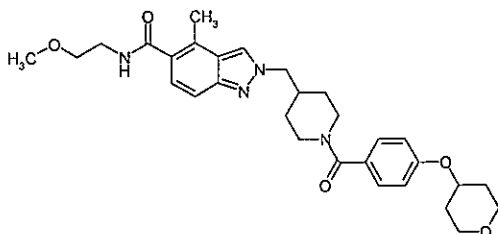
Example 352: N-(2-methoxyethyl)-4-methyl-2-((1-[(3-phenyl-1,2,4-oxadiazol-5-yl)carbonyl]-piperidin-4-yl)methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 12 mg of the title compound was obtained from 100 mg of 71a and 78 mg of 3-phenyl-1,2,4-oxadiazol-5-carboxylic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.33 (2H), 1.50 - 1.72 (2H), 2.31 - 2.45 (1H), 2.53 (3H), 2.86 - 3.04 (1H), 3.16 - 3.26 (1H), 3.28 (3H), 3.41 (2H), 3.43 - 3.51 (2H), 3.94 - 4.13 (1H), 4.28 - 4.55 (3H), 7.19 (1H), 7.43 (1H), 7.52 - 7.72 (3H), 7.96 - 8.09 (2H), 8.13 (1H), 8.52 (1H).

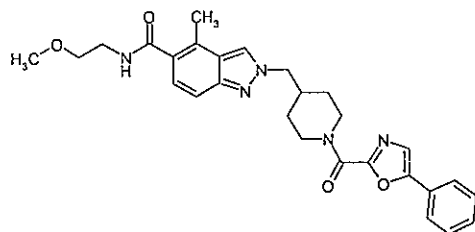
Example 353: N-(2-methoxyethyl)-4-methyl-2-((1-[4-(3,4,5,6-tetrahydro-2H-pyran-4-yloxy)-benzoyl]piperidin-4-yl)methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 21 mg of the title compound was obtained from 100 mg of 71a and 91 mg of 4-(3,4,5,6-tetrahydro-2H-pyran-4-yloxy)benzoic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.38 - 1.70 (4H), 1.97 (2H), 2.18 - 2.41 (1H), 2.53 (3H), 2.75 - 3.02 (2H), 3.28 (3H), 3.35 - 3.54 (8H), 3.84 (2H), 4.35 (2H), 4.52 - 4.77 (1H), 7.00 (2H), 7.18 (1H), 7.30 (2H), 7.42 (1H), 8.13 (1H), 8.50 (1H).

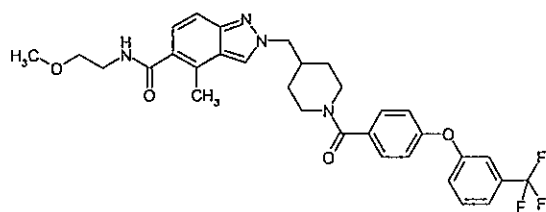
Example 354: N-(2-methoxyethyl)-4-methyl-2-({1-[(5-phenyloxazol-2-yl)carbonyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 10 mg of the title compound was obtained from 100 mg of 71a and 77 mg of 5-phenyl-oxazol-2-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.41 (2H), 1.59 (2H), 2.28 - 2.45 (1H), 2.52 - 2.56 (3H), 2.85 (1H), 3.20 (1H), 3.28 (3H), 3.36 - 3.43 (2H), 3.44 - 3.50 (2H), 4.38 (3H), 4.57 - 4.74 (1H), 7.19 (1H), 7.39 - 7.57 (4H), 7.74 - 7.82 (2H), 7.88 (1H), 8.13 (1H), 8.52 (1H).

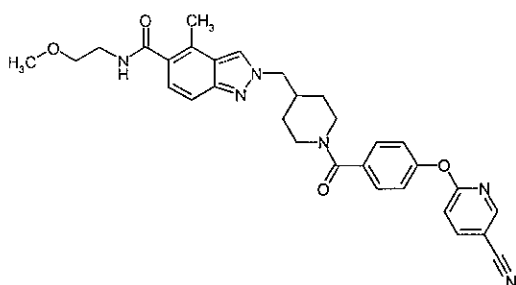
Example 355: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[3-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 40 mg of the title compound was obtained from 100 mg of 71a and 115 mg of 4-[3-(trifluoromethyl)phenoxy]benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.35 - 1.65 (2H), 2.19 - 2.38 (1H), 2.52 - 2.56 (3H), 2.71 - 3.12 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.53 - 3.87 (1H), 4.35 (3H), 7.07 - 7.15 (2H), 7.18 (1H), 7.30 - 7.48 (5H), 7.54 (1H), 7.64 (1H), 8.13 (1H), 8.50 (1H).

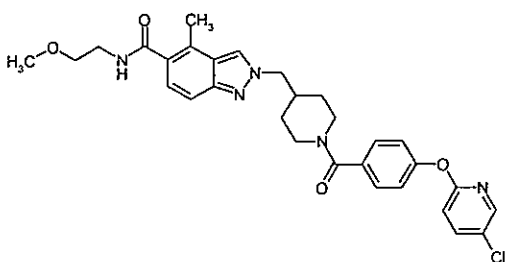
Example 356: 2-[(1-{4-[(5-cyanopyridin-2-yl)oxy]benzoyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 15 mg of the title compound was obtained from 100 mg of 71a and 98 mg of 4-[(5-cyanopyridin-2-yl)oxy]benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.21 - 1.35 (2H), 1.36 - 1.75 (2H), 2.23 - 2.39 (1H),
 5 2.53 (3H), 2.69 - 3.19 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.46 (2H), 3.53 - 3.85 (1H), 4.36 (3H),
 7.18 (1H), 7.23 - 7.33 (3H), 7.36 - 7.53 (3H), 8.13 (1H), 8.34 (1H), 8.51 (1H), 8.66 (1H).

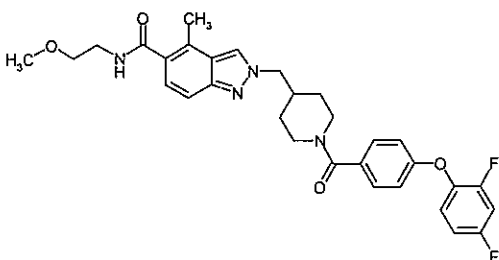
Example 357: 2-[(1-{4-[(5-chloropyridin-2-yl)oxy]benzoyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



10 Analogously to Example 1b, Version B, 40 mg of the title compound was obtained from 100 mg of 71a and 102 mg of 4-[(5-chloropyridin-2-yl)oxy]benzoic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.25 (2H), 1.36 - 1.70 (2H), 2.20 - 2.39 (1H), 2.53
 (3H), 2.66 - 3.15 (2H), 3.28 (3H), 3.40 (2H), 3.43 - 3.50 (2H), 3.55 - 3.90 (1H), 4.36 (3H), 7.08 -
 7.28 (4H), 7.42 (3H), 7.99 (1H), 8.10 - 8.30 (2H), 8.51 (1H).

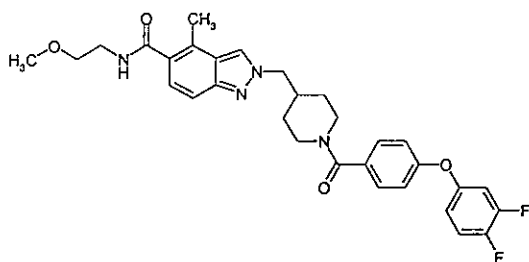
15 **Example 358:** 2-({1-[4-(2,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 22 mg of the title compound was obtained from 100 mg of 71a and 102 mg of 4-(2,4-difluorophenoxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.21 (2H), 1.33 - 1.76 (2H), 2.20 - 2.39 (1H), 2.52 (3H), 2.69 - 3.13 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.53 - 3.90 (1H), 4.35 (3H),
5 6.91 - 7.04 (2H), 7.11 - 7.22 (2H), 7.30 - 7.46 (4H), 7.51 (1H), 8.13 (1H), 8.50 (1H).

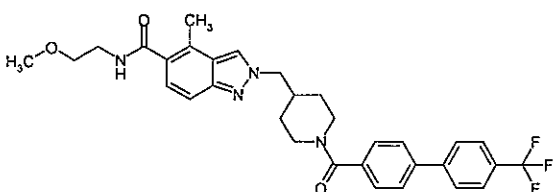
Example 359: 2-({1-[4-(3,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 23 mg of the title compound was obtained from 100 mg
10 of 71a and 102 mg of 4-(3,4-difluorophenoxy)benzoic acid in DMF.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.30 (2H), 1.35 - 1.70 (2H), 2.20 - 2.40 (1H), 2.52 (3H), 2.74 - 3.14 (2H), 3.28 (3H), 3.40 (2H), 3.43 - 3.51 (2H), 3.52 - 3.89 (1H), 4.35 (3H), 6.94 (1H), 7.05 (2H), 7.18 (1H), 7.30 (1H), 7.36 - 7.59 (4H), 8.15 (1H), 8.51 (1H).

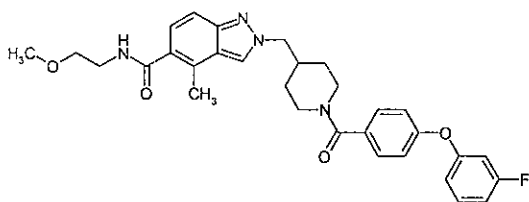
Example 360: N-(2-methoxyethyl)-4-methyl-2-[(1-{[4'-(trifluoromethyl)biphenyl-4-yl]-carbonyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
15



Analogously to Example 1b, Version B, 44 mg of the title compound was obtained from 100 mg of 71a and 109 mg of 4'-(trifluoromethyl)biphenyl-4-carboxylic acid in DMF

LC-MS: R_t = 1.28 min, MS (ES⁺): m/z = 579 (M+H)⁺.

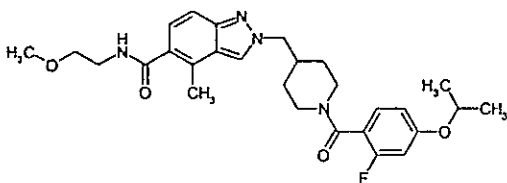
Example 361: 2-({1-[4-(3-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
20



Analogously to Example 1b, Version B, 43 mg of the title compound was obtained from 100 mg of 71a and 95 mg of 4-(3-fluorophenoxy)benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.31 (2H), 1.33 - 1.69 (2H), 2.19 - 2.38 (1H),
 5 2.52 (3H), 2.70 - 3.11 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.49 (2H), 3.54 - 3.85 (1H), 4.35 (3H), 6.85 - 7.13 (5H), 7.18 (1H), 7.35 - 7.55 (4H), 8.15 (1H), 8.51 (1H).

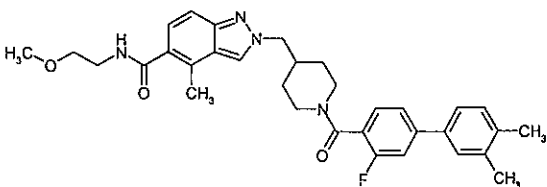
Example 362: 2-([1-(2-fluoro-4-isopropoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



10 Analogously to Example 1b, Version B, 31 mg of the title compound was obtained from 75 mg of 71a and 41 mg of 2-fluoro-4-isopropoxybenzoic acid.

¹H-NMR (300 MHz, chloroform-d): δ [ppm]= 1.21 - 1.42 (8H), 1.51 (1H), 1.69 (1H), 2.38 (1H),
 2.56 - 2.80 (4H), 2.87 - 3.08 (1H), 3.36 (3H), 3.48 - 3.73 (4H), 4.28 (2H), 4.51 (1H), 4.74 (1H),
 5.28 (1H), 6.19 (1H), 6.54 (1H), 6.67 (1H), 7.13 - 7.38 (2H), 7.50 (1H), 7.94 (1H).

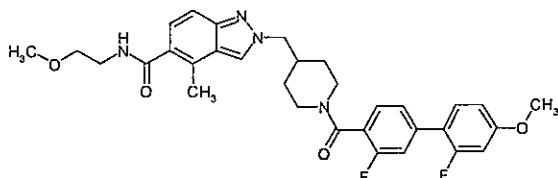
15 **Example 363:** 2-([1-((3-fluoro-3',4'-dimethylbiphenyl-4-yl)carbonyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 35 mg of the title compound was obtained from 75 mg of 71a and 50 mg of 3-fluoro-3',4'-dimethylbiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, chloroform-d): δ [ppm]= 1.25 - 1.51 (2H), 1.57 (1H), 1.76 (1H), 2.25 - 2.37 (6H), 2.45 (1H), 2.61 - 2.71 (3H), 2.79 (1H), 3.06 (1H), 3.40 (3H), 3.53 - 3.77 (5H), 4.34 (2H), 4.83 (1H), 6.04 - 6.30 (1H), 7.15 - 7.47 (7H), 7.55 (1H), 7.98 (1H).

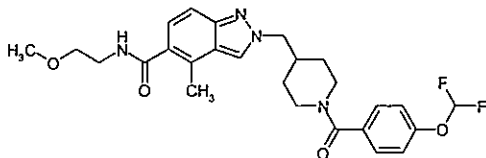
Example 364: 2-({1-[(2',3-difluoro-4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 28 mg of the title compound was obtained from 90 mg of 71a and 65 mg of 3,2'-difluoro-4'-methoxybiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, chloroform-d): δ [ppm]= 1.25 - 1.50 (2H), 1.50 - 1.84 (2H), 2.44 (1H), 2.66 (3H), 2.78 (1H), 3.03 (1H), 3.39 (3H), 3.53 - 3.76 (5H), 3.85 (3H), 4.23 - 4.45 (2H), 4.83 (1H), 6.17 (1H), 6.64 - 6.92 (2H), 7.18 - 7.48 (5H), 7.54 (1H), 7.97 (1H).

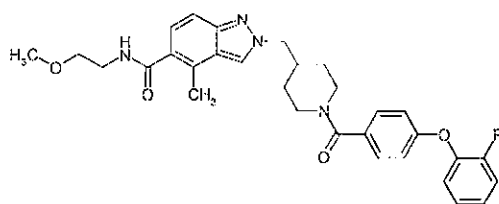
Example 365: 2-({1-[4-(difluoromethoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 23 mg of the title compound was obtained from 75 mg of 71a and 38 mg of 4-(difluoromethoxy)benzoic acid.

¹H-NMR (300 MHz, chloroform-d): δ [ppm]= 1.26 (2H), 1.59 (2H), 2.26 - 2.49 (1H), 2.62 (3H), 2.84 (2H), 3.36 (3H), 3.43 (1H), 3.50 - 3.96 (4H), 4.29 (2H), 4.46 - 5.02 (1H), 6.13 - 6.25 (1H), 6.26 - 6.82 (1H), 7.11 (2H), 7.22 - 7.43 (3H), 7.50 (1H), 7.93 (1H).

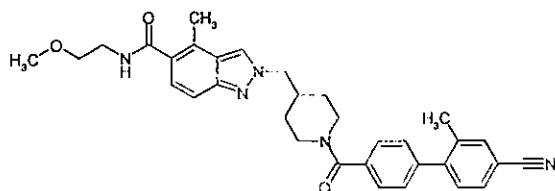
Example 366: 2-({1-[4-(2-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



120 mg of compound 54, 26 mg of o-fluorophenol, 152 mg of caesium carbonate, 6.7 mg of copper(I) bromide and 3.2 mg of (2-pyridyl)acetone were suspended in 6 ml dimethyl sulphoxide and stirred for 3 days at 80°C. The reaction mixture was filtered, the filter cake
 5 washed with water and ethyl acetate, and the aqueous phase extracted several times with ethyl acetate. The combined organic phases were dried with sodium sulphate and concentrated. The residue was purified by HPLC and 12 mg of the title compound was obtained.

¹H-NMR (300 MHz, chloroform-d): δ [ppm]= 1.16 - 1.47 (2H), 1.62 (2H), 2.43 (1H), 2.66 (3H), 2.85 (2H), 3.39 (3H), 3.53 - 3.74 (4H), 3.79 - 4.19 (1H), 4.33 (2H), 4.47 - 5.02 (1H), 6.15 (1H),
 10 6.96 (2H), 7.07 - 7.23 (3H), 7.30 - 7.45 (3H), 7.54 (1H), 7.96 (1H).

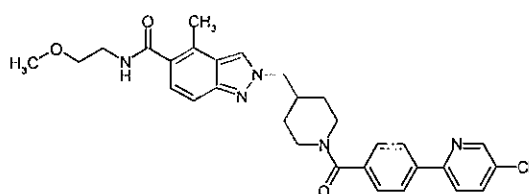
Example 367: 2-({1-[(4'-cyano-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 40 mg of the title compound was obtained from 80 mg of 54 and
 15 38 mg of (2-methyl-4-cyanophenyl)boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.20 - 1.34 (2H), 1.36 - 1.69 (2H), 2.28 (4H), 2.53 (3H), 2.69 - 3.19 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.54 - 3.77 (1H), 4.37 (3H), 7.18 (1H), 7.37 - 7.50 (6H), 7.74 (1H), 7.83 (1H), 8.15 (1H), 8.52 (1H).

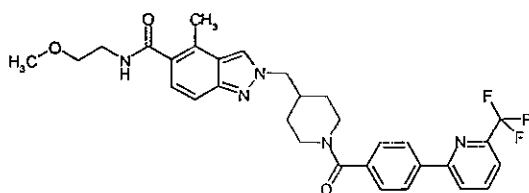
Example 368: 2-({1-[4-(5-chloropyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
 20



200 mg of the aryl bromide prepared in Example 54 together with 187 mg of (5-chloropyridin-2-yl)-2-boronic acid pinacol ester were first placed in 3 ml DMF, treated with 48 mg of 1,1'-bis(diphenylphosphino)ferrocenodichloropalladium(II), 39 mg of copper(I)chloride 254 mg of caesium carbonate and 22 mg of 1,1'-bis(diphenylphosphino)ferrocene and heated for ca. 1
 5 day at 70°C, until further progress of the reaction could no longer be observed. The reaction mixture was treated with water and saturated sodium hydrogen carbonate solution, treated several times with ethyl acetate, and the combined organic phases dried with sodium sulphate and concentrated. After HPLC purification, this yielded 92 mg of the title compound.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.36 (2H), 1.37 - 1.72 (2H), 2.23 - 2.41 (1H),
 10 2.52 - 2.57 (3H), 2.70 - 3.13 (2H), 3.28 (3H), 3.39 (2H), 3.45 (2H), 3.51 - 3.80 (1H), 4.36 (3H), 7.18 (1H), 7.36 - 7.52 (3H), 8.05 (2H), 8.09 - 8.22 (3H), 8.51 (1H), 8.73 (1H).

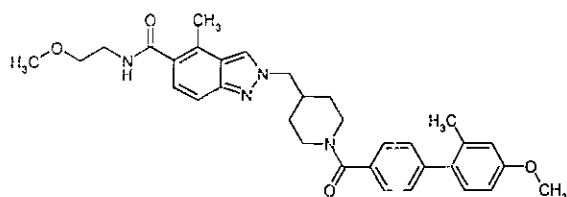
Example 369: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[6-(trifluoromethyl)pyridin-2-yl]-benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



15 Analogously to Example 317, 33 mg of the title compound was obtained from 100 mg of 54 and 80 mg of [6-(trifluoromethyl)pyridin-2-yl]boronic acid pinacol ester under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.18 - 1.35 (2H), 1.36 - 1.67 (2H), 2.25 - 2.41 (1H),
 2.53 (3H), 2.70 - 3.15 (2H), 3.28 (3H), 3.35 - 3.43 (2H), 3.43 - 3.51 (2H), 3.53 - 3.73 (1H), 4.37
 (3H), 7.18 (1H), 7.42 (1H), 7.53 (2H), 7.89 (1H), 8.06 - 8.26 (4H), 8.33 (1H), 8.51 (1H).

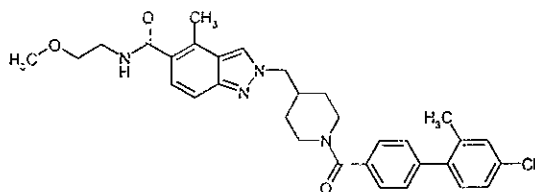
20 **Example 370:** N-(2-methoxyethyl)-2-({1-[(4'-methoxy-2'-methylbiphenyl-4-yl)carbonyl]-piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 52 mg of the title compound was obtained from 100 mg of 54 and 48 mg of (4-methoxy-2-methylphenyl)boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.16 - 1.35 (2H), 1.37 - 1.67 (2H), 2.23 (3H), 2.26 - 2.42 (1H), 2.53 (3H), 2.70 - 3.12 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.77 (4H), 4.36 (3H), 6.77 - 6.92 (2H), 7.10 - 7.22 (2H), 7.39 (5H), 8.15 (1H), 8.52 (1H).

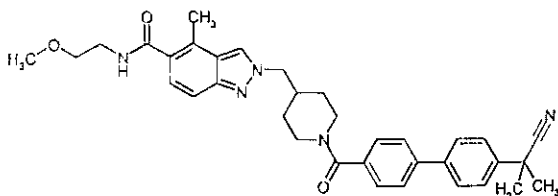
Example 371: 2-({1-[(4'-chloro-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 43 mg of the title compound was obtained from 100 mg of 54 and 50 mg of (4-chloro-2-methylphenyl)boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.27 (2H), 1.38 - 1.70 (2H), 2.23 (4H), 2.53 (3H), 2.70 - 3.15 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.49 (2H), 3.55 - 3.84 (1H), 4.36 (3H), 7.12 - 7.27 (2H), 7.28 - 7.35 (1H), 7.36 - 7.47 (6H), 8.15 (1H), 8.52 (1H).

