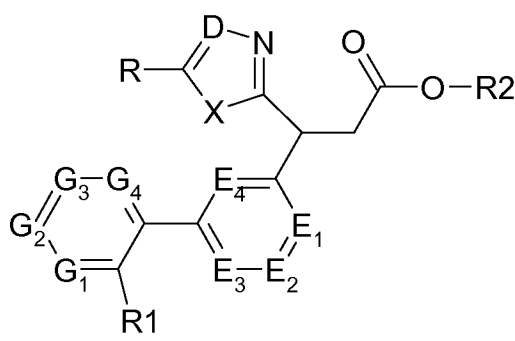


Biaryl-propionic acid derivatives and their use as pharmaceuticals

5 The present invention relates to compounds of the formula I,



10 wherein X, R, R1, R2, D, E1, E2, E3, E4, G1, G2, G3 and G4 have the meanings indicated below, which are valuable pharmaceutical active compounds. They are inhibitors of the protease cathepsin A, and are useful for the treatment of diseases such as atherosclerosis, heart failure, renal diseases, liver diseases or inflammatory diseases, for example. The invention furthermore relates to processes for the preparation of the compounds of the formula I, their use and pharmaceutical
15 compositions comprising them.

Cathepsin A (EC = 3.4.16.5; gene symbol CTSA) is a protease also known as lysosomal carboxypeptidase A or protective protein. It belongs to a family of serine carboxypeptidases which contains only two other mammalian representatives,
20 retinoid-inducible serine carboxypeptidase and vitellogenic carboxypeptidase-like protein. Within the cell cathepsin A resides in lysosomes where it forms a high molecular weight complex with beta-galactosidase and neuraminidase. The interaction of cathepsin A with these glycosidases is essential for their correct routing to the lysosome and protects them from intralysosomal proteolysis. A deficiency of
25 cathepsin A resulting from various mutations in the ctsa gene leads to a secondary deficiency of beta-galactosidase and neuraminidase that is manifest as the

autosomal recessive lysosomal storage disorder galactosialidosis (cf. A. d'Azzo et al., in "The Metabolic and Molecular Bases of Inherited Disease", vol. 2 (1995), 2835-2837). The majority of identified mutations in *ctsa* are missense mutations affecting the folding or the stability of the protein. None of them was shown to occur in the active site of the enzyme (G. Rudenko et al., Proc. Natl. Acad. Sci. USA 95 (1998), 621-625). Accordingly, the lysosomal storage disorder can be corrected with catalytically inactive cathepsin A mutants (N. J. Galjart et al., J. Biol. Chem. 266 (1991), 14754-14762). The structural function of cathepsin A is therefore separable from its catalytic activity. This is also underscored by the observation that in contrast to mice deficient in the *ctsa* gene, mice carrying a catalytically inactivating mutation in the *ctsa* gene do not develop signs of the human disease galactosialidosis (R. J. Rottier et al., Hum. Mol. Genet. 7 (1998), 1787-1794; V. Seyrantepe et al., Circulation 117 (2008), 1973-1981).

Cathepsin A displays carboxypeptidase activity at acidic pH and deamidase and esterase activities at neutral pH against various naturally occurring bioactive peptides. In vitro studies have indicated that cathepsin A converts angiotensin I to angiotensin 1-9 and bradykinin to bradykinin 1-8, which is the ligand for the bradykinin B1 receptor. It hydrolyzes endothelin-1, neurokinin and oxytocin, and deamidates substance P (cf. M. Hiraiwa, Cell. Mol. Life Sci. 56 (1999), 894-907). High cathepsin A activity has been detected in urine, suggesting that it is responsible for tubular bradykinin degradation (M. Saito et al., Int. J. Tiss. Reac. 17 (1995), 181-190). However, the enzyme can also be released from platelets and lymphocytes and is expressed in antigen-presenting cells where it might be involved in antigen processing (W. L. Hanna et al., J. Immunol. 153 (1994), 4663-4672; H. Ostrowska, Thromb. Res. 86 (1997), 393-404; M. Reich et al., Immunol. Lett. (online Nov. 30, 2009)). Immunohistochemistry of human organs revealed prominent expression in renal tubular cells, bronchial epithelial cells, Leydig's cells of the testis and large neurons of the brain (O. Sohma et al., Pediatr. Neurol. 20 (1999), 210-214). It is upregulated during differentiation of monocytes to macrophages (N. M. Stamatatos et al., FEBS J. 272 (2005), 2545-2556). Apart from structural and enzymatic functions, cathepsin A has been shown to associate with neuraminidase and an alternatively

spliced beta-galactosidase to form the cell-surface laminin and elastin receptor complex expressed on fibroblasts, smooth muscle cells, chondroblasts, leukocytes and certain cancer cell types (A. Hinek, *Biol. Chem.* 377 (1996), 471-480).

- 5 The importance of cathepsin A for the regulation of local bradykinin levels has been demonstrated in animal models of hypertension. Pharmacological inhibition of cathepsin A activity increased renal bradykinin levels and prevented the development of salt-induced hypertension (H. Ito et al., *Br. J. Pharmacol.* 126 (1999), 613-620). This could also be achieved by antisense oligonucleotides suppressing the
- 10 expression of cathepsin A (I. Hajashi et al., *Br. J. Pharmacol.* 131 (2000), 820-826). Besides in hypertension, beneficial effects of bradykinin have been demonstrated in various further cardiovascular diseases and other diseases (cf. J. Chao et al., *Biol. Chem.* 387 (2006), 665-75; P. Madeddu et al., *Nat. Clin. Pract. Nephrol.* 3 (2007), 208-221). Key indications of cathepsin A inhibitors therefore include atherosclerosis,
- 15 heart failure, cardiac infarction, cardiac hypertrophy, vascular hypertrophy, left ventricular dysfunction, in particular left ventricular dysfunction after myocardial infarction, renal diseases such as renal fibrosis, renal failure and kidney insufficiency; liver diseases such as liver fibrosis and liver cirrhosis, diabetes complications such as nephropathy, as well as organ protection of organs such as the heart and the
- 20 kidney.

- As indicated above, cathepsin A inhibitors can prevent the generation of the bradykinin B1 receptor ligand bradykinin 1-8 (M. Saito et al., *Int. J. Tiss. Reac.* 17 (1995), 181-190). This offers the opportunity to use cathepsin A inhibitors for the
- 25 treatment of pain, in particular neuropathic pain, and inflammation, as has been shown for bradykinin B1 receptor antagonists (cf. F. Marceau et al., *Nat. Rev. Drug Discov.* 3 (2004), 845-852). Cathepsin A inhibitors can further be used as anti-platelet agents as has been demonstrated for the cathepsin A inhibitor ebelactone B, a propiolactone derivative, which suppresses platelet aggregation in hypertensive
- 30 animals (H. Ostrowska et al., *J. Cardiovasc. Pharmacol.* 45 (2005), 348-353).

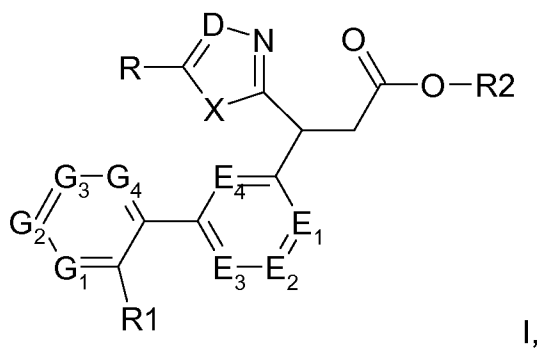
Further, like other serine proteases such as prostatic, elastase or matriptase, cathepsin A can stimulate the amiloride-sensitive epithelial sodium channel (ENaC) and is thereby involved in the regulation of fluid volumes across epithelial membranes (cf. C. Planes et al., Curr. Top. Dev. Biol. 78 (2007), 23-46). Thus, respiratory diseases can be ameliorated by the use of cathepsin A inhibitors, such as cystic fibrosis, chronic bronchitis, chronic obstructive pulmonary disease, asthma, respiratory tract infections and lung carcinoma. Cathepsin A modulation in the kidney could be used to promote diuresis and thereby induce a hypotensive effect. Also beneficial effects of Cathepsin A inhibitors on atrial fibrillation are reported in J. Med. Chem. 2012, 55, no. 17, 7636-7649.

Besides for the above-mentioned compound ebelactone B, an inhibitory effect on cathepsin A has been found for certain dipeptidic phenylalanine derivatives which are described in JP 2005/145839. Also in WO2011/092187, WO2012/101197, WO2012/101199, WO2013/014204, WO2013/014205, WO2013/072327, WO2013/072328 compounds with inhibitory activity on Cathepsin A are disclosed. There is a need for further compounds which inhibit cathepsin A and offer an opportunity for the treatment of the mentioned diseases and further diseases in which cathepsin A plays a role. The present invention satisfies this need by providing the oxygen-substituted 3-heteroaroylamino-propionic acid derivatives of the formula I defined below.

Certain compounds in which a 3-heteroaroylamino-propionic acid moiety can be present, have already been described. For example, in WO 2006/076202 amine derivatives, which modulate the activity of steroid nuclear receptors, are described which carry on the nitrogen atom of the amine function a heteroaroyl group and a further group which is defined very broadly. In US 2004/0072802 broadly-defined beta-amino acid derivatives are described which carry an acyl group on the beta-amino group and are inhibitors of matrix metalloproteases and/or tumor necrosis factor. In WO 2009/080226 and WO 2009/080227, which relate to antagonists of the platelet ADP receptor P2Y₁₂ and inhibit platelet aggregation, pyrazoloylamino-substituted carboxylic acid derivatives are described which, however, additionally

carry a carboxylic acid derivative group on the carbon atom carrying the pyrazoloylamino group. Other pyrazoloylamino-substituted compounds, in which the nitrogen atom of the amino group is connected to a ring system and which are inhibitors of the blood clotting enzymes factor Xa and/or factor VIIa, are described in WO 2004/056815.

A subject of the present invention is a compound of the formula I, in any of its stereoisomeric forms or a mixture of stereoisomeric forms in any ratio, or a physiologically acceptable salt thereof, or a physiologically acceptable solvate of any of them,



wherein the meanings are:

X is S or O;

D is N or $-\text{C}(\text{R}_3)=$;

R is H, $(\text{C}_1\text{-C}_6)$ -alkyl, $(\text{C}_3\text{-C}_8)$ -cycloalkyl;

R1 is H, F, Cl, CF_3 , CN, CO-H, CO- $(\text{C}_1\text{-C}_6)$ -alkyl, CO-NR20R21; wherein R20 and

R21 are independently from each other H or $(\text{C}_1\text{-C}_6)$ -alkyl;

R3 is H, methyl or ethyl;

R2 is Hydrogen or $(\text{C}_1\text{-C}_6)$ -alkyl;

E1 is N or $-\text{C}(\text{R}_4)=$;

E2 is N or $-\text{C}(\text{R}_5)=$;

E3 is N or $-\text{C}(\text{R}_6)=$;

E₄ is N or -C(R7)=; wherein none or one of E₁, E₂, E₃ or E₄ is N;

R₄ is H or O-(C₁-C₆)-alkyl;

R₅ is H, F, Cl, CF₃, (C₁-C₆)-alkyl, (C₁-C₆)-cycloalkyl;

5 R₆ is H;

R₇ is H;

G₁ is N or -C(R8)=;

G₂ is N or -C(R9)=;

10 G₃ is N or -C(R10)=;

G₄ is N or -C(R11)=; wherein none or one of G₁, G₂, G₃ or G₄ is N;

or G₃ and G₄ are -C(R10)= and -C(R11)=, wherein R10 and R11 form a 4 to 7 membered saturated carbocycle- or heterocycle with one or two oxygen atoms; which is optionally mono- or disubstituted by (C₁-C₃)-alkyl;

15

R₈ is H, F, Cl, (C₁-C₆)-alkyl, O-(C₁-C₆)-alkyl, CF₃, -CH₂CO-NH₂ or OCF₃;

R₉ is H, F, Cl, OH, O-(C₁-C₆)-alkyl, CH₂OH, CO-NH₂, (C₁-C₆)-alkyl, O-(C₁-C₆)-alkyl, CF₃ or OCF₃;

R₁₀ is H, F, Cl, OH, (C₁-C₆)-alkyl, CH₂OH, CO-O-(C₁-C₆)-alkyl, SO₂-(C₁-C₆)-alkyl,

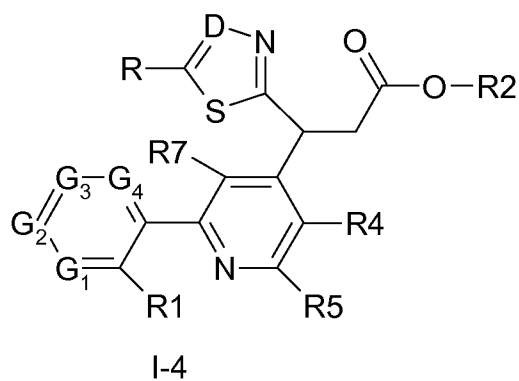
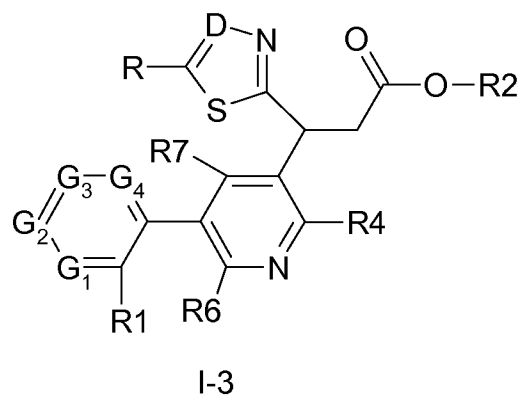
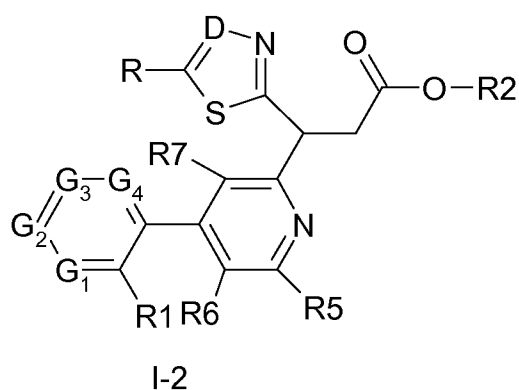
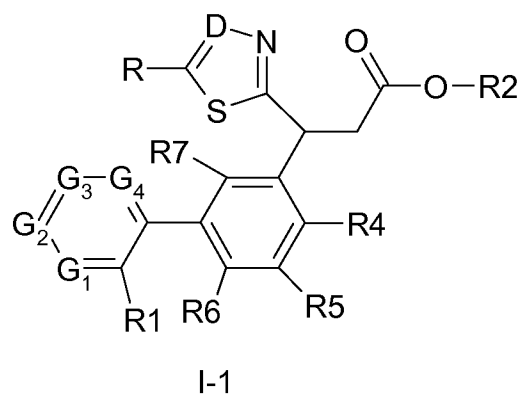
20 CN, O-(C₁-C₆)-alkyl, CF₃ or OCF₃;

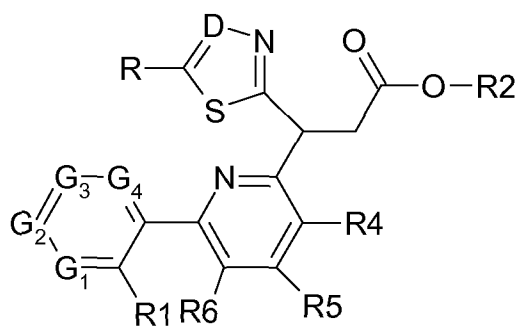
R₁₁ is H, F, Cl, OH, O-(C₁-C₆)-alkyl, CH₂OH, CO-(C₁-C₆)-methyl, CO-N(R₂₀R₂₁), CO-O-(C₁-C₆)-alkyl, CN, (C₁-C₆)-alkyl, O-(C₁-C₆)-alkyl or OCF₃; wherein R₂₀ and R₂₁ are independently from each other H or (C₁-C₃)-alkyl or form together with the nitrogen to which they are attached a 5 or 6 membered saturated ring.

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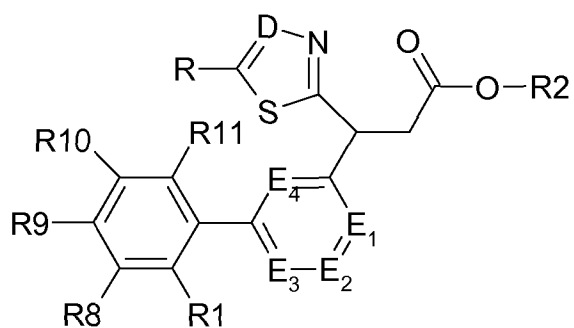
In terms of formulae resulting from formula I by incorporation of meanings of D, X, E₁, E₂, E₃, and E₄, or G₁, G₂, G₃ and G, in one embodiment of the invention a compound of the formula I is a compound of any one or more of formulae I-1 to I-20, for example a compound of formula I-1, or a compound of formula I-2, or a
30 compound of formula I-3, or a compound of formula I-4, or a compound of formula I-5, or a compound of formula I-6, or a compound of formula I-7, or a compound of formula I-8, or a compound of formula I-9 or formula I-10, or a compound of formula

I-11 or formula I-12, or a compound of formula I-13 or formula I-14 or formula I-15, or formula I-16, or formula I-17, or formula I-18, or formula I-19, or formula I-20, in any of its stereoisomeric forms or a mixture of stereoisomeric forms in any ratio, or a physiologically acceptable salt thereof, or a physiologically acceptable solvate of any of them, wherein in the compounds of formulae I-1 to I-20 the groups D, R, R1, R2, R4, R5, R6, R7, R8, R9, R10, R11, E1, E2, E3, E4, G1, G2, G3 and G4, are defined as in the compounds of formula I in general or in any embodiment specified above or below.