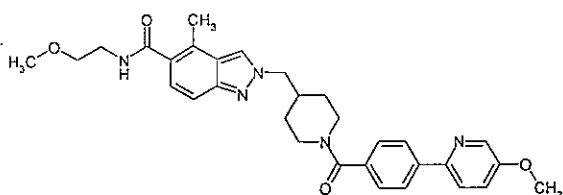
Example 372: 2-[(1-[(4'-(2-cyanopropan-2-yl)biphenyl-4-yl)carbonyl]piperidin-4-yl)-methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 67 mg of the title compound was obtained from 100 mg of 54 and 55 mg of [4-(1-cyano-1-methylethyl)phenyl]boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.28 (2H), 1.36 - 1.65 (2H), 1.72 (6H), 2.24 - 2.40 (1H), 2.53 (3H), 2.70 - 3.14 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.49 (2H), 3.54 - 3.77 (1H), 4.36 (3H), 7.18 (1H), 7.38 - 7.50 (3H), 7.62 (2H), 7.75 (4H), 8.15 (1H), 8.52 (1H).

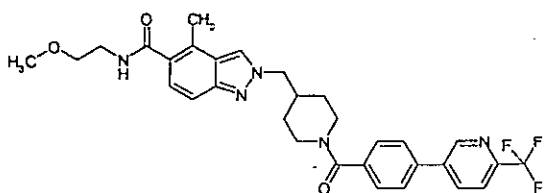
Example 373: N-(2-methoxyethyl)-2-({1-[4-(5-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}-methyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 368, 12 mg of the title compound was obtained from 200 mg of 54 and 183 mg of (5-methoxypyridin-2-yl)boronic acid pinacol ester under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.22 - 1.33 (2H), 1.36 - 1.66 (2H), 2.23 - 2.38 (1H), 2.53 (3H), 2.68 - 3.12 (2H), 3.28 (3H), 3.35 - 3.42 (2H), 3.43 - 3.50 (2H), 3.54 - 3.78 (1H), 3.88 (3H), 4.36 (3H), 7.18 (1H), 7.36 - 7.55 (4H), 7.96 (1H), 8.07 (2H), 8.13 (1H), 8.39 (1H), 8.51 (1H).

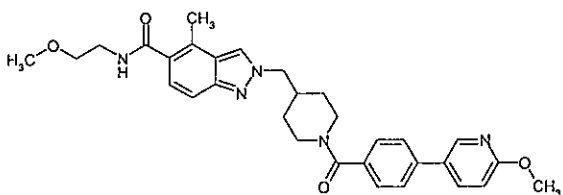
Example 374: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[6-(trifluoromethyl)pyridin-3-yl]-benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 317, 37 mg of the title compound was obtained from 100 mg of 54 and 56 mg of [2-(trifluoromethyl)pyridin-5-yl]boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.36 (2H), 1.36 - 1.69 (2H), 2.21 - 2.42 (1H), 2.53 (3H), 2.70 - 3.14 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.53 - 3.72 (1H), 4.36 (3H), 7.18 (1H), 7.42 (1H), 7.53 (2H), 7.89 (2H), 8.01 (1H), 8.15 (1H), 8.41 (1H), 8.52 (1H), 9.13 (1H).

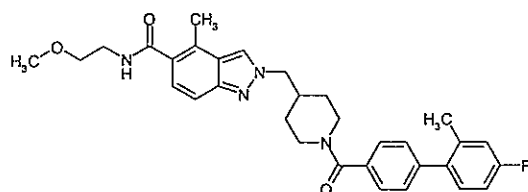
Example 375: N-(2-methoxyethyl)-2-({1-[4-(6-methoxypyridin-3-yl)benzoyl]piperidin-4-yl)-methyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 50 mg of the title compound was obtained from 100 mg of 54 and 47 mg of (2-methoxypyridin-5-yl)boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.26 (2H), 1.34 - 1.70 (2H), 2.21 - 2.40 (1H), 2.53 (3H), 2.69 - 3.15 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.49 (2H), 3.51 - 3.79 (1H), 3.90 (3H), 4.36 (3H), 6.93 (1H), 7.18 (1H), 7.36 - 7.52 (3H), 7.72 (2H), 8.05 (1H), 8.15 (1H), 8.46 - 8.57 (2H).

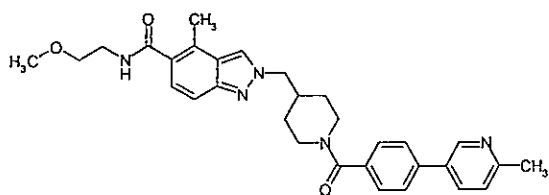
- 5 **Example 376:** 2-({1-[(4'-fluoro-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 261 mg of the title compound was obtained from 300 mg of 54 and 135 mg of (4-fluoro-2-methylphenyl)boronic acid under reflux.

- 10 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.27 (2H), 1.37 - 1.70 (2H), 2.24 (4H), 2.53 (3H), 2.71 - 3.14 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.55 - 3.83 (1H), 4.37 (3H), 7.05 - 7.12 (1H), 7.18 (2H), 7.22 - 7.30 (1H), 7.40 (5H), 8.14 (1H), 8.52 (1H).

Example 377: N-(2-methoxyethyl)-4-methyl-2-({1-[4-(6-methylpyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

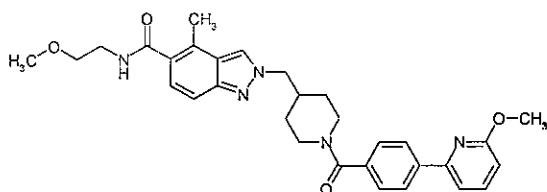


15

Analogously to Example 317, 45 mg of the title compound was obtained from 80 mg of 54 and 32 mg of (2-methylpyridin-5-yl)boronic acid under reflux.

- 20 ¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.26 (2H), 1.37 - 1.71 (2H), 2.18 - 2.39 (1H), 2.52 - 2.58 (3H), 2.73 - 3.16 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.45 (2H), 3.54 - 3.79 (1H), 4.36 (3H), 7.18 (1H), 7.30 - 7.54 (4H), 7.76 (2H), 8.00 (1H), 8.15 (1H), 8.51 (1H), 8.78 (1H).

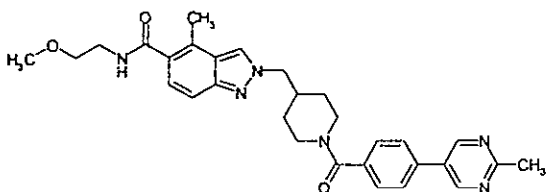
Example 378: N-(2-methoxyethyl)-2-({1-[4-(6-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 30 mg of the title compound was obtained from 100 mg of 54 and 45 mg of (6-methoxypyridin-2-yl)boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.26 (2H), 1.35 - 1.68 (2H), 2.32 (1H), 2.52 (3H),
 5 2.70 - 3.13 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.53 - 3.74 (1H), 3.96 (3H), 4.36 (3H), 6.81 (1H), 7.18 (1H), 7.37 - 7.67 (4H), 7.81 (1H), 8.15 (3H), 8.51 (1H).

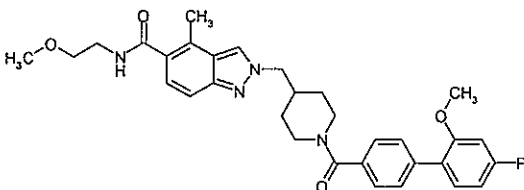
Example 379: N-(2-methoxyethyl)-4-methyl-2-({1-[4-(2-methylpyrimidin-5-yl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



10 Analogously to Example 317, 36 mg of the title compound was obtained from 100 mg of 54 and 64 mg of (2-methylpyrimidin-5-yl)boronic acid pinacol ester under reflux.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.27 (2H), 1.38 - 1.70 (2H), 2.21 - 2.40 (1H), 2.53 (3H), 2.67 (3H), 2.73 - 3.16 (2H), 3.28 (3H), 3.40 (2H), 3.43 - 3.50 (2H), 3.53 - 3.76 (1H), 4.36 (3H), 7.18 (1H), 7.42 (1H), 7.51 (2H), 7.84 (2H), 8.13 (1H), 8.51 (1H), 9.05 (2H).

15 **Example 380:** 2-({1-[4-(4'-fluoro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

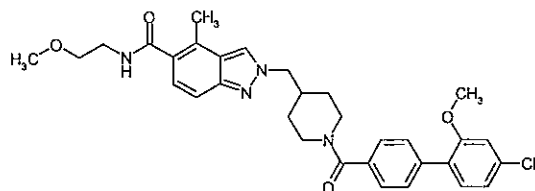


Analogously to Example 317, 47 mg of the title compound was obtained from 100 mg of 54 and 50 mg of (4-fluoro-2-methoxyphenyl)boronic acid under reflux.

20 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.34 (2H), 1.37 - 1.71 (2H), 2.22 - 2.40 (1H), 2.53 (3H), 2.70 - 3.15 (2H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.56 - 3.75 (1H), 3.79

(3H), 4.36 (3H), 6.87 (1H), 6.99 - 7.10 (1H), 7.18 (1H), 7.30 - 7.46 (4H), 7.50 (2H), 8.13 (1H), 8.51 (1H).

Example 381: 2-({1-[(4'-chloro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



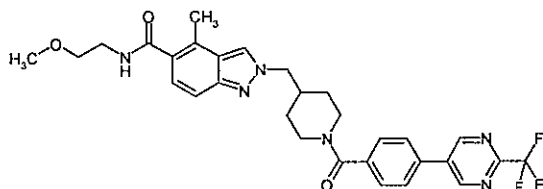
5

Analogously to Example 317, 87 mg of the title compound was obtained from 100 mg of 54 and 54 mg of (4-fluoro-2-methoxyphenyl)boronic acid under reflux.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.26 (2H), 1.39 - 1.69 (2H), 2.19 - 2.39 (1H), 2.53 (3H), 2.75 - 3.14 (2H), 3.28 (3H), 3.40 (2H), 3.45 (2H), 3.57 - 3.74 (1H), 3.81 (3H), 4.36 (3H), 7.04 - 7.25 (3H), 7.28 - 7.62 (6H), 8.13 (1H), 8.51 (1H).

10

Example 382: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[2-(trifluoromethyl)pyrimidin-5-yl]-benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



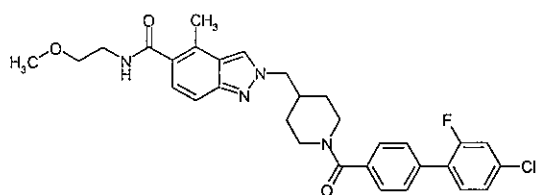
15

Analogously to Example 317, 14 mg of the title compound was obtained from 100 mg of 54 and 56 mg of [2-(trifluoromethyl)pyrimidin-5-yl]boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.24 (2H), 1.38 - 1.73 (2H), 2.22 - 2.41 (1H), 2.53 (3H), 2.70 - 2.89 (1H), 2.94 - 3.18 (1H), 3.28 (3H), 3.36 - 3.43 (2H), 3.44 - 3.49 (2H), 3.51 - 3.70 (1H), 4.37 (3H), 7.18 (1H), 7.42 (1H), 7.57 (2H), 7.97 (2H), 8.13 (1H), 8.51 (1H), 9.43 (2H).

20

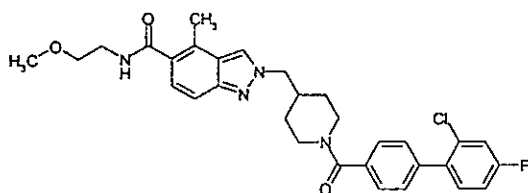
Example 383: 2-({1-[(4'-chloro-2'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 51 mg of the title compound was obtained from 100 mg of 54 and 51 mg of (4-chloro-2-fluorophenyl)boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.27 (2H), 1.38 - 1.68 (2H), 2.32 (1H), 2.53 (3H),
 5 2.71 - 2.91 (1H), 2.93 - 3.16 (1H), 3.28 (3H), 3.40 (2H), 3.45 (2H), 3.53 - 3.78 (m, 1H), 4.37 (3H), 7.18 (1H), 7.33 - 7.52 (4H), 7.53 - 7.74 (4H), 8.13 (1H), 8.51 (1H).

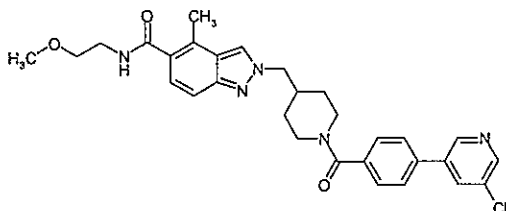
Example 384: 2-({1-[(2'-chloro-4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



10 Analogously to Example 317, 26 mg of the title compound was obtained from 100 mg of 54 and 51 mg of (2-chloro-4-fluorophenyl)boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.28 (2H), 1.38 - 1.67 (2H), 2.23 - 2.39 (1H), 2.53 (3H), 2.71 - 2.91 (1H), 2.94 - 3.15 (1H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.54 - 3.74 (1H), 4.37 (3H), 7.18 (1H), 7.33 (1H), 7.39 - 7.63 (7H), 8.13 (1H), 8.51 (1H).

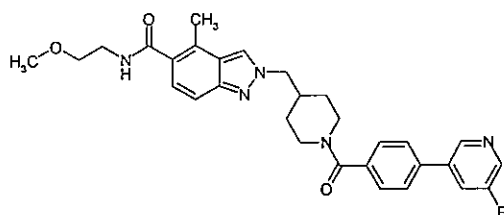
15 **Example 385:** 2-({1-[4-(5-chloropyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 22 mg of the title compound was obtained from 100 mg of 54 and 46 mg of (5-chloropyridin-3-yl)boronic acid under reflux.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.21 - 1.35 (2H), 1.36 - 1.69 (2H), 2.23 - 2.40 (1H), 2.53 (3H), 2.69 - 2.90 (1H), 2.92 - 3.16 (1H), 3.28 (3H), 3.40 (2H), 3.45 (2H), 3.53 - 3.73 (1H), 4.36 (3H), 7.18 (1H), 7.35 - 7.57 (3H), 7.86 (2H), 8.15 (1H), 8.30 (1H), 8.52 (1H), 8.65 (1H), 8.90 (1H).

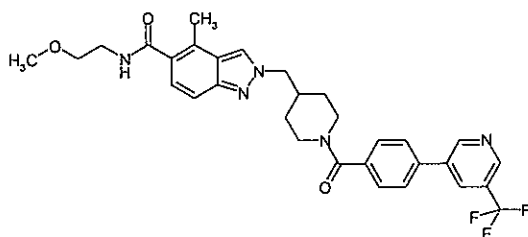
- 5 **Example 386:** 2-({1-[4-(5-fluoropyridin-3-yl)benzoyl]piperidin-4-yl)methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 57 mg of the title compound was obtained from 100 mg of 54 and 41 mg of (5-fluoropyridin-3-yl)boronic acid under reflux.

- 10 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.27 (2H), 1.37 - 1.70 (2H), 2.24 - 2.40 (1H), 2.52 - 2.55 (3H), 2.70 - 2.89 (1H), 2.93 - 3.15 (1H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.53 - 3.73 (1H), 4.36 (3H), 7.18 (1H), 7.42 (1H), 7.50 (2H), 7.86 (2H), 8.07 - 8.18 (2H), 8.51 (1H), 8.61 (1H), 8.84 (1H).

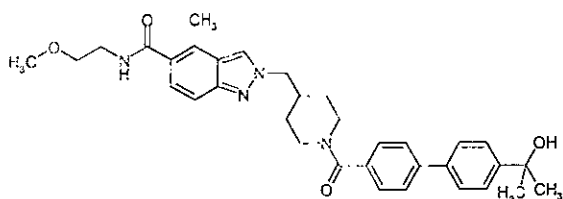
- 15 **Example 387:** N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[5-(trifluoromethyl)pyridin-3-yl]-benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 317, 73 mg of the title compound was obtained from 100 mg of 54 and 56 mg of [5-(trifluoromethyl)pyridin-3-yl]boronic acid under reflux.

- 20 ¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.20 - 1.35 (2H), 1.36 - 1.67 (2H), 2.23 - 2.40 (1H), 2.53 (3H), 2.69 - 2.92 (1H), 2.93 - 3.17 (1H), 3.28 (3H), 3.35 - 3.43 (2H), 3.43 - 3.49 (2H), 3.53 - 3.75 (1H), 4.37 (3H), 7.18 (1H), 7.42 (1H), 7.52 (2H), 7.93 (2H), 8.07 - 8.20 (1H), 8.45 - 8.58 (2H), 9.00 (1H), 9.24 (1H).

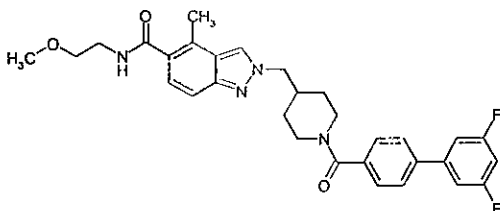
Example 388: 2-[(1-{[4'-(2-hydroxypropan-2-yl)biphenyl-4-yl]carbonyl}piperidin-4-yl)-methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 46 mg of the title compound was obtained from 100 mg of 54 and 77 mg of (4-hydroxy-tert-butylphenyl)boronic acid pinacol ester under reflux.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.26 (2H), 1.45 (8H), 2.22 - 2.42 (1H), 2.53 (3H), 2.71 - 2.90 (1H), 2.91 - 3.14 (1H), 3.28 (3H), 3.40 (2H), 3.43 - 3.50 (2H), 3.56 - 3.80 (1H), 4.36 (3H), 5.06 (1H), 7.18 (d, 1H), 7.37 - 7.50 (3H), 7.50 - 7.67 (4H), 7.71 (2H), 8.15 (1H), 8.52 (1H).

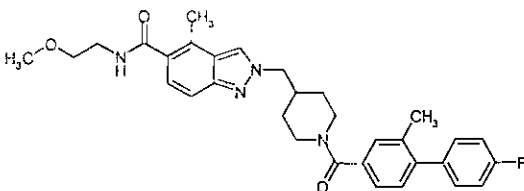
Example 389: 2-[(1-{[3',5'-difluorobiphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 22 mg of the title compound was obtained from 100 mg of 54 and 46 mg of (3,5-difluorophenyl)boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.18 - 1.35 (2H), 1.36 - 1.70 (2H), 2.19 - 2.39 (1H), 2.52 - 2.56 (3H), 2.71 - 2.92 (1H), 2.93 - 3.17 (1H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.49 (2H), 3.52 - 3.71 (1H), 4.36 (3H), 7.11 - 7.33 (2H), 7.37 - 7.54 (5H), 7.81 (2H), 8.13 (1H), 8.51 (1H).

Example 390: 2-[(1-{[4'-fluoro-2-methylbiphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

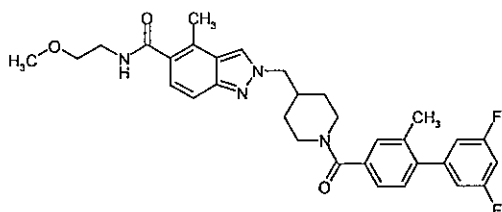


- 270 -

Analogously to Example 317, 25 mg of the title compound was obtained from 60 mg of 332 and 19 mg of (4-fluorophenyl)boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.25 (2H), 1.37 - 1.67 (2H), 2.24 (4H), 2.53 (3H), 2.69 - 2.87 (1H), 2.92 - 3.14 (1H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.50 (2H), 3.56 - 3.81 (1H), 4.36 (3H), 7.11 - 7.34 (6H), 7.36 - 7.46 (3H), 8.15 (1H), 8.52 (1H).

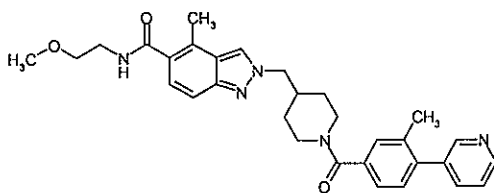
Example 391: 2-({1-[(3',5'-difluoro-2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 317, 30 mg of the title compound was obtained from 60 mg of 332 and 27 mg of (3,5-difluorophenyl)boronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.23 (2H), 1.36 - 1.69 (2H), 2.27 (4H), 2.53 (3H), 2.69 - 2.90 (1H), 2.92 - 3.15 (1H), 3.28 (3H), 3.36 - 3.43 (2H), 3.45 (2H), 3.53 - 3.76 (1H), 4.36 (3H), 7.08 - 7.21 (3H), 7.22 - 7.34 (4H), 7.42 (1H), 8.15 (1H), 8.52 (1H).

Example 392: N-(2-methoxyethyl)-4-methyl-2-({1-[3-methyl-4-(3-pyridyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

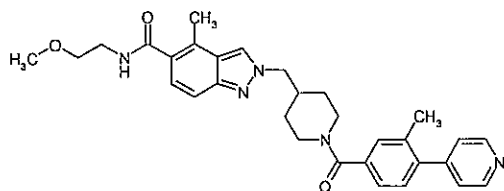


Analogously to Example 317, 19 mg of the title compound was obtained from 60 mg of 332 and 21 mg of pyridine-3-ylboronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.20 - 1.33 (2H), 1.37 - 1.71 (2H), 2.26 (4H), 2.53 (3H), 2.70 - 2.88 (1H), 2.94 - 3.17 (1H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.49 (2H), 3.56 - 3.82 (1H), 4.36 (3H), 7.18 (1H), 7.23 - 7.37 (3H), 7.39 - 7.56 (2H), 7.83 (1H), 8.15 (1H), 8.52 (1H), 8.56 - 8.65 (2H).

- 271 -

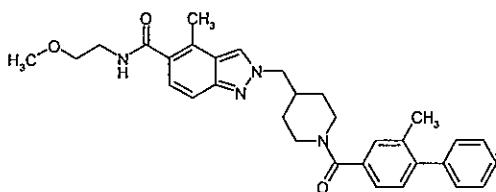
Example 393: N-(2-methoxyethyl)-4-methyl-2-({1-[3-methyl-4-(4-pyridyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 317, 19 mg of the title compound was obtained from 60 mg of 332 and 21 mg of pyridine-4-ylboronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.18 - 1.33 (2H), 1.37 - 1.67 (2H), 2.27 (4H), 2.52 - 2.56 (3H), 2.70 - 2.86 (1H), 2.93 - 3.13 (1H), 3.28 (3H), 3.36 - 3.43 (2H), 3.43 - 3.49 (2H), 3.55 - 3.75 (1H), 4.36 (3H), 7.18 (1H), 7.23 - 7.36 (3H), 7.38 - 7.48 (3H), 8.15 (1H), 8.52 (1H), 8.60 - 8.69 (2H).

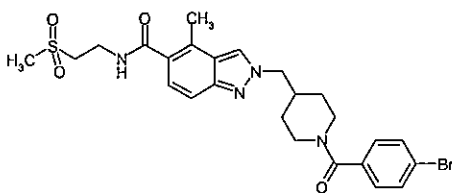
Example 394: N-(2-methoxyethyl)-4-methyl-2-({1-[(2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide



Analogously to Example 317, 57 mg of the title compound was obtained from 60 mg of 332 and 21 mg of phenylboronic acid under reflux.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.34 (2H), 1.37 - 1.67 (2H), 2.25 (4H), 2.53 (3H), 2.68 - 2.88 (1H), 2.92 - 3.17 (1H), 3.28 (3H), 3.36 - 3.43 (2H), 3.45 (2H), 3.59 - 3.81 (1H), 4.36 (3H), 7.13 - 7.32 (4H), 7.33 - 7.50 (6H), 8.15 (1H), 8.52 (1H).

Example 395: 2-{{1-[4-(4-bromobenzoyl)piperidin-4-yl]methyl}-N-(2-mesyloethyl)-4-methyl-2H-indazole-5-carboxamide

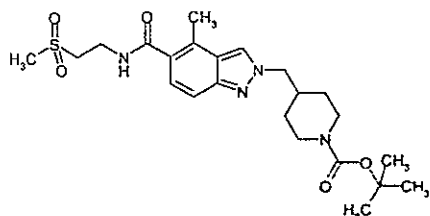


Analogously to Example 1b, Version B, 383 mg of the title compound was obtained from 1 g of 395b and 797 mg of 4-bromobenzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.21 - 1.32 (2H), 1.34 - 1.67 (2H), 2.21 - 2.38 (1H), 2.55 (3H), 2.68 - 2.87 (1H), 3.05 (4H), 3.39 (2H), 3.45 - 3.59 (1H), 3.66 (2H), 4.35 (3H), 7.23 (1H), 7.28 - 7.37 (2H), 7.43 (1H), 7.59 - 7.70 (2H), 8.34 (1H), 8.52 (1H).

The starting material was prepared as follows:

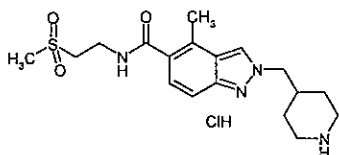
Example 395a: Tert-butyl 4-({5-[N-(2-mesyloethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}-methyl)piperidin-1-carboxylate



10 Analogously to Example 266a, 1.26 g of the title compound was obtained from 2.506 g of 1c and 3.022 g of 2-mesyloethylamine hydrochloride.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.02 - 1.17 (2H), 1.34 - 1.54 (11H), 2.08 - 2.24 (1H), 2.55 (3H), 2.73 (2H), 3.06 (3H), 3.34 - 3.47 (2H), 3.57 - 3.76 (2H), 3.91 (2H), 4.32 (2H), 7.23 (1H), 7.43 (1H), 8.33 (1H), 8.52 (1H).

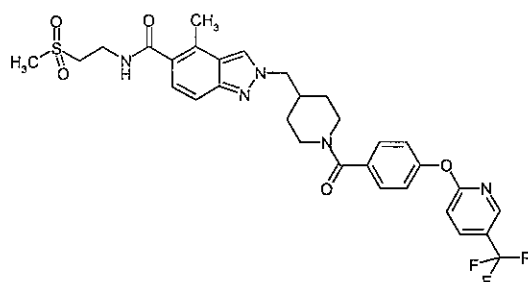
15 **Example 395b:** N-(2-mesyloethyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 258b from 1.26 g of compound 395a, 1.29 g of the title compound was obtained, which was reacted without further purification.

20 **Example 396:** N-(2-mesyloethyl)-4-methyl-2-{[1-(4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}-benzoyl)piperidin-4-yl]methyl}-2H-indazole-5-carboxamide

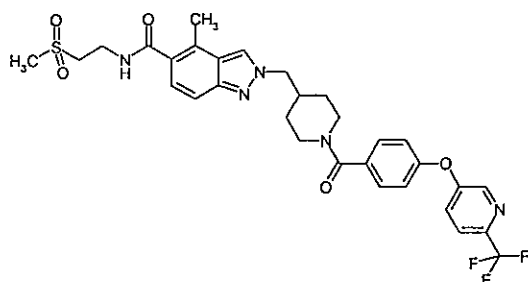
- 273 -



Analogously to Example 1b, Version B, 30 mg of the title compound was obtained from 80 mg of 395b and 90 mg of 4-{{[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.35 (2H), 1.36 - 1.72 (2H), 2.22 - 2.41 (1H),
 5 2.55 (3H), 3.06 (5H), 3.39 (2H), 3.49 - 3.89 (3H), 4.37 (3H), 7.20 - 7.32 (4H), 7.40 - 7.49 (3H),
 8.25 (1H), 8.34 (1H), 8.53 (1H), 8.58 (1H).

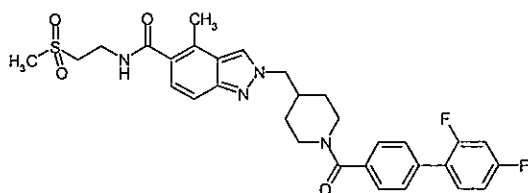
Example 397: N-(2-mesylethyl)-4-methyl-2-{{[1-(4-{{[6-(trifluoromethyl)pyridin-3-yl]oxy}-benzoyl]piperidin-4-yl]methyl}-2H-indazole-5-carboxamide



10 Analogously to Example 1b, Version B, 30 mg of the title compound was obtained from 80 mg of 395b and 90 mg of 303b in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.24 (2H), 1.35 - 1.70 (2H), 2.22 - 2.40 (1H), 2.55
 (3H), 3.06 (5H), 3.39 (2H), 3.55 - 3.75 (3H), 4.36 (3H), 7.20 - 7.28 (3H), 7.40 - 7.49 (3H), 7.65
 (1H), 7.92 (1H), 8.34 (1H), 8.53 (1H), 8.60 (1H).

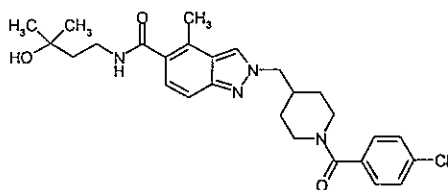
15 **Example 398:** 2-({[1-((2',4'-difluorobiphenyl-4-yl)carbonyl)piperidin-4-yl]methyl}-N-(2-mesylethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 25 mg of the title compound was obtained from 80 mg of 395b and 74 mg of 2',4'-difluoro-biphenyl-4-carboxylic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.21 - 1.35 (2H), 1.37 - 1.70 (2H), 2.22 - 2.41 (1H), 2.55 (3H), 3.05 (5H), 3.38 (2H), 3.50 - 3.83 (3H), 4.37 (3H), 7.15 - 7.29 (2H), 7.31 - 7.52 (4H), 7.55 - 7.73 (3H), 8.34 (1H), 8.53 (1H).

Example 399: 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-N-(3-hydroxy-3-methylbutyl)-4-methyl-2H-indazole-5-carboxamide

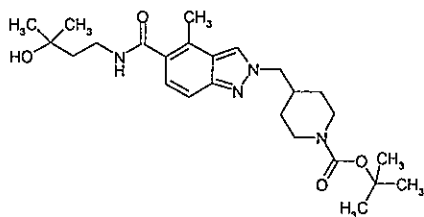


Analogously to Example 1b, Version B, 37 mg of the title compound was obtained from 160 mg of 399b and 105 mg of 4-chlorobenzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.14 (6H), 1.17 - 1.32 (2H), 1.33 - 1.71 (4H), 2.23 - 2.37 (1H), 2.51 (3H), 2.68 - 2.87 (1H), 2.87 - 3.13 (1H), 3.31 (2H), 3.47 - 3.55 (1H), 4.34 (s, 4H), 7.17 (1H), 7.33 - 7.45 (3H), 7.50 (2H), 7.97 - 8.09 (1H), 8.50 (1H).

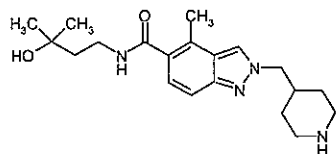
The starting material was prepared as follows:

Example 399a: Tert-butyl 4-({5-[N-(3-hydroxy-3-methylbutyl)carbamoyl]-4-methyl-2H-indazol-2-yl}methyl)piperidin-1-carboxylate



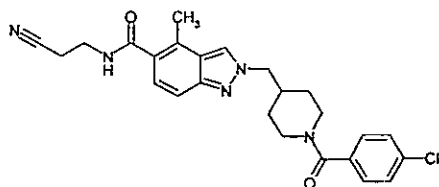
Analogously to Example 1b, Version B, 150 mg of 1d and 69 mg of 4-amino-2-methylbutan-2-ol in DMF were reacted at 60°C. After phase separation and extraction, the combined organic phases were additionally washed with 0.1 N hydrochloric acid and saturated sodium hydrogen carbonate solution. This yielded 132 mg of the title compound, which was used in the next step without further purification.

Example 399b: N-(3-hydroxy-3-methylbutyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide



Analogously to Example 258b, from 132 mg of 399a, 162 mg of the title compound was obtained, which was used in the next step without further purification.

Example 400: 2-[[1-(4-chlorobenzoyl)piperidin-4-yl]methyl]-N-(2-cyanoethyl)-4-methyl-2H-indazole-5-carboxamide

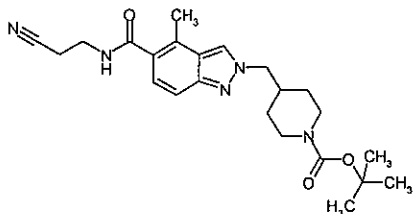


Analogously to Example 1b, Version B, 55 mg of the title compound was obtained from 177 mg of 400b and 128 mg of 4-chlorobenzoic acid in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm] = 1.23 (2H), 1.33 - 1.70 (2H), 2.20 - 2.38 (1H), 2.56 (3H), 2.78 (3H), 2.90 - 3.11 (1H), 3.47 (3H), 4.35 (3H), 7.22 (1H), 7.34 - 7.56 (5H), 8.44 - 8.57 (2H).

The starting material was prepared as follows:

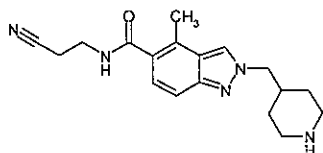
Example 400a: Tert-butyl 4-({5-[N-(2-cyanoethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}-methyl)-piperidin-1-carboxylate



Analogously to Example 1b, Version B, 150 mg of 1d and 42 mg of 3-aminopropanitrile in DMF were reacted at 60°C. After phase separation and extraction, the combined organic phases were additionally washed with 0.1 N hydrochloric acid and saturated sodium hydrogen carbonate

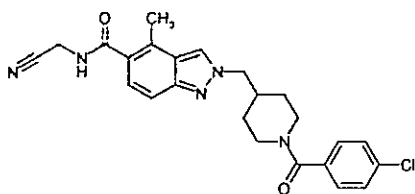
solution. This yielded 137 mg of the title compound, which was used in the next step without further purification.

Example 400b: N-(2-cyanoethyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide



- 5 Analogously to Example 258b, from 137 mg of 400a 177 mg of the title compound was obtained, which was used in the next step without further purification.

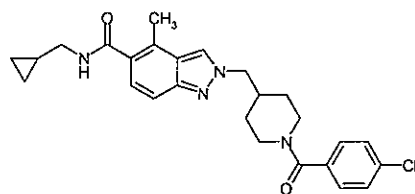
Example 401: 2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(cyanomethyl)-4-methyl-2H-indazole-5-carboxamide



- 10 Analogously to Example 1b, Version B, 75 mg of the title compound was obtained from 90 mg of 238c and 29 mg of aminoacetonitrile in DMF.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.21 - 1.33 (23H), 1.34 - 1.69 (2H), 2.15 - 2.41 (1H), 2.57 (3H), 2.75 - 2.86 (1H), 2.93 - 3.14 (1H), 3.39 - 3.69 (1H), 4.29 (2H), 4.33 - 4.55 (3H), 7.18 - 7.28 (1H), 7.39 (2H), 7.43 - 7.55 (3H), 8.57 (1H), 8.77 - 8.90 (1H).

- 15 **Example 402:** 2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(cyclopropylmethyl)-4-methyl-2H-indazole-5-carboxamide



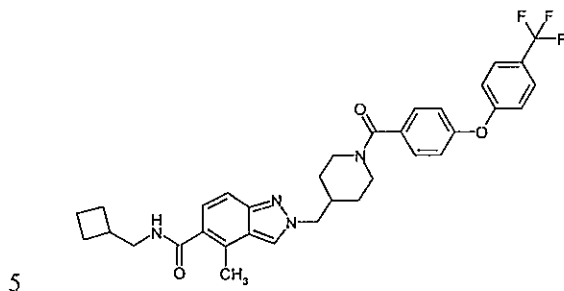
Analogously to Example 1b, Version B, 44 mg of the title compound was obtained from 90 mg of 238c) and 19 mg of cyclopropylmethylamine in DMF.

- 20 ¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.23 (2H), 0.33 - 0.53 (2H), 0.94 - 1.11 (1H), 1.20 - 1.33 (2H), 1.34 - 1.68 (2H), 2.21 - 2.41 (1H), 2.54 (3H), 2.69 - 2.85 (1H), 2.93 - 3.08 (1H), 3.13

- 277 -

(2H), 3.44 - 3.68 (1H), 4.35 (3H), 7.18 (1H), 7.33 - 7.46 (3H), 7.47 - 7.60 (2H), 8.07 - 8.30 (1H), 8.51 (1H).

Example 403: N-(cyclobutylmethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



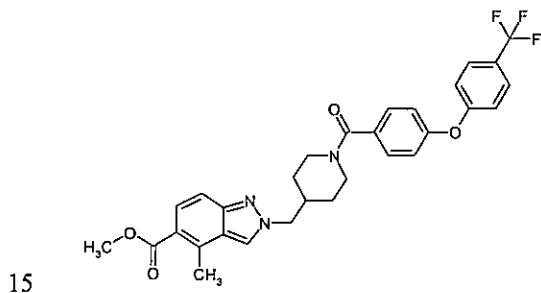
Analogously to Example 1b, Version B, 48 mg of the title compound was obtained from 137 mg of 403b and 31 mg of (cyclobutylamino)methylamine hydrochloride.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.17 - 1.32 (2H), 1.35 - 1.64 (2H), 1.67 - 1.90 (4H), 1.93 - 2.08 (2H), 2.22 - 2.38 (1H), 2.51 - 2.52 (3H), 2.69 - 2.88 (1H), 2.90 - 3.13 (1H), 3.20 - 3.30 (2H), 3.51 - 3.83 (1H), 4.36 (3H), 7.09 - 7.27 (5H), 7.35 - 7.52 (3H), 7.76 (2H), 8.13 (1H), 8.51 (1H).

10

The starting material was prepared as follows:

Example 403a: Methyl 4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazol-5-carboxylate

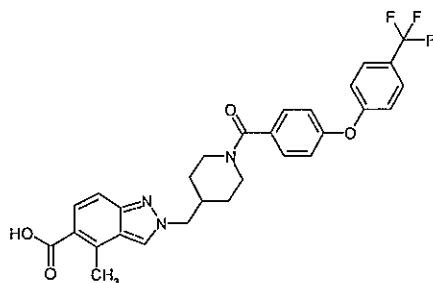


Analogously to Example 1b, Version B, 2.73 g of the title compound was obtained from 3.76 g of 238a and 3.28 g of 4-[4-(trifluoromethyl)phenoxy]benzoic acid.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.35 (2H), 1.38 - 1.74 (2H), 2.17 - 2.41 (1H), 2.76 (3H), 2.78 - 3.14 (2H), 3.44 - 3.76 (1H), 3.82 (3H), 4.37 (3H), 7.06 - 7.28 (4H), 7.38 - 7.50 (3H), 7.67 (1H), 7.76 (2H), 8.71 (1H).

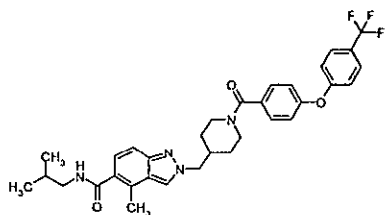
20

Example 403b: 4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)-methyl]-2H-indazol-5-carboxylic acid



Analogously to Example 1d, from 1.05 g of 403a, 970 mg of the title compound was obtained,
5 which was used in the next step without further purification.

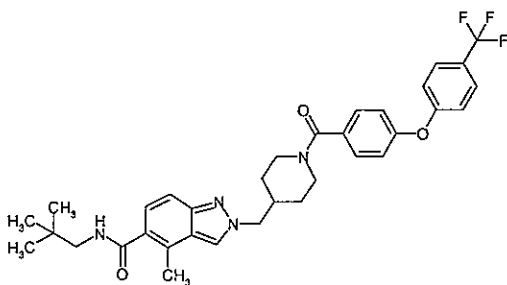
Example 404: N-isobutyl-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 35 mg of the title compound was obtained from 100 mg
10 of 403b and 14 mg of isobutylamine.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.91 (6H), 1.24 (2H), 1.34 - 1.69 (2H), 1.82 (1H), 2.19 - 2.42 (1H), 2.52 (3H), 2.71 - 2.90 (1H), 3.06 (3H), 3.51 - 3.85 (1H), 4.36 (3H), 7.05 - 7.27 (5H), 7.37 - 7.52 (3H), 7.76 (2H), 8.16 (1H), 8.51 (1H).

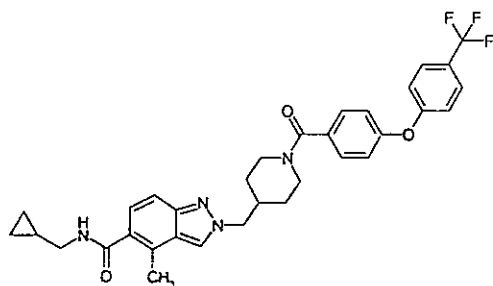
Example 405: 4-methyl-N-neopentyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 36 mg of the title compound was obtained from 100 mg of 403b and 16 mg of neopentylamine.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 0.92 (9H), 1.17 - 1.32 (2H), 1.34 - 1.69 (2H), 2.21 - 2.38 (1H), 2.52 - 2.57 (3H), 2.70 - 2.88 (1H), 3.08 (3H), 3.53 - 3.85 (1H), 4.36 (3H), 7.04 - 7.28 (5H), 7.35 - 7.52 (3H), 7.76 (2H), 8.12 (1H), 8.51 (1H).

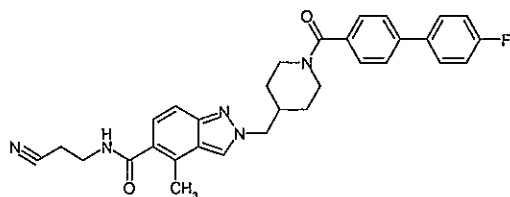
Example 406: N-(cyclopropylmethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]-benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 33 mg of the title compound was obtained from 100 mg of 403b and 26 mg of cyclopropylmethylamine.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 0.23 (2H), 0.35 - 0.54 (2H), 0.93 - 1.11 (1H), 1.23 (2H), 1.36 - 1.67 (2H), 2.18 - 2.41 (1H), 2.54 (3H), 2.66 - 3.22 (4H), 3.47 - 3.85 (1H), 4.36 (3H), 7.03 - 7.29 (5H), 7.35 - 7.53 (3H), 7.76 (2H), 8.22 (1H), 8.52 (1H).