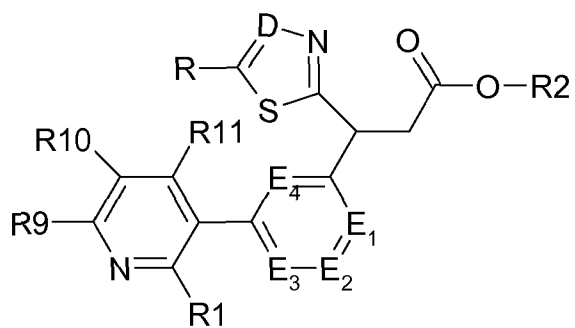




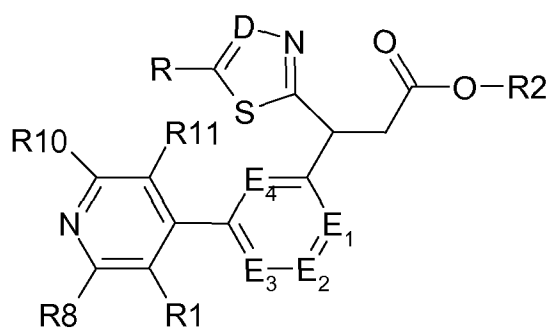
I-5



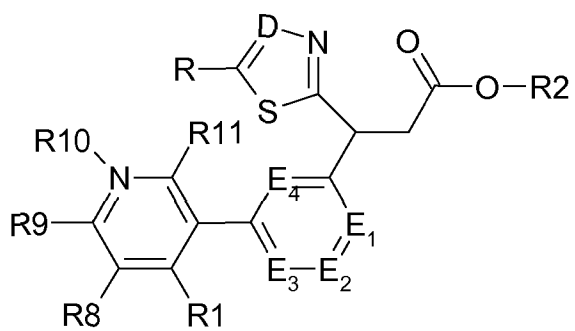
I-6



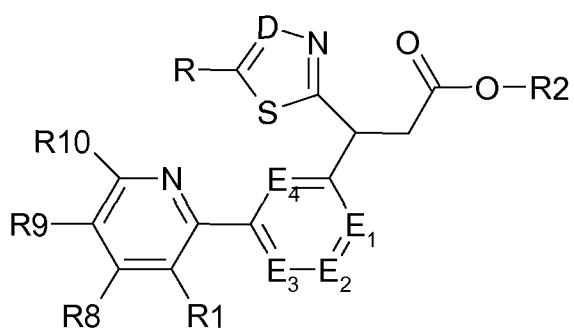
I-7



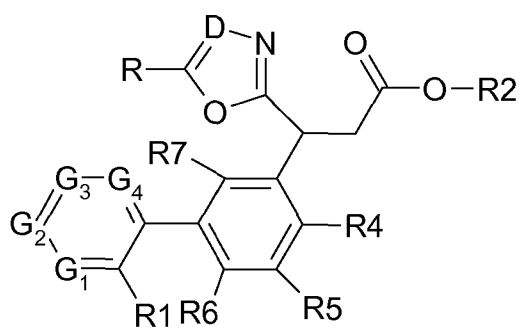
I-8



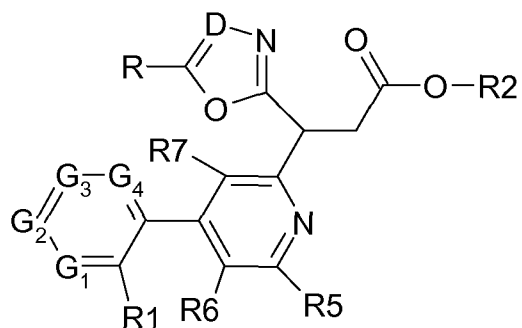
I-9



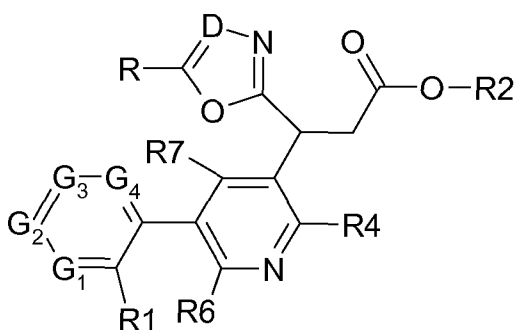
I-10



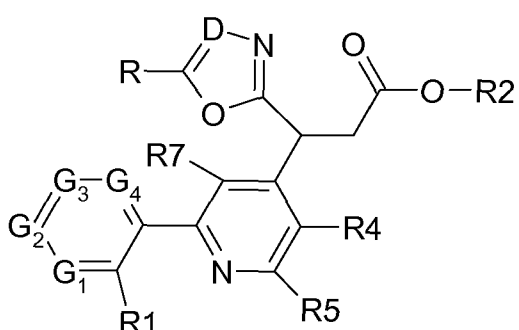
I-11



I-12

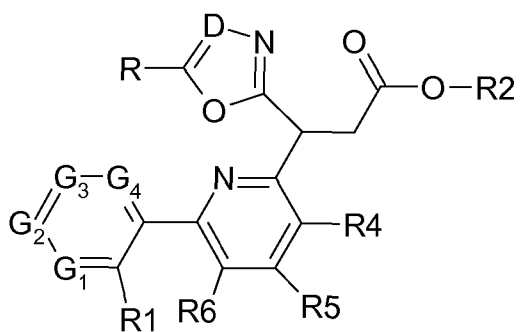


I-13

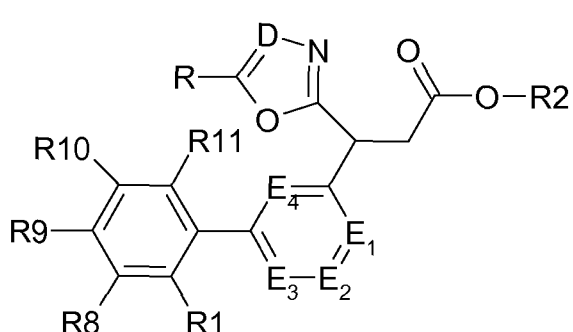


I-14

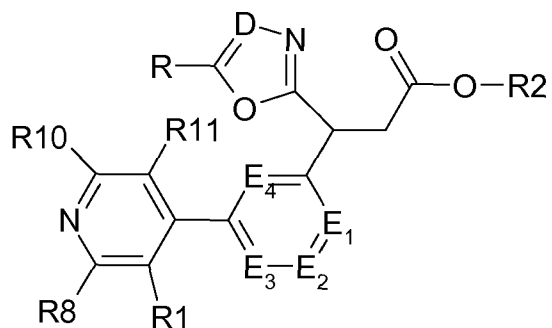
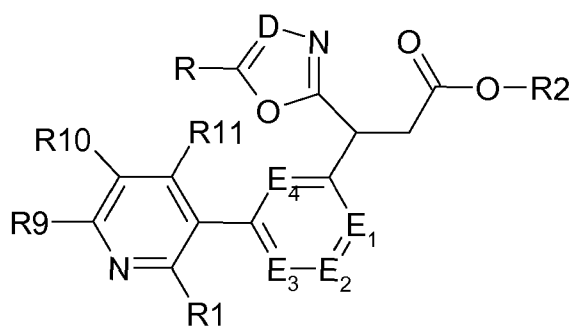
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I-15



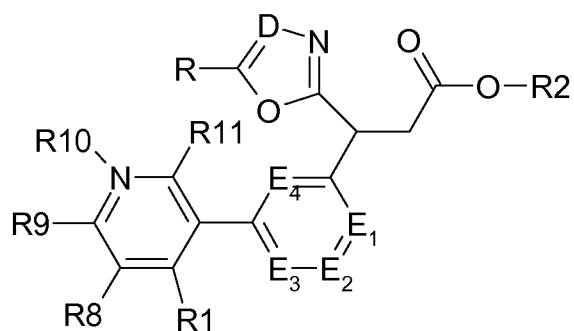
I-16



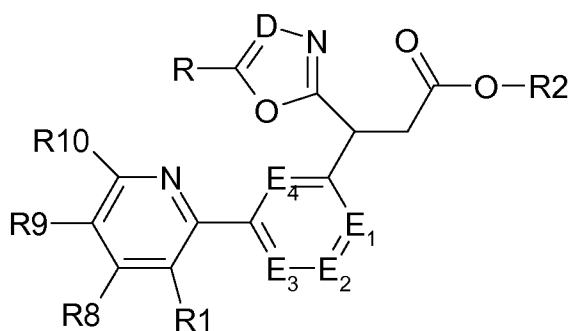
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I-17

I-18



I-19



I-20.

Another embodiment of the invention are compounds of formula I-1, I-2, I-3, I-4, I-5, I-11, I-12, I-13, I-14 or I-15, the groups D, R, R1, R2, R4, R5, R6, R7, R8, R9, R10, R11, E₁, E₂, E₃, E₄, defined as in the compounds of formula I in general or in any embodiment specified above or below,

wherein

G₁ is -C(R8)=;

G₂ is -C(R9)=;

G₃ is -C(R10)=;

G₄ -C(R11)=.

Another embodiment of the invention are compounds of formula I-1, I-2, I-3, I-4, I-5, I-11, I-12, I-13, I-14 or I-15, the groups D, R, R1, R2, R4, R5, R6, R7, R8, R9, R10, R11, E₁, E₂, E₃, E₄, defined as in the compounds of formula I in general or in any embodiment specified above or below, wherein

G₁ is N;

G₂ is -C(R9)=;

G₃ is -C(R10)=;

G₄ -C(R11)=.

Another embodiment of the invention are compounds of formula I-1, I-2, I-3, I-4, I-5, I-11, I-12, I-13, I-14 or I-15, the groups D, R, R1, R2, R4, R5, R6, R7, R8, R9, R10, R11, E₁, E₂, E₃, E₄, defined as in the compounds of formula I in general or in any embodiment specified above or below, wherein

- 5 G₁ is—C(R8)=;
G₂ is N;
G₃ is—C(R10)=;
G₄ is —C(R11)=.

- 10 Another embodiment of the invention are compounds of formula I-1, I-2, I-3, I-4, I-5, I-11, I-12, I-13, I-14 or I-15, the groups D, R, R1, R2, R4, R5, R6, R7, R8, R9, R10, R11, E₁, E₂, E₃, E₄, defined as in the compounds of formula I in general or in any embodiment specified above or below, wherein

- G₁ is—C(R8)=;
15 G₂ is—C(R9)=;
G₃ is N;
G₄ —C(R11)=.

Another embodiment of the invention are compounds of formula I-1, I-2, I-3, I-4, I-5, I-11, I-12, I-13, I-14 or I-15, the groups D, R, R1, R2, R4, R5, R6, R7, R8, R9, R10, R11, E₁, E₂, E₃, E₄, defined as in the compounds of formula I in general or in any embodiment specified above or below, wherein

- 20 G₁ is—C(R8)=;
G₂ is—C(R9)=;
25 G₃ is—C(R10)=;
G₄ is N.

- Another embodiment of the invention are compounds of formula I-6, I-7, I-8, I-9, I-10, I-16, I-17, I-18, I-19 or I-20, the groups D, R, R1, R2, R4, R5, R6, R7, R8, R9, R10, R11, G₁, G₂, G₃ and G₄, defined as in the compounds of formula I in general or in any embodiment specified above or below, wherein
- 30 E₁ is —C(R4)=;

E₂ is -C(R5)=;

E₃ is -C(R6)=;

E₄ is -C(R7)=.

- 5 Another embodiment of the invention are compounds of formula I-6, I-7, I-8, I-9, I-10, I-16, I-17, I-18, I-19 or I-20, the groups D, R, R1, R2, R4, R5, R6, R7, R8, R9, R10, R11, G₁, G₂, G₃ and G₄, defined as in the compounds of formula I in general or in any embodiment specified above or below, wherein

E₁ is N;

10 E₂ is -C(R5)=;

E₃ is -C(R6)=;

E₄ is -C(R7)=.

- Another embodiment of the invention are compounds of formula I-6, I-7, I-8, I-9, I-10,
15 I-16, I-17, I-18, I-19 or I-20, the groups D, R, R1, R2, R4, R5, R6, R7, R8, R9, R10, R11, G₁, G₂, G₃ and G₄, defined as in the compounds of formula I in general or in any embodiment specified above or below, wherein

E₁ is -C(R4)=;

E₂ is N;

20 E₃ is -C(R6)=;

E₄ is -C(R7)=.

- Another embodiment of the invention are compounds of formula I-6, I-7, I-8, I-9, I-10,
I-16, I-17, I-18, I-19 or I-20, the groups D, R, R1, R2, R4, R5, R6, R7, R8, R9, R10,
25 R11, G₁, G₂, G₃ and G₄, defined as in the compounds of formula I in general or in any embodiment specified above or below, wherein

E₁ or -C(R4)=;

E₂ is -C(R5)=;

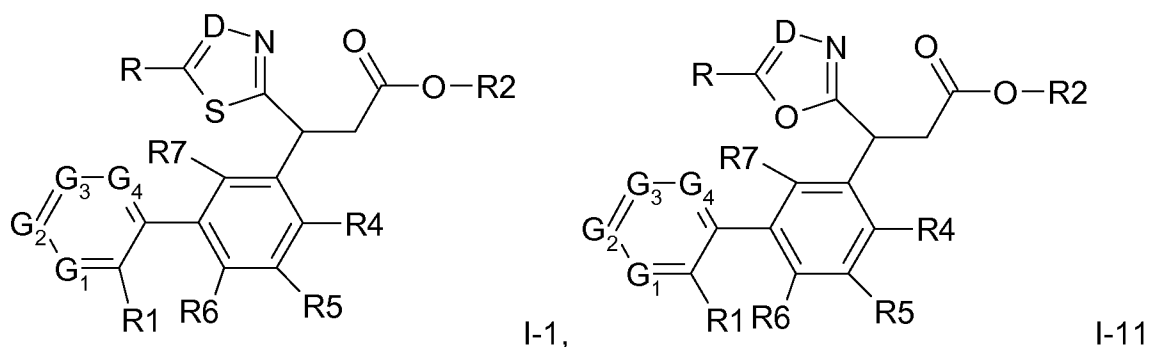
E₃ is N;

30 E₄ is -C(R7)=.

Another embodiment of the invention are compounds of formula I-6, I-7, I-8, I-9, I-10, I-16, I-17, I-18, I-19 or I-20, the groups D, R, R₁, R₂, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁, G₁, G₂, G₃ and G₄, defined as in the compounds of formula I in general or in any embodiment specified above or below, wherein

- 5 E₁ is -C(R₄)=;
 E₂ is -C(R₅)=;
 E₃ is -C(R₆)=;
 E₄ is N.

- 10 Another embodiment of the invention are compounds of formula I-1 or I-11



wherein the groups D, R, R₁, R₂, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ are defined as in the compounds of formula I in general or in any embodiment specified above or below, wherein

- 15 G₁ is -C(R₈)=;
 G₂ is -C(R₉)=;
 G₃ is -C(R₁₀)=;
 G₄ is -C(R₁₁)=.

- 20 One embodiment of the invention are compounds of formula I wherein
 R is Hydrogen, methyl or ethyl.

One embodiment of the invention are compounds of formula I wherein

- R₁ is H, F, Cl, CF₃, CN, CO-H, CO-(C₁-C₂)-alkyl, CO-NH₂, CO-NHR₂₁, CO-N((C₁-
 25 C₂)-alkyl)₂, wherein R₂₁ is H or (C₁-C₂)-alkyl.

One embodiment of the invention are compounds of formula I wherein

R1 is H, F, Cl, CF₃, CN, CO-H, CO-methyl, CO-NH₂, CO-N(CH₃)₂.

One embodiment of the invention are compounds of formula I wherein
R2 is Hydrogen or (C₁-C₆-)-alkyl.

- 5 One embodiment of the invention are compounds of formula I wherein
R2 is Hydrogen.

One embodiment of the invention are compounds of formula I wherein
R4 is H or O-methyl.

- 10 One embodiment of the invention are compounds of formula I wherein
R5 is H F, Cl, CF₃, (C₁-C₆-)-alkyl, cyclopropyl.

One embodiment of the invention are compounds of formula I wherein
R8 is H, F, Cl, (C₁-C₆-)-alkyl, O-(C₁-C₆-)-alkyl, CF₃ or OCF₃.

- 15 One embodiment of the invention are compounds of formula I wherein
R8 is H, F, Cl, methyl, O-methyl, CF₃ or OCF₃.

One embodiment of the invention are compounds of formula I wherein
R8 is H, F, Cl, methyl or O-methyl.

- 20 One embodiment of the invention are compounds of formula I wherein
R9 is H, F, Cl, OH, O-propyl, CH₂OH, CO-NH₂, methyl, O-methyl, CF₃ or OCF₃;

One embodiment of the invention are compounds of formula I wherein
R9 is H, F, Cl, OH, O-propyl, CH₂OH, CO-NH₂, methyl, O-methyl.

- 25 One embodiment of the invention are compounds of formula I wherein
R10 is H, F, Cl, OH, i-propyl, t-butyl, CH₂OH, CO-O-methyl, SO₂-methyl, CN, methyl,
O-methyl, CF₃ or OCF₃.

One embodiment of the invention are compounds of formula I wherein
R10 is H, F, Cl, OH, methyl, i-propyl, t-butyl, CH₂OH, CO-O-methyl, O-methyl.

30

One embodiment of the invention are compounds of formula I wherein

R11 is H, F, Cl, OH, O-methyl, O-i-propyl, CH₂OH, CO-methyl, CO-N(methyl)₂, CO-pyrrolidin, CO-O-methyl, CN, methyl or OCF₃;

One embodiment of the invention are compounds of formula I wherein

R11 is H, F, Cl, OH, methyl, O-i-propyl, CH₂OH, CO-methyl, CO-N(methyl)₂, CO-O-methyl, O-methyl.

One embodiment of the invention are compounds of formula I wherein

10 G₃ and G₄ are –C(R10)= and –C(R11)=, wherein R10 and R11 form a 5 or 6 membered saturated carbocycle or heterocycle with one or two oxygen atoms; which is optionally mono- or disubstituted by halogen and or (C₁-C₃)-alkyl.

One embodiment of the invention are compounds of formula I wherein

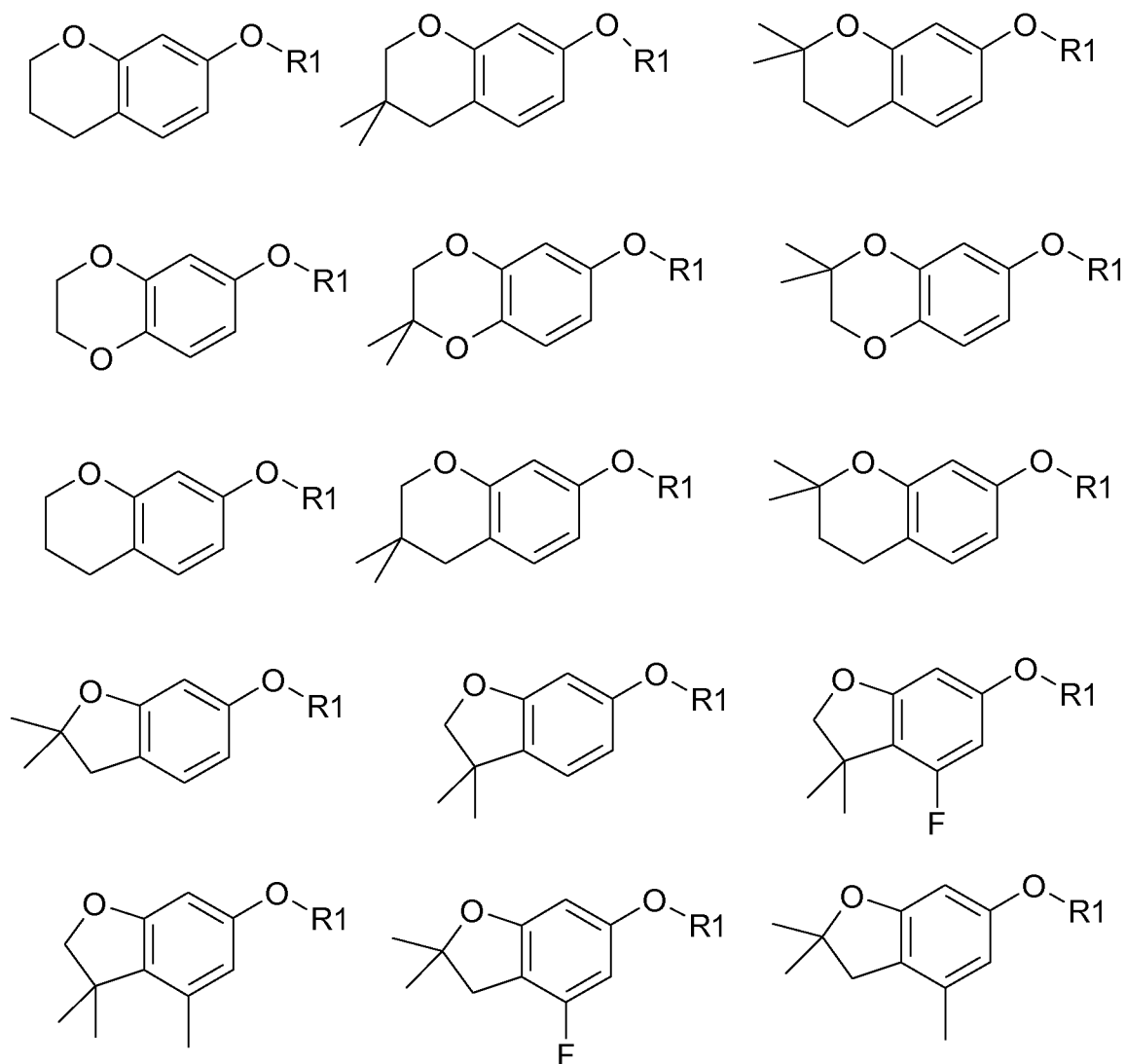
15 G₁ is –C(R8)=;

G₂ is –C(R9)=;

G₃ is –C(R10)=;

G₄ is –C(R11)=;

20 wherein R10 and R11 form a 5 or 6 membered saturated carbocycle or heterocycle with one or two oxygen atoms; which is optionally mono- or disubstituted by (C₁-C₃)-alkyl forming a bicyclic ringstructure selected from



One embodiment of the invention are compounds of formula I wherein

5 G_1 is $-C(R_8)=$;

G_2 is $-C(R_9)=$;

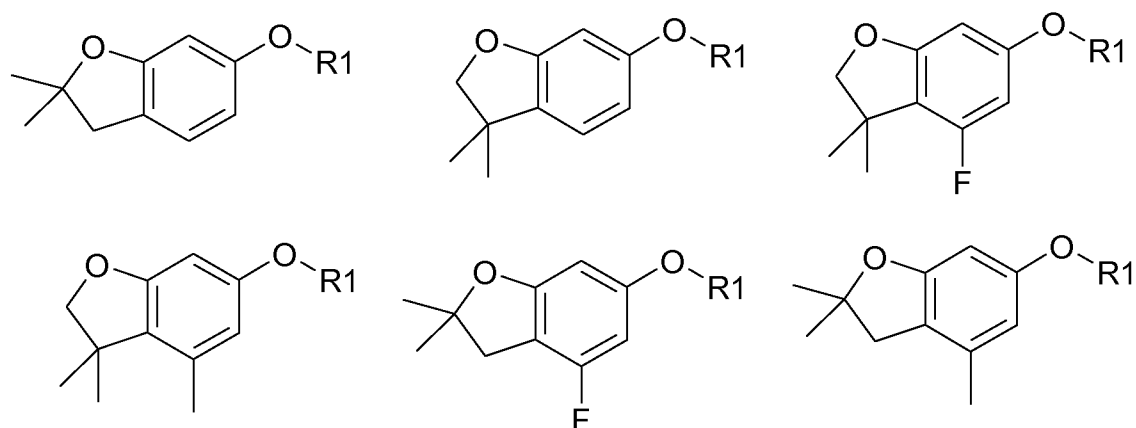
G_3 is $-C(R_{10})=$;

G_4 is $-C(R_{11})=$;

wherein R_{10} and R_{11} form a 5 membered saturated carbo- or heterocycle with one

10 or two oxygen atoms which is optionally mono- or disubstituted by (C_1-C_3) -alkyl

forming a bicyclic ringstructure selected from



One embodiment of the invention are compounds of formula I wherein

G_1 is $-\text{C}(\text{R}_8)=$;

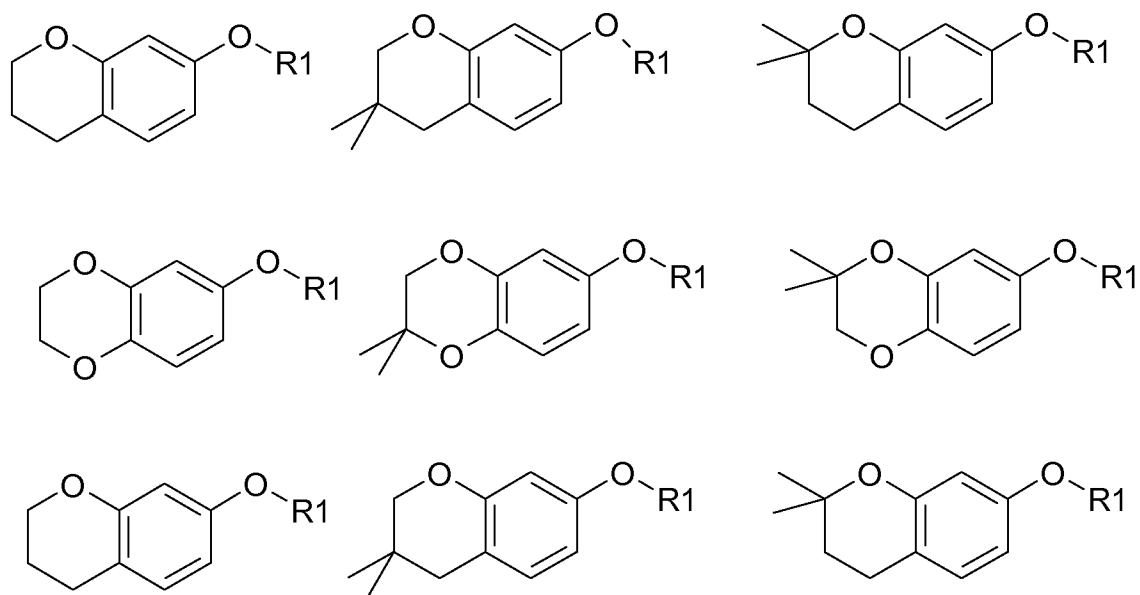
5 G_2 is $-\text{C}(\text{R}_9)=$;

G_3 is $-\text{C}(\text{R}_{10})=$;

G_4 is $-\text{C}(\text{R}_{11})=$;

wherein R_{10} and R_{11} form a 6 membered saturated carbo- or heterocycle with one or two oxygen atoms which is optionally mono- or disubstituted by $(\text{C}_1\text{-C}_3)$ -alkyl

10 forming a bicyclic ringstructure selected from



Another embodiment of the invention are compounds of formula I wherein

G_1 is $-C(R_8)=$;

G_2 is $-C(R_9)=$;

5 G_3 and G_4 are $-C((\text{methyl})_2)-CH_2-O-$;

R_8 is H;

R_9 is H.