Example 407: N-(2-cyanoethyl)-2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}-methyl)-4-methyl-2H-indazole-5-carboxamide

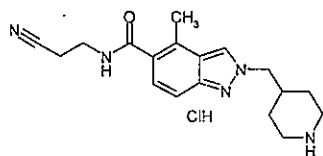


Analogously to Example 1b, Version B, 37 mg of the title compound was obtained from 140 mg of 407a and 84 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.18 - 1.37 (2H), 1.37 - 1.68 (2H), 2.22 - 2.41 (1H), 2.56 (3H), 2.78 (4H), 3.39 - 3.53 (2H), 3.56 - 3.82 (1H), 4.37 (3H), 7.14 - 7.37 (3H), 7.39 - 7.53 (3H), 7.63 - 7.83 (4H), 8.36 - 8.60 (2H).

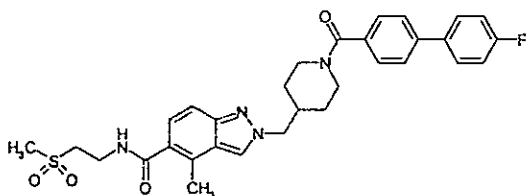
The starting material was prepared as follows:

Example 407a: N-(2-cyanoethyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 165 mg of 400a, 140 mg of the title compound was obtained, which was used in the next step without further purification.

Example 408: 2-(1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl)methyl)-N-(2-mesylyl)-4-methyl-2H-indazole-5-carboxamide

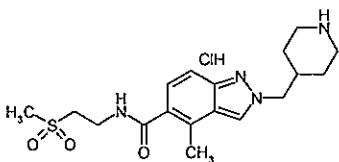


Analogously to Example 1b), Version B, 126 mg of the title compound was obtained from 300 mg of 408a) and 156 mg of 4'-fluorobiphenyl-4-carboxylic acid.

LC-MS: $R_t = 1.20$ min, MS (ES⁺): $m/z = 577$ (M+H)⁺.

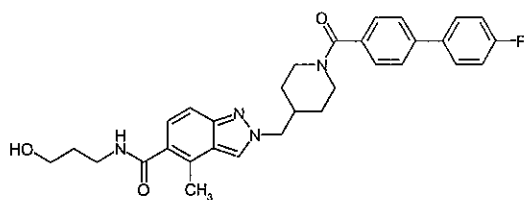
The starting material was prepared as follows:

Example 408a: N-(2-mesylyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 346 mg of 395a, 300 mg of the title compound was obtained, which was used in the next step without further purification.

Example 409: 2-(1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl)methyl)-N-(3-hydroxypropyl)-4-methyl-2H-indazole-5-carboxamide

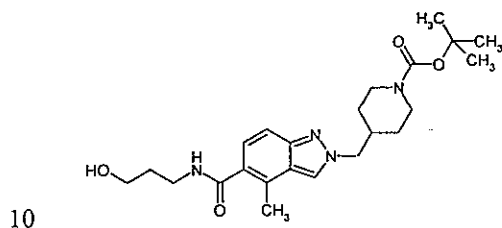


Analogously to Example 1b, Version B, 20 mg of the title compound was obtained from 124 mg of 409b) and 73 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.26 (2H), 1.38 - 1.59 (2H), 1.67 (2H), 2.23 - 2.42 (1H), 2.52 - 2.57 (3H), 2.69 - 3.17 (2H), 3.19 - 3.94 (6H), 4.36 (3H), 7.18 (1H), 7.31 (2H), 7.38 - 7.51 (3H), 7.63 - 7.86 (4H), 8.08 (1H), 8.51 (1H).

The starting material was prepared as follows:

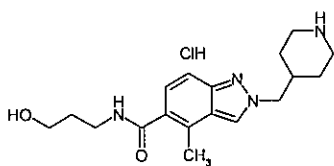
Example 409a: Tert-butyl 4-({5-[N-(3-hydroxypropyl)carbamoyl]-4-methyl-2H-indazol-2-yl}-methyl)piperidin-1-carboxylate



Analogously to Example 1b, Version B, 151 mg of the title compound was obtained from 150 mg of 1d and 30 mg of 3-amino-propan-1-ol.

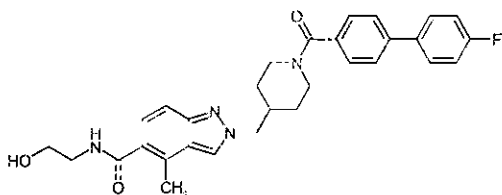
¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.01 - 1.30 (3H), 1.32 - 1.49 (10H), 1.67 (2H), 2.08 - 2.27 (1H), 2.52 (3H), 2.57 - 2.80 (2H), 3.29 (2H), 3.41 - 3.54 (2H), 3.81 - 4.00 (2H), 4.32 (2H), 4.48 (1H), 7.18 (1H), 7.42 (1H), 8.10 (1H), 8.50 (1H).

Example 409b: N-(3-hydroxypropyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazol-5-carboxamide



Analogously to Example 1a, from 246 mg of 409a, 124 mg of the title compound was obtained, which was used in the next step without further purification.

Example 410: 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-hydroxyethyl)-4-methyl-2H-indazole-5-carboxamide

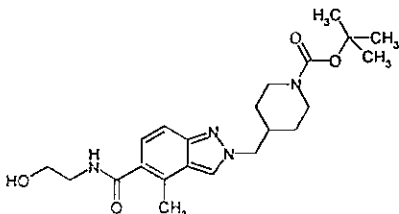


Analogously to Example 1b, Version B, 36 mg of the title compound was obtained from 117 mg of 410b and 72 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.26 (2H), 1.39 - 1.74 (2H), 2.21 - 2.40 (1H), 2.53 (3H), 2.75 - 3.16 (2H), 3.20 - 3.37 (2H), 3.43 - 3.74 (4H), 4.36 (3H), 7.21 (1H), 7.31 (2H), 7.38 - 7.52 (3H), 7.65 - 7.81 (4H), 8.03 (1H), 8.51 (1H).

The starting material was prepared as follows:

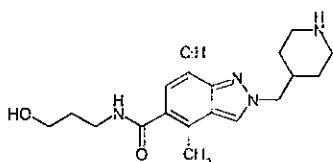
Example 410a: Tert-butyl 4-({5-[N-(2-hydroxyethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}-methyl)piperidin-1-carboxylate



Analogously to Example 1b, Version B, 143 mg of the title compound was obtained from 150 mg of 1d) and 25 mg of 2-aminoethanol.

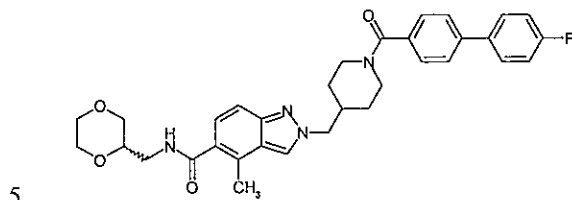
¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.01 - 1.30 (3H), 1.32 - 1.51 (10H), 2.07 - 2.26 (1H), 2.52 - 2.55 (3H), 2.58 - 2.82 (2H), 3.30 (2H), 3.45 - 3.58 (2H), 3.83 - 3.98 (2H), 4.32 (2H), 4.69 (1H), 7.21 (1H), 7.41 (1H), 8.05 (1H), 8.50 (1H).

Example 410b: N-(3-hydroxypropyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazol-5-carboxamide hydrochloride



Analogously to Example 1a) from 138 mg of 410a, 117 mg of the title compound was obtained, which was used in the next step without further purification.

Example 411: (+/-)-N-(1,4-dioxan-2-ylmethyl)-2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]-piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide



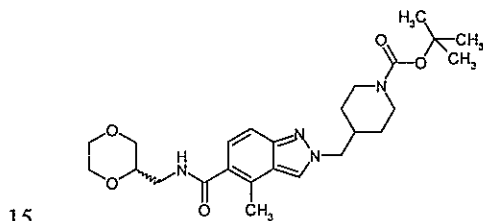
Analogously to Example 1b, Version B, 62 mg of the title compound was obtained from 226 mg of 411b) and 119 mg of 4'-fluorobiphenyl-4-carboxylic acid.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.34 (2H), 1.37 - 1.73 (2H), 2.23 - 2.42 (1H), 2.53 (3H), 2.69 - 2.90 (1H), 2.90 - 3.14 (1H), 3.17 - 3.30 (3H), 3.47 (1H), 3.58 (1H), 3.59 - 3.70 (3H), 3.71 - 3.82 (2H), 4.37 (3H), 7.19 (1H), 7.31 (2H), 7.38 - 7.53 (3H), 7.66 - 7.81 (4H), 8.18 (1H), 8.52 (1H).

10

The starting material was prepared as follows:

Example 411a: (+/-)-tert-butyl 4-({5-[N-(1,4-dioxan-2-ylmethyl)carbamoyl]-4-methyl-2H-indazol-2-yl}methyl)piperidin-1-carboxylate

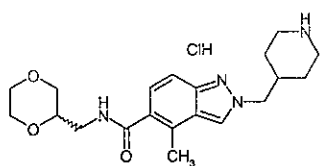


Analogously to Example 1b, Version B, 267 mg of the title compound was obtained from 250 mg of 1d and 78 mg of (+/-)-1,4-dioxan-2-ylmethylamine.

¹H-NMR (400 MHz, DMSO-d₆): δ [ppm]= 0.99 - 1.18 (2H), 1.38 (1H), 2.09 - 2.24 (1H), 2.53 (3H), 3.10 - 3.29 (4H), 3.41 - 3.52 (1H), 3.53 - 3.70 (4H), 3.70 - 3.81 (2H), 3.84 - 3.98 (2H), 4.32 (2H), 7.18 (1H), 7.41 (1H), 8.12 - 8.24 (1H), 8.51 (1H).

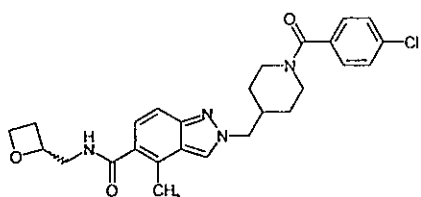
20

Example 411b: (+/-)-N-(1,4-dioxan-2-ylmethyl)-4-methyl-2-(4-piperidylmethyl)-2H-indazole-5-carboxamide hydrochloride



Analogously to Example 1a, from 261 mg of 411a, 226 mg of the title compound was obtained, which was used in the next step without further purification.

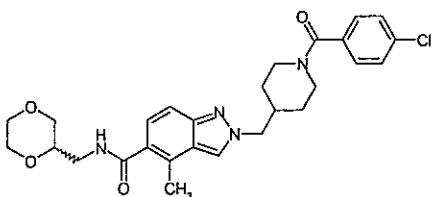
Example 412: (+/-)-2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methyl-N-(oxetan-2-ylmethyl)-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 55 mg of the title compound was obtained from 350 mg of 238c) and 83 mg of (+/-)-oxetan-2-ylmethylamine.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.16 - 1.31 (2H), 1.33 - 1.69 (2H), 2.19 - 2.38 (2H), 2.53 (3H), 2.58 - 2.66 (1H), 2.66 - 2.86 (1H), 2.90 - 3.14 (1H), 3.38 - 3.64 (3H), 4.35 (5H), 4.71 - 4.96 (1H), 7.19 (1H), 7.33 - 7.45 (3H), 7.47 - 7.55 (2H), 8.21 - 8.37 (1H), 8.51 (1H).

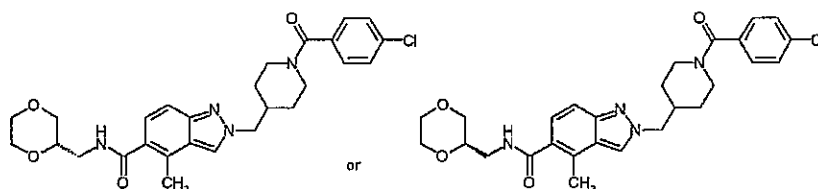
Example 413: (+/-)-2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(1,4-dioxan-2-ylmethyl)-4-methyl-2H-indazole-5-carboxamide



Analogously to Example 1b, Version B, 185 mg of the title compound was obtained from 376 mg of 238c and 111 mg of (+/-)-1,4-dioxan-2-ylmethylamine hydrochloride.

¹H-NMR (300 MHz, DMSO-d₆): δ [ppm]= 1.19 - 1.32 (2H), 1.33 - 1.66 (2H), 2.20 - 2.40 (1H), 2.53 (3H), 2.71 - 2.86 (1H), 2.90 - 3.11 (1H), 3.16 - 3.32 (3H), 3.40 - 3.59 (3H), 3.60 - 3.70 (2H), 3.71 - 3.83 (2H), 4.35 (3H), 7.18 (1H), 7.31 - 7.46 (3H), 7.46 - 7.60 (2H), 8.21 (1H), 8.51 (1H).

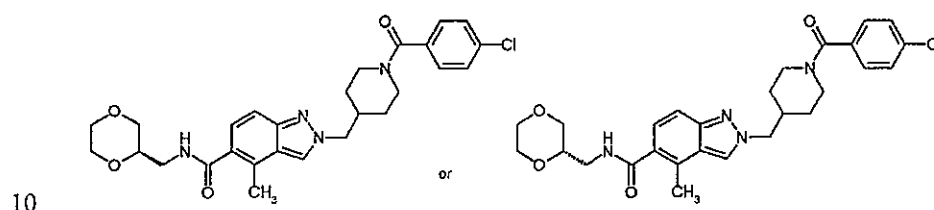
Example 414: (R or S)-2-{{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(1,4-dioxan-2-ylmethyl)-4-methyl-2H-indazole-5-carboxamide



From 185 mg of the racemate prepared in Example 413, 51 mg of the title compound together with 58 mg of the slower-eluting enantiomer (Example 415) were obtained by racemate separation by means of preparative chiral HPLC (Method D).

Analytical chiral HPLC: 12.62 min.

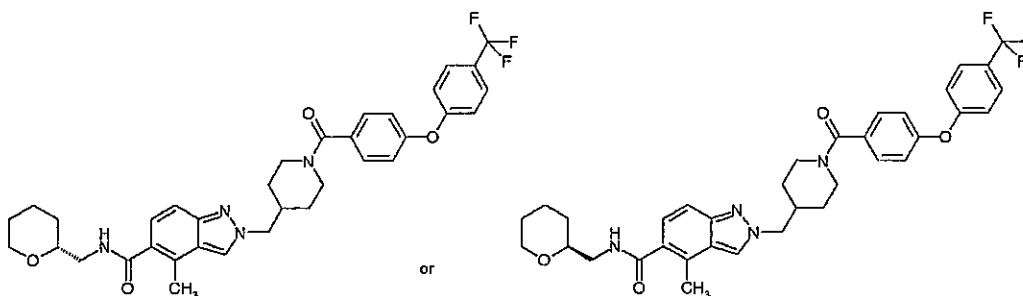
Example 415: (S or R)-2-{{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(1,4-dioxan-2-ylmethyl)-4-methyl-2H-indazole-5-carboxamide



From 185 mg of the racemate prepared in Example 413, 58 mg of the title compound together with 51 mg of the faster-eluting enantiomer (Example 414) were obtained by racemate separation by means of preparative chiral HPLC (Method D).

Analytical chiral HPLC: 13.68 min.

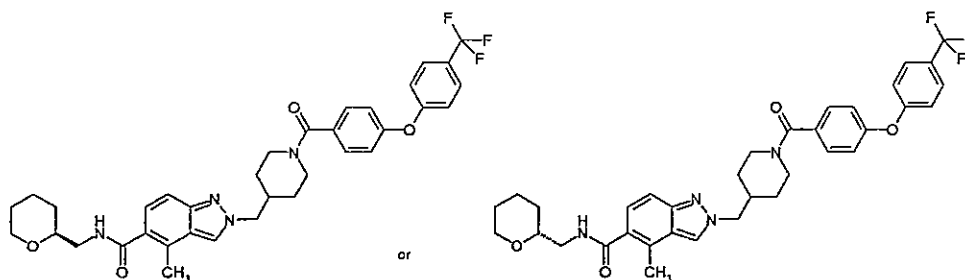
Example 416: (R or S)-4-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



From 231 mg of the racemate prepared in Example 248, 24 mg of the title compound together with 24 mg of the slower-eluting enantiomer (Example 417) were obtained by racemate separation by means of preparative chiral HPLC (Method E).

Analytical chiral HPLC: 7.08 min.

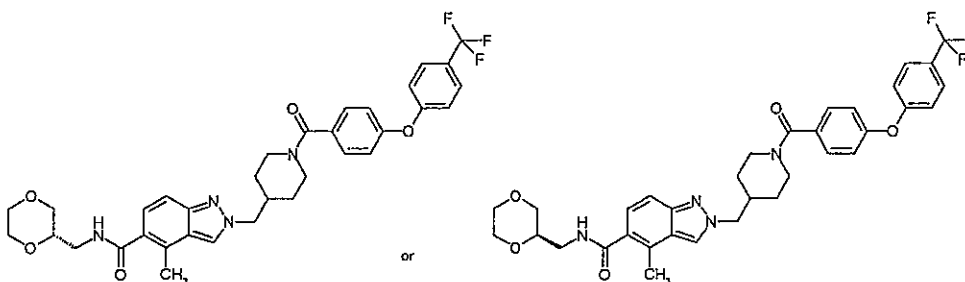
- 5 **Example 417:** (S or R)-4-methyl-N-(3,4,5,6-tetrahydro-2H-pyran-2-ylmethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



- From 231 mg of the racemate prepared in Example 248, 24 mg of the title compound together with 24 mg of the faster-eluting enantiomer (Example 416) were obtained by racemate separation by means of preparative chiral HPLC (Method E).
- 10

Analytical chiral HPLC: 8.98 min.

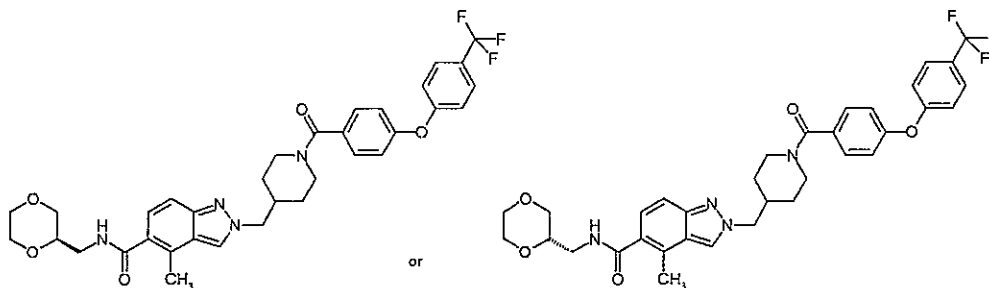
- Example 418:** (R or S)-N-(1,4-dioxan-2-ylmethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



- 15 From 280 mg of the racemate prepared in Example 249, 65 mg of the title compound together with 75 mg of the slower-eluting enantiomer (Example 419) were obtained by racemate separation by means of preparative chiral HPLC (Method D (injection volume: 0.1 ml; detection: UV 210 nM)).

Analytical chiral HPLC: 13.66 min.

Example 419: (R or S)-N-(1,4-dioxan-2-ylmethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)-phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide



From 280 mg of the racemate prepared in Example 249, 75 mg of the title compound together
5 with 65 mg of the faster-eluting enantiomer (Example 418) were obtained by racemate separation by means of preparative chiral HPLC (Method D (injection volume: 0.1 ml; Detection: UV 210 nm)).

Analytical chiral HPLC: 14.90 min.

Biological Examples:

The following biological examples which are described in sections 1.-5 and have only been carried out using EP2 receptor antagonists are already contained in patent application PCT/EP2012/073556, unpublished at the priority date of the present specification.

5 1. Detection of antagonism to the human prostaglandin E₂ (subtype EP2) receptor signal**1. 1 Detection principle**

The binding of PGE₂ to the EP2 subtype of the human PGE₂ receptor induces the activation of membrane-located adenylate cyclases and leads to the formation of cAMP. In the presence of the phosphodiesterase inhibitor IBMX, the cAMP accumulated owing to this stimulation and
10 released by cell lysis is used in a competitive detection method. In this test, the cAMP present in the lysate competes with a fluorescence-labelled cAMP (cAMP-d2) for binding to an Eu cryptate-labelled anti-cAMP antibody.

In the absence of cellular cAMP, a maximal signal is produced, which is attributable to the binding of this cAMP-d2 molecule to the antibody. After excitation of the cAMP-d2 molecule at
15 337 nm, there is a fluorescence resonance energy transfer (FRET) to the Eu cryptate molecules of the anti-cAMP antibody (labelled therewith), followed by a long-lasting emission signal at 665 nm (and at 620 nm). Both signals are measured in a suitable measurement instrument with a time delay, i.e. after decay of the background fluorescence. Any increase in the low FRET signal caused by prostaglandin E₂ administration (measured as well ratio change =
20 $\text{emission}_{665\text{nm}}/\text{emission}_{620\text{nm}} * 10000$) indicates the action of antagonists.

1.2. Detection method**1.2.1. Test for antagonism (data per well of a 384-well plate):**

To a test plate with the substance solutions already added (0.05 µl; 100% DMSO, concentration range from 0.8 nM – 16.5 µM) were added 4 µl of a cAMP-d2/cell suspension (625000
25 cells/ml). After a 20-minute preincubation at room temperature (RT), 2 µl of a 3xPGE2 solution (1.5 nM, in PBS-IBMX) were added and incubated in the presence of the agonist for a further 60 mins at RT (volume: ~ 6 µl). Next the reaction was stopped by addition of 2 µl of lysis buffer and incubated for a further 20 mins at RT before the actual measurement (volume: ~ 8 µl).

2. Detection of antagonism to the human prostaglandin E₂ (subtype EP₄) receptor signal

2.1 Detection principle

The binding of PGE₂ to the EP₄ subtype of the human PGE₂ receptor induces the activation of membrane-located adenylate cyclases and leads to the formation of cAMP. In the presence of the phosphodiesterase inhibitor IBMX, the cAMP accumulated owing to this stimulation and released by cell lysis is used in a competitive detection method. In this test, the cAMP present in the lysate competes with a fluorescence-labelled cAMP (cAMP-d2) for binding to an Eu cryptate-labelled anti-cAMP antibody.

In the absence of cellular cAMP a maximal signal is produced, which is attributable to the binding of this cAMP-d2 molecule to the antibody. After excitation of the cAMP-d2 molecule at 337 nm, there is a fluorescence resonance energy transfer (FRET) to the Eu cryptate molecules of the anti-cAMP antibody (labelled therewith), followed by a long-lasting emission signal at 665 nm (and at 620 nm). Both signals are measured in a suitable measurement instrument with a time delay, i.e. after decay of the background fluorescence. Any increase in the low FRET signal caused by prostaglandin E₂ administration (measured as well ratio change = $\text{emission}_{665\text{nm}}/\text{emission}_{620\text{nm}} * 10000$) indicates the action of antagonists.

2.2. Detection method

2.2.1. Test for antagonism (data per well of a 384-well plate):

To a test plate with the substance solutions already added (0.05 µl; 100% DMSO, concentration range from 0.8 nM - 16.5 µM) were added 4 µl of a cAMP-d2/cell suspension (312500 cells/ml). After a 20-minute preincubation at room temperature (RT), 2 µl of a 3xPGE₂ solution (0.3 nM, in PBS-IBMX) were added and incubated in the presence of the agonist for a further 60 mins at RT (volume: ~ 6 µl). Next the reaction was stopped by addition of 2 µl of lysis buffer and incubated for a further 20 mins at RT before the actual measurement (volume: ~ 8 µl).

3. Detection of antagonism to the human prostaglandin D receptor signal

3.1 Detection principle

The binding of prostaglandin D₂ to the human PGD receptor induces the activation of membrane-located adenylate cyclases and leads to the formation of cAMP. In the presence of the phosphodiesterase inhibitor IBMX, the cAMP accumulated owing to this stimulation and released by cell lysis is used in a competitive detection method. In this test, the cAMP present in the lysate competes with a fluorescence-labelled cAMP (cAMP-d2) for binding to an Eu cryptate-labelled anti-cAMP antibody.

In the absence of cellular cAMP, a maximal signal is produced, which is attributable to the binding of this cAMP-d2 molecule to the antibody. After excitation of the cAMP-d2 molecule at 337 nm, there is a fluorescence resonance energy transfer (FRET) to the Eu cryptate molecules of the anti-cAMP antibody (labelled therewith), followed by a long-lasting emission signal at 665 nm (and at 620 nm). Both signals are measured in a suitable measurement instrument with a time delay, i.e. after decay of the background fluorescence. Any increase in the low FRET signal caused by prostaglandin E₂ administration (measured as well ratio change = $\text{emission}_{665\text{nm}}/\text{emission}_{620\text{nm}} * 10000$) indicates the action of antagonists.

3.2. Detection method

3.2.1. Test for antagonism (data per well of a 384-well plate):

To a test plate with the substance solutions already added (0.05 µl; 100% DMSO, concentration range from 0.8 nM – 16.5 µM) were added 4 µl of a cAMP-d2/cell suspension (625000 cells/ml). After a 20-minute preincubation at room temperature (RT), 2 µl of a 3xPGD2 solution (6 nM, in PBS-IBMX) were added and incubated in the presence of the agonist for a further 30 mins at RT (volume: ~ 6 µl). Next the reaction was stopped by addition of 2 µl of lysis buffer and incubated for a further 20 mins at RT before the actual measurement (volume: ~ 8 µl).

4. The EP2 subtype of the PGE₂ receptor and the pre-ovulatory cumulus expansion

4.1. Background:

In the pre-ovulatory antral follicle, the oocyte is surrounded by cumulus cells, which form a dense cell crown around the oocyte. After the LH peak (luteinizing hormone), a series of processes is activated, which results in a marked morphological change in this cell crown of cumulus cells. During this, the cumulus cells form an extracellular matrix, which leads to the so-called cumulus expansion (Vanderhyden *et al.* Dev Biol.; (1990); 140(2): 307-317) This cumulus expansion is an important component of the ovulatory process and the subsequent possibility of fertilization.

During the cumulus expansion, prostaglandins and here prostaglandin E₂, synthesis whereof is induced by the LH peak, are of decisive importance. Prostanoid EP2 knockout mice (Hizaki *et al.*, 1999, Proc Natl Acad Sci U. S. A.; (1999) 96(18):10501-10506) show a markedly decreased cumulus expansion and severe subfertility, which demonstrates the importance of the prostanoid EP2 receptor for this process.

4.2. Cumulus expansions test *in vitro*

In immature female mice (strain: B6D2F1 from Charles River), at an age of 14 - 18 days, the folliculogenesis was induced by a single administration (intraperitoneal) of 10 IU of PMSG (pregnant mare serum gonadotropin; Sigma G-4877, Lot 68H0909). 47-50 hours after the injection, the ovaries were removed and the cumulus-oocyte complexes removed. At this stage, the cumulus complex is not yet expanded.

The cumulus-oocyte complexes were now incubated for 20-24 hours with prostaglandin E₂ (PGE₂) (0.3 µM), vehicle control (ethanol) or test substances. Medium: alpha-MEM medium with 0.1 mM IBMX, pyruvate (0.23 mM), glutamine (2 mM), pen/strep 100 IU/ml pen. and 100 µg/ml strep.), HSA (8 mg/ml) and foetal bovine serum (FBS, 10%). The cumulus expansion was then established through the subdivision into four stages (after Vanderhyden *et al.* Dev Biol.; (1990); 140(2):307-317).

4.3 *In-vivo* action on post-ovulatory *in-vitro* fertilization :

Substances can exert an influence on fertility by decreasing the fertilizability of oocytes or cumulus-oocyte complexes. In order to study such effects, substances can be administered *in vivo* and subjected to *in vitro* fertilization after ovulation of cumulus-oocyte complexes has taken

place. The *in-vitro* fertilization rate, where no test substance is any longer present, allows conclusions as to the *in-vivo* effects of the test substances.

Immature female mice (strain: B6D2F1, Charles River, Suelzfeld, age: 19-25 days) were kept in Macrolon cages in rooms with controlled illumination (12 hrs darkness: 12 hrs light), fed with a standard diet and provided with drinking water ad libitum.

The mice were primed with PMSG (pregnant mare serum gonadotropin) (10 IU/animal i.p.). After 48 hours, a stimulus triggering ovulation was created in the animals by one administration of 10 IU/animal i.p. (hCG, human chorionic gonadotropin). The test substances were dissolved in benzyl benzoate/castor oil (1+4 v/v) and administered in a volume of 0.1 ml s.c. 1 hour before hCG (n=5 animals per group). Fourteen hours after hCG administration, the animals were killed. Ovulated oocytes and cumulus-oocyte complexes were obtained from the bursa ovarii and/or oviduct and subjected to *in-vitro* fertilization, wherein for the fertilization a sperm count of 40000 sperms/0.5 ml was used for 1 hour. Twenty-four hours after the incubation with the sperms, the number of fertilized oocytes is established and the percentage fertilization rate determined.

The results in Table 4 show that Example 17 according to the invention has a dose-dependent influence on the fertilizability of ovulated cumulus-oocyte complexes.

4.4 *In-vivo* action on fertility in non-human primates (Cynomolgus):

In order to study the effect of substances on fertility, mating studies can be performed in monkeys (Jensen *et al.* Contraception 81; (2010) 82(2): 165–171). For this, the test substances are administered to female cynomolgus monkeys (*Macaca Fascicularis*) which are being kept in groups, then the animals are mated with a male animal. Matings are checked for by sperm detection in daily vaginal smears. Pregnancies resulting therefrom are identified by hormone determinations and ultrasound examinations. Through the changes in the serum oestradiol concentrations during the cycle, (rise before the midcycle LH peak), the fertile phase within a cycle of the individual animals can be determined. Matings in this fertile period are described as “timed matings” (matings in the fertile phase). Apart from the absolute number of pregnancies occurring, the effect on the fertility can also be expressed as pregnancies per “timed matings”.

To test the action of EP2 receptor antagonists on fertility in monkeys, Examples 17 and 56 according to the invention were administered over 6 months, dissolved in 0.5 ml castor oil. Example 17 was administered once daily at a dosage of 10 mg/kg (n= 10 animals), while Example 56 was administered twice daily at a dosage of 10 mg/kg (n=9 animals). Only the vehicle was administered to a control group (n=10 animals). In the first month of the treatment,

no male animal was placed with the female animals. After this, female and male animals were kept together over 5 months, with detection of pregnancies and cycle monitoring.

Table 5 shows that both substances result in a marked reduction in the number of pregnancies occurring. These data for the first time show the strong contraceptive effect of EP2 receptor antagonists in the primate. Further results of the study show that the substances have no effects on hormone levels and cycle length. This confirms that the substances produce a contraceptive effect by non-hormonal mechanisms. The reversibility of the contraceptive effect was demonstrated on some animals after discontinuation of the treatment.

5. Determination of the pharmacokinetic parameters after intravenous administration

5.1. Intravenous administration:

For this, the substances were administered in dissolved form, compatible solubilizers such as PEG400 and/or ethanol being used in tolerable quantity. The substances were administered at a dosage of 0.1 -1 mg/kg. The administration was effected in the female rat as a bolus injection. Here, at various times after bolus injection ca. 100-150 μ l blood samples were withdrawn from the jugular vein via a catheter. The blood samples were treated with lithium-heparin as anticoagulant and stored refrigerated until further work-up. After centrifugation of the samples for 15 mins at 3000 rpm, an aliquot of 100 μ l was withdrawn from the supernatant (plasma) and precipitated by addition of 400 μ l of cold acetonitrile or methanol (absolute). The precipitated samples were frozen overnight at -20°C, then centrifuged once again for 15 mins at 3000 rpm, before 150 μ l of the clear supernatant was withdrawn for the concentration determination. The analysis was performed with an Agilent 1200 HPLC system with attached LCMS/MS detection.

5.2 Calculation of the PK parameters

The calculation was performed by means of the PK calculation software WinNonLin[®], where $t_{1/2}$ means half-life within a specified interval (here: terminal $t_{1/2}$, in hrs).

Table 1: Examples for the biological activity of the compounds according to the invention on the hEP2 receptor (IC_{50} measured by cAMP antagonism test), selectivity towards hDP & hEP₄ (IC_{50} measured by cAMP antagonism test): x: 1 - 10, xx: 10 - 100, xxx > 100 :

Example	Antagonism hEP2 IC_{50} [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
1	1.03E-07	xxx	xxx
2	2.17E-08	xxx	xxx
3	4.91E-09	xxx	xxx
4	6.81E-08	xxx	xxx
5	5.11E-09	xxx	xxx
6	3.13E-07	xx	xx
7	2.58E-08	xxx	xxx
8	9.37E-09	xxx	xxx
9	1.72E-08	xxx	xxx
10	8.67E-09	xxx	xxx
11	8.14E-09	xxx	xxx
12	2.03E-08	xxx	xxx
13	3.59E-08	xxx	xxx
14	3.67E-09	xxx	xxx
15	4.17E-09	xxx	xxx
16	5.85E-09	xxx	xxx
17	3.44E-09	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
18	7.66E-09	xxx	xxx
19	8.57E-09	xxx	xxx
20	7.33E-09	xxx	xxx
21	3.58E-08	xxx	xxx
22	3.03E-08	xxx	xxx
23	1.61E-08	xxx	xxx
24	6.01E-08	xxx	xxx
25	2.52E-08	xxx	xxx
26	2.42E-08	xxx	xxx
27	1.93E-08	xxx	xxx
28	3.39E-08	xxx	xxx
29	9.31E-09	xxx	xxx
30	6.43E-08	xxx	xxx
31	3.08E-09	xxx	xxx
32	1.04E-08	xxx	xxx
33	3.42E-08	xxx	xxx
34	1.34E-07	xxx	xxx
35	6.92E-08	xxx	xxx
36	6.16E-08	xxx	xxx
37	1.04E-07	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
38	2.81E-08	xxx	xxx
39	1.13E-07	xxx	xxx
40	1.15E-08	xxx	xxx
41	5.62E-09	xxx	xxx
42	4.75E-09	xxx	xxx
43	4.82E-09	xxx	xxx
44	3.85E-09	xxx	xxx
45	5.51E-09	xxx	xxx
46	4.91E-09	xxx	xxx
47	5.61E-09	xxx	xxx
48	5.59E-09	xxx	xxx
49	6.44E-09	xxx	xxx
50	3.18E-09	xxx	xxx
51	6E-08	xxx	xxx
52	8.3E-07	xx	xx
53	4.42E-08	xxx	xxx
54	4.74E-09	xxx	xxx
55	9.67E-09	xxx	xxx
56	1.31E-08	xxx	xxx
57	5.29E-09	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
58	1.08E-06	xx	xx
59	7.58E-07	xx	xx
60	1.75E-07	xx	xx
61	1.5E-06	xx	xx
62	6.74E-07	xx	xx
63	1.44E-06	xx	xx
64	4.13E-06	x	x
65	2.79E-06	x	x
66	3.21E-07	xx	xx
67	4.05E-07	xx	xx
68	1.77E-08	xxx	xxx
69	2.9E-08	xxx	
70	4.9E-08	xxx	
71	8.29E-08	xxx	xxx
72	1.2E-07	xxx	xxx
73	7.78E-08	xxx	xxx
74	9.54E-08	xxx	xxx
75	6.7E-08	xxx	xxx
76	1.79E-07	xx	xx
77	5.39E-08	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
78	3.17E-08	xxx	xxx
79	1.25E-07	xxx	xxx
80	8.59E-08	xxx	xxx
81	5.13E-07	xx	xx
82	2.18E-07	xx	xx
83	3.26E-08	xxx	xxx
84	1.92E-08	xxx	xxx
85	3.55E-08	xxx	xxx
86	1.54E-08	xxx	xxx
87	2.4E-08	xxx	xxx
88	1.62E-08	xxx	xxx
89	9.4E-09	xxx	xxx
90	1.48E-08	xxx	xxx
91	2.03E-08	xxx	xxx
92	1.27E-08	xxx	xxx
93	1.15E-08	xxx	xxx
94	1.28E-08	xxx	xxx
95	6.18E-09	xxx	xxx
96	9.42E-09	xxx	xxx
97	1.27E-08	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
98	7.07E-09	xxx	xxx
99	1.04E-08	xxx	xxx
100	6.21E-09	xxx	xxx
101	6.58E-09	xxx	xxx
102	4.1E-07	xx	xx
103	5.95E-07	xx	xx
104	5.13E-07	xx	xx
105	2.85E-07	xx	xx
106	1.5E-07	xxx	xxx
107	1.14E-08	xxx	xxx
108	6.46E-09	xxx	xxx
109	9.53E-09	xxx	xxx
110	4.4E-09	xxx	xxx
111	1.37E-08	xxx	xxx
112	1.19E-08	xxx	xxx
113	1.39E-08	xxx	xxx
114	6.08E-09	xxx	xxx
115	9.05E-09	xxx	xxx
116	5.52E-09	xxx	xxx
117	1.78E-08	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
118	8.17E-09	xxx	
119	1.32E-08	xxx	
120	1.55E-08	xxx	
121	1.55E-08	xxx	xxx
122	2.86E-08	xxx	xxx
123	8.92E-09	xxx	xxx
124	7.99E-09	xxx	xxx
125	3.66E-09	xxx	xxx
126	2.12E-08	xxx	xxx
127	5.56E-08	xxx	xxx
128	4.71E-07	xx	xx
129	3.09E-07	xx	xx
130	2.41E-07	xx	xx
131	2.28E-07	xx	xx
132	1.09E-07	xxx	xxx
133	4.19E-08	xxx	xxx
134	5.65E-08	xxx	xxx
135	1.95E-08	xxx	xxx
136	6.59E-09	xxx	xxx
137	7.65E-09	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
138	1.6E-07	xxx	xxx
139	4.19E-07	xx	xx
140	5.53E-08	xxx	xxx
141	1.43E-07	xxx	xxx
142	7.13E-07	xx	xx
143	3.68E-08	xxx	xxx
144	5.08E-08	xxx	xxx
145	1.88E-08	xxx	xxx
146	5.26E-08	xxx	xxx
147	5.74E-08	xxx	xxx
148	2.74E-08	xxx	xxx
149	2.96E-08	xxx	xxx
150	1.28E-08	xxx	xxx
151	4.85E-08	xxx	xxx
152	3.33E-08	xxx	xxx
153	6.08E-08	xxx	xxx
154	8.08E-08	xxx	xxx
155	2.94E-08	xxx	xxx
156	1.84E-07	xx	xx
157	1.36E-08	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
158	6.21E-09	xxx	xxx
159	6.24E-09	xxx	xxx
160	8.29E-08	xxx	xxx
161	9.82E-08	xxx	xxx
162	9.33E-09	xxx	xxx
163	6.8E-09	xxx	xxx
164	1.09E-08	xxx	xxx
165	3.34E-08	xxx	xxx
166	2.87E-09	xxx	xxx
167	1.35E-08	xxx	xxx
168	2.93E-08	xxx	xxx
169	2.77E-08	xxx	xxx
170	4.25E-09	xxx	xxx
171	3.88E-09	xxx	xxx
172	3.16E-09	xxx	xxx
173	2.71E-09	xxx	xxx
174	6.58E-09	xxx	xxx
175	8.05E-09	xxx	xxx
176	7.77E-09	xxx	xxx
177	1.18E-07	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
178	6.83E-08	xxx	xxx
179	2.61E-06	x	x
180	1.73E-08	xxx	xxx
181	3.52E-08	xxx	xxx
182	6.42E-08	xxx	xxx
183	4.66E-08	xxx	xxx
184	7.68E-08	xxx	xxx
185	2.48E-08	xxx	xxx
186	5.19E-08	xxx	xxx
187	2.35E-08	xxx	xxx
188	1.49E-07	xxx	xxx
189	3.36E-08	xxx	xxx
190	2.72E-07	xx	xx
191	4.71E-08	xxx	xxx
192	2.06E-07	xx	xx
193	1.45E-07	xxx	xxx
194	4.09E-08	xxx	xxx
195	2.27E-07	xx	xx
196	5.08E-08	xxx	xxx
197	6.32E-08	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
198	1.53E-07	xxx	xxx
199	9.71E-08	xxx	xxx
200	1.51E-08	xxx	xxx
201	2.49E-08	xxx	xxx
202	5.06E-09	xxx	xxx
203	8.57E-08	xxx	xxx
204	1E-08	xxx	xxx
205	4.33E-08	xxx	xxx
206	4.13E-08	xxx	xxx
207	3.03E-08	xxx	xxx
208	8.06E-08	xxx	xxx
209	2.51E-08	xxx	xxx
210	1.95E-08	xxx	xxx
211	5.01E-08	xxx	xxx
212	1.06E-07	xxx	xxx
213	8.24E-08	xxx	xxx
214	6.49E-08	xxx	xxx
215	3.9E-08	xxx	xxx
216	8.61E-08	xxx	xxx
217	1.14E-07	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
218	1.28E-07	xxx	xxx
219	2.73E-07	xx	xx
220	1.47E-07	xxx	xxx
221	2.35E-07	xx	xx
222	4.09E-08	xxx	xxx
223	2E-08	xxx	xxx
224	5.62E-09	xxx	xxx
225	1.43E-08	xxx	xxx
226	1.19E-08	xxx	xxx
227	9.89E-09	xxx	xxx
228	8.09E-09	xxx	xxx
229	1.02E-08	xxx	xxx
230	1.37E-08	xxx	xxx
231	1.32E-08	xxx	xxx
232	3.83E-08	xxx	xxx
233	3.97E-08	xxx	xxx
234	4.77E-08	xxx	xxx
235	1.67E-08	xxx	xxx
236	5.75E-08	xxx	xxx
237	3.1E-08	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
238	3.24E-08	xxx	xxx
239	2.7E-08	xxx	xxx
240	2.51E-08	xxx	xxx
241	9.03E-09	xxx	xxx
242	3.05E-07	xx	xx
243	3.35E-07	xx	xx
244	4.42E-08	xxx	xxx
245	7.79E-09	xxx	xxx
246	9.23E-09	xxx	xxx
247	1.26E-08	xxx	xxx
248	1.47E-08	xxx	xxx
249	1.12E-08	xxx	xxx
250	3.06E-08	xxx	xxx
251	3.81E-06	x	x
252	2.95E-07	xx	xx
253	1.68E-07	xx	xx
254	5.52E-08	xxx	xxx
255	8.35E-08	xxx	xxx
256	1.07E-07		xxx
257	1.02E-07	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
258	1.09E-07	xxx	xxx
259	2.56E-08	xxx	xxx
260	8.02E-09	xxx	xxx
261	8.1E-08	xxx	xxx
262	3.34E-08	xxx	xxx
263	1.81E-07	xx	xx
264	1.37E-08	xxx	xxx
265	3.15E-08	xxx	xxx
266	1.89E-08	xxx	xxx
267	4.55E-08	xxx	xxx
268	2.29E-08	xxx	xxx
269	7.54E-09	xxx	xxx
270	5.31E-08	xxx	xxx
271	3.35E-08	xxx	xxx
272	2.37E-08	xxx	xxx
273	1.85E-08	xxx	xxx
274	4.46E-08	xxx	xxx
275	3.45E-08	xxx	xxx
276	2.26E-07	xx	xx
277	4.79E-08	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
278	1.04E-07	xxx	xxx
279	5.74E-08	xxx	xxx
280	3.07E-08	xxx	xxx
281	2.3E-08	xxx	xxx
282	2.01E-08	xxx	xxx
283	6.55E-08	xxx	xxx
284	5.64E-08	xxx	xxx
285	1E-07	xxx	xxx
286	1E-07	xxx	xxx
287	1.05E-06	xx	xx
288	4E-08	xxx	xxx
289	5.16E-08	xxx	xxx
290	4.26E-08	xxx	xxx
291	5.94E-09	xxx	xxx
292	2E-08	xxx	xxx
293	2.83E-08	xxx	xxx
294	1.32E-08	xxx	xxx
295	7.08E-08	xxx	xxx
296	2.97E-08	xxx	xxx
297	4.6E-07	xx	xx