Another embodiment of the invention are compounds of formula I wherein

10 X is S;

D is $-C(R_3)=$;

R is Hydrogen or methyl or ethyl;

R_3 is H, methyl or ethyl.

15 Another embodiment of the invention are compounds of formula I wherein

X is O;

D is $-C(R_3)=$;

R is Hydrogen or methyl or ethyl;

R_3 is H, methyl or ethyl.

20

Another embodiment of the invention are compounds of formula I wherein

X is O;

D is N;

R is Hydrogen or methyl or ethyl;

25 R_3 is H, methyl or ethyl.

Alkyl groups, i.e. saturated hydrocarbon residues, can be linear (straight-chain) or branched. This also applies if these groups are substituted or are part of another group, for example an alkyl-O- group (alkyloxy group, alkoxy group) or an HO-

30 substituted alkyl group (hydroxyalkyl group). Depending on the respective definition, the number of carbon atoms in an alkyl group can be 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10, or 1, 2, 3, 4, 5, 6, 7 or 8, or 1, 2, 3, 4, 5 or 6, or 1, 2, 3 or 4, or 1, 2 or 3, or 1 or 2, or 1,

for example. In one embodiment of the invention, a (C₁-C₁₀)-alkyl group present in the compounds of the formula I is a (C₁-C₈)-alkyl group, in another embodiment a (C₁-C₆)-alkyl group, in another embodiment a (C₁-C₄)-alkyl group, in another embodiment a (C₁-C₃)-alkyl group, in another embodiment a (C₁-C₂)-alkyl group, in another embodiment a (C₂-C₃)-alkyl group, in another embodiment a methyl group. In one embodiment of the invention, a (C₁-C₈)-alkyl group present in any position of the compounds of the formula I is a (C₁-C₆)-alkyl group, in another embodiment a (C₁-C₄)-alkyl group, in another embodiment a (C₁-C₃)-alkyl group, in another embodiment a (C₁-C₂)-alkyl group, in another embodiment a (C₂-C₃)-alkyl group, in another embodiment a methyl group, where any (C₁-C₈)-alkyl group present in the compounds of the formula I can independently of each other (C₁-C₈)-alkyl group be a group of any of these embodiments. In one embodiment of the invention, a (C₁-C₆)-alkyl group present in any position of the compounds of the formula I is a (C₁-C₄)-alkyl group, in another embodiment a (C₁-C₃)-alkyl group, in another embodiment a (C₁-C₂)-alkyl group, in another embodiment a (C₂-C₃)-alkyl group, in another embodiment a methyl group, where any (C₁-C₆)-alkyl group present in the compounds of the formula I can independently of each other (C₁-C₆)-alkyl group be a group of any of these embodiments. In one embodiment of the invention, a (C₁-C₄)-alkyl group present in any position of the compounds of the formula I is a (C₁-C₃)-alkyl group, in another embodiment a (C₁-C₂)-alkyl group, in another embodiment a (C₂-C₃)-alkyl group, in another embodiment a methyl group, where any (C₁-C₄)-alkyl group present in the compounds of the formula I can independently of each other (C₁-C₄)-alkyl group be a group of any of these embodiments. Examples of alkyl groups are methyl, ethyl, propyl groups including propyl (i.e. n-propyl) and isopropyl, butyl groups including butyl (i.e. n-butyl), sec-butyl, isobutyl and tert-butyl, pentyl groups including pentyl (i.e. n-pentyl), 1-methylbutyl, isopentyl, neopentyl and tert-pentyl, hexyl groups including hexyl (i.e. n-hexyl), 3,3-dimethylbutyl and isohexyl, heptyl groups including heptyl (i.e. n-heptyl), octyl groups including octyl (i.e. n-octyl), nonyl groups including nonyl (i.e. n-nonyl), and decyl groups including decyl (i.e. n-decyl). Examples of alkyl-O- groups are methoxy, ethoxy, propoxy (i.e. n-propoxy), isopropoxy, butoxy (i.e. n-butoxy), isobutoxy, tert-butoxy, pentoxy (i.e. n-pentoxy). Examples of alkyl-S(O)_m- are methylsulfanyl- (CH₃-S-), methanesulfinyl- (CH₃-S(O)-),

methanesulfonyl ($\text{CH}_3\text{-S(O)}_2\text{-}$), ethylsulfanyl- ($\text{CH}_3\text{-CH}_2\text{-S-}$), ethanesulfinyl- ($\text{CH}_3\text{-CH}_2\text{-S(O)-}$), ethanesulfonyl ($\text{CH}_3\text{-CH}_2\text{-S(O)}_2\text{-}$), 1-methylethylsulfanyl- ($(\text{CH}_3)_2\text{CH-S-}$), 1-methylethanesulfinyl- ($(\text{CH}_3)_2\text{CH-S(O)-}$), 1-methylethanesulfonyl ($(\text{CH}_3)_2\text{CH-S(O)}_2\text{-}$). In one embodiment of the invention the number m is chosen from 0 and 2, wherein all numbers m are independent of each other and can be identical or different. In another embodiment the number m in any of its occurrences is, independently of its meaning in other occurrences, 0. In another embodiment the number m in any of its occurrences is, independently of its meaning in other occurrences, 2.

A substituted alkyl group can be substituted in any positions, provided that the respective compound is sufficiently stable and is suitable as a pharmaceutical active compound. The prerequisite that a specific group and a compound of the formula I are sufficiently stable and suitable as a pharmaceutical active compound, applies in general with respect to the definitions of all groups in the compounds of the formula I. In one embodiment of the invention, an individual carbon atom in any alkyl group in the compounds of the formula I, as well as in other groups such as cycloalkyl groups and heterocyclic groups, for example, independently of any other carbon atom does not carry more than one substituent which is bonded via an oxygen atom, nitrogen atom or sulfur atom, such as HO- , $(\text{C}_1\text{-C}_4)\text{-alkyl-O-}$ or $(\text{C}_1\text{-C}_4)\text{-alkyl-S(O)}_m\text{-}$ substituents, for example. An alkyl group which is optionally substituted by one or more fluorine substituents can be unsubstituted, i.e. not carry fluorine substituents, or substituted, for example by one, two, three, four, five, six, seven, eight, nine, ten or eleven fluorine substituents, or by one, two, three, four, five, six or seven fluorine substituents, or by one, two, three, four or five fluorine substituents, or by one, two or three fluorine substituents, which can be located in any positions. For example, in a fluoro-substituted alkyl group one or more methyl groups can carry three fluorine substituents each and be present as trifluoromethyl groups, and/or one or more methylene groups (CH_2) can carry two fluorine substituents each and be present as difluoromethylene groups. The explanations with respect to the substitution of a group by fluorine also apply if the group additionally carries other substituents and/or is part of another group, for example of an alkyl-O- group. Examples of fluoro-

substituted alkyl groups are trifluoromethyl, 2-fluoroethyl, 1-fluoroethyl, 1,1-difluoroethyl, 2,2,2-trifluoroethyl, pentafluoroethyl, 3,3,3-trifluoropropyl, 2,2,3,3,3-pentafluoropropyl, 4,4,4-trifluorobutyl and heptafluoroisopropyl. Examples of fluoro-substituted alkyl-O- groups are trifluoromethoxy, 2,2,2-trifluoroethoxy, pentafluoroethoxy and 3,3,3-trifluoropropoxy. Examples of fluoro-substituted alkyl-S(O)_m- groups are trifluoromethylsulfanyl- (CF₃-S-), trifluoromethanesulfinyl- (CF₃-S(O)-) and trifluoromethanesulfonyl (CF₃-S(O)₂-).

- 10 The number of ring carbon atoms in a (C₃-C₇)-cycloalkyl group can be 3, 4, 5, 6 or 7. Examples of cycloalkyl are cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and cycloheptyl. As regards the optional substitution of cycloalkyl groups by one or more (C₁-C₄)-alkyl substituents, they be unsubstituted, i.e. not carry alkyl substituents, or substituted, for example by one, two, three or four, or by one or two, identical or
- 15 different (C₁-C₄)-alkyl substituents, for example by methyl groups, which substituents can be located in any positions. Examples of such alkyl-substituted cycloalkyl groups are 1-methylcyclopropyl, 2,2-dimethylcyclopropyl, 1-methylcyclopentyl, 2,3-dimethylcyclopentyl, 1-methylcyclohexyl, 4-methylcyclohexyl, 4-isopropylcyclohexyl, 4-tert-butylcyclohexyl and 3,3,5,5-tetramethylcyclohexyl. As regards the optional
- 20 substitution of cycloalkyl groups by one or more fluorine substituents, they can be unsubstituted, i.e. not carry fluorine substituents, or substituted, for example by one, two, three, four, five, six, seven, eight, nine, ten or eleven fluorine substituents, or by one, two, three, four, five or six fluorine substituents, or by one, two, three or four fluorine substituents, or by one or two fluorine substituents. The fluorine substituents
- 25 can be located in any positions of the cycloalkyl group and can also be located in an alkyl substituent on the cycloalkyl group. Examples of fluoro-substituted cycloalkyl groups are 1-fluorocyclopropyl, 2,2-difluorocyclopropyl, 3,3-difluorocyclobutyl, 1-fluorocyclohexyl, 4,4-difluorocyclohexyl and 3,3,4,4,5,5-hexafluorocyclohexyl. Cycloalkyl groups can also be substituted simultaneously by fluorine and alkyl.
- 30 Halogen is fluorine, chlorine, bromine or iodine. In one embodiment of the invention, halogen in any occurrence in the compounds of the formula I, independently of all

other occurrences, is fluorine, chlorine or bromine, in another embodiment fluorine or chlorine, in another embodiment fluorine.

An oxo substituent, i.e. an oxygen atom which is bonded via a double bond, when bonded to a carbon atom, replaces two hydrogen atoms on the carbon atom of the parent system to which it is bonded. Thus, if a CH₂ group is substituted by oxo, it becomes a carbonyl group (C(O), C=O). An oxo substituent cannot be present on a carbon atom in an aromatic ring.

The present invention comprises all stereoisomeric forms of the compounds of the formula I, for example all enantiomers and diastereomers including cis/trans isomers. The invention likewise comprises mixtures of two or more stereoisomeric forms, for example mixtures of enantiomers and/or diastereomers including cis/trans isomers, in all ratios. Asymmetric centers contained in the compounds of the formula I, for example in unsubstituted or substituted alkyl groups, can all independently of each other have the S configuration or the R configuration. The invention relates to enantiomers, both the levorotatory and the dextrorotatory antipode, in enantiomerically pure form and essentially enantiomerically pure form, for example with a molar ratio of the two enantiomers of 99 : 1 or greater, and in the form of racemates and in the form of mixtures of the two enantiomers in all ratios. The invention likewise relates to diastereomers in the form of pure and essentially pure diastereomers and in the form of mixtures of two or more diastereomers in all ratios. The invention also comprises all cis/trans isomers of the compounds of the formula I in pure form and essentially pure form, for example with a molar ratio of the cis/trans isomers of 99 : 1 or greater, and in the form of mixtures of the cis isomer and the trans isomer in all ratios. Cis/trans isomerism can occur in substituted rings. The preparation of individual stereoisomers, if desired, can be carried out by resolution of a mixture according to customary methods, for example, by chromatography or crystallization, or by use of stereochemically uniform starting compounds in the synthesis or by stereoselective reactions. Optionally, before a separation of stereoisomers a derivatization can be carried out. The separation of a mixture of stereoisomers can be carried out at the stage of the compound of the formula I or at

the stage of an intermediate in the course of the synthesis. The invention also comprises all tautomeric forms of the compounds of the formula I.

5 Physiologically acceptable salts, including pharmaceutically utilizable salts, of the compounds of the formula I generally comprise a nontoxic salt component. They can contain inorganic or organic salt components. Such salts can be formed, for example, from compounds of the formula I which contain an acidic group, for example a carboxylic acid group (hydroxycarbonyl group, HO-C(O)-), and nontoxic inorganic or organic bases. Suitable bases are, for example, alkali metal compounds or alkaline earth metal compounds, such as sodium hydroxide, potassium hydroxide, sodium carbonate or sodium hydrogencarbonate, or ammonia, organic amino compounds and quaternary ammonium hydroxides. Reactions of compounds of the formula I with bases for the preparation of the salts are in general carried out according to customary procedures in a solvent or diluent. Examples of salts of acidic groups thus are sodium, potassium, magnesium or calcium salts or ammonium salts which can also carry one or more organic groups on the nitrogen atom. Compounds of the formula I which contain a basic, i.e. protonatable, group, for example an amino group or a basic heterocycle, can be present in the form of their acid addition salts with physiologically acceptable acids, for example as salt with hydrogen chloride, hydrogen bromide, phosphoric acid, sulfuric acid, acetic acid, benzoic acid, methanesulfonic acid, p-toluenesulfonic acid, which in general can be prepared from the compounds of the formula I by reaction with an acid in a solvent or diluent according to customary procedures. If the compounds of the formula I simultaneously contain an acidic and a basic group in the molecule, the invention also includes internal salts (betaines, zwitterions) in addition to the salt forms mentioned. The present invention also comprises all salts of the compounds of the formula I which, because of low physiological tolerability, are not directly suitable for use as a pharmaceutical, but are suitable as intermediates for chemical reactions or for the preparation of physiologically acceptable salts, for example by means of anion exchange or cation exchange. The present invention also comprises all solvates of the compounds of the formula I and their salts, including physiologically acceptable solvates, such as hydrates, i.e. adducts with water, and adducts with alcohols like

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(C₁-C₄)-alkanols, as well as active metabolites of compounds of the formula I and prodrugs of the compounds of the formula I, i.e. compounds which in vitro may not necessarily exhibit pharmacological activity but which in vivo are converted into pharmacologically active compounds of the formula I, for example compounds which are converted by metabolic hydrolysis into a compound of the formula I, such as compounds in which a carboxylic acid group is present in esterified form or in the form of an amide.

A subject of the invention are all compounds of the formula I wherein any one or more structural elements such as groups, substituents and numbers are defined as in any of the specified embodiments or definitions of the elements or have one or more of the specific meanings which are mentioned herein as examples of elements, wherein all combinations of one or more specified embodiments and/or definitions and/or specific meanings of the elements are a subject of the present invention. Also with respect to all such compounds of the formula I, all their stereoisomeric forms and mixtures of stereoisomeric forms in any ratios, and their physiologically acceptable salts, and the physiologically acceptable solvates of any of them, are a subject of the present invention.

A subject of the invention also is a compound of the formula I which is chosen from any of the specific compounds of the formula I which are disclosed herein, or is any one of the specific compounds of the formula I which are disclosed herein, irrespective thereof whether they are disclosed as a free compound and/or as a specific salt, or a physiologically acceptable salt thereof, or a physiologically acceptable solvate of any of them, wherein the compound of the formula I is a subject of the invention in any of its stereoisomeric forms or a mixture of stereoisomeric forms in any ratio.

Another subject of the present invention are the novel starting compounds and intermediates occurring in the synthesis of the compounds of the formula I, in any of their stereoisomeric forms or a mixture of stereoisomeric forms in any ratio, and their salts, and solvates of any of them, and their use as synthetic intermediates or starting compounds. All general explanations, specifications of embodiments and definitions of numbers and groups given above with respect to the compounds of the formula I apply correspondingly to the said intermediates and starting compounds. A subject of the invention are in particular the novel specific starting compounds and intermediates described herein. Independently thereof whether they are described as a free compound and/or as a specific salt, they are a subject of the invention both in the form of the free compounds and in the form of their salts, and if a specific salt is described, additionally in the form of this specific salt.

The compounds of the formula I inhibit the protease cathepsin A as can be demonstrated in the pharmacological test described below and in other tests which are known to a person skilled in the art. The compounds of the formula I and their physiologically acceptable salts and solvates therefore are valuable pharmaceutical active compounds. The compounds of the formula I and their physiologically acceptable salts and solvates can be used for the treatment of cardiovascular diseases such as heart failure including systolic heart failure, diastolic heart failure, diabetic heart failure and heart failure with preserved ejection fraction, cardiomyopathy, myocardial infarction, left ventricular dysfunction including left ventricular dysfunction after myocardial infarction, cardiac hypertrophy, myocardial remodeling including myocardial remodeling after infarction or after cardiac surgery, valvular heart diseases, vascular hypertrophy, vascular remodeling including vascular stiffness, hypertension including pulmonary hypertension, portal hypertension and systolic hypertension, atherosclerosis, peripheral arterial occlusive disease (PAOD), restenosis, thrombosis and vascular permeability disorders, ischemia and/or reperfusion damage including ischemia and/or reperfusion damage of the heart and ischemia and/or reperfusion damage of the retina, inflammation and inflammatory diseases such as rheumatoid arthritis and osteoarthritis, renal diseases such as renal papillary necrosis and renal failure including renal failure after

ischemia/reperfusion, pulmonary diseases such as cystic fibrosis, chronic bronchitis, chronic obstructive pulmonary disease (COPD), asthma, acute respiratory distress syndrome (ARDS), respiratory tract infections and lung carcinoma, immunological diseases, diabetic complications including diabetic nephropathy and diabetic

5 cardiomyopathy, fibrotic diseases such as pulmonary fibrosis including idiopathic lung fibrosis, cardiac fibrosis, vascular fibrosis, perivascular fibrosis, renal fibrosis including renal tubulointerstitial fibrosis, fibrosing skin conditions including keloid formation, collagenosis and scleroderma, and liver fibrosis, liver diseases such as liver cirrhosis, pain such as neuropathic pain, diabetic pain and inflammatory pain,

10 macular degeneration, neurodegenerative diseases or psychiatric disorders, or for cardioprotection including cardioprotection after myocardial infarction and after cardiac surgery, or for renoprotection, for example. The compounds of the formula I and their physiologically acceptable salts and solvates can be used as a diuretic (stand-alone treatment or in combination with established diuretics). The compounds

15 of the formula I and their physiologically acceptable salts and solvates can also be used for treatment and/or prevention of atrial fibrillation. The treatment of diseases is to be understood as meaning both the therapy of existing pathological changes or malfunctions of the organism or of existing symptoms with the aim of relief, alleviation or cure, and the prophylaxis or prevention of pathological changes or malfunctions of

20 the organism or of symptoms in humans or animals which are susceptible thereto and are in need of such a prophylaxis or prevention, with the aim of a prevention or suppression of their occurrence or of an attenuation in the case of their occurrence. For example, in patients who on account of their disease history are susceptible to myocardial infarction, by means of the prophylactic or preventive medicinal treatment

25 the occurrence or re-occurrence of a myocardial infarction can be prevented or its extent and sequelae decreased, or in patients who are susceptible to attacks of asthma, by means of the prophylactic or preventive medicinal treatment such attacks can be prevented or their severity decreased. The treatment of diseases can occur both in acute cases and in chronic cases. The efficacy of the compounds of the

30 formula I can be demonstrated in the pharmacological test described below and in other tests which are known to a person skilled in the art.