- 310 -

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
298	3.45E-07	xx	xx
299	5.41E-07	xx	xx
300	7.98E-08	xxx	xxx
301	2.45E-08	xxx	xxx
302	1.31E-07	xxx	xxx
303	1.39E-08	xxx	xxx
304	9.98E-08	xxx	xxx
305	3.01E-07	xx	xx
306	7.66E-08	xxx	xxx
307	3.06E-08	xxx	xxx
308	2.22E-08	xxx	xxx
309	6.6E-08	xxx	xxx
310	8.06E-07	xx	xx
311	8.41E-08	xxx	xxx
312	1.58E-07	xxx	xxx
313	8.94E-08	xxx	xxx
314	2.78E-08	xxx	xxx
315	4.3E-08	xxx	xxx
316	7.9E-08	xxx	xxx
317	1.81E-07	xx	xx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
318	2.63E-08	xxx	xxx
319	3.15E-07	xx	xx
320	1.56E-08	xxx	xxx
321	1.01E-07	xxx	xxx
322	7.62E-08	xxx	xxx
323	1.28E-07	xxx	xxx
324	4.6E-07	xx	xx
325	3.21E-07	xx	xx
326	4.85E-07	xx	xx
327	1.21E-06	xx	xx
328	2.56E-07	xx	xx
329	2.32E-07	xx	xx
330	9.53E-09	xxx	xxx
331	9.47E-09	xxx	xxx
332	5.47E-09	xxx	xxx
333	6.42E-09	xxx	xxx
334	7.42E-08	xxx	xxx
335	5.09E-09	xxx	xxx
336	5.59E-09	xxx	xxx
337	1.81E-08	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
338	2.8E-08	xxx	xxx
339	7.15E-09	xxx	xxx
340	1.99E-08	xxx	xxx
341	5.53E-08	xxx	xxx
342	1.03E-07	xxx	xxx
343	4.24E-08	xxx	xxx
344	1.03E-08	xxx	xxx
345	6.26E-09	xxx	xxx
346	1.49E-08	xxx	xxx
347	7.88E-09	xxx	xxx
348	8.88E-09	xxx	xxx
349	4.82E-09	xxx	xxx
350	8.41E-09	xxx	xxx
351	6.84E-09	xxx	xxx
352	2.17E-08	xxx	xxx
353	5.02E-08	xxx	xxx
354	5.87E-09	xxx	xxx
355	5.39E-09	xxx	xxx
356	6.36E-08	xxx	xxx
357	7.36E-09	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
358	8.01E-09	xxx	xxx
359	6.99E-09	xxx	xxx
360	6.27E-09	xxx	xxx
361	5.11E-09	xxx	xxx
362	1.51E-08	xxx	xxx
363	5.48E-09	xxx	xxx
364	6.61E-09	xxx	xxx
365	5.3E-09	xxx	xxx
366	1.89E-08	xxx	xxx
367	5.84E-09	xxx	xxx
368	1.2E-08	xxx	xxx
369	5.66E-09	xxx	xxx
370	3.07E-09	xxx	xxx
371	6.11E-09	xxx	xxx
372	7.29E-09	xxx	xxx
373	4.07E-08	xxx	xxx
374	6.58E-09	xxx	xxx
375	5.8E-09	xxx	xxx
376	3.59E-09	xxx	xxx
377	2.73E-08	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
378	7.77E-09	xxx	xxx
379	6.26E-08	xxx	xxx
380	5.35E-09	xxx	xxx
381	5.84E-09	xxx	xxx
382	1.94E-08	xxx	xxx
383	5.78E-09	xxx	xxx
384	3.95E-09	xxx	xxx
385	1.31E-08	xxx	xxx
386	2.57E-08	xxx	xxx
387	1.32E-08	xxx	xxx
388	2.05E-08	xxx	xxx
389	4.51E-09	xxx	xxx
390	3.14E-09	xxx	xxx
391	6.07E-09	xxx	xxx
392	2.18E-08	xxx	xxx
393	5E-08	xxx	xxx
394	5.7E-09	xxx	xxx
395	3.04E-08	xxx	xxx
396	1.8E-08	xxx	xxx
397	1.25E-08	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
398	1.82E-08	xxx	xxx
399	5.93E-07	xx	xx
400	5.61E-08	xxx	xxx
401	1.55E-07	xxx	xxx
402	7.77E-08	xxx	xxx
403	9.59E-09	xxx	xxx
404	2.02E-08	xxx	xxx
405	2.14E-06	x	x
406	8.19E-09	xxx	xxx
407	1.16E-08	xxx	xxx
408	2.35E-08	xxx	xxx
409	3.18E-08	xxx	xxx
410	9.04E-08	xxx	xxx
411	1.65E-08	xxx	xxx
412	6.35E-08	xxx	xxx
413	3.78E-08	xxx	xxx
414	2.23E-07	xx	xx
415	2.13E-08	xxx	xxx
416	1.07E-08	xxx	xxx
417	9.75E-09	xxx	xxx

Example	Antagonism hEP2 IC ₅₀ [M]	Selectivity hEP2/hDP	Selectivity hEP2/hEP4
418	1.58E-08	xxx	xxx
419	1.09E-08	xxx	xxx

Table 2: Reduction in the cumulus expansion in percent with use of 0.3 μ M PGE2 for stimulation, reference 1: example 62 from DE102009049662A1

Example	% reduction at 1 μ M	% reduction at 0.5 μ m
Reference 1*	79% (*reduction at 10 μ M)	
Example 5	59	56
Example 17	76	71
Example 42	79	48
Example 117	54	30
Example 283	26	25
Example 291	73	76
Example 303	47	38
Example 345	36	18

- 5 Table 3: Terminal half-life ($t_{1/2}$) after intravenous administration in the female rat, reference 1: example 62 from DE102009049662A1

Example	$t_{1/2}$ [hr]
Reference 1	0.4
Example 17	2.5

Example 42	3.2
Example 117	2.0
Example 283	5.8

Table 4: Reduction in *in-vitro* fertilization rate after *in-vivo* administration of the test substance:

Example	% fertilization with vehicle	% fertilization with 1 mg/kg	% fertilization with 5 mg/kg	% fertilization with 10 mg/kg
Example 17	30 ± 9	20 ± 10	9 ± 1	8 ± 4

5 Table 5: Reduction in the pregnancy rates in the non-human primate (*Cynomolgus*)

Group	Animals per group	Number of “timed matings”*	Pregnancies	% Pregnancies/ “timed mating”
Control	10	21	8	38
Example 17	10	25	3	6
Example 56	9	34	2	12

*Detection of vaginal sperms in the fertile phase of the cycle

6. Effect of a single administration of EP2 receptor antagonists and COX inhibitors alone or in combination on the fertility of the rat

The rat is a particularly suitable animal model for demonstrating the effect of substances on fertility, since the cycle can be observed easily by vaginal swabs and mating reliably leads to a high number of offspring.

In the experiments below, female Han Wistar rats having a weight of 200-270 g were used. The animals were kept in Makrolon cages in rooms with controlled lighting (12 hours of darkness; 12 hours of light), fed with a standard diet and provided with water ad libitum.

5 EP2 receptor antagonists were dissolved in benzyl benzoate/castor oil (1+4 v/v), and the dose was administered in a volume of 1 ml/kg of body weight s.c. Optionally, EP2 receptor antagonists were also administered orally in the same vehicle as the COX inhibitors (see below). Here, the oral administrations were divided to three points in time (12.30, 16.00 and 17.00 hours) (see Table 6.3).

10 The COX inhibitors were suspended in a carrier liquid (85 mg of Myrj®53 polyoxyl(50)stearate; CAS No. 9004-99-3) in 100 ml of 0.9% w/v NaCl solution), and the dose corresponding to the treatment group was administered in a volume of 2 ml/kg of body weight.

Prior to the start of the experiment, two cycles were observed using vaginal swabs. Only animals having a regular 4-day cycle were used in the experiment. The animals were randomized to the treatment groups.

15 The animals were treated once in the proestrus, the day in the cycle when the LH peak occurs in the evening, with the test substances either alone or in combination. On the evening of the proestrus, the female animals mated with a male rat. Successive mating was confirmed by detection of vaginal sperms in the vaginal swab the next morning. The effect on fertility was determined 9-15 days after mating by the number of implantation sites of the animals. The group
20 size was n = 6 per group.

As can be seen from Tables 6.1 – 6.6, in these experiments piroxicam, indomethacin and meloxicam in combination with various EP2 receptor antagonists had a strong antifertility effect. In particular, it should be noted that the individual administrations showed no or only a weak effect, whereas the combination showed a markedly increased contraceptive efficacy.

25 **Table 6.1:** Effect of a single administration of the EP2 receptor antagonist E16: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide (Example 17) or piroxicam alone or in combination on the fertility of the female rat.

Treatment group	Time of treatment (o'clock)	Absolute implantations number of 6 animals	Implantations per animal (mean $\pm$ SD)
Vehicle	7:00	74	12.3 $\pm$ 0.8
E16 4 mg/animal	7:00	54	9.0 $\pm$ 3.4
Piroxicam 2 mg/animal	7:00	81	13.5 $\pm$ 1.0
E16 4 mg/animal + Piroxicam 2 mg/animal	7:00	22	3.7 $\pm$ 4.3
E16 4 mg/animal	13:00	49*	8.2 $\pm$ 3.1
Piroxicam 2 mg/animal	13:00	68	11.3 $\pm$ 1.6
E16 4 mg/animal + Piroxicam 2 mg/animal	13:00	8*	1.4 $\pm$ 1.1

* Data of n = 5 animals since one animal did not mate (no sperms detected in the vaginal swab)

Table 6.2: Effect of a single administration of the EP2 receptor antagonist E161: N-(2-methoxyethyl)-4-methyl-2-[[1-(4-[[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoyl)piperidin-4-yl]methyl]-2H-indazole-5-carboxamide (Example 345) or piroxicam alone or in combination on the fertility of the female rat.

Treatment group	Time of treatment (o'clock)	Absolute implantations number of 6 animals	Implantations per animal (mean $\pm$ SD)
Vehicle	18:00	77	12.8 $\pm$ 2.2
E161 2 mg/animal	18:00	55	9.0 $\pm$ 3.4
Piroxicam 2 mg/animal	18:00	53*	10.6 $\pm$ 1.5
E161 2 mg/animal + Piroxicam 2 mg/animal	18:00	24	4.0 $\pm$ 2.1

5 * Data of n = 5 animals since one animal did not mate (no sperms detected in the vaginal swab)

Table 6.3: Effect of an oral administration of EP2 receptor antagonist E16: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide (Example 17) or E161: N-(2-methoxyethyl)-4-methyl-2-[[1-(4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoyl)piperidin-4-yl]methyl]-2H-indazole-5-carboxamide (Example 345) or piroxicam alone or in combination on the fertility of the female rat.

Treatment group	Time of treatment (o'clock)#	Absolute implantations number of 6 animals	Implantations per animal (mean $\pm$ SD)
Vehicle	12:30; 16:00 and 17:00	73	12.3 $\pm$ 1.2
Piroxicam 2 mg/animal	12:30	71	11.8 $\pm$ 3.7
E16 2 mg/animal	16:00 and 17:00	63	10.5 $\pm$ 1.5
E161 2 mg/animal	16:00 and 17:00	69	11.5 $\pm$ 3.1
E16 2 mg/animal + Piroxicam 2 mg/animal	12:30; 16:00 and 17:00	21	3.5 $\pm$ 2.9
E161 2 mg/animal + Piroxicam 2 mg/animal	12:30; 16:00 and 17:00	32	5.3 $\pm$ 4.1

#Dosage of the EP2 receptor antagonists was divided to two points in time after administration of the COX inhibitor

Table 6.4: Effect of a single administration of the EP2 receptor antagonist E16: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide (Example 17) or meloxicam in various dosages alone or in combination on the fertility of the female rat.

Treatment group	Time of treatment (o'clock)#	Absolute implantations number of 6 animals	Implantations per animal (mean $\pm$ SD)
Vehicle	17:00	31***	10.3 $\pm$ 0.6
Meloxicam 0.25 mg/animal	17:00	32*	6.4 $\pm$ 5.9
Meloxicam 0.5 mg/animal	17:00	70	11.7 $\pm$ 6.4
Meloxicam 1 mg/animal	17:00	66	11 $\pm$ 5.5
Meloxicam 2 mg/animal	17:00	54*	10.8 $\pm$ 4.4
E16 4 mg/animal	17:00	22**	5.5 $\pm$ 5.4
E16 4 mg/animal + Meloxicam 0.25 mg/animal	17:00	31*	6.2 $\pm$ 6.3
E16 4 mg/animal + Meloxicam 0.5 mg/animal	17:00	15***	5.0 $\pm$ 1.7
E16 4 mg/animal + Meloxicam 1 mg/animal	17:00	11*	2.2 $\pm$ 3.3
E16 4 mg/animal + Meloxicam 2 mg/animal	17:00	7**	1.8 $\pm$ 2.4

5 * Data of n = 5 animals since one animal did not mate (no sperms detected in the vaginal swab)

** Data of n = 4 animals since two animals did not mate (no sperms detected in the vaginal swab)

*** Data of n = 3 animals since three animals did not mate (no sperms detected in the vaginal swab)

Table 6.5: Effect of a single dose-dependent administration of the EP2 receptor antagonist E16: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide (Example 17) or meloxicam alone or in combination on the fertility of the female rat.

Treatment group	Time of treatment (o'clock)#	Absolute implantations number of 6 animals	Implantations per animal (mean $\pm$ SD)
Vehicle	17:00	61*	12.2 $\pm$ 1.9
E16 0.25 mg/animal	17:00	78	13 $\pm$ 2.3
E16 0.5 mg/animal	17:00	60	10 $\pm$ 2.8
E16 1 mg/animal	17:00	42	7 $\pm$ 5.4
E16 2 mg/animal	17:00	30**	7.5 $\pm$ 1.7
E16 4 mg/animal	17:00	32	5.3 $\pm$ 4.5
Meloxicam 2 mg/animal	17:00	63*	12.6 $\pm$ 3.4
E16 0.25 mg/animal + Meloxicam 2 mg/animal	17:00	52	8.7 $\pm$ 2.5
E16 0.5 mg/animal + Meloxicam 2 mg/animal	17:00	36	6.0 $\pm$ 4.3
E16 1 mg/animal + Meloxicam 2 mg/animal	17:00	35	5.8 $\pm$ 3.4
E16 2 mg/animal + Meloxicam 2 mg/animal	17:00	19	3.2 $\pm$ 2.8
E16 4 mg/animal + Meloxicam 2 mg/animal	17:00	11***	3.7 $\pm$ 4.7

5 * Data of n = 5 animals since one animal did not mate (no sperms detected in the vaginal swab)

** Data of n = 4 animals since two animals did not mate (no sperms detected in the vaginal swab)

*** Data of n = 3 animals since three animals did not mate (no sperms detected in the vaginal swab)

Table 6.6: Effect of a single dose-dependent administration of the EP2 receptor antagonist E16: N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide (Example 17) or indomethacin alone or in combination on the fertility of the female rat.

Treatment group	Time of treatment (o'clock)#	Absolute implantations number of 6 animals	Implantations per animal (mean $\pm$ SD)
Vehicle	16:00	67	11.2 $\pm$ 2.6
E16 1 mg/animal	16:00	62	10.3 $\pm$ 2.5
E16 2 mg/animal	16:00	71	11.8 $\pm$ 2.1
E16 4 mg/animal	16:00	50	8.3 $\pm$ 3.4
E16 8 mg/animal	16:00	43*	8.6 $\pm$ 2.8
Indomethacin 1 mg/animal	16:00	67	11.2 $\pm$ 2.9
E16 1 mg/animal + Indomethacin 1 mg/animal	16:00	58	9.7 $\pm$ 1.9
E16 2 mg/animal + Indomethacin 1 mg/animal	16:00	38	6.3 $\pm$ 3.7
E16 4 mg/animal + Indomethacin 1 mg/animal	16:00	27	4.5 $\pm$ 1.6
E16 8 mg/animal + Indomethacin 1 mg/animal	16:00	26	4.3 $\pm$ 1.9

5 * Data of n = 5 animals since one animal did not mate (no sperms detected in the vaginal swab)

7. *In-vivo* effect on postovulatory *in-vitro* fertilization in the mouse

Substances can influence fertility by lowering fertilizability of oocytes or cumulus-oocyte complexes. To examine such effects, substances can be administered *in vivo* and, after ovulation of cumulus-oocyte complexes, the latter are subjected to *in-vitro* fertilization. The *in-vitro* fertilization rate in the absence of test substance allows conclusions to be drawn on the *in-vivo* effects of the test substances.

Immature female mice (strain: B6D2F1, Charles River, Suelzfeld, age: 19-25 days) were kept in Makrolon cages in rooms with controlled lighting (12 hours of darkness: 12 hours of light), fed with a standard diet and provided with water ad libitum.

The mice were primed with PMSG (Pregnant Mare Serum Gonadotropin) (10 IU/animal i.p.). After 48 hours, by administration of 10 IU/animal i.p., an ovulation-triggering stimulus (hCG, human Chorion Gonadotropin) was generated in the animals. EP2 receptor antagonists were dissolved in benzyl benzoate/castor oil (1+4 v/v) and administered in a volume of 0.1 ml s.c. 5 1 hour prior to hCG (n = 5 animals per group). The COX inhibitor was suspended in a carrier liquid (85 mg of Myrj®53 polyoxyl(50)stearate; CAS No. 9004-99-3) in 100 ml of 0.9% w/v NaCl solution), and the appropriate dose was administered in a volume of 2 ml/kg of body weight orally 1 hour prior to hCG. Fourteen hours after administration of hCG, the animals were sacrificed. Ovulated oocytes and cumulus-oocyte complexes were obtained from the ovarii bursa 10 and/or the oviduct and subjected to *in-vitro* fertilization, with a sperm number of 40 000 sperms/0.5 ml for 1 hour being used for fertilization. Twenty-four hours after incubation with the sperms, the number of fertilized oocytes and the number of 2-cell embryos per animal were determined.

Table 7.1: The results in this table show that the EP2 receptor antagonist according to the invention E126: 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide (Example 291) has a dose-dependent effect on the number of 2-cell embryos per animal.

Treatment group	2-Cell embryos per animal (mean $\pm$ SD)
Vehicle*	36.3 $\pm$ 9.5
Piroxicam 0.1 mg/animal	10.8 $\pm$ 3.8
E126 0.1 mg/animal	30.8 $\pm$ 7.3
Piroxicam 0.1 mg/animal + E126 0.1 mg/animal	6.2 $\pm$ 4.5
E126 0.5 mg/animal	12.6 $\pm$ 3.6
Piroxicam 0.1 mg/animal + E126 0.5 mg/animal*	3.3 $\pm$ 2.1

5 * Data of n = 4 animals since there was no follicle maturation

Preparation Example 1:

Combination tablets comprising 60 mg of piroxicam and 600 mg of N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide (E16)

- 5 In a fluidized-bed granulator, 1667 g of E16, 167 g of piroxicam, 444 g of microcrystalline cellulose and 125 g of croscarmellose sodium are initially charged. This powder mixture is granulated using a granulation liquid consisting of 75 g of hydroxypropylmethylcellulose 5cP, 2.5 g of sodium lauryl sulphate and 1800 g of water. The granules are dried in the fluidized-bed granulator and then sieved and subsequently mixed with 20 g of magnesium stearate for
- 10 5 minutes. The powder mixture is then compressed on a rotary tablet press to oval tablets having a mass of 900 mg, a length of 18 mm and a width of 8 mm. The tablets are then coated with a dispersion of 55.6 g of ready-to-use film powder and 315 g of water. The ready-to-use film powder consists of 50.56% hypromellose 5cP, 10.12% Macrogol3350, 10.12% talc, 23.20% titanium dioxide and 6.00% yellow iron oxide.
- 15 Combination tablets comprising other EP2 receptor antagonists according to the invention and COX inhibitors can be prepared analogously. For alternative doses, the ratios can be adapted appropriately.

Alternatively, it is possible to formulate the EP2 receptor antagonist and COX inhibitor separately in separate tablets.

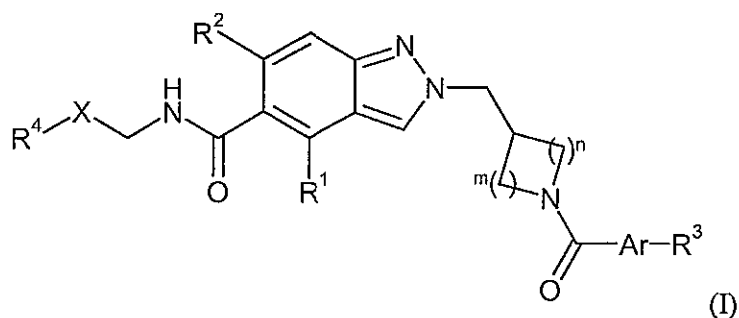
20 **Preparation Example 2:**

Kit comprising 2 combination tablets comprising 60 mg of piroxicam and 600 mg of N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide (E16)

- A folded box is manufactured which contains 3 × 2 combination tablets according to Preparation
- 25 Example 1 and can thus be used for three expected sexual intercourses. The composition of the combination tablets is identical. The folded box furthermore contains a package insert which describes that the woman has to take one tablet orally 6 to 12 hours before sexual intercourse and a further tablet within 24 hours after sexual intercourse. Furthermore, the package insert indicates that intake of the second tablet can be dispensed with if there has been no sexual
- 30 intercourse. If there has been no sexual intercourse after two administrations, the two tablets not used because of this may be used for a further administration.

Claims:

1. Use of a pharmaceutical composition comprising (a) at least one EP2 receptor antagonist according to formula (I)



wherein

R^1 : means H, C_1 - C_2 alkyl or C_1 - C_2 alkyloxy;

R^2 : H or methyl;

subject to the proviso that one of the two residues R^1 or R^2 equals H;

10 X : $-(CH_2)_l$ -, $-(CH_2)_k-O$ -, $-CH_2-S$ -, $CH_2-S(O)_2$ -, $-CH(CH_3)$ -, $-CH(CH_3)-O$ - or $-C(CH_3)_2-O$ -;

k : 1 or 2;

l : 0, 1 or 2;

R^4 : H, C_1 - C_4 alkyl, C_3 - C_4 cycloalkyl or CH_2 - C_3 - C_4 cycloalkyl;

15 and in the case that X means $-(CH_2)_l$ - with l meaning 0 or 1 or $-CH(CH_3)$ -

R^4 : additionally means a 4-6-membered heterocyclyl residue;

and in the case that X means $-(CH_2)_l$ - or $-CH(CH_3)$ -

R^4 : additionally means CN;

m : 1 or 2;

n: 1 or 2;

Ar: a 6-10-membered aryl or 5-10-membered hetaryl residue,

R³: halogen, CN, SF₅, C₁-C₄ alkyl, C₃-C₆ cycloalkyl, C₄-C₆ heterocyclyl, O-C₁-C₄ alkyl, O-C₃-C₆ cycloalkyl, O-C₄-C₆ heterocyclyl, S-C₁-C₄ alkyl, S(O)₂-C₁-C₄ alkyl, Ar', O-Ar', C(CH₃)₂-CN or C(CH₃)₂-OH; and

Ar': an optionally singly or doubly substituted 6-membered aryl or 5-6-membered heteroaryl residue;

wherein the substituents are selected from F, Cl, CN, C₁-C₄ alkyl, O-C₁-C₄ alkyl, C(CH₃)₂-CN, C(CH₃)₂-OH and C(O)NH₂;

and comprising (b) at least one COX inhibitor selected from the list below:

C1 indomethacin;

C2 diclofenac;

C4 naproxen;

C5 ibuprofen;

C6 meloxicam;

C7 piroxicam;

C8 nimesulide;

C9 ketoprofen;

C10 tenoxicam;

C11 mefanemic acid;

C12 ketoralac;

C13 celecoxib (4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl]benzenesulphonamide);

C14 parecoxib(N-[4-(5-methyl-3-phenyl-4-isoxazolyl)phenyl]sulphonylpropionamide);

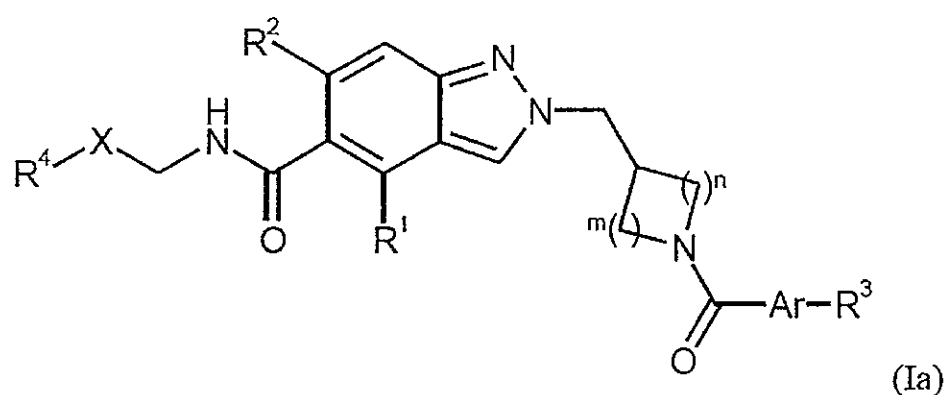
C15 rofecoxib (4-(4-mesylphenyl)-3-phenylfuran-2(5H)-one);

C16 valdecoxib (4-[5-methyl-3-phenyl-4-isoxazolyl]benzenesulphonamide);

C17 etoricoxib

for on-demand contraception.

- 5 2. Pharmaceutical composition comprising (a) at least one EP2 receptor antagonist according to formula (Ia)



wherein

R^1, R^2 : mean H or methyl;

10 subject to the proviso that one of the two residues R^1 or R^2 equals H;

X: $-(CH_2)_l-$ or $-(CH_2)_k-O-$;

k: 1 or 2;

l: 0, 1 or 2;

R^4 : H, C_1-C_4 alkyl, C_3-C_4 cycloalkyl or $CH_2-C_3-C_4$ cycloalkyl;

15 m: 1 or 2;

n: 1 or 2;

Ar: a 6-10-membered aryl or 5-10-membered hetaryl residue,

R³: halogen, CN, SF₅, C₁-C₄ alkyl, C₃-C₆ cycloalkyl, C₄-C₆ heterocyclyl, O-C₁-C₄ alkyl, O-C₃-C₆ cycloalkyl, O-C₄-C₆ heterocyclyl, S-C₁-C₄ alkyl, S(O)₂-C₁-C₄ alkyl, Ar', O-Ar', C(CH₃)₂-CN or C(CH₃)₂-OH; and

Ar': an optionally singly or doubly substituted 6-membered aryl or 5-6-membered heteroaryl residue;

wherein the substituents are selected from F, Cl, CN, C₁-C₄ alkyl, O-C₁-C₄ alkyl, C(CH₃)₂-CN, C(CH₃)₂-OH and C(O)NH₂;

and isomers, diastereomers, enantiomers and salts or cyclodextrin clathrates thereof

and comprising

(b) at least one COX inhibitor selected from the list

C1 indomethacin;

C2 diclofenac;

C4 naproxen;

C5 ibuprofen;

C6 meloxicam;

C7 piroxicam;

C8 nimesulide;

C9 ketoprofen;

C10 tenoxicam;

C11 mefanemic acid;

C12 ketoralac;

C13 celecoxib (4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl]benzenesulphonamide);

C14 parecoxib (N-[4-(5-methyl-3-phenyl-4-isoxazolyl)phenyl]sulphonylpropionamide);

- 332 -

C15 rofecoxib (4-(4-mesyphenyl)-3-phenylfuran-2(5H)-one);

C16 valdecoxib (4-[5-methyl-3-phenyl-4-isoxazolyl]benzenesulphonamide);

C17 etoricoxib

as active compounds and at least one pharmaceutically acceptable carrier.

5 3. Pharmaceutical composition according to Claim 2, wherein

R¹: means a methyl group;

R²: H;

X: -(CH₂)_l- or -(CH₂)_k-O-;

k: 1;

10 l: 0 or 1;

R⁴: C₁-C₄ alkyl, C₃-C₄ cycloalkyl or CH₂-C₃-C₄ cycloalkyl;

m, n: 1; and

Ar: a phenyl residue;

and isomers, diastereomers, enantiomers and salts or cyclodextrin clathrates thereof and

15 R³ and Ar' have the meanings stated in Claim 2.

4. Pharmaceutical composition according to Claim 2, wherein

R¹: means a methyl group;

R²: H;

X: -(CH₂)_l- or -(CH₂)_k-O-;

20 k: 1;

l: 0 or 1;

R⁴: C₁-C₄ alkyl, C₃-C₄ cycloalkyl or CH₂-C₃-C₄ cycloalkyl;

m, n: 2; and

Ar: a phenyl residue;

and isomers, diastereomers, enantiomers and salts or cyclodextrin clathrates thereof and R³ and Ar' have the meanings stated in Claim 2.

- 5 5. Pharmaceutical composition according to Claim 3 or 4, wherein the EP2 receptor antagonist is selected from the following group:
 - E1 2-{{1-(4-cyano-2-fluorobenzoyl)piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
 - E2 2-{{1-(4-tert-butoxybenzoyl)piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
 - 10 E3 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
 - E4 N-(2-methoxyethyl)-4-methyl-2-{{1-(4-morpholinobenzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
 - E5 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
 - 15 E6 2-{{1-(2-fluoro-4-mesylbenzoyl)piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
 - E7 2-{{1-(2-fluoro-4-methoxybenzoyl)piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
 - 20 E8 2-{{1-(4-bromo-2-fluorobenzoyl)piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
 - E9 2-{{1-(2-fluoro-4-methylbenzoyl)piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
 - E10 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
 - 25 E11 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pentafluoro- λ^6 -sulphonyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

- E12 N-(2-methoxyethyl)-4-methyl-2-([1-(4-methylbenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E13 2-([1-[4-(4-chlorophenoxy)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E14 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(4-methylphenoxy)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E15 2-([1-[4-(4-tert-butylphenoxy)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E16 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(4-(trifluoromethyl)phenoxy)benzoyl]piperidin-10 4-yl]methyl)-2H-indazole-5-carboxamide
- E17 N-(2-methoxyethyl)-2-([1-[4-(4-methoxyphenoxy)benzoyl]piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- E18 N-(2-methoxyethyl)-4-methyl-2-([1-(4-phenoxybenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 15 E19 2-([1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E20 2-([1-(4-methoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E21 2-([1-(4-fluorobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-20 indazole-5-carboxamide
- E22 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E23 2-([1-(2-methoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E24 N-(2-methoxyethyl)-4-methyl-2-([1-[4-[(trifluoromethyl)sulphonyl]benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E25 N-(2-methoxyethyl)-4-methyl-2-([1-[3-(trifluoromethyl)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide

- E26 N-(2-methoxyethyl)-4-methyl-2-([1-(3-methylbenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E27 2-([1-(3-chlorobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E28 2-([1-[4-(4-carbamoylphenoxy)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E29 2-([1-[4-(cyclopentyloxy)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E30 2-([1-[4-(difluoromethyl)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E31 2-([1-(4-cyanobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E32 2-([1-[4-(1H-imidazol-1-yl)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E33 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(oxazol-2-yl)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E34 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(oxazol-5-yl)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E35 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(isoxazol-5-yl)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 20 E36 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(1H-pyrazol-1-yl)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E37 N-(2-methoxyethyl)-4-methyl-2-([1-[4-(1H-1,2,4-triazol-1-yl)benzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 25 E38 2-([1-[4-(difluoromethoxy)-2-fluorobenzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E39 2-([1-[2-fluoro-4-(pyrrolidin-1-yl)benzoyl]piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E40 2-({1-[(3,4'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E41 2-({1-[(3-fluoro-4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E42 2-({1-[(3-fluoro-4'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E43 2-[(1-{[3-fluoro-3'-(trifluoromethyl)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E44 2-[(1-{[3-fluoro-2'-(trifluoromethoxy)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-
10 (2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E45 2-({1-[(2'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E46 2-({1-[(2',4'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E47 2-({1-[(2-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E48 N-(2-methoxyethyl)-4-methyl-2-({1-[(2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide
- E49 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(4-pyridyloxy)benzoyl]piperidin-4-yl}methyl)-
20 2H-indazole-5-carboxamide
- E50 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(4H-1,2,4-triazol-4-yl)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide
- E51 2-({1-(2-fluoro-4-(morpholin-4-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E52 2-({1-[4-bromobenzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E53 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide

- E54 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E55 N-(2-methoxyethyl)-4-methyl-2-([1-(4-[(trifluoromethyl)sulphonyl]benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 5 E56 4-methyl-2-([1-(4-(pentafluoro- λ^6 -sulphonyl)benzoyl)piperidin-4-yl]methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E57 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E58 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-(trifluoromethoxy)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 10 E59 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide
- E60 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-([1-(4-methylbenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 15 E61 N-[2-(cyclopropylmethoxy)ethyl]-2-([1-(4-(4-fluorophenoxy)benzoyl)piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- E62 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-([1-(4-(trifluoromethyl)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E63 N-[2-(cyclopropylmethoxy)ethyl]-2-([1-(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E64 2-([1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl)-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide
- E65 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- 25 E66 4-methyl-2-([1-(4-methylbenzoyl)piperidin-4-yl]methyl)-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E67 2-([1-(4-(4-fluorophenoxy)benzoyl)piperidin-4-yl]methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide

- E68 4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2-({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E69 2-({1-[4-(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- 5 E70 2-({1-[4-(cyclopropylbenzoyl)piperidin-4-yl]methyl}-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E71 2-({1-[4-(chlorobenzoyl)piperidin-4-yl]methyl}-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E72 N-(2-isopropoxyethyl)-4-methyl-2-({1-[4-(methylbenzoyl)piperidin-4-yl]methyl}-2H-indazole-5-carboxamide
- 10 E73 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E74 N-(2-isopropoxyethyl)-4-methyl-2-({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- 15 E75 2-({1-[4-(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E76 2-({1-[4-(cyclopropylbenzoyl)piperidin-4-yl]methyl}-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E77 2-({1-[4-(chlorobenzoyl)piperidin-4-yl]methyl}-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- 20 E78 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- E79 2-({1-[4-(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- 25 E80 4-methyl-N-[2-(trifluoromethoxy)ethyl]-2-({1-[4-[4-(trifluoromethyl)phenoxy]benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E81 N-(2-tert-butoxyethyl)-2-({1-[4-(chlorobenzoyl)piperidin-4-yl]methyl}-4-methyl-2H-indazole-5-carboxamide

- E82 N-(2-tert-butoxyethyl)-2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E83 N-(2-tert-butoxyethyl)-2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E84 N-(2-tert-butoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E85 2-({[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methyl-N-{2-[(²H3)methyloxy]-(²H4)ethyl}-2H-indazole-5-carboxamide
- 10 E86 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-{2-[(²H3)methyloxy](²H4)ethyl}-2H-indazole-5-carboxamide
- E87 2-({[1-(4-chlorobenzoyl)azetidin-3-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E88 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[(trifluoromethyl)sulphonyl]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- 15 E89 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide
- E90 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E91 2-({[1-(4-chloro-2-fluorobenzoyl)azetidin-3-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E92 2-({1-[3-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E93 2-({1-[4-chloro-3-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E94 2-({[1-(4-cyclopropylbenzoyl)azetidin-3-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E95 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide

- E96 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E97 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl]-methyl)-2H-indazole-5-carboxamide
- 5 E98 2-([1-[2-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E99 4-methyl-2-([1-[4-(pentafluoro- λ^6 -sulphanyl)benzoyl]azetidin-3-yl]methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 10 E100 N-[2-(cyclopropyloxy)ethyl]-4-methyl-2-([1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl]methyl)-2H-indazole-5-carboxamide
- E101 N-[2-(cyclobutyloxy)ethyl]-4-methyl-2-([1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl]methyl)-2H-indazole-5-carboxamide
- E102 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide
- 15 E103 N-[2-(cyclopropylmethoxy)ethyl]-2-([1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl]-methyl)-4-methyl-2H-indazole-5-carboxamide
- E104 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-([1-(4-methylbenzoyl)azetidin-3-yl]-methyl)-2H-indazole-5-carboxamide
- E105 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- 20 E106 N-(2-tert-butoxyethyl)-2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- E107 N-[2-(cyclopropylmethoxy)ethyl]-2-([1-[(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E108 2-([1-[(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E109 2-([1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E110 2-({1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E111 2-({1-[4-(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 5 E112 2-{{1-[4-(4-chlorobenzoyl)azetidin-3-yl]methyl}-4-methyl-N-[2-(2,2,2-trifluoroethoxy)-ethyl]-2H-indazole-5-carboxamide
- E113 4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- E114 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}-
10 azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- E115 2-({1-[4-(4-chlorophenoxy)benzoyl]azetidin-3-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E116 2-({1-[4-(4-chlorophenoxy)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E117 N-(2-methoxyethyl)-4-methyl-2-{{1-[4-{{5-(trifluoromethyl)pyridin-2-yl}oxy}benzoyl]-azetidin-3-yl}methyl}-2H-indazole-5-carboxamide
- E118 4-methyl-N-(2,2,2-trifluoroethyl)-2-{{1-[4-{{5-(trifluoromethyl)pyridin-2-yl}oxy}benzoyl]azetidin-3-yl}methyl}-2H-indazole-5-carboxamide
- E119 2-{{1-[4-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-ethyl-4-methyl-2H-indazole-5-
20 carboxamide
- E120 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E121 2-({1-[4-(4-chlorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 25 E122 4-methyl-2-({1-[4-(4-methylphenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E123 2-({1-[4-(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide

- E124 4-methyl-2-([1-(4-morpholinobenzoyl)piperidin-4-yl]methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E125 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-[(trifluoromethyl)sulphonyl]benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 5 E126 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-[4-(trifluoromethyl)phenoxy]benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E127 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-[[5-(trifluoromethyl)pyridin-2-yl]oxy]benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E128 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-[[6-(trifluoromethyl)pyridin-3-yl]oxy]benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 10 E129 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-[[6-(trifluoromethyl)pyridin-2-yl]oxy]benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E130 2-([1-(4-(4-cyanophenoxy)benzoyl)piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 15 E131 2-([1-(4-(3-fluorophenoxy)benzoyl)piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E132 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-[3-(trifluoromethyl)phenoxy]benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E133 2-([1-(4-[[5-cyanopyridin-2-yl]oxy]benzoyl)piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 20 E134 2-([1-(4-[[5-chloropyridin-2-yl]oxy]benzoyl)piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E135 4-methyl-N-(2,2,2-trifluoroethyl)-2-([1-(4-[5-(trifluoromethyl)pyridin-2-yl]benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 25 E136 2-([1-(4-(2,4-difluorophenoxy)benzoyl)piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E137 2-([1-(4-(3,4-difluorophenoxy)benzoyl)piperidin-4-yl]methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide

- E138 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{[4'-(trifluoromethyl)biphenyl-4-yl]carbonyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E139 2-[[1-(4-bromobenzoyl)piperidin-4-yl]methyl]-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 5 E140 2-({1-[4-(5-chloropyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E141 2-({1-[(4'-methoxy-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E142 4-methyl-2-({1-[4-(6-methylpyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 10 E143 2-({1-[(4'-fluoro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E144 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[6-(trifluoromethyl)pyridin-3-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 15 E145 2-({1-[4-(6-methoxypyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E146 2-({1-[4-(6-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E147 4-methyl-2-({1-[4-(5-methylpyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 20 E148 2-({1-[4-(5-fluoropyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E149 2-({1-[4-(5-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 25 E150 4-methyl-2-({1-[4-(2-methylpyrimidin-5-yl)benzoyl]piperidin-4-yl}methyl)-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E151 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[2-(trifluoromethyl)pyrimidin-5-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide

- E152 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[6-(trifluoromethyl)pyridin-2-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E153 2-({1-[4-(4-cyanophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E154 2-({1-[4-(4-bromo-3-methylbenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E155 2-({1-[4-(4-tert-butylbenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E156 2-({1-[4-(1-hydroxy-1-methylethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E157 2-({1-[4-(4-cyclohexylbenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E158 2-({1-[4-(1-cyano-1-methylethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E159 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyrimidin-2-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E160 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyridin-3-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E161 N-(2-methoxyethyl)-4-methyl-2-({1-[4-{5-(trifluoromethyl)pyridin-2-yl}oxy}benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- 20 E162 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyridin-2-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E163 N-(2-methoxyethyl)-4-methyl-2-({1-[4-{4-(trifluoromethyl)pyrimidin-2-yl}oxy}benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- 25 E164 N-(2-methoxyethyl)-4-methyl-2-({1-[4-{6-(trifluoromethyl)pyridin-3-yl}oxy}benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E165 N-(2-methoxyethyl)-4-methyl-2-({1-[4-{6-(trifluoromethyl)pyridin-2-yl}oxy}benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