The compounds of the formula I and their physiologically acceptable salts and solvates can therefore be used in animals, in particular in mammals and specifically in humans, as a pharmaceutical or medicament on their own, in mixtures with one another or in the form of pharmaceutical compositions. A subject of the present invention also are the compounds of the formula I and their physiologically acceptable salts and solvates for use as a pharmaceutical, as well as pharmaceutical compositions and medicaments which comprise an efficacious dose of at least one compound of the formula I and/or a physiologically acceptable salt thereof and/or solvate thereof as an active ingredient and a pharmaceutically acceptable carrier, i.e. one or more pharmaceutically innocuous, or nonhazardous, vehicles and/or excipients, and optionally one or more other pharmaceutical active compounds. A subject of the present invention furthermore are the compounds of the formula I and their physiologically acceptable salts and solvates for use in the treatment of the diseases mentioned above or below, including the treatment of any one of the mentioned diseases, for example the treatment of heart failure, myocardial infarction, cardiac hypertrophy, diabetic nephropathy, diabetic cardiomyopathy, cardiac fibrosis, or ischemia and/or reperfusion damage, or for cardioprotection, the use of the compounds of the formula I and their physiologically acceptable salts and solvates for the manufacture of a medicament for the treatment of the diseases mentioned above or below, including the treatment of any one of the mentioned diseases, for example the treatment of heart failure, myocardial infarction, cardiac hypertrophy, diabetic nephropathy, diabetic cardiomyopathy, cardiac fibrosis, or ischemia and/or reperfusion damage, or for cardioprotection, wherein the treatment of diseases comprises their therapy and prophylaxis as mentioned above, as well as their use for the manufacture of a medicament for the inhibition of cathepsin A. A subject of the invention also are methods for the treatment of the diseases mentioned above or below, including the treatment of any one of the mentioned diseases, for example the treatment of heart failure, myocardial infarction, cardiac hypertrophy, diabetic nephropathy, diabetic cardiomyopathy, cardiac fibrosis, or ischemia and/or reperfusion damage, or for cardioprotection, which comprise administering an efficacious amount of at least one compound of the formula I and/or a physiologically acceptable salt thereof and/or solvate thereof to a human or an animal which is in

need thereof. The compounds of the formula I and pharmaceutical compositions and medicaments comprising them can be administered enterally, for example by oral, sublingual or rectal administration, parenterally, for example by intravenous, intramuscular, subcutaneous or intraperitoneal injection or infusion, or by another
5 type of administration such as topical, percutaneous, transdermal, intra-articular or intraocular administration.

The compounds of the formula I and their physiologically acceptable salts and solvates can also be used in combination with other pharmaceutical active
10 compounds, wherein in such a combination use the compounds of the formula I and/or their physiologically acceptable salts and/or solvates and one or more other pharmaceutical active compounds can be present in one and the same pharmaceutical composition or in two or more pharmaceutical compositions for separate, simultaneous or sequential administration. Examples of such other
15 pharmaceutical active compounds are diuretics, aquaretics, angiotensin converting enzyme (ACE) inhibitors, angiotensin receptor blockers, renin inhibitors, beta blockers, digoxin, aldosterone antagonists, NO donors, nitrates, hydralazines, ionotropes, vasopressin receptor antagonists, soluble guanylate cyclase activators, statins, peroxisome proliferator-activated receptor-alpha (PPAR- α) activators,
20 peroxisome proliferator-activated receptor-gamma (PPAR- γ) activators, rosiglitazone, pioglitazone, metformin, sulfonylureas, glucagon-like peptide 1 (GLP-1) agonists, dipeptidyl peptidase IV (DPPIV) inhibitors, insulins, anti-arrhythmics, endothelin receptor antagonists, calcium antagonists, phosphodiesterase inhibitors, phosphodiesterase type 5 (PDE5) inhibitors, factor II/factor IIa inhibitors, factor
25 IX/factor IXa inhibitors, factor X/factor Xa inhibitors, factor XIII/factor XIIIa inhibitors, heparins, glycoprotein IIb/IIIa antagonists, P2Y₁₂ receptor antagonists, clopidogrel, coumarins, cyclooxygenase inhibitors, acetylsalicylic acid, RAF kinase inhibitors and p38 mitogen-activated protein kinase inhibitors. A subject of the present invention also is the said combination use of any one or more of the compounds of the formula
30 I disclosed herein and their physiologically acceptable salts and solvates, with any one or more, for example one or two, of the mentioned other pharmaceutical active compounds.

The pharmaceutical compositions and medicaments according to the invention normally contain from about 0.5 to about 90 percent by weight of compounds of the formula I and/or physiologically acceptable salts and/or solvates thereof, and an amount of active ingredient of the formula I and/or its physiologically acceptable salt and/or solvate which in general is from about 0.2 mg to about 1.5 g, particularly from about 0.2 mg to about 1 g, more particularly from about 0.5 mg to about 0.5 g, for example from about 1 mg to about 0.3 g, per unit dose. Depending on the kind of the pharmaceutical composition and other particulars of the specific case, the amount may deviate from the indicated ones. The production of the pharmaceutical compositions and medicaments can be carried out in a manner known per se. For this, the compounds of the formula I and/or their physiologically acceptable salts and/or solvates are mixed together with one or more solid or liquid vehicles and/or excipients, if desired also in combination with one or more other pharmaceutical active compounds such as those mentioned above, and brought into a suitable form for dosage and administration, which can then be used in human medicine or veterinary medicine.

As vehicles, which may also be looked upon as diluents or bulking agents, and excipients suitable organic and inorganic substances can be used which do not react in an undesired manner with the compounds of the formula I. As examples of types of excipients, or additives, which can be contained in the pharmaceutical compositions and medicaments, lubricants, preservatives, thickeners, stabilizers, disintegrants, wetting agents, agents for achieving a depot effect, emulsifiers, salts, for example for influencing the osmotic pressure, buffer substances, colorants, flavorings and aromatic substances may be mentioned. Examples of vehicles and excipients are water, vegetable oils, waxes, alcohols such as ethanol, isopropanol, 1,2-propanediol, benzyl alcohols, glycerol, polyols, polyethylene glycols or polypropylene glycols, glycerol triacetate, polyvinylpyrrolidone, gelatin, cellulose, carbohydrates such as lactose or starch like corn starch, sodium chloride, stearic acid and its salts such as magnesium stearate, talc, lanolin, petroleum jelly, or mixtures thereof, for example saline or mixtures of water with one or more organic

solvents such as mixtures of water with alcohols. For oral and rectal use, pharmaceutical forms such as, for example, tablets, film-coated tablets, sugar-coated tablets, granules, hard and soft gelatin capsules, suppositories, solutions, including oily, alcoholic or aqueous solutions, syrups, juices or drops, furthermore suspensions or emulsions, can be used. For parenteral use, for example by injection or infusion, pharmaceutical forms such as solutions, for example aqueous solutions, can be used. For topical use, pharmaceutical forms such as ointments, creams, pastes, lotions, gels, sprays, foams, aerosols, solutions or powders can be used. Further suitable pharmaceutical forms are, for example, implants and patches and forms adapted to inhalation. The compounds of the formula I and their physiologically acceptable salts can also be lyophilized and the obtained lyophilizates used, for example, for the production of injectable compositions. In particular for topical application, also liposomal compositions are suitable. The pharmaceutical compositions and medicaments can also contain one or more other active ingredients and/or, for example, one or more vitamins.

As usual, the dosage of the compounds of the formula I depends on the circumstances of the specific case and is adjusted by the physician according to the customary rules and procedures. It depends, for example, on the compound of the formula I administered and its potency and duration of action, on the nature and severity of the individual syndrome, on the sex, age, weight and the individual responsiveness of the human or animal to be treated, on whether the treatment is acute or chronic or prophylactic, or on whether further pharmaceutical active compounds are administered in addition to a compound of the formula I. Normally, in the case of administration to an adult weighing about 75 kg, a dose from about 0.1 mg to about 100 mg per kg per day, in particular from about 1 mg to about 20 mg per kg per day, for example from about 1 mg to about 10 mg per kg per day (in each case in mg per kg of body weight), is administered. The daily dose can be administered in the form of a single dose or divided into a number of individual doses, for example two, three or four individual doses. The administration can also be carried out continuously, for example by continuous injection or infusion.

Depending on the individual behavior in a specific case, it may be necessary to deviate upward or downward from the indicated dosages.

Besides as a pharmaceutical active compound in human medicine and veterinary medicine, the compounds of the formula I can also be employed as an aid in biochemical investigations or as a scientific tool or for diagnostic purposes, for example in in-vitro diagnoses of biological samples, if an inhibition of cathepsin A is intended. The compounds of the formula I and their salts can also be used as intermediates, for example for the preparation of further pharmaceutical active substances.

The following examples illustrate the invention.

Abbreviations

ACN	acetonitrile
DCM	dichloromethane
DMF	N,N-dimethylformamide
DMSO	dimethyl sulfoxide
EA	ethyl acetate
EDIA	N-ethyl-diisopropylamine
FA	formic acid
MOH	methanol
NEM	N-ethyl-morpholine
TFA	trifluoroacetic acid
THF	tetrahydrofuran
TOTU	O-(cyano(ethoxycarbonyl)methyleneamino)-N,N,N',N'-tetramethyluronium tetrafluoroborate

When example compounds containing a basic group were purified by preparative high pressure liquid chromatography (HPLC) on reversed phase (RP) column material and, as customary, the eluent was a gradient mixture of water and

acetonitrile containing trifluoroacetic acid, they were in part obtained in the form of their acid addition salts with trifluoroacetic acid, depending on the details of the work-up such as evaporation or lyophilization conditions. In the names of the example compounds and the structural formulae such contained trifluoroacetic acid is not specified. Likewise are other acid components of example compounds obtained in the form of an acid addition salt in general not specified in the name and the formula.

The prepared compounds were in general characterized by spectroscopic data and chromatographic data, in particular mass spectra (MS) and HPLC retention times (Rt; in min) which were obtained by combined analytical HPLC/MS characterization (LC/MS), and/or nuclear magnetic resonance (NMR) spectra. Unless specified otherwise, ^1H -NMR spectra were recorded at 500 MHz in D_6 -DMSO as solvent at 298 K. In the NMR characterization, the chemical shift δ (in ppm), the number of hydrogen atoms (H), and the multiplicity (s: singlet, d: doublet, dd: doublet of doublets, t: triplet, q: quartet, m: multiplet) of the peaks as determined from the graphically depicted spectra are given. In the MS characterization, in general the mass number (m/z) of the peak of the molecular ion [M], for example $[\text{M}^+]$, or of a related ion such as the ion $[\text{M}+1]$, for example $[(\text{M}+1)^+]$, i.e. the protonated molecular ion $[(\text{M}+\text{H})^+]$, or the ion $[\text{M}-1]$, for example $[(\text{M}-1)^-]$, i.e. the deprotonated molecular ion $[(\text{M}-\text{H})^-]$, which was formed depending on the ionization method used, is given. Generally, the ionization method was electrospray ionization (ES). The particulars of the LC/MS methods used are as follows.

Method LC1

Column: Waters UPLC BEH C18, 50 x 2.1 mm, 1.7 μm ; flow: 0.9 ml/min; 55 $^\circ\text{C}$; eluent A: water + 0.05 % FA; eluent B: ACN + 0.035 % FA; gradient: from 98 % A + 2 % B to 5 % A + 95 % B within 2.0 min, then 5 % A + 95 % B for 0.6 min, then to 95 % A + 5 % B within 0.1 min, then 95 % A + 5 % B for 0.3 min; MS ionization method: ES^+

Method LC2

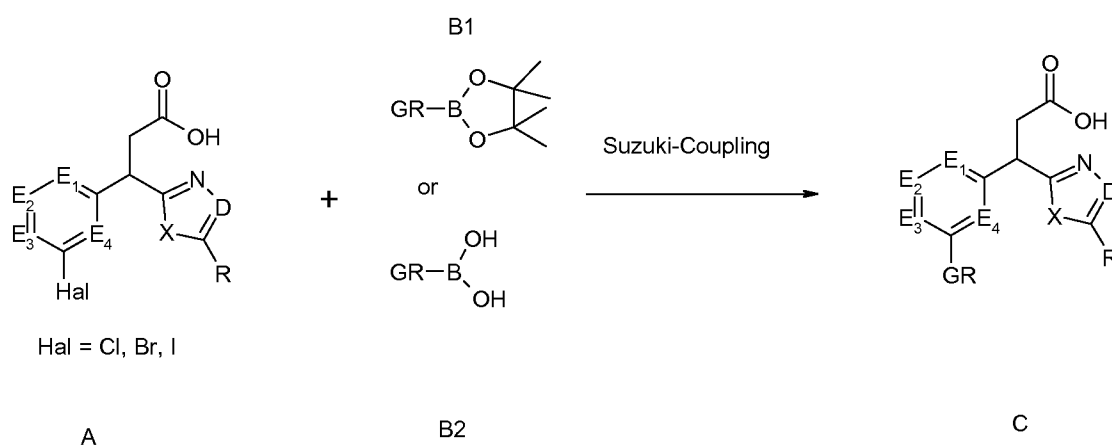
Column: Waters XBridge C18, 50 x 4.6 mm, 2.5 μ m; flow: 1.7 ml/min; 40 °C; eluent A: water + 0.05 % TFA; eluent B: ACN + 0.05 % TFA; gradient: 95 % A + 5 % B for 0.2 min, then to 5 % A + 95 % B within 2.2 min, then 5 % A + 95 % B for 0.8 min, then to 95 % A + 5 % B within 0.1 min, then 95 % A + 5 % B for 0.7 min; MS ionization method: ES⁺

Method LC3

Column: Waters XBridge C18, 50 x 4.6 mm, 2.5 μ m; flow: 1.7 ml/min; 40 °C; eluent A: water + 0.1 % TFA; eluent B: ACN + 0.1 % TFA; gradient: 97 % A + 3 % B for 0.2 min, then to 40 % A + 60 % B within 3.5 min, then to 2 % A + 98 % B within 0.5 min, then 2 % A + 98 % B for 0.5 min, then to 97 % A + 3 % B within 0.2 min, then 97 % A + 3 % B for 0.3 min; MS ionization method: ES⁺

Experimental

In general the compounds of formula I are synthesized according to one of the general schemes below:



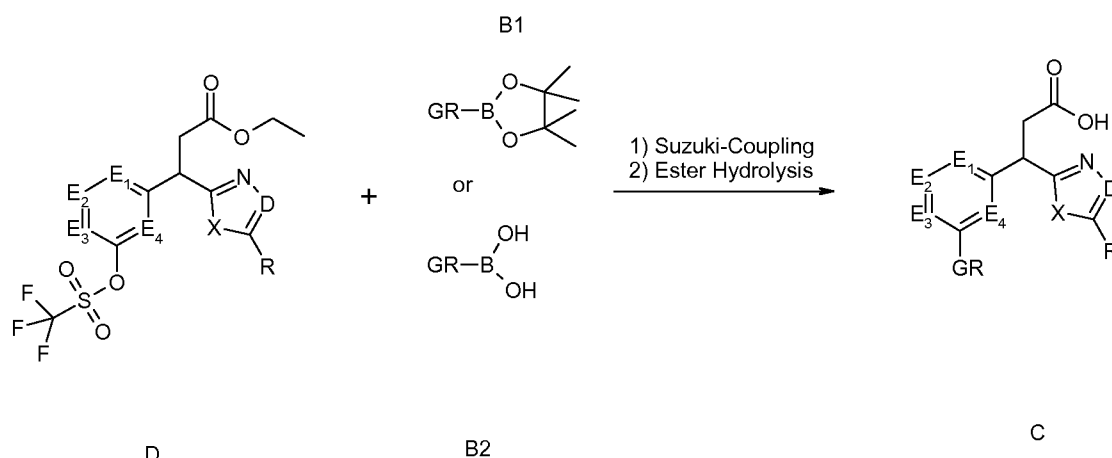
Scheme A and General Procedure A:

0,3 mmol boronic acid is placed in a reaction tube, 0,2mmol of the intermediate A dissolved in 2ml of DMF and 0,8mmol Cs₂CO₃ dissolved in 1ml water is added and

the reaction tube purged with Ar. After the addition of 0,02 mmol mmol Pd[PPh₃]Cl₂ is reaction mixture is heated overnight at 95°C

The reaction mixture is adjusted to pH = 5 with 2N HCl, 15ml EtOAc and 5ml 5% NaCl solution are added, the organic phase is separated and dried over Na₂SO₄.

- 5 The solvent is removed in vacuo and the crude product subjected to HPLC chromatography. Yields are in the range from 10 to 95%.



Scheme B and General Procedure B

0,3 mmol boronic acid is placed in a reaction tube, 0,2mmol of the intermediate D dissolved in 2ml of DMF and 0,8mmol Cs₂CO₃ dissolved in 1ml water is added and the reaction tube purged with Ar. After the addition of 0,02 mmol mmol Pd[PPh₃]Cl₂ is reaction mixture is heated overnight at 95°C

The reaction mixture is adjusted to pH = 5 with 2N HCl, 15ml EtOAc and 5ml 5% NaCl solution are added, the organic phase is separated and dried over Na₂SO₄.

The solvent is removed in vacuo, the crude product is dissolved in 2ml EtOH and 0,5 ml 1N NaOH is added. After stirring overnight at RT the solvent is removed and the isolated material is subjected to HPLC chromatography. Yields are in the range from 10 to 95%

The Suzuki reactions described above can be carried out by all procedures well known to a person skilled in the art and described for example in M. Mora et. al. Current Organic Chemistry 2012, 1128-1150

Solvents other than DMF like isopropanol, toluene, dioxane, THF, acetonitrile, water or any combinations of them might be used, the reaction is also possible by using ionic liquids. Any combination of a palladium salt and ligand or any preformed palladium catalyst system can be applied for these reactions. Polymer-bound Pd-catalysts or polymer-bound ligands may also be used. As base one might use Na_2CO_3 , K_2CO_3 , AgCO_3 or CsF instead of Cs_2CO_3 . The reaction temperatures are in the range from 40°C to 150°C and may be reached by thermal heating or by using a microwave reactor for a period from 5 minutes to 48 hours.

In scheme B one can isolate the product of the Suzuki coupling from the first step prior to the formation of the final product by ester hydrolysis or one can run both reactions in a one pot procedure without isolating the ester material. Instead of the ethyl ester in scheme B any other ester known to people skilled in the art may be used. Although instead of the triflate in scheme B one might use any other suitable leaving groups for Suzuki reactions.

Intermediates A are synthesized according to the general scheme C by addition of a metallated five-membered heterocycle F to an aromatic acid derivative E: E can be an aromatic acid ester, an aromatic acid chloride or a Weinreb amide of an aromatic carboxylic acid E. The resulting ketone G is subjected to a Wittig reaction and the Wittig condensation product H is reduced by applying Zn/HOAc and the ester residue is converted into the free acid A.

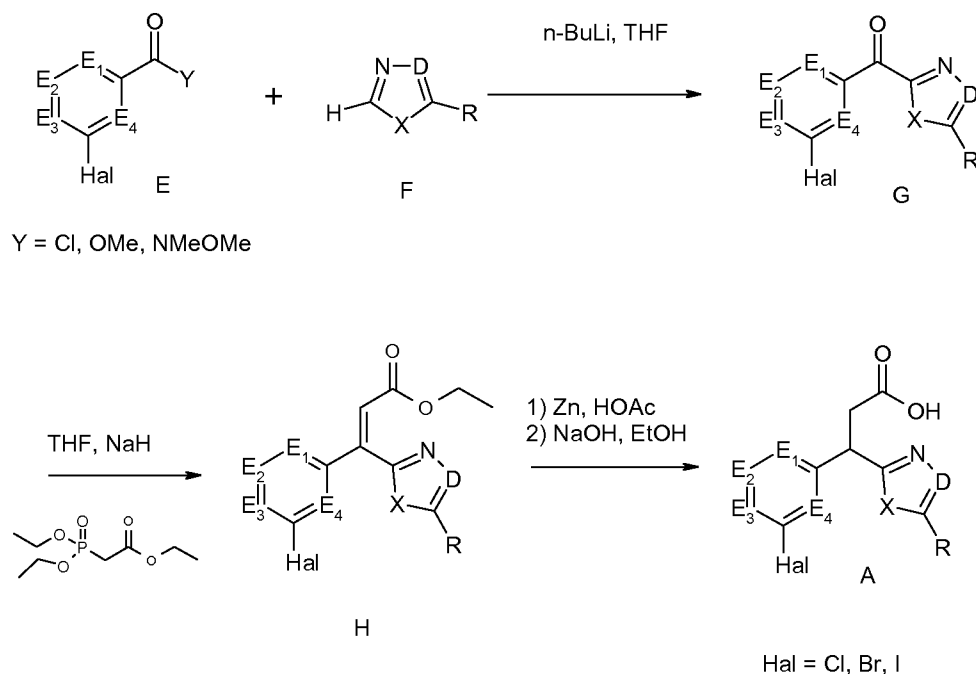
In the metallation step instead of n-BuLi tert-BuLi or MeLi a Grignard reagent like MeMgBr, iso-propyl-MgBr or iso-propyl-MgCl*LiCl may be used. To facilitate the addition to the intermediate E a transmetallation to another metal ion like copper or zinc might be necessary.

For the Wittig reaction one might use other strong bases instead of NaH like NaOtBu or KOtBu or phosphazene bases. And the reduction might be carried out using Zn in

aq. HCl using a concentration of aq. HCl ranging from 0,01M to 12M or in diluted H₂SO₄.