- E166 2-({1-[(4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E167 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(3,4,5,6-tetrahydro-2H-pyran-4-yloxy)benzoyl]piperidin-4-yl} methyl)-2H-indazole-5-carboxamide
- 5 E168 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[3-(trifluoromethyl)phenoxy]benzoyl} piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E169 2-[(1-{4-[(5-cyanopyridin-2-yl)oxy]benzoyl} piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E170 2-[(1-{4-[(5-chloropyridin-2-yl)oxy]benzoyl} piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E171 2-({1-[4-(2,4-difluorophenoxy)benzoyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E172 2-({1-[4-(3,4-difluorophenoxy)benzoyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E173 N-(2-methoxyethyl)-4-methyl-2-[(1-{[4'-(trifluoromethyl)biphenyl-4-yl]carbonyl} piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E174 2-({1-[4-(3-fluorophenoxy)benzoyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E175 2-[[1-(2-fluoro-4-isopropoxybenzoyl)piperidin-4-yl]methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E176 2-({1-[(3-fluoro-3',4'-dimethylbiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E177 2-({1-[(2',3-difluoro-4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E178 2-({1-[4-(difluoromethoxy)benzoyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E179 2-({1-[4-(2-fluorophenoxy)benzoyl]piperidin-4-yl} methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E180 2-({1-[(4'-cyano-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E181 2-({1-[4-(5-chloropyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E182 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[6-(trifluoromethyl)pyridin-2-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E183 N-(2-methoxyethyl)-2-({1-[(4'-methoxy-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E184 2-({1-[(4'-chloro-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E185 2-[(1-{4'-(2-cyanopropan-2-yl)biphenyl-4-yl}carbonyl)piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E186 N-(2-methoxyethyl)-2-({1-[4-(5-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E187 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[6-(trifluoromethyl)pyridin-3-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E188 N-(2-methoxyethyl)-2-({1-[4-(6-methoxypyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E189 2-({1-[(4'-fluoro-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E190 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(6-methylpyridin-3-yl)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide
- E191 N-(2-methoxyethyl)-2-({1-[4-(6-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E192 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(2-methylpyrimidin-5-yl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E193 2-({1-[(4'-fluoro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E194 2-({1-[(4'-chloro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E195 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[2-(trifluoromethyl)pyrimidin-5-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 5 E196 2-({1-[(4'-chloro-2'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E197 2-({1-[(2'-chloro-4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E198 2-({1-[4-(5-chloropyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E199 2-({1-[4-(5-fluoropyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E200 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[5-(trifluoromethyl)pyridin-3-yl]benzoyl}-piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- 15 E201 2-[(1-{[4'-(2-hydroxypropan-2-yl)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E202 2-({1-[(3',5'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E203 2-({1-[(4'-fluoro-2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E204 2-({1-[(3',5'-difluoro-2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E205 N-(2-methoxyethyl)-4-methyl-2-({1-[3-methyl-4-(3-pyridyl)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide
- 25 E206 N-(2-methoxyethyl)-4-methyl-2-({1-[3-methyl-4-(4-pyridyl)benzoyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide and
- E207 N-(2-methoxyethyl)-4-methyl-2-({1-[(2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}-methyl)-2H-indazole-5-carboxamide

6. Pharmaceutical composition according to Claim 5, where the EP2 receptor antagonist is selected from the list below:

- E2 2-([1-(4-tert-butoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E3 2-([1-(4-(4-fluorophenoxy)benzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E5 2-([1-(4'-(4-fluorobiphenyl-4-yl)carbonyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E7 2-([1-(2-fluoro-4-methoxybenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E8 2-([1-(4-bromo-2-fluorobenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E9 2-([1-(2-fluoro-4-methylbenzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E10 2-([1-(2-fluoro-4-(trifluoromethyl)benzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E11 N-(2-methoxyethyl)-4-methyl-2-([1-(4-(pentafluoro- λ^6 -sulphanyl)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 20 E12 N-(2-methoxyethyl)-4-methyl-2-([1-(4-methylbenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E13 2-([1-(4-(4-chlorophenoxy)benzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E14 N-(2-methoxyethyl)-4-methyl-2-([1-(4-(4-methylphenoxy)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 25 E15 2-([1-(4-(4-tert-butylphenoxy)benzoyl)piperidin-4-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E16 N-(2-methoxyethyl)-4-methyl-2-([1-(4-(4-(trifluoromethyl)phenoxy)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide

- E17 N-(2-methoxyethyl)-2-({1-[4-(4-methoxyphenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E18 N-(2-methoxyethyl)-4-methyl-2-({1-[4-phenoxybenzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- 5 E19 2-({1-[4-(cyclopropylbenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E20 2-({1-[4-(4-methoxybenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E21 2-({1-[4-(4-fluorobenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E22 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E24 N-(2-methoxyethyl)-4-methyl-2-({1-[4-({trifluoromethyl)sulphonyl]benzoyl}piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 15 E25 N-(2-methoxyethyl)-4-methyl-2-({1-[3-(trifluoromethyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E26 N-(2-methoxyethyl)-4-methyl-2-({1-[3-methylbenzoyl]piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E27 2-({1-[3-chlorobenzoyl]piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E29 2-({1-[4-(cyclopentyloxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E30 2-({1-[4-(difluoromethyl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E31 2-({1-[4-(cyanobenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E36 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(1H-pyrazol-1-yl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

- E38 2-({1-[4-(difluoromethoxy)-2-fluorobenzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E39 2-({1-[2-fluoro-4-(pyrrolidin-1-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E40 2-({1-[(3,4'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E41 2-({1-[(3-fluoro-4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E42 2-({1-[(3-fluoro-4'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E43 2-[(1-{[3-fluoro-3'-(trifluoromethyl)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E44 2-[(1-{[3-fluoro-2'-(trifluoromethoxy)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E45 2-({1-[(2'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E46 2-({1-[(2',4'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E47 2-({1-[(2-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E48 N-(2-methoxyethyl)-4-methyl-2-({1-[(2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E51 2-({1-[2-fluoro-4-(morpholin-4-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E52 2-{[1-(4-bromobenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E53 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

- E54 2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E55 N-(2-methoxyethyl)-4-methyl-2-([1-(4-[(trifluoromethyl)sulphonyl]benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 5 E59 2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide
- E60 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-([1-(4-methylbenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E61 N-[2-(cyclopropylmethoxy)ethyl]-2-([1-(4-(4-fluorophenoxy)benzoyl)piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E62 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-([1-(4-(trifluoromethyl)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E63 N-[2-(cyclopropylmethoxy)ethyl]-2-([1-(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E64 2-{[1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl}-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide
- E65 2-{[1-(4-chlorobenzoyl)piperidin-4-yl]methyl}-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E66 4-methyl-2-([1-(4-methylbenzoyl)piperidin-4-yl]methyl)-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- 20 E67 2-([1-(4-(4-fluorophenoxy)benzoyl)piperidin-4-yl]methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E68 4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2-([1-(4-(trifluoromethyl)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 25 E69 2-([1-(4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl]methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- E70 2-([1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl)-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide

- E71 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E72 N-(2-isopropoxyethyl)-4-methyl-2-([1-(4-methylbenzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 5 E73 2-([1-(4-(4-fluorophenoxy)benzoyl)piperidin-4-yl]methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E74 N-(2-isopropoxyethyl)-4-methyl-2-([1-(4-(trifluoromethyl)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- E75 2-([1-(4'-(4-fluorobiphenyl-4-yl)carbonyl)piperidin-4-yl]methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E76 2-([1-(4-cyclopropylbenzoyl)piperidin-4-yl]methyl)-N-(2-isopropoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E77 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- 15 E78 2-([1-(4-(4-fluorophenoxy)benzoyl)piperidin-4-yl]methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- E79 2-([1-(4'-(4-fluorobiphenyl-4-yl)carbonyl)piperidin-4-yl]methyl)-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- E80 4-methyl-N-[2-(trifluoromethoxy)ethyl]-2-([1-(4-(4-(trifluoromethyl)phenoxy)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide
- 20 E81 N-(2-tert-butoxyethyl)-2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- E82 N-(2-tert-butoxyethyl)-2-([1-(4-(4-fluorophenoxy)benzoyl)piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E83 N-(2-tert-butoxyethyl)-2-([1-(4'-(4-fluorobiphenyl-4-yl)carbonyl)piperidin-4-yl]methyl)-4-methyl-2H-indazole-5-carboxamide
- E84 N-(2-tert-butoxyethyl)-4-methyl-2-([1-(4-(4-(trifluoromethyl)phenoxy)benzoyl)piperidin-4-yl]methyl)-2H-indazole-5-carboxamide

- E85 2-([1-(4-chlorobenzoyl)piperidin-4-yl]methyl)-4-methyl-N-{2-[(2H3)methyloxy](2H4)ethyl}-2H-indazole-5-carboxamide
- E86 2-({1-[4-(4-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-{2-[(2H3)methyloxy](2H4)ethyl}-2H-indazole-5-carboxamide
- 5 E87 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E88 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[(trifluoromethyl)sulphonyl]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- E89 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide
- 10
- E90 2-({1-[2-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E91 2-([1-(4-chloro-2-fluorobenzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E92 2-({1-[3-fluoro-4-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E93 2-({1-[4-chloro-3-(trifluoromethyl)benzoyl]azetidin-3-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E94 2-([1-(4-cyclopropylbenzoyl)azetidin-3-yl]methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20
- E95 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- E100 N-[2-(cyclopropyloxy)ethyl]-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide
- 25 E101 N-[2-(cyclobutyloxy)ethyl]-4-methyl-2-({1-[4-(trifluoromethoxy)benzoyl]azetidin-3-yl}methyl)-2H-indazole-5-carboxamide
- E102 2-([1-(4-chlorobenzoyl)azetidin-3-yl]methyl)-N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2H-indazole-5-carboxamide

- E103 N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl)methyl}-4-methyl-2H-indazole-5-carboxamide
- E104 N-[2-(cyclopropylmethoxy)ethyl]-4-methyl-2-({1-[4-(4-methylbenzoyl)azetidin-3-yl)methyl}-2H-indazole-5-carboxamide
- 5 E105 2-({1-[4-(4-chlorobenzoyl)azetidin-3-yl)methyl}-4-methyl-N-[2-(trifluoromethoxy)ethyl]-2H-indazole-5-carboxamide
- E107 N-[2-(cyclopropylmethoxy)ethyl]-2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl)methyl}-4-methyl-2H-indazole-5-carboxamide
- E108 2-({1-[(4'-fluorobiphenyl-4-yl)carbonyl]azetidin-3-yl)methyl)-N-(2-methoxyethyl)-4-
10 methyl-2H-indazole-5-carboxamide
- E109 2-({1-[4-(4-fluorophenoxy)benzoyl]azetidin-3-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E112 2-({1-[4-(4-chlorobenzoyl)azetidin-3-yl)methyl}-4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2H-indazole-5-carboxamide
- 15 E113 4-methyl-N-[2-(2,2,2-trifluoroethoxy)ethyl]-2-({1-[4-(4-(trifluoromethyl)phenoxy]benzoyl}azetidin-3-yl)methyl)-2H-indazole-5-carboxamide
- E114 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}azetidin-3-yl)methyl]-2H-indazole-5-carboxamide
- E115 2-({1-[4-(4-chlorophenoxy)benzoyl]azetidin-3-yl)methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
20
- E116 2-({1-[4-(4-chlorophenoxy)benzoyl]azetidin-3-yl)methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E117 N-(2-methoxyethyl)-4-methyl-2-({1-[4-{5-(trifluoromethyl)pyridin-2-yl}oxy}benzoyl]azetidin-3-yl)methyl)-2H-indazole-5-carboxamide
- 25 E125 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[(trifluoromethyl)sulphanyl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E126 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide

- E127 4-methyl-N-(2,2,2-trifluoroethyl)-2-{{1-(4-{{5-(trifluoromethyl)pyridin-2-yl}oxy}benzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- E128 4-methyl-N-(2,2,2-trifluoroethyl)-2-{{1-(4-{{6-(trifluoromethyl)pyridin-3-yl}oxy}benzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- 5 E131 2-({1-[4-(3-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E132 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[3-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E137 2-({1-[4-(3,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 10 E138 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4'-(trifluoromethyl)biphenyl-4-yl}carbonyl)piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E141 2-({1-[(4'-methoxy-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- 15 E143 2-({1-[(4'-fluoro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-N-(2,2,2-trifluoroethyl)-2H-indazole-5-carboxamide
- E153 2-({1-[4-(4-cyanophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E154 2-{{1-[4-bromo-3-methylbenzoyl]piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E155 2-{{1-[4-tert-butylbenzoyl]piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E157 2-{{1-[4-cyclohexylbenzoyl]piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E160 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyridin-3-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E161 N-(2-methoxyethyl)-4-methyl-2-{{1-(4-{{5-(trifluoromethyl)pyridin-2-yl}oxy}benzoyl)piperidin-4-yl}methyl}-2H-indazole-5-carboxamide

- E162 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(pyridin-2-yloxy)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E163 N-(2-methoxyethyl)-4-methyl-2-{{1-[4-{[4-(trifluoromethyl)pyrimidin-2-yl]oxy}benzoyl]piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- 5 E164 N-(2-methoxyethyl)-4-methyl-2-{{1-[4-{[6-(trifluoromethyl)pyridin-3-yl]oxy}benzoyl]piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- E165 N-(2-methoxyethyl)-4-methyl-2-{{1-[4-{[6-(trifluoromethyl)pyridin-2-yl]oxy}benzoyl]piperidin-4-yl}methyl}-2H-indazole-5-carboxamide
- E166 2-({1-[(4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-
10 4-methyl-2H-indazole-5-carboxamide
- E168 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[3-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E170 2-[(1-{4-[(5-chloropyridin-2-yl)oxy]benzoyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E171 2-({1-[4-(2,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E172 2-({1-[4-(3,4-difluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E173 N-(2-methoxyethyl)-4-methyl-2-[(1-{[4'-(trifluoromethyl)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
20
- E174 2-({1-[4-(3-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E175 2-{{1-[2-fluoro-4-isopropoxybenzoyl]piperidin-4-yl}methyl}-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E176 2-({1-[(3-fluoro-3',4'-dimethylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E177 2-({1-[(2',3-difluoro-4'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide

- E178 2-({1-[4-(difluoromethoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E179 2-({1-[4-(2-fluorophenoxy)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E180 2-({1-[4-(4'-cyano-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E181 2-({1-[4-(5-chloropyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E182 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[6-(trifluoromethyl)pyridin-2-yl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
10
- E183 N-(2-methoxyethyl)-2-({1-[(4'-methoxy-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E184 2-({1-[(4'-chloro-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E185 2-[(1-{[4'-(2-cyanopropan-2-yl)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E186 N-(2-methoxyethyl)-2-({1-[4-(5-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E187 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[6-(trifluoromethyl)pyridin-3-yl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
20
- E188 N-(2-methoxyethyl)-2-({1-[4-(6-methoxypyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide
- E189 2-({1-[(4'-fluoro-2'-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E190 N-(2-methoxyethyl)-4-methyl-2-({1-[4-(6-methylpyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E191 N-(2-methoxyethyl)-2-({1-[4-(6-methoxypyridin-2-yl)benzoyl]piperidin-4-yl}methyl)-4-methyl-2H-indazole-5-carboxamide

- E193 2-({1-[(4'-fluoro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E194 2-({1-[(4'-chloro-2'-methoxybiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 5 E195 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[2-(trifluoromethyl)pyrimidin-5-yl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E196 2-({1-[(4'-chloro-2'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E197 2-({1-[(2'-chloro-4'-fluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 10 E198 2-({1-[4-(5-chloropyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E199 2-({1-[4-(5-fluoropyridin-3-yl)benzoyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 15 E200 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[5-(trifluoromethyl)pyridin-3-yl]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide
- E201 2-[(1-{[4'-(2-hydroxypropan-2-yl)biphenyl-4-yl]carbonyl}piperidin-4-yl)methyl]-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E202 2-({1-[(3',5'-difluorobiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 20 E203 2-({1-[(4'-fluoro-2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- E204 2-({1-[(3',5'-difluoro-2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl}methyl)-N-(2-methoxyethyl)-4-methyl-2H-indazole-5-carboxamide
- 25 E205 N-(2-methoxyethyl)-4-methyl-2-({1-[3-methyl-4-(3-pyridyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide
- E206 N-(2-methoxyethyl)-4-methyl-2-({1-[3-methyl-4-(4-pyridyl)benzoyl]piperidin-4-yl}methyl)-2H-indazole-5-carboxamide

E207 N-(2-methoxyethyl)-4-methyl-2-({1-[(2-methylbiphenyl-4-yl)carbonyl]piperidin-4-yl)methyl}-2H-indazole-5-carboxamide

7. Pharmaceutical composition according to Claim 6, where the EP2 receptor antagonist is selected from the list below:

5 E16 N-(2-methoxyethyl)-4-methyl-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide

E126 4-methyl-N-(2,2,2-trifluoroethyl)-2-[(1-{4-[4-(trifluoromethyl)phenoxy]benzoyl}piperidin-4-yl)methyl]-2H-indazole-5-carboxamide

10 E161 N-(2-methoxyethyl)-4-methyl-2-{{1-(4-{[5-(trifluoromethyl)pyridin-2-yl]oxy}benzoyl)piperidin-4-yl)methyl}-2H-indazole-5-carboxamide

8. Pharmaceutical composition according to any of Claims 2 - 7, characterized in that it comprises the COX inhibitor in the following ranges:

	C1	indomethacin:	19 – 800 mg/day;
	C2	diclofenac:	37.5 – 800 mg/day;
15	C4	naproxen:	125 – 4000 mg/day;
	C5	ibuprofen:	200 – 4800 mg/day;
	C6	meloxicam:	2 – 60 mg/day;
	C7	piroxicam:	5 – 80 mg/day;
	C8	nimesulide:	25 – 800 mg/day;
20	C9	ketoprofen:	25 – 1200 mg/day;
	C10	tenoxicam:	10 – 160 mg/day;
	C11	mefanemic acid:	125 – 2000 mg/day;
	C12	ketoralac:	2.5 – 160 mg/day;
	C13	celecoxib:	25 – 800 mg/day;
25	C14	parecoxib	12.5 – 200 mg/day;

- 360 -

C15 rofecoxib: 6 - 200 mg/day;

C16 valdecoxib: 10 - 160 mg/day;

C17 etoricoxib: 15 - 360 mg/day.

9. Pharmaceutical composition according to any of Claims 2 - 7, where the COX inhibitor is

5 C1 indomethacin;

C2 diclofenac;

C4 naproxen;

C5 ibuprofen;

C6 meloxicam;

10 C7 piroxicam;

C8 nimesulide.

10. Pharmaceutical composition according to Claim 9, where the COX inhibitor is:

C1 indomethacin;

C2 diclofenac;

15 C6 meloxicam;

C7 piroxicam.

11. Use of the pharmaceutical composition according to one of Claims 2-10 for the production of a drug containing suitable formulation and carrier substances.

12. Use of the pharmaceutical composition according to one of Claims 2-10 and the drugs
20 according to Claim 11 for fertility control/contraception.

13. Use of the pharmaceutical composition according to one of Claims 2-10 and the drugs according to Claim 11 for emergency contraception.

14. Use according to Claim 1 or 13, where the pharmaceutical composition is administered in a single administration dose in a period of up to 24 hours, preferably 6 - 12 hours, before expected sexual intercourse or up to 2 hours after sexual intercourse.
15. Method for on-demand contraception, characterized in that the pharmaceutical composition according to any of Claims 2 - 10 is administered on demand and once in a period of 0 - 24, preferably 6 - 12, hours prior to expected sexual intercourse or up to 2 hours after sexual intercourse.
16. Method for on-demand contraception, characterized in that initially a first pharmaceutical composition according to any of Claims 2 - 10 is taken in a period of up to 24 hours, preferably 6 - 12 hours, prior to expected sexual intercourse, and a second pharmaceutical composition according to any of Claims 2 - 10 is taken in a period of up to 48 hours after sexual intercourse, where the preparations taken before and after intercourse are identical.
17. Method for on-demand contraception, characterized in that initially a first pharmaceutical composition according to any of Claims 2 - 10 is taken in a period of up to 24 hours, preferably 6 - 12 hours, prior to expected sexual intercourse, and a second pharmaceutical composition according to any of Claims 2 - 10 is taken in a period of up to 48 hours after sexual intercourse, where the preparations taken before and after intercourse are not identical.
18. Administration protocol for on-demand contraception, characterized in that a pharmaceutical composition according to any of Claims 2 - 10 is taken in a period of up to 24 hours, preferably 6 - 12 hours, prior to expected sexual intercourse and a further pharmaceutical composition according to any of Claims 2 - 10 is taken in a period of up to 48 hours after sexual intercourse.
19. Kit for providing an administration protocol according to Claim 18 or for carrying out a method according to either of Claims 16 - 17 for on-demand contraception, comprising in each case at least one tablet comprising a pharmaceutical composition according to any of Claims 2 - 10, for administration before, and at least one tablet comprising a pharmaceutical composition according to any of Claims 2 - 10, for administration after expected sexual intercourse.
20. Use of the pharmaceutical composition according to one of Claims 2-10, in the form of a pharmaceutical preparation for oral, pulmonary, nasal, dermal or transdermal administration.
21. Use of a pharmaceutical composition according to Claim 1, wherein

- 362 -

in the case that X means $\cdot(\text{CH}_2)_l\cdot$ with l meaning 0

R^4 : is additionally a 4-6-membered heterocyclyl radical linked via a ring carbon atom.