Intermediates E might be commercially available or might be prepared in one step from the commercially available carboxylic acids

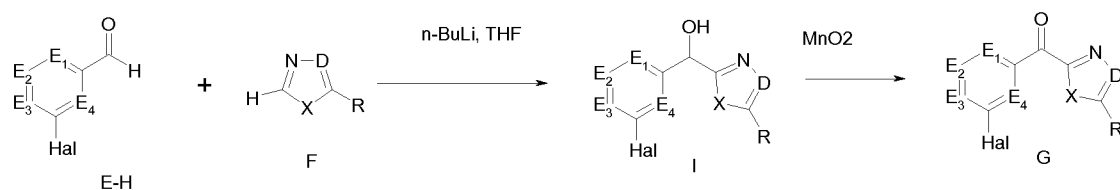
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Scheme C

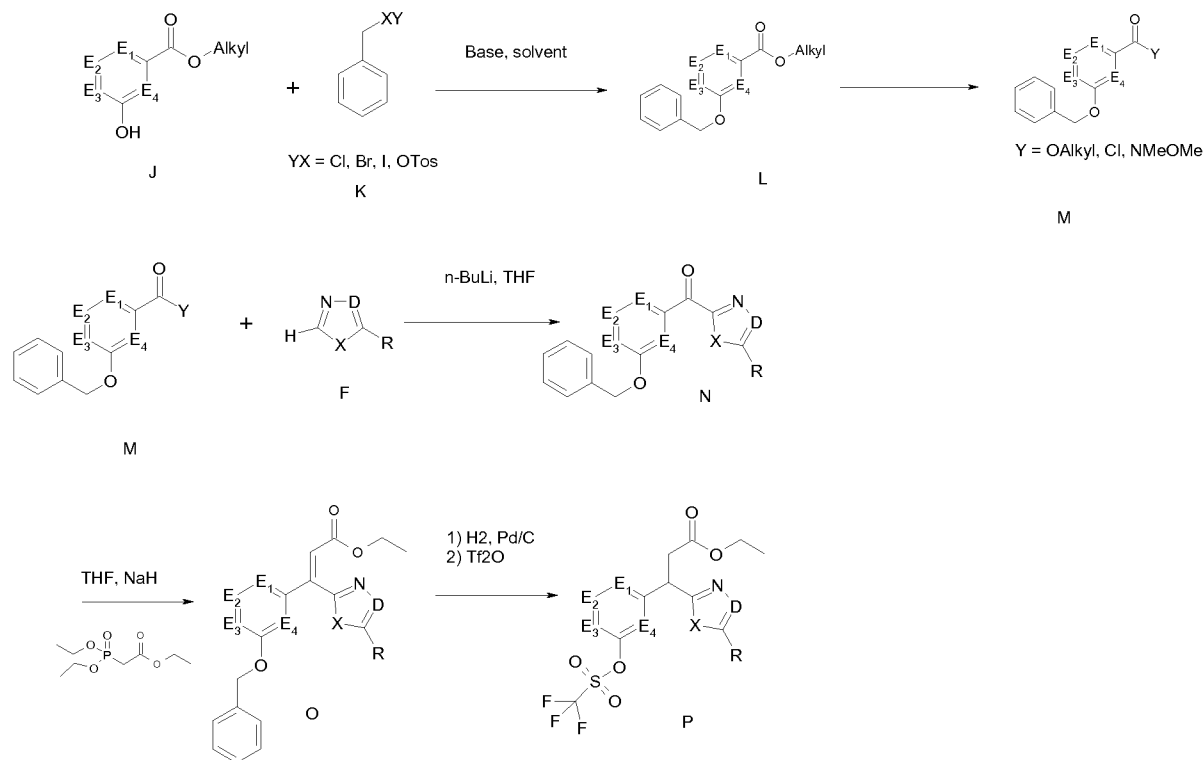
- 10 Under certain circumstances the group Y might represent a hydrogen atom forming the aromatic aldehyde intermediate E-H. After the addition of the metallated heterocycle F the corresponding alcohol I can be oxidized to the desired intermediate G by using MnO₂ or other oxidants like KMnO₄, CrO₃, and H₂O₂.

15



Scheme D

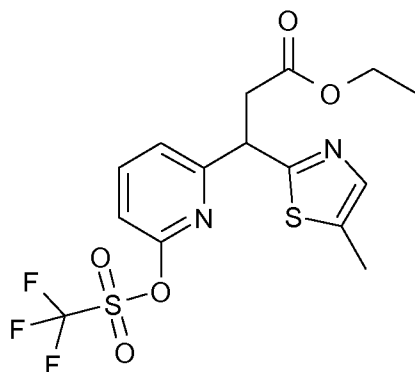
In an alternative approach to the synthesis described in scheme C, the residue Hal in building block E might be replaced by a benzyloxy-residue leading to building block M. When the reduction of the CC-double bond of the Wittig condensation product O is carried out with Pd/C and H₂ the benzyl residue is removed at the same time. Conversion of the newly formed aromatic hydroxy group into the triflate furnishes intermediate P, a suitable starting material for palladium catalyzed cross-couplings. The starting materials M can be obtained from aromatic carboxylic acid esters having a free hydroxy substituent like J. The hydroxy substituent is alkylated with benzylbromide, benzyl chloride or benzyl iodide or any other benzyl derivative capable of undergoing a nucleophilic substitution reaction with an aromatic hydroxy group. Depending on the reactivity of the intermediate L towards the metallation reaction, L might be converted into the corresponding derivatives M by standard procedures well known to a person skilled in the art.



Scheme E

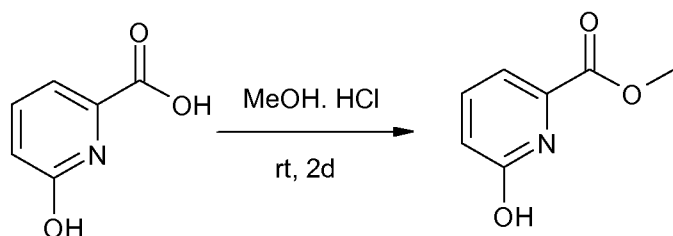
Synthesis of intermediate P1:

3-(5-methylthiazol-2-yl)-3-(6-(trifluoromethyl-sulfonyloxy)pyridin-2-yl)propanoate



Step 1: Synthesis of methyl 6-hydroxypicolinate

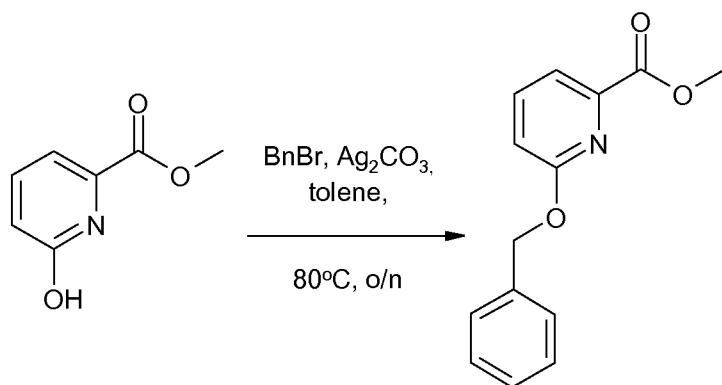
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To a solution of 6-hydroxypicolinic acid (13.0 g, 93.5 mmol) in methanol (150 mL) at room temperature was added HCl in dioxane (4N, 10 mL). The resulting mixture was stirred at room temperature for 48 hours. The reaction mixture was concentrated to give methyl 6-hydroxypicolinate (13 g, 90%) as a white solid.

10

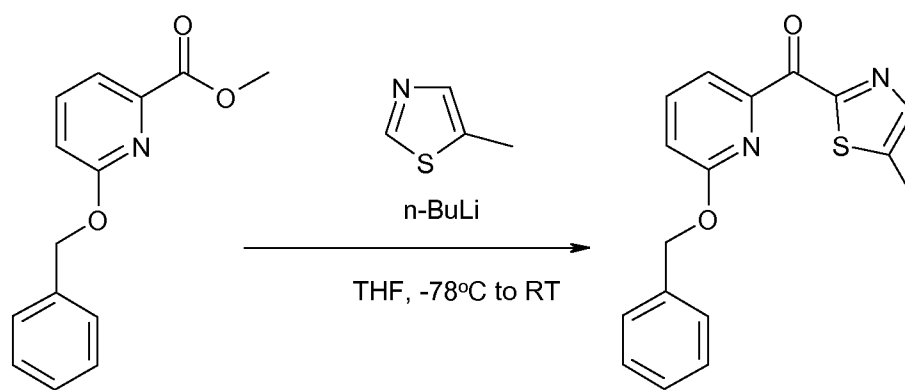
Synthesis of methyl 6-(benzyloxy)picolinate



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A mixture of methyl 6-hydroxypicolinate (3.06 g, 20.0 mmol), (bromomethyl)benzene (6.84 g, 40.0 mmol), and silver carbonate (11 g, 40 mmol) in toluene (150 mL) was stirred at 80 °C overnight. After cooling and filtration, the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography eluting with petroleum ether/ethyl acetate (20:1) to give methyl 6-(benzyloxy)picolinate (2.6 g, 53%) as a white solid.

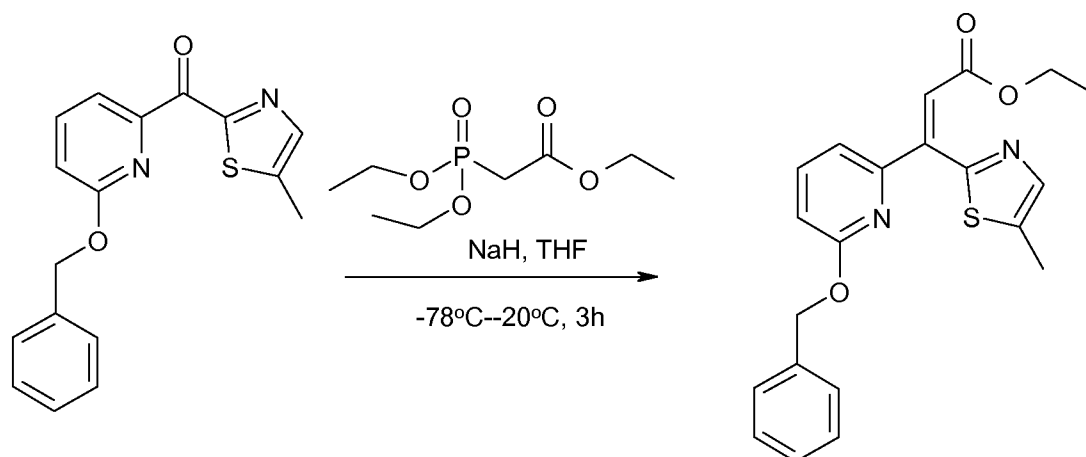
Synthesis of (6-(benzyloxy)pyridin-2-yl)(5-methylthiazol-2-yl)methanone



- 10 To a solution of n-BuLi (2.5 M in hexane, 16 mL, 40 mmol) in dry THF (120 mL) at -78 °C was added 5-methylthiazole (4.17 g, 40 mmol) drop wise over 20 minutes under nitrogen atmosphere. The mixture was stirred at -78 °C for 2 hrs and a solution of methyl 6-(benzyloxy)picolinate (4.7 g, 20 mmol) in THF (30 mL) was added drop wise over 20 minutes. After being stirred at -78 °C for 1 hr, the mixture was acidified with HCl (1N) to pH around 6 and extracted with ethyl acetate. The organic layer was concentrated and then diluted with MTBE. The white precipitate was collected by filtration to give A030-4. The remaining filtrate was concentrated. The residue was purified by silica gel chromatography eluting with petroleum ether/ethyl acetate (10/1) to afford A030-4. Total 6.2 g of (6-(benzyloxy)pyridin-2-yl)(5-methylthiazol-2-yl)methanone (85% yield) was obtained as a white solid.

¹H NMR (400 MHz, CDCl₃) δ 8.00 (d, *J* = 7.2 Hz, 1H), 7.89 (s, 1H), 7.82 (t, *J* = 8.0 Hz, 1H), 7.51-7.36 (m, 5H), 7.01 (d, *J* = 8.0 Hz, 1H), 5.56 (s, 2H), 2.60 (s, 3H).

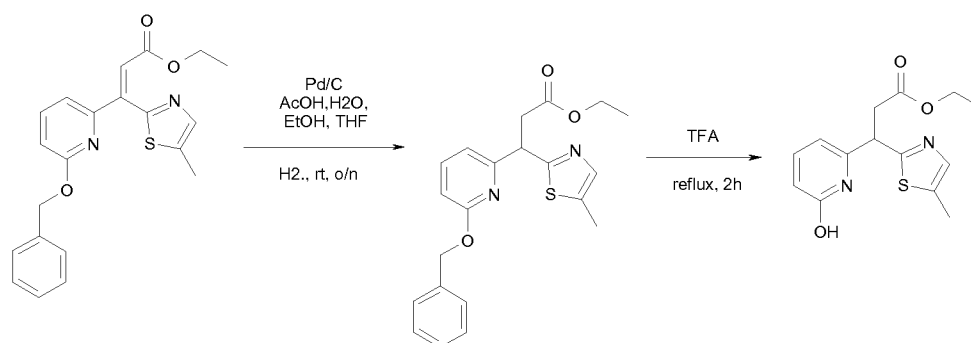
Synthesis of (Z)-ethyl 3-(6-(benzyloxy)pyridin-2-yl)-3-(5-methylthiazol-2-yl)acrylate



To a suspension of NaH (60%, 1.3 g, 33 mmol) in dry THF (200 mL) at -78°C was added ethyl 2-(diethoxyphosphoryl)acetate (7.5 g, 33 mmol). After being stirred at -78°C for 1 hour, a solution of (6-(benzyloxy)pyridin-2-yl)(5-methylthiazol-2-yl)methanone (3.47 g, 11.2 mmol) in THF (30 mL) was added drop wise over 30 minutes. After being stirred at -20 to -30°C for 2 hours, the mixture was acidified with HCl (0.5 N) and then extracted with ethyl acetate. The organic layer was dried and concentrated. The residue was washed with MTBE and petroleum ether to afford (6-(benzyloxy)pyridin-2-yl)(5-methylthiazol-2-yl)methanone (6.2 g, 85% yield) as a white solid.

^1H NMR (400 MHz, CDCl_3) δ 7.58-7.25 (m, 8H), 6.80 (d, $J = 8.4$ Hz, 1H), 6.75 (d, $J = 7.2$ Hz, 1H), 5.44 (s, 2H), 4.15 (q, $J = 3.2$ Hz, 2H), 2.54 (s, 3H), 1.21 (t, $J = 3.2$ Hz, 3H).

Synthesis of ethyl 3-(6-hydroxypyridin-2-yl)-3-(5-methylthiazol-2-yl)propanoate

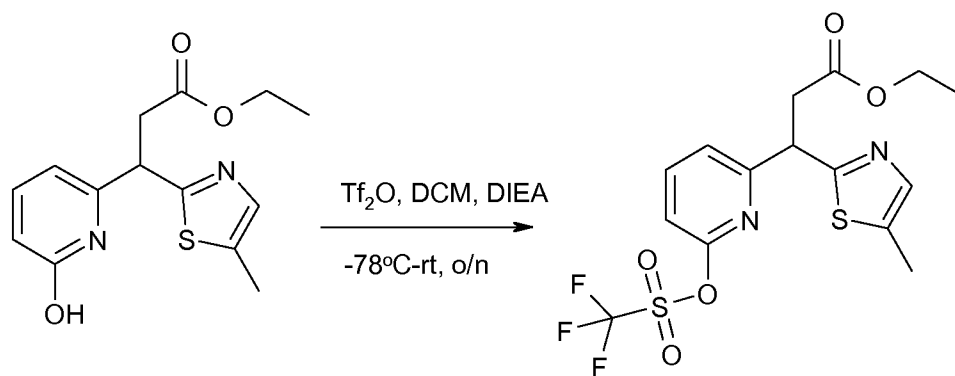


A mixture of (6-(benzyloxy)pyridin-2-yl)(5-methylthiazol-2-yl)methanone (8.0 g, 21 mmol), acetic acid (1.5 mL), H_2O (3 mL), and Pd/C (10%, 4 g) in ethanol (90 mL) and

THF (60 mL) was stirred under H₂ atmosphere at room temperature overnight. After filtration through a pad of Celite and evaporation of the solvent, the residue was washed with MTBE and petroleum ether to afford the product (3g) as a white solid. The filtration was mixed with TFA (20 mL) and the resulting mixture was refluxed for 2h. After cooling down, the mixture was treated with water and extracted with ethyl acetate. The organic layer was dried and concentrated. The residue was washed with MTBE and petroleum ether to afford ethyl 3-(6-hydroxypyridin-2-yl)-3-(5-methylthiazol-2-yl)propanoate (1.4 g, total 71%) as a white solid.

¹H NMR (400 MHz, CDCl₃) δ 7.40 (s, 1H), 7.34-7.30 (m, 1H), 6.46 (dd, *J* = 9.2, 1.2 Hz, 1H), 6.14 (d, *J* = 6.0 Hz, 1H), 4.63 (t, *J* = 7.2 Hz, 1H), 4.13-4.07 (m, 2H), 3.25-3.19 (m, 1H), 3.05-2.97 (m, 1H), 2.43 (s, 3H), 1.19 (t, *J* = 3.2 Hz, 3H).

Synthesis of ethyl 3-(5-methylthiazol-2-yl)-3-(6-(trifluoromethylsulfonyloxy)pyridin-2-yl)propanoate



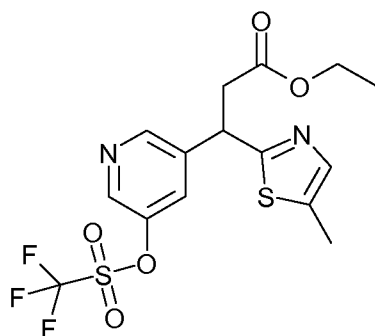
To a solution of ethyl 3-(6-hydroxypyridin-2-yl)-3-(5-methylthiazol-2-yl)propanoate (3.45 g, 11.8 mmol) in DCM (200 mL) at -78 °C was added trifluoromethanesulfonic anhydride (5.00 g, 17.7 mmol) and DIPEA (5.00 mL, 23.6 mmol) drop wise over 30 minutes. The resulting mixture was stirred at -78 °C for 30 minutes and warmed to room temperature overnight. The mixture was cooled to 0 °C and treated with water (100 mL). The organic layer was dried over Na₂SO₄ and concentrated. The oily residue was treated with ether and the solid was filtered off. The filtrate was concentrated to afford ethyl 3-(5-methylthiazol-2-yl)-3-(6-(trifluoromethylsulfonyloxy)pyridin-2-yl)propanoate (4.6 g, 93%) as brown oil.

¹H NMR (400 MHz, CDCl₃) δ 7.83 (t, *J* = 7.6 Hz, 1H), 7.42 (d, *J* = 7.6 Hz, 1H), 7.33 (s, 1H), 7.05 (d, *J* = 8.4 Hz, 1H), 5.02-4.99 (m, 1H), 4.08 (q, *J* = 3.2 Hz, 2H), 3.41-

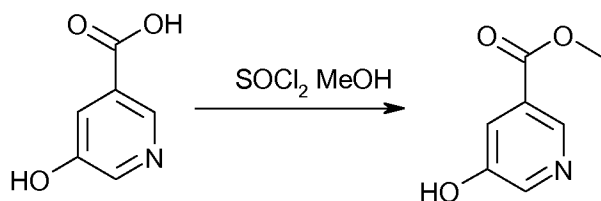
3.35 (m, 1H), 3.22-3.16 (m, 1H), 2.41 (s, 3H), 1.18 (t, $J = 3.2$ Hz, 3H).

Synthesis of intermediate P2:

Ethyl 3-(5-methylthiazol-2-yl)-3-(5-(trifluoromethylsulfonyloxy)pyridin-3-yl)-propanoate



5 Synthesis of methyl 5-hydroxynicotinate

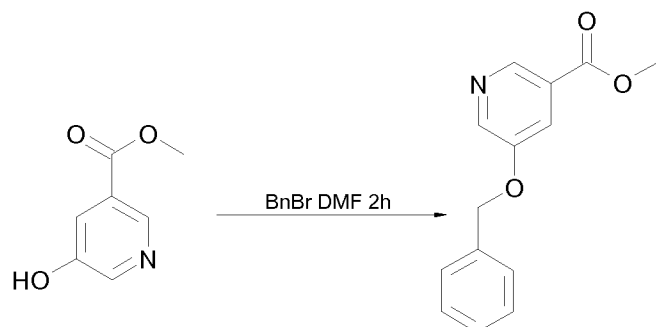


To a stirred solution of 5-hydroxynicotinic acid (10 g, 71.9 mmol) in MeOH (100 mL) was added sulfurous dichloride (1 mL) drop wise over 5 min. The resulting mixture was stirred at room temperature overnight. The resulted solution was added 100 mL of NaHCO₃. The precipitate was filtrated and washed with MeOH for several cycles to give 8.5 g of A031-2 (75% yields), which was used for the nest step without further purification.

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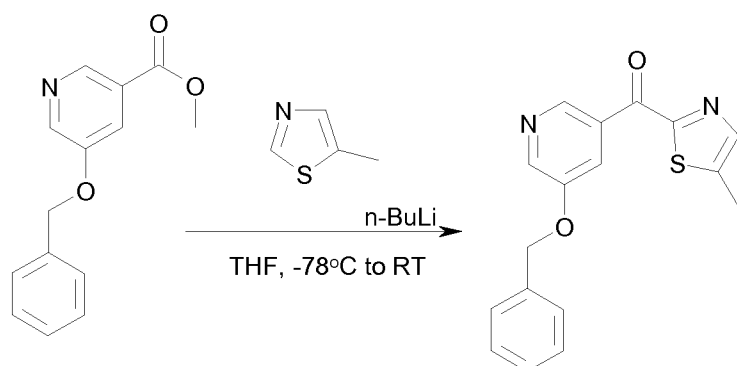
Synthesis of methyl 5-(benzyloxy)nicotinate

44



To a stirred solution of methyl 5-hydroxynicotinate (8.5 g, 55.6 mmol) and K_2CO_3 (11.5 g, 83.3 mmol) in DMF (20 mL) at 0 °C was added (bromomethyl)benzene (8 mL, 66.7 mmol) drop wise over 5 min. The resulting mixture was stirred at room temperature for 2 hours. The mixture was quenched with water (100 mL) and extracted with EA (100mL x 3). The combined organic layers were washed with brine (100 mL), dried over Na_2SO_4 . After filtration and evaporation of the solvent, the residue was purified by silica gel column chromatography eluting with petroleum ether/ethyl acetate (4:1) to give methyl 5-(benzyloxy)nicotinate (4 g, 30.7%) as a yellow oil.

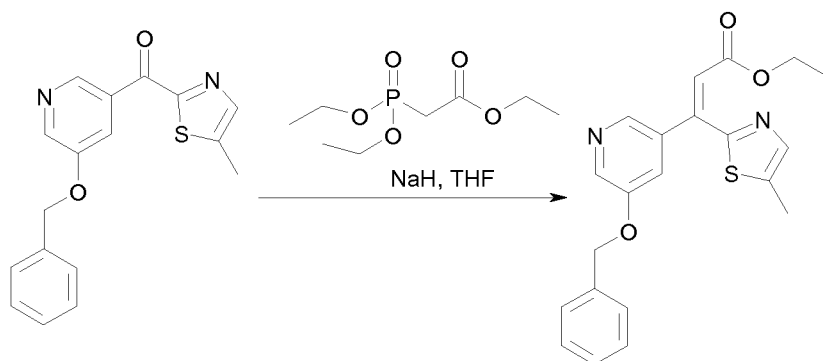
Synthesis of (5-(benzyloxy)pyridin-3-yl)(5-methylthiazol-2-yl)methanone



To a stirring mixture of 5-methylthiazole (3.3 g, 33 mmol) in dry THF (20 mL) was added $n-BuLi$ (33 mmol 13.2 mL solution in hexanes,) drop wise over 10 minutes at -78 °C under nitrogen atmosphere. The mixture was stirred between -78 °C and -60 °C for 1.5 h and then cooled to -78 °C. A solution of methyl 5-(benzyloxy)nicotinate

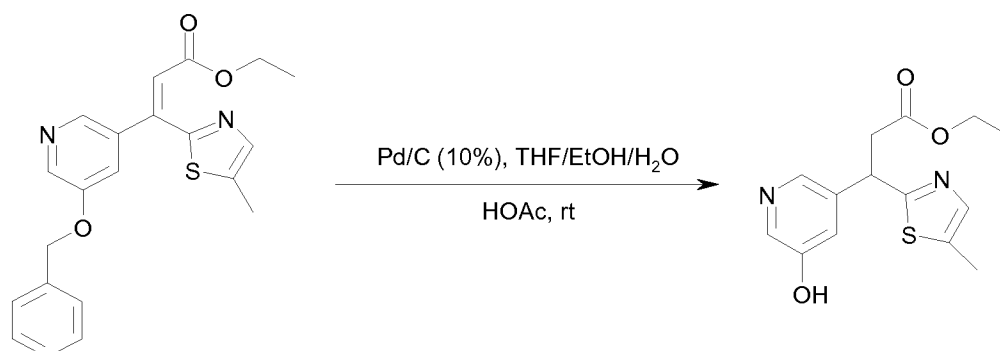
(4.0 g, 16.5 mmol) in THF (10 mL) was added drop wise over 10 minutes. The resulting mixture was stirred at -78 °C for 30 minutes and warmed to room temperature with stirring overnight. The mixture was cooled to 0 °C and treated with water (50 mL). The resulting mixture was adjusted to pH around 6 with HCl (1N) and extracted with EA (100 mL x 3). The combined organic layers were dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography eluting with petroleum ether/EtOAc (3:1) to afford (5-(benzyloxy)-2-methoxyphenyl)(thiazol-2-yl)methanone (3.2 g, 42%) as a yellow gel.

Synthesis of (Z)-ethyl 3-(5-(benzyloxy)pyridin-3-yl)-3-(5-methylthiazol-2-yl)acrylate



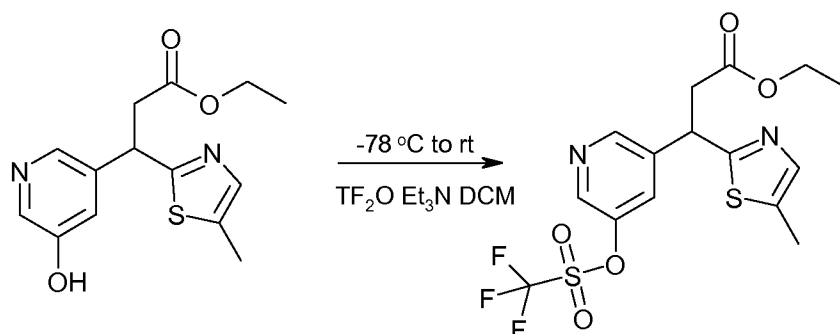
To a suspension of NaH (60%, 1.2 g, 30.9 mmol) in dry THF (20 mL) was added ethyl 2-(diethoxyphosphoryl)acetate (6.9 g, 30.9 mmol) at -78 °C. After being stirred at -78 °C for 1 hour, a solution of (5-(benzyloxy)pyridin-3-yl)(5-methylthiazol-2-yl)methanone (3.2 g, 10.3 mmol) in THF (5 mL) was added drop wise over 30 min. The mixture was stirred at -78 °C for 30 min and then warmed to room temperature with stirring overnight. The mixture was cooled to 0 °C and treated with water (50 mL), pH value was adjusted to 6 with HCl (0.5 N) and extracted with EA (50 mL x 3). The combined organic layers were dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by silica gel chromatography eluting with petroleum ether/EtOAc (3: 1) to afford (Z)-ethyl 3-(5-(benzyloxy)-2-methoxyphenyl)-3-(thiazol-2-yl)acrylate (3.1 g, 80.0%) as yellow gel.

Synthesis of ethyl 3-(5-hydroxypyridin-3-yl)-3-(5-methylthiazol-2-yl)propanoate



To a mixture of (Z)-ethyl 3-(5-(benzyloxy)pyridin-3-yl)-3-(5-methylthiazol-2-yl)acrylate (3.1 g, 8.2 mmol), HOAc (1.0 mL), and H₂O (2.0 mL) in EtOH (20 mL) and THF (10 mL) was added Pd/C (10%, dry, 820 mg) under H₂ atmosphere. The mixture was stirred at room temperature under H₂ atmosphere for 48 hrs. After filtration through a pad of Celite and evaporation of the solvent, the residue was purified by silica gel chromatography eluting with petroleum ether/EtOAc (3:1) to afford 3-(5-hydroxypyridin-3-yl)-3-(5-methylthiazol-2-yl)propanoate (2.2 g, 92%) as a yellow-green gel.

Synthesis of ethyl 3-(5-methylthiazol-2-yl)-3-(5-(trifluoromethylsulfonyloxy)pyridin-3-yl)propanoate



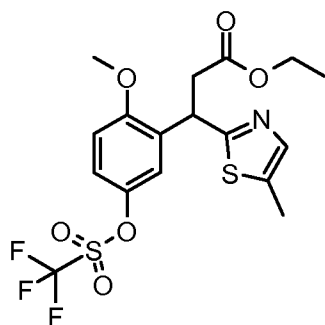
To a stirred solution of ethyl 3-(5-hydroxypyridin-3-yl)-3-(5-methylthiazol-2-yl)propanoate (2.2 g, 7.5 mmol) and Et₃N (1.52 g, 15.1 mmol) in DCM (20 mL) at -78 °C was added trifluoromethanesulfonic anhydride (2.3 g, 15.1 mmol) drop wise over 5 min. The resulting mixture was stirred at -78 °C for 30 minutes and warmed to

room temperature with stirring overnight. The mixture was cooled to 0°C, treated with water (50 mL) and extracted with DCM (50 mL x 3). The combined organic layers were washed with brine (50 mL), dried over Na₂SO₄, and concentrated. The residue was purified by silica gel column chromatography eluting with petroleum ether/ethyl acetate (4:1) to give ethyl 3-(2-methoxy-5-(trifluoromethylsulfonyloxy)phenyl)-3-(thiazol-2-yl)propanoate (900 mg, 28.0%) as yellow oil.

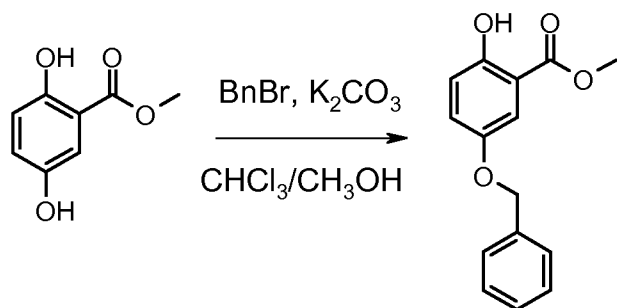
¹H NMR (400 MHz, MeOD) δ 8.71 (d, *J* = 1.2 Hz, 1H), 8.58 (d, *J* = 2.4 Hz, 1H), 7.96 (s, 1H), 7.43 (s, 1H), 5.02 (t, 1H), 4.10-4.05 (m, 2H), 3.43-3.37 (m, 1H), 3.22-3.15 (m, 1H), 2.45 (s, 3H), 1.18-1.14 (m, 3H).

Synthesis of intermediate P3:

3-(2-Methoxy-5-trifluoromethanesulfonyloxy-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid ethyl ester



Methyl 5-(benzyloxy)-2-hydroxybenzoate

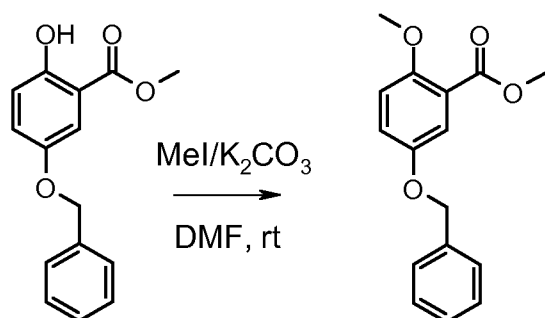


To a refluxed suspension of K₂CO₃ (32.8 g, 238 mmol) in methanol (180 mL) and CHCl₃ (350 mL) was then added a mixture of methyl 2,5-dihydroxybenzoate (10.0 g, 59.5 mmol) and bromomethylbenzene (7.10 mL, 59.5 mmol) in methanol/CHCl₃ (50

mL/25 mL) drop wise over 30 minutes. The resulting mixture was stirred at reflux for another 4 hours. After cooling down and filtration, the filter cake was washed with CHCl_3 (20 mL). The combined filtrate was concentrated under reduced pressure. The residue was redissolved in CHCl_3 (200 mL), washed with HCl (1N) (100 mL x 2). The organic was then washed with brine (100 mL), dried over Na_2SO_4 , filtered, and concentrated. The residue was purified by silica gel column chromatography eluting with petroleum ether/EtOAc (8: 1) to give methyl 5-(benzyloxy)-2-hydroxybenzoate (10.6 g, 69.0%) as a white solid.

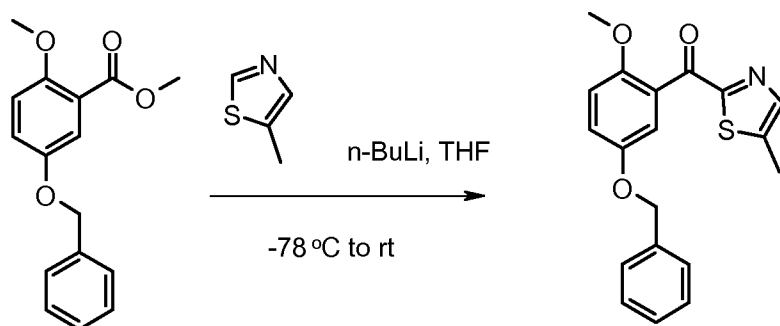
^1H NMR (500 MHz, $\text{DMSO}-d_6$) δ 10.09 (s, 1H), 7.43 (d, J = 7.2 Hz, 2H), 7.40-7.36 (m, 2H), 7.34-7.32 (2H, m), 7.23 (dd, J = 9.2, 3.2 Hz, 1H), 6.93 (d, J = 9.2 Hz, 1H), 5.06 (s, 2H), 3.88 (s, 3H). NOESY showed the desired product as well.

Methyl 5-(benzyloxy)-2-methoxybenzoate



To a mixture of methyl 5-(benzyloxy)-2-hydroxybenzoate (10.6 g, 41.1 mmol) and K_2CO_3 (11.3 g, 82.2 mmol) in DMF (100 mL) was added iodomethane (2.60 mL, 49.3 mmol) drop wise over 5 minutes. The resulting mixture was stirred at room temperature overnight. The reaction mixture was poured into water (400 mL), filtered, and collected the solid. The solid was dissolved in EtOAc (300 mL), washed with water (50 mL) and brine (100 mL), dried over Na_2SO_4 , filtered, and concentrated under reduced pressure to give methyl 5-(benzyloxy)-2-methoxybenzoate (10.4 g, 98%) as a yellow solid.

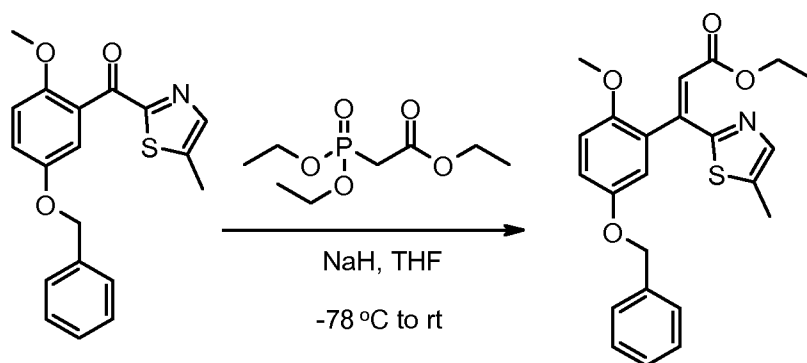
(5-(benzyloxy)-2-methoxyphenyl)(5-methylthiazol-2-yl)methanone



To a stirring mixture of 5-methylthiazole (3.80 g, 38.2 mmol) in dry THF (200 mL) was
 5 added n-BuLi (2.5 M in hexane, 15.3 mL, 38.2 mmol) drop wise over 20 minutes at -
 78 °C under nitrogen atmosphere. The mixture was stirred between -78 °C and -60
 °C for 1.5 h and then cooled to -78 °C. A solution of methyl 5-(benzyloxy)-2-
 methoxybenzoate (10.4 g, 38.2 mmol) in THF (50 mL) was added drop wise over 30
 minutes. The resulting mixture was stirred at -78 °C for 30 minutes and warmed to
 10 room temperature with stirring overnight. The mixture was cooled to 0 °C and treated
 with water (50 mL). The resulting mixture was adjusted to pH around 6 with HCl (1N).
 The organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced
 pressure. The residue was purified by silica gel column chromatography eluting with
 petroleum ether/EtOAc (3:1) to afford (5-(benzyloxy)-2-methoxyphenyl)(5-
 15 methylthiazol-2-yl)methanone (2.5 g, 19%) as a yellow gel. 3.0 g of starting material
 methyl 5-(benzyloxy)-2-methoxybenzoate was recovered.

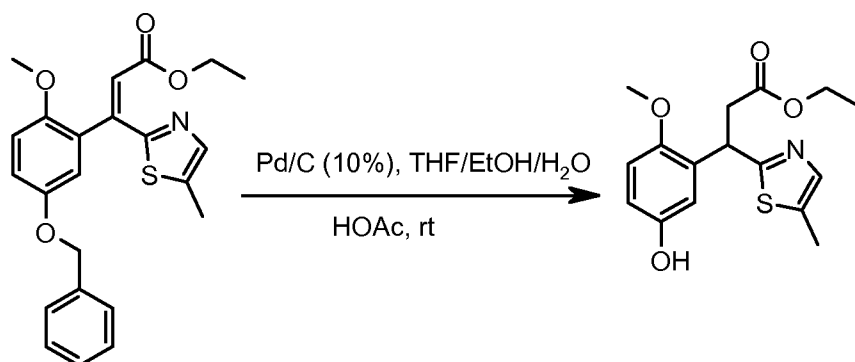
Synthesis of (Z)-ethyl 3-(5-(benzyloxy)-2-methoxyphenyl)-3-(5-methylthiazol-2-
 yl)acrylate

50



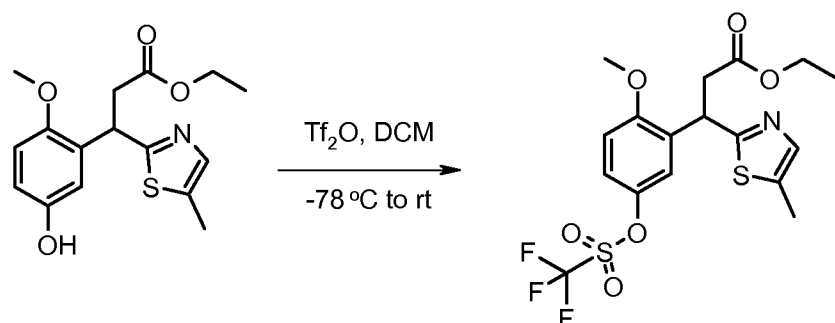
To a suspension of NaH (60%, 884 mg, 22.1 mmol) in dry THF (120 mL) was added ethyl 2-(diethoxyphosphoryl)acetate (4.96 g, 22.1 mmol) at -78 °C. After being stirred at -78 °C for 1 hour, a solution of (5-(benzyloxy)-2-methoxyphenyl)(5-methylthiazol-2-yl)methanone (2.50 g, 7.37 mmol) in THF (30 mL) was added drop wise over 30 min. The mixture was stirred at -78 °C for 30 min and then warmed to room temperature with stirring overnight. The mixture was cooled to 0 °C and treated with water (50 mL), pH value was adjusted to 6 with HCl (0.5 N). The organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by silica gel chromatography eluting with petroleum ether/EtOAc (3: 1) to afford (Z)-ethyl 3-(5-(benzyloxy)-2-methoxyphenyl)-3-(5-methylthiazol-2-yl)acrylate (1.75 g, 58.0%) as yellow gel.

Synthesis of ethyl 3-(5-hydroxy-2-methoxyphenyl)-3-(5-methylthiazol-2-yl)propanoate



To a mixture of (Z)-ethyl 3-(5-(benzyloxy)-2-methoxyphenyl)-3-(5-methylthiazol-2-yl)acrylate (4.0 g, 9.8 mmol), HOAc (2.0 mL), and H₂O (5.0 mL) in EtOH (100 mL) and THF (60 mL) was added Pd/C (10%, dry, 5.0 g) under nitrogen atmosphere. The mixture was stirred at room temperature under H₂ atmosphere for 48 hrs. After filtration through a pad of Celite and evaporation of the solvent, the residue was purified by silica gel chromatography eluting with petroleum ether/EtOAc (3:1) to afford ethyl 3-(5-hydroxy-2-methoxyphenyl)-3-(5-methylthiazol-2-yl)propanoate (1.9 g, 60%) as a yellow-green gel.

10 Ethyl-3-(2-methoxy-5-(trifluoromethylsulfonyloxy)phenyl)-3-(5-methylthiazol-2-yl)propanoate



15 To a stirred solution of ethyl 3-(5-hydroxy-2-methoxyphenyl)-3-(5-methylthiazol-2-yl)propanoate (4.50 g, 14.0 mmol) and DIPEA (4.8 mL, 28 mmol) in DCM (150 mL) at -78 °C was added trifluoromethanesulfonic anhydride (4.74 g, 16.8 mmol) drop wise over 30 min. The resulting mixture was stirred at -78 °C for 30 minutes and warmed to room temperature with stirring overnight. The mixture was cooled to 0 °C and treated with water (50 mL). The organic layer was washed with brine (50 mL), dried over Na₂SO₄, and concentrated. The residue was purified by silica gel column chromatography eluting with petroleum ether/ethyl acetate (4:1) to give ethyl 3-(2-methoxy-5-(trifluoromethylsulfonyloxy)phenyl)-3-(5-methylthiazol-2-yl)propanoate (4.63 g, 73.0%) as yellow oil.

25 ¹H NMR (400 MHz, DMSO-*d*₆) δ 7.41 (dd, *J* = 9.1, 3.1 Hz, 1H), 7.37 (d, *J* = 1.2 Hz, 1H), 7.35 (d, *J* = 3.1 Hz, 1H), 7.18 (d, *J* = 9.2 Hz, 1H), 5.07 (dd, *J* = 8.3, 7.1 Hz, 1H),

3.99 (q, $J = 7.1$ Hz, 2H), 3.86 (s, 3H), 3.32 (dd, $J = 16.3, 6.9$ Hz, 1H), 3.02 (dd, $J = 16.3, 6.9$ Hz, 1H), 2.35 (d, $J = 1.0$ Hz, 3H), 1.08 (dd, $J = 9.2, 5.1$ Hz, 3H).

5

Synthesis of intermediate P4:

3-(2-Methoxy-5-trifluoromethanesulfonyloxy-phenyl)-3-thiazol-2-yl-propionic acid ethyl ester

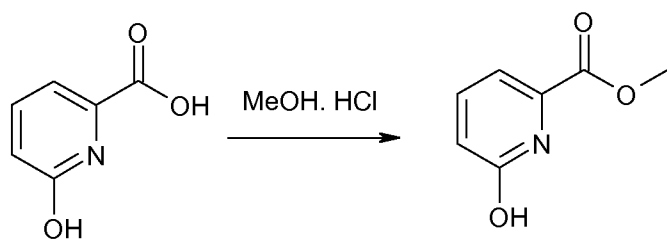
- 10 The synthesis is carried out as described for the synthesis of intermediate P3 using 1,3-thiazole instead of 5-methyl-1,3-thiazole as starting material.

Synthesis of intermediate P5

- 15 Ethyl 3-(5-methyl-1, 3, 4-oxadiazol-2-yl)-3-(6-(trifluoromethyl sulfonyloxy)pyridin-yl)-propanoate

Methyl 6-hydroxypicolinate

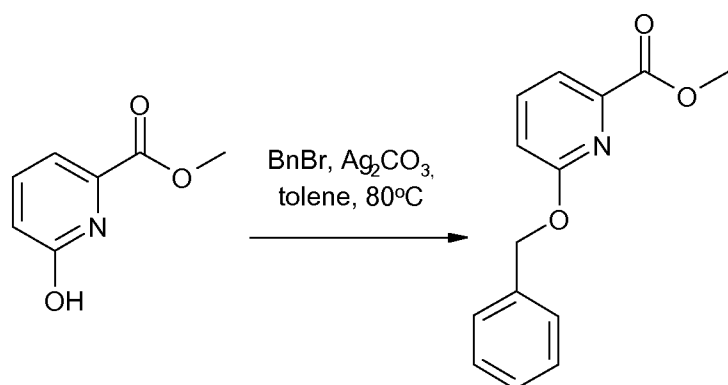
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- To a solution of 6-hydroxypicolinic acid (13.0 g, 93.5 mmol) in methanol (150 mL) at room temperature was added HCl in dioxane (4N, 10 mL). The resulting mixture was stirred at room temperature for 48 h. The reaction mixture was concentrated to give
- 25 methyl 6-hydroxypicolinate (13.0 g, 90 %) as a white solid.

6-(Benzyloxy)picolinate

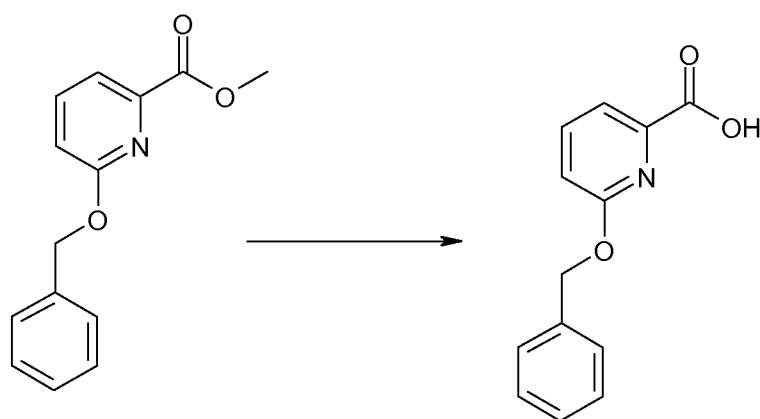
53



A solution of methyl 6-hydroxypicolinate (3.06 g, 20 mmol), (bromomethyl)benzene (6.84 g, 40 mmol) and silver carbonate (11 g, 40 mmol) in toluene (150 mL) was stirred at 80°C overnight. Cooled and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel chromatography (eluted with petroleum ether: ethyl acetate = 20: 1) to give methyl 6-(benzyloxy)picolinate (2.6 g, 53 %) as a white solid.

10

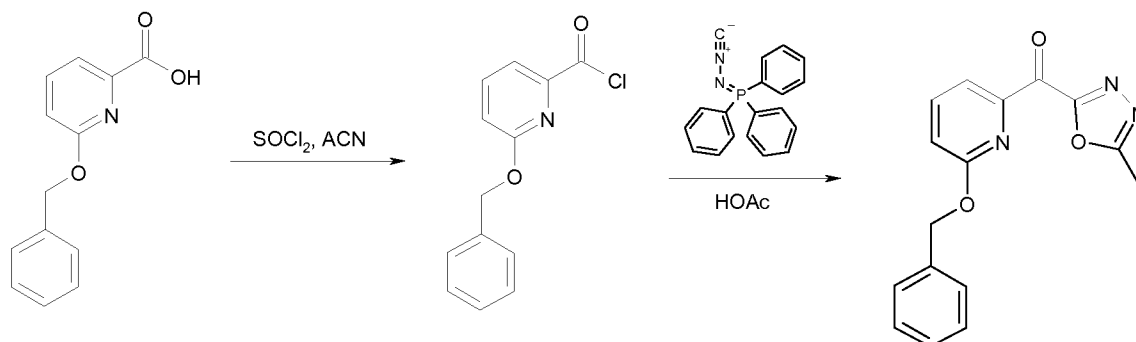
6-(Benzyloxy)pyridine-2-carboxylic acid



To a solution of methyl 6-(benzyloxy)picolinate (6 g, 24.7 mmol) in THF (75 mL) and water (15 mL) was added $\text{LiOH} \cdot \text{H}_2\text{O}$ (3.12 g, 74 mmol). The reaction solution was stirred at rt overnight. Workup gave 6-(benzyloxy)pyridine-2-carboxylic acid (5 g, 100%) as a white foam.

15

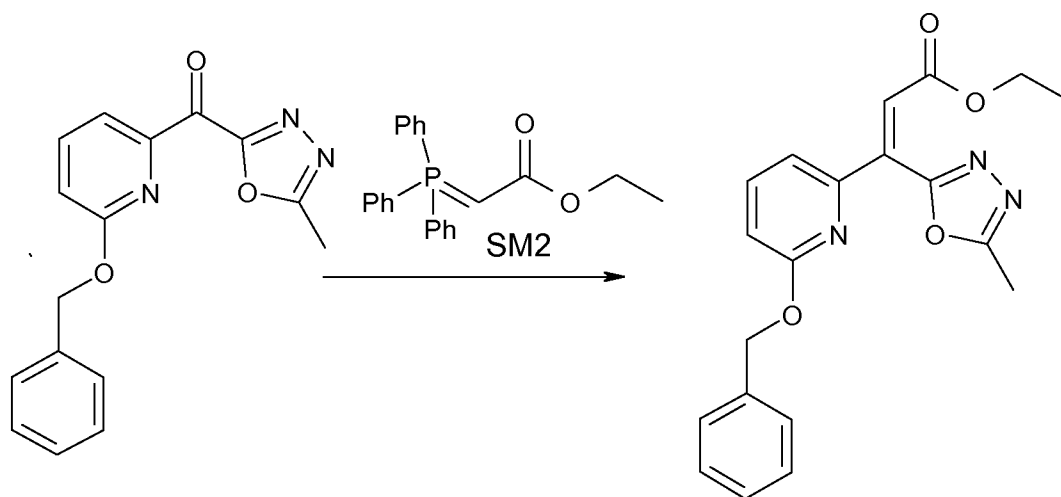
[6-(Benzyloxy)pyridin-2-yl](5-methyl-1,3,4-oxadiazol-2-yl)methanone



To a solution of 6-(benzyloxy)pyridine-2-carboxylic acid (2.28 g, 10 mmol) in
 5 acetonitrile (20 mL) was added SOCl_2 (5 mL) at rt and the reaction was slowly heated
 to reflux for 20 min. After cooled and concentrated, the residue acid chloride was
 added dichloromethane (30 mL), followed by the addition of (N-
 isocyanimino)triphenylphosphorane (3.12 g, 10 mmol) at rt. After 2h, acetic acid (1.2
 g, 20 mmol) was added before a solution of triethylamine (2.02 g, 20 mmol) in
 10 dichloromethane (10 mL) was added dropwise. The solution was stirred for an
 additional 8 h, and quenched with water. The organic layer was washed with brine,
 dried and concentrated *in vacuo*. The residue was purified by flash column
 chromatography (petroleum ether: ethyl acetate=8; 1) to give [6-(benzyloxy)pyridin-2-
 yl](5-methyl-1,3,4-oxadiazol-2-yl)methanone (200 mg, 7%) as a brown oil.

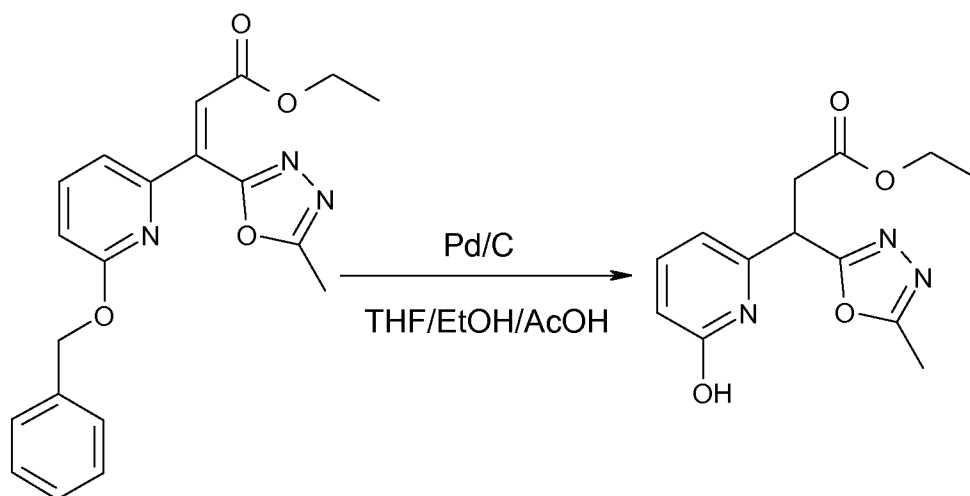
15

Ethyl 3-(6-(benzyloxy) pyridin-2-yl)-3-(5-methyl-1, 3, 4-oxadiazol-2-yl) acrylate



A mixture of [6-(benzyloxy)pyridin-2-yl](5-methyl-1,3,4-oxadiazol-2-yl)methanone (2.2 g, 7.5 mmol) and ethyl (triphenylphosphoranylidene)acetate (2.6 g, 7.5 mmol) in benzene (15 mL) was stirred at 100 °C for 48 h. The solution was evaporated and the residue was purified on column chromatography (silica gel, EA/PE = 1: 8) to give ethyl 3-(6-(benzyloxy)pyridin-2-yl)-3-(5-methyl-1, 3, 4-oxadiazol-2-yl)acrylate (2.3 g, 84%) as a brown oil.

10 Ethyl 3-(6-hydroxypyridin-2-yl)-3-(5-methyl-1, 3 4-oxadiazol-2-yl)propanoate

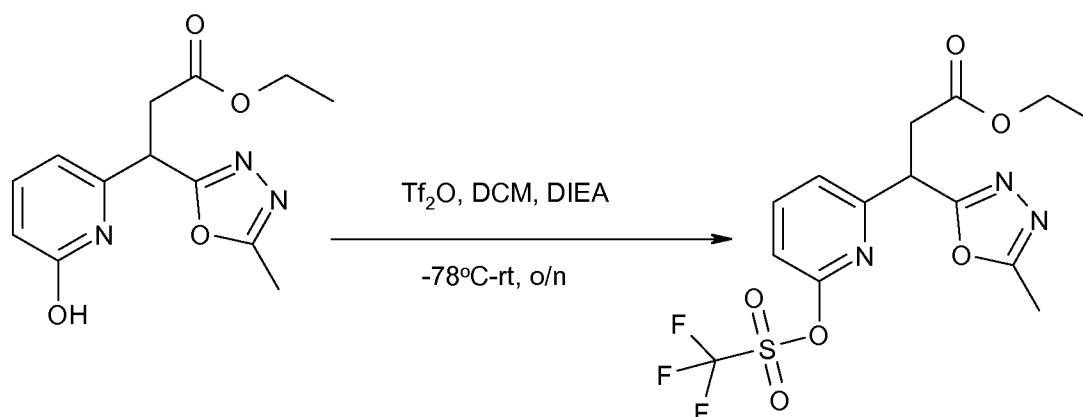


To a solution of ethyl 3-(6-(benzyloxy)pyridin-2-yl)-3-(5-methyl-1, 3, 4-oxadiazol-2-yl)acrylate (2.2 g, 6.0 mmol) in EtOH (30 mL), THF (30 mL) and AcOH (1 mL) was added Pd/C (220 mg, 10%). The system was evacuated and then refilled with H₂ for

3 times. The mixture solution was stirred at rt overnight under H₂ atmosphere. The mixture was filtrated and the filtration was evaporated *in vacuo* to give ethyl 3-(6-hydroxypyridin-2-yl)-3-(5-methyl-1, 3 4-oxadiazol-2-yl)propanoate (1.4 g, 84%).

5

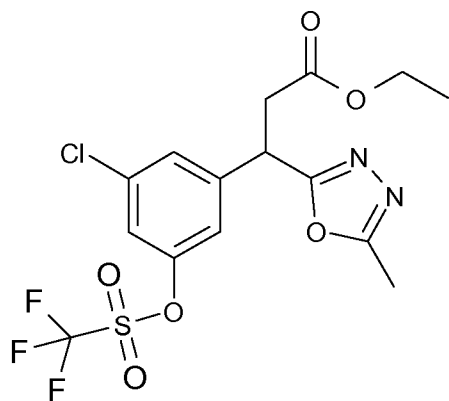
3-(5-Methyl-1, 3, 4-oxadiazol-2-yl)-3-(6-(trifluoromethyl sulfonyloxy)pyridin-2-yl)propanoate



A mixture of ethyl 3-(6-hydroxypyridin-2-yl)-3-(5-methyl-1, 3 4-oxadiazol-2-yl) propanoate (1.0 g, 3.6 mmol) and DIPEA (1.4 g, 10.8 mmol) in DCM (40 mL) was stirred at -78°C for 0.5h. Then trifluoromethanesulfonic anhydride (1.5g, 5.4 mmol) in DCM (6mL) was added dropwise at -78 °C. The mixture solution was stirred at -78°C for 1h then warmed to room temperature overnight. The mixture was concentrated in vacuum to give a residue which was purified on column chromatography (elution was 12% EA in PE) to give ethyl 3-(5-methyl-1, 3, 4-oxadiazol-2-yl)-3-(6-(trifluoromethylsulfonyloxy) pyridin-2-yl) propanoate (1.47 g, 99%) as a yellow liquid.

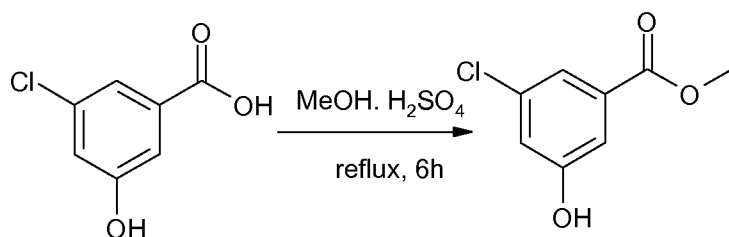
20 Synthesis of intermediate P6

Ethyl 3-(3-chloro-5-[(trifluoromethyl)sulfonyl]oxy)phenyl)-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoate



3-Chloro-5-hydroxybenzoate

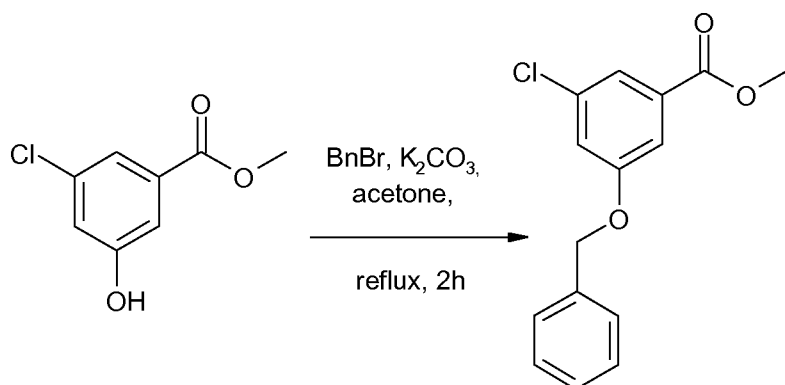
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- 10 To a solution of 3-chloro-5-hydroxybenzoic acid (1.72 g, 10 mmol) in methanol (20 mL) at room temperature was added conc. H_2SO_4 (0.5 mL). And then the resulting mixture was refluxed for 6 hours. The reaction mixture was concentrated and the residue was added NaHCO_3 solution and ethyl acetate. The organic layer was separated, dried and concentrated to give methyl 3-chloro-5-hydroxybenzoate (1.8 g,
- 15 97 %) as a brown solid, which was used in the next step without further purification.

Methyl 3-(benzyloxy)-5-chlorobenzoate

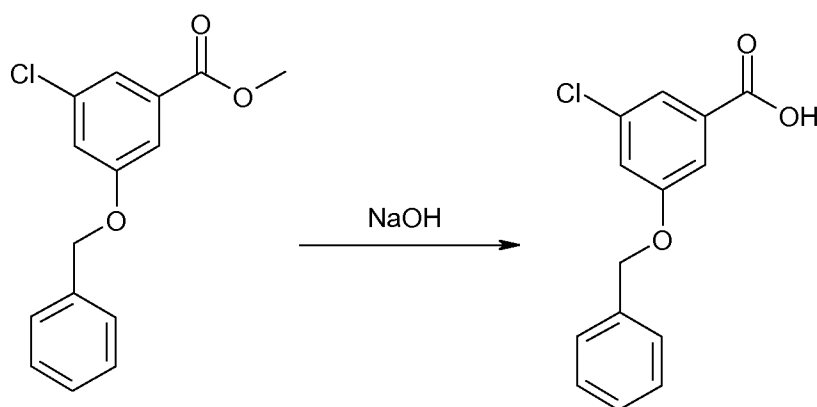
58



A solution of methyl 3-chloro-5-hydroxybenzoate (12.75 g, 68.5 mmol), (bromomethyl)benzene (11 mL, 82.2 mmol) and potassium carbonate (19 g, 137 mmol) in acetone (300 mL) was refluxed for 2h. Cooled and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified on silica gel chromatography (eluted with petroleum ether: ethyl acetate =20: 1) to give methyl 3-(benzyloxy)-5-chlorobenzoate (18 g, 95 %) as a light yellow oil.

10

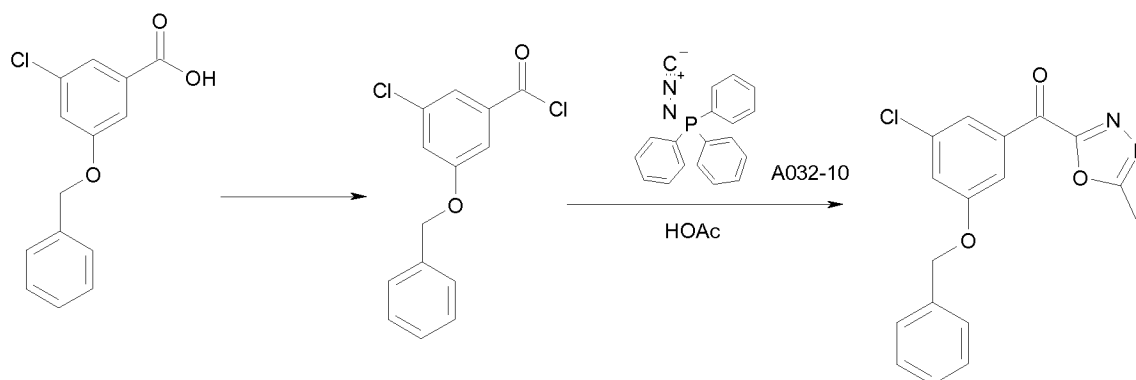
3-(Benzyloxy)-5-chlorobenzoic acid



To a solution of methyl 3-(benzyloxy)-5-chlorobenzoate (5 g, 18 mmol) in methanol (60 mL) and water (20 mL) was added NaOH (2.18 g, 54 mmol) at rt. The mixture was refluxed overnight. Workup gave 3-(benzyloxy)-5-chlorobenzoic acid (4.8 g, 100%) as a white solid.

15

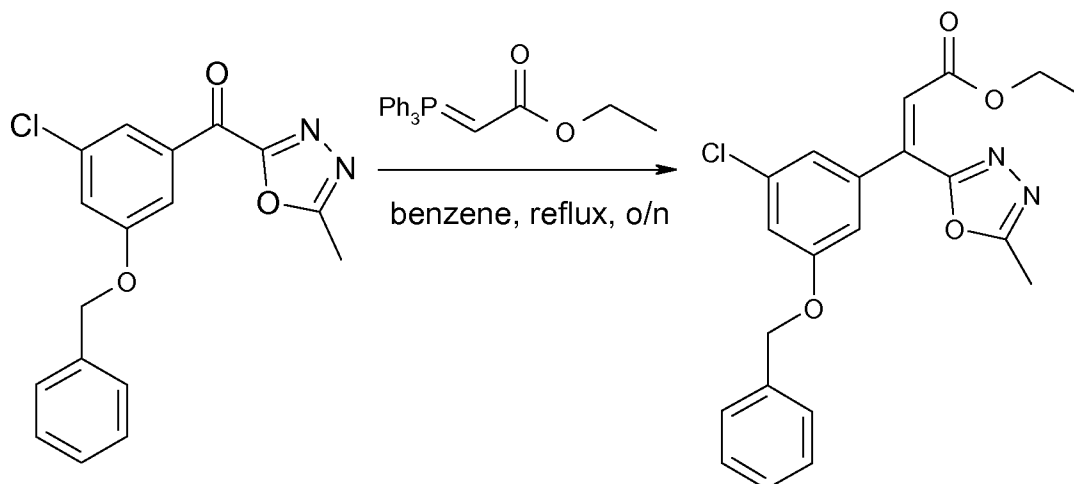
[3-(Benzyloxy)-5-chlorophenyl](5-methyl-1,3,4-oxadiazol-2-yl)methanone



- 5 To a solution of 3-(benzyloxy)-5-chlorobenzoic acid (1.31 g, 5 mmol) in acetonitrile (20 mL) was added SOCl_2 (2.5 mL) at rt, then slowly heated to reflux for 20 min. After cooled and concentrated, the residue acid chloride was added dichloromethane (25 mL), followed by (N-isocyanimine)triphenylphosphorane (1.51 g, 5 mmol) at rt. After 2h, acetic acid (600 mg, 10 mmol) was added before a solution of triethylamine (1.01 g, 10 mmol) in dichloromethane (10 mL) was added dropwise. The solution was
- 10 stirred for an additional 8 h, and washed with brine, dried and concentrated on rotary evaporator. The residue was purified by flash column chromatography (petroleum ether: ethyl acetate=10/1) to give [3-(benzyloxy)-5-chlorophenyl](5-methyl-1,3,4-oxadiazol-2-yl)methanone (700 mg, 42%) as a brown oil.

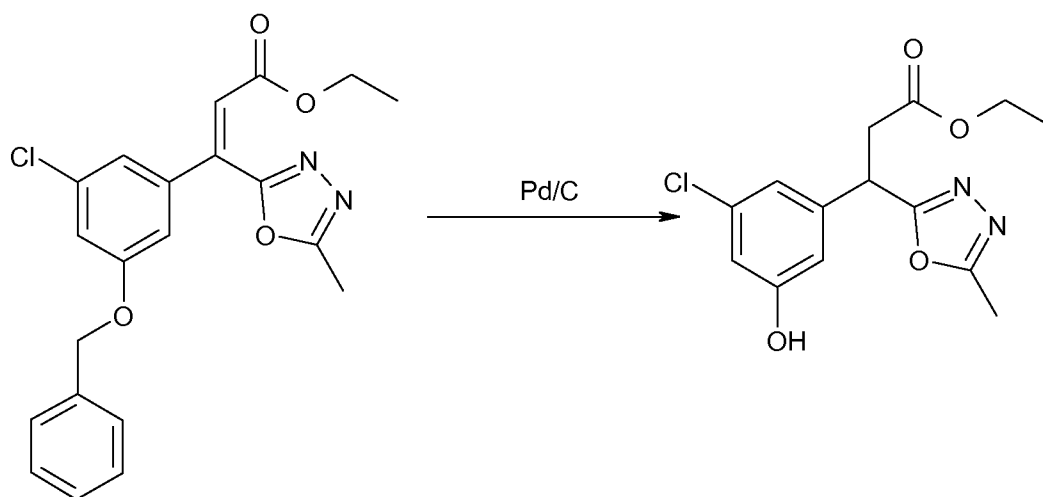
15

Ethyl 3-(3-(benzyloxy)-5-chlorophenyl)-3-(5-methyl-1,3,4-oxadiazol-2-yl)acrylate



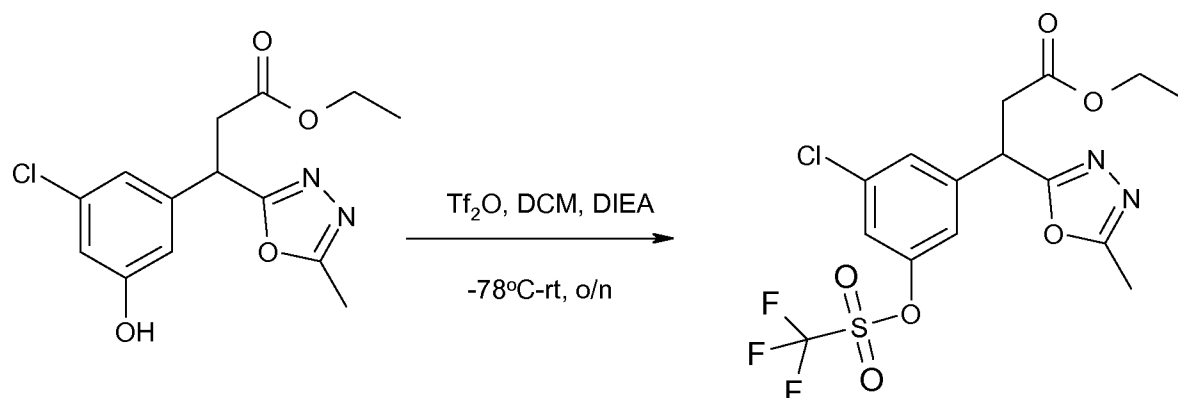
A solution of [3-(benzyloxy)-5-chlorophenyl](5-methyl-1,3,4-oxadiazol-2-yl)methanone (700 mg, 2.13 mmol) and ethyl (triphenylphosphoranylidene)acetate (800 mg, 2.13 mmol) in benzene (20 mL) was refluxed overnight. After concentration, the resulted residue was purified by flash column chromatography (petroleum ether: ethyl acetate=10/1 to 4/1) to give ethyl 3-(3-(benzyloxy)-5-chlorophenyl)-3-(5-methyl-1,3,4-oxadiazol-2-yl)acrylate (780 mg, 93%) as a brown oil.

Ethyl 3-(3-chloro-5-hydroxyphenyl)-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoate



To a mixture of ethyl 3-(3-(benzyloxy)-5-chlorophenyl)-3-(5-methyl-1,3,4-oxadiazol-2-yl)acrylate (700 mg, 1.76 mmol) in ethanol (20 mL) was added Pd/C (10%, 200 mg) under nitrogen atmosphere. The mixture was stirred at room temperature under H₂ atmosphere overnight. After filtration through a pad of Celite, the filtrate was concentrated to give crude ethyl 3-(3-chloro-5-hydroxyphenyl)-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoate (650 mg, 100%) as a yellow oil.

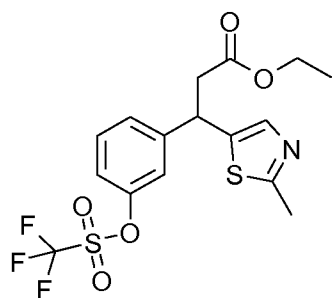
Ethyl 3-(3-chloro-5-[(trifluoromethyl)sulfonyl]oxyphenyl)-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoate



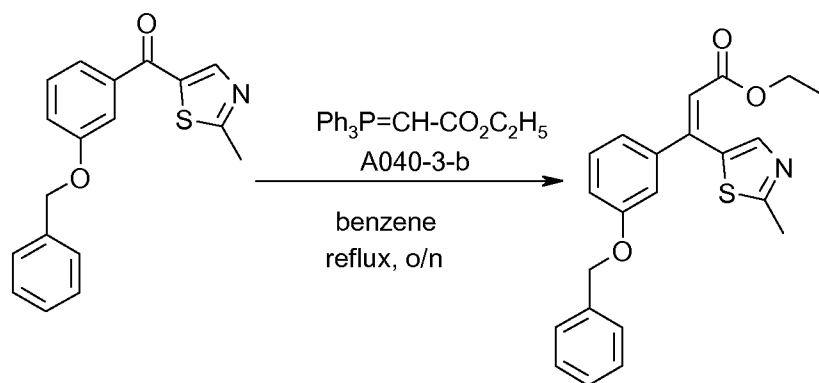
To a solution of ethyl 3-(3-chloro-5-hydroxyphenyl)-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoate (600 mg, 2 mmol) in DCM (30 mL) at -78°C was added trifluoromethanesulfonic anhydride (846 mg, 3 mmol) and DIPEA (516 mg, 4 mmol) dropwise. The resulting mixture was stirred at -78°C for 30 minutes and warmed to room temperature overnight. The mixture was cooled to 0°C and quenched with water (20 mL), and then the organic layer was dried over Na_2SO_4 and concentrated. The oily residue was added ether, the solid was filtered and the filtration was concentrated. The resulted residue was purified by flash column chromatography (petroleum ether: ethyl acetate= 3/1) to give ethyl 3-(3-chloro-5-
 5 {{{(trifluoromethyl)sulfonyl}oxy}phenyl)-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoate (375 mg, 42%) as a yellow oil.

Synthesis of intermediate P7

15 Ethyl 3-(2-methylthiazol-5-yl)-3-(3-(trifluoromethylsulfonyloxy)phenyl) propanoate

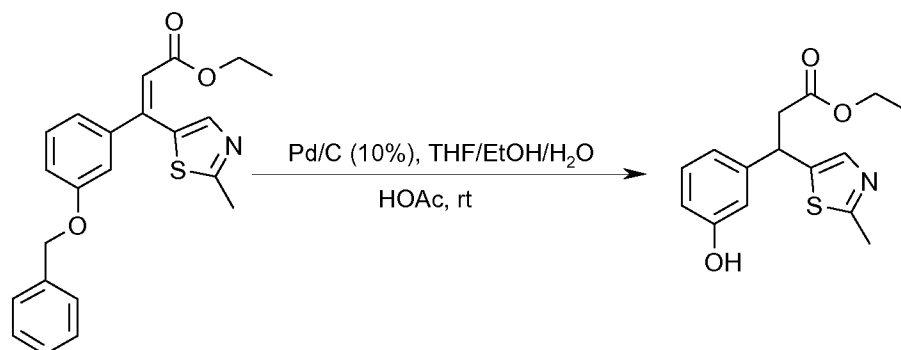


(Z)-Ethyl 3-(3-(benzyloxy)phenyl)-3-(2-methylthiazol-5-yl)acrylate



- 5 The solution of 3-(3-(benzyloxy)phenyl)(2-methylthiazol-5-yl)methanone (1.9 g, 6.1 mmol) in benzene (20 mL) was added A040-3-b (2.1 g 6.1 mmol). The resulting mixture was refluxed overnight. The reaction was cooled to rt and diluted with water (50 mL) and EA (100 mL). The organic layer was separated and the aqueous washed with EA (2 x 100 mL). The combined organic layers were dried over Na_2SO_4 .
- 10 After filtration and evaporation of the solvent, the crude residue was purified over silica gel (4:1 PE/EA) to give a clear oil (1.6g, 70%).

Ethyl 3-(3-(benzyloxy)phenyl)-3-(2-methylthiazol-5-yl)propanoate



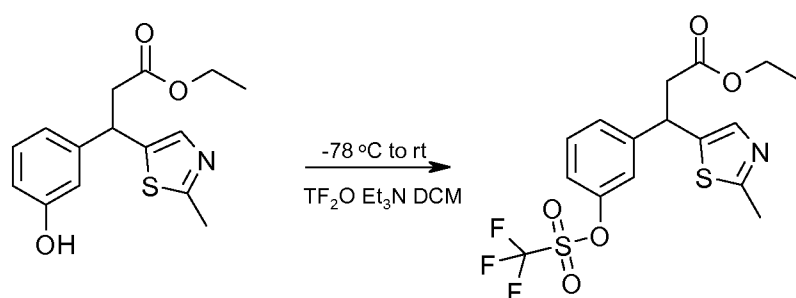
15

To a mixture of (Z)-ethyl 3-(3-(benzyloxy)phenyl)-3-(2-methylthiazol-5-yl)acrylate (1.6 g, 4.22 mmol), HOAc (1.0 mL), and H_2O (2.0 mL) in EtOH (20 mL) and THF (10 mL) was added Pd/C (10%, dry, 1.6g) under N_2 atmosphere. The mixture was stirred at

room temperature under H₂ atmosphere for 48 hrs. After filtration through a pad of celite and evaporation of the solvent, the residue was purified by silica gel chromatography eluting with petroleum ether/EtOAc (3:1) to afford 3-(3-(benzyloxy)phenyl)-3-(2-methylthiazol-5-yl)propanoate (1.1 g, 90%) as a yellow oil.

5

Ethyl 3-(2-methylthiazol-5-yl)-3-(3-(trifluoromethylsulfonyloxy)phenyl)propanoate



10

To a stirred solution of ethyl 3-(3-(benzyloxy)phenyl)-3-(2-methylthiazol-5-yl)propanoate (1.1 g, 3.4 mmol) and Et₃N (0.7 g, 6.8 mmol) in DCM (10 mL) at -78 °C was added trifluoromethanesulfonic anhydride (1.9 g, 6.8 mmol) dropwise over 5 min. The resulting mixture was stirred at -78 °C for 30 minutes and warmed to room temperature with stirring overnight. The mixture was cooled to 0 °C and treated with water (50 mL). The organic layer was washed with brine (50 mL), dried over Na₂SO₄. After filtration and evaporation of the solvent, the residue was purified by silica gel column chromatography eluting with petroleum ether/ethyl acetate (4:1) to give ethyl 3-(2-methoxy-5-(trifluoromethylsulfonyloxy)phenyl)-3-(thiazol-2-yl)propanoate (450 mg, 32%) as yellow oil.

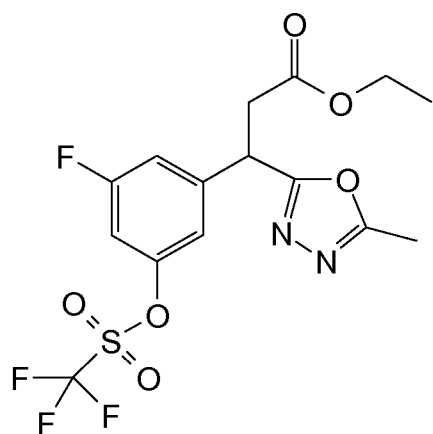
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Synthesis of intermediate P8

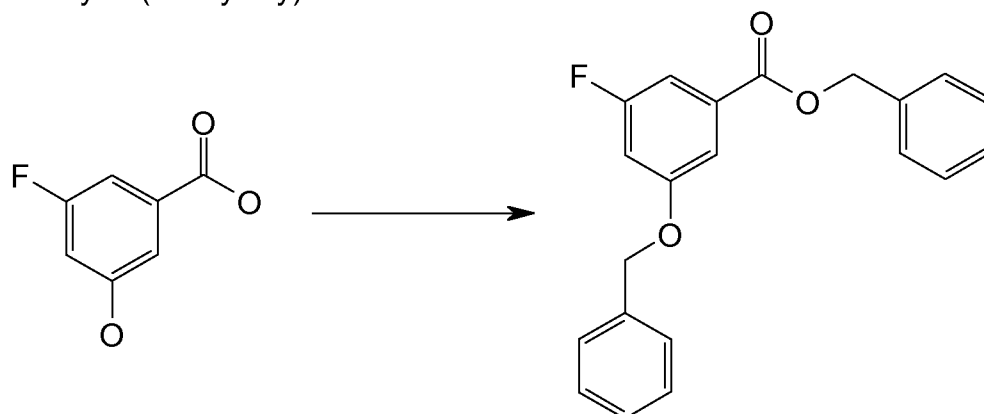
Ethyl 3-(3-fluoro-5-(trifluoromethylsulfonyloxy)phenyl)-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoate

25



A033

Benzyl 3-(benzyloxy)-5-fluorobenzoate



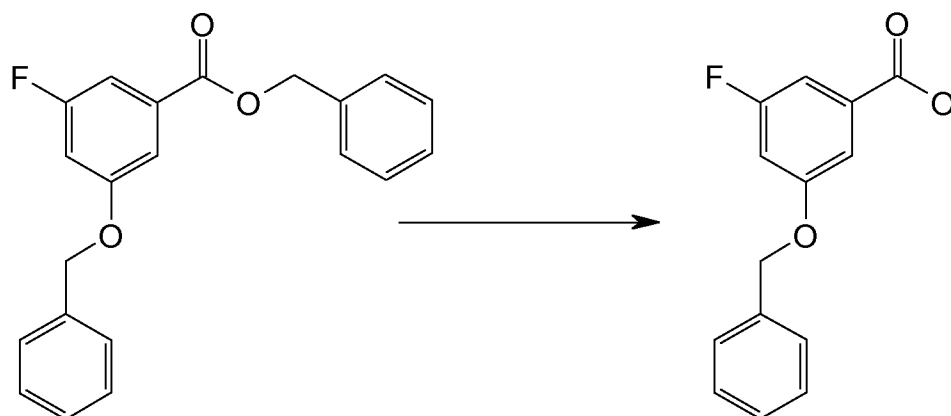
5 The mixture of the starting material (10 g, 64.1 mmol), K_2CO_3 (26.5 g, 192.3 mmol) and BnBr (24.1 g, 141.0 mmol) in DMF (150 mL) was stirred at r.t. for 12 h. After the reaction was completed, the mixture was added water (200 mL), extracted with EA (150 mL x 3). The combined organic layers were washed with LiCl aq., brine, dried

10 over Na_2SO_4 . After filtration and evaporation of the solvent, the residue was purified by silica gel chromatography (EA/PE = 1:30) to give product (17g, 78.9%) as color less oil.

15

20 3-(Benzyloxy)-5-fluorobenzoic acid

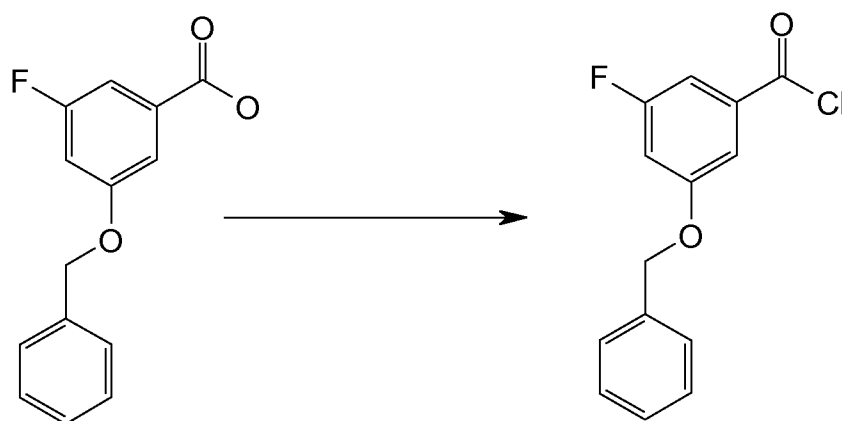
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To the solution of the starting material (17 g, 50.6 mmol) in MeOH (50 mL) was added NaOH aq (50mL, 2.0mmol/L). The reaction solution was stirred and heated to 50°C overnight. After the reaction was completed, 40 mL of HCl (3N) was added. The mixture was extracted with EA (100 mL x 3). The combined organic layers were washed with brine and dried over Na₂SO₄. After filtration and evaporation of the solvent, the product (12.5g, 99.6 %) was obtained.

10

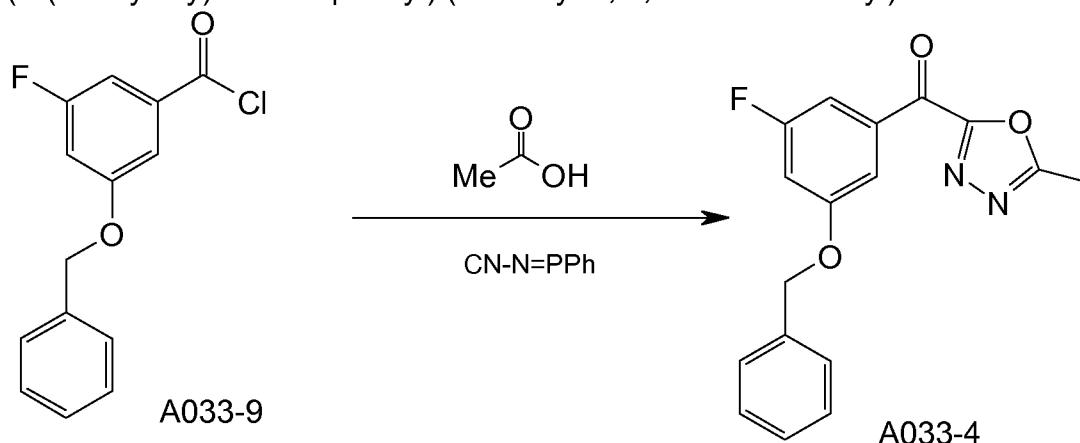
3-(Benzyloxy)-5-fluorobenzoyl chloride



15

To a mixture of the starting material (4.5 g, 18.3 mmol) in MeCN (40 mL) was added 10mL of SOCl₂. The mixture was stirred and heated to 80°C for 0.5h. After cooling, the solution was evaporated and the residue (4.6g, crude) was used for next step directly.

(3-(Benzyloxy)-5-fluorophenyl) (5-methyl-1, 3, 4-oxadiazol-2-yl)methanone

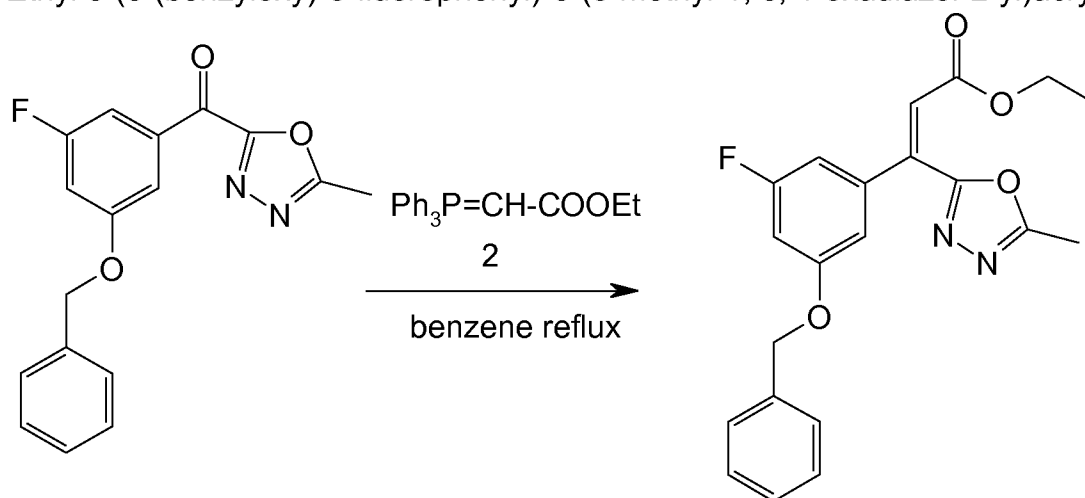


The mixture of starting material (4.6 g, 17.4 mmol), $\text{Ph}_3\text{P}=\text{N-CN}$ (5.3 g, 17.5 mmol) in CH_2Cl_2 (40 mL) was stirred at r.t. for 2h. Acetic acid (2.1 g, 34.8 mmol) was added.

- 5 After the addition, NEt_3 (3.5 g, 34.8 mmol) in 5mL of CH_2Cl_2 was added drop wise.

The mixture was stirred at r.t. overnight. After the reaction was completed, the solution was evaporated and the residue was purified on column chromatography (silica gel, EA/PE = 1: 10) to give the product (3.0g, 55.2 %) as a yellow oil.

- 10 Ethyl 3-(3-(benzyloxy)-5-fluorophenyl)-3-(5-methyl-1, 3, 4-oxadiazol-2-yl)acrylate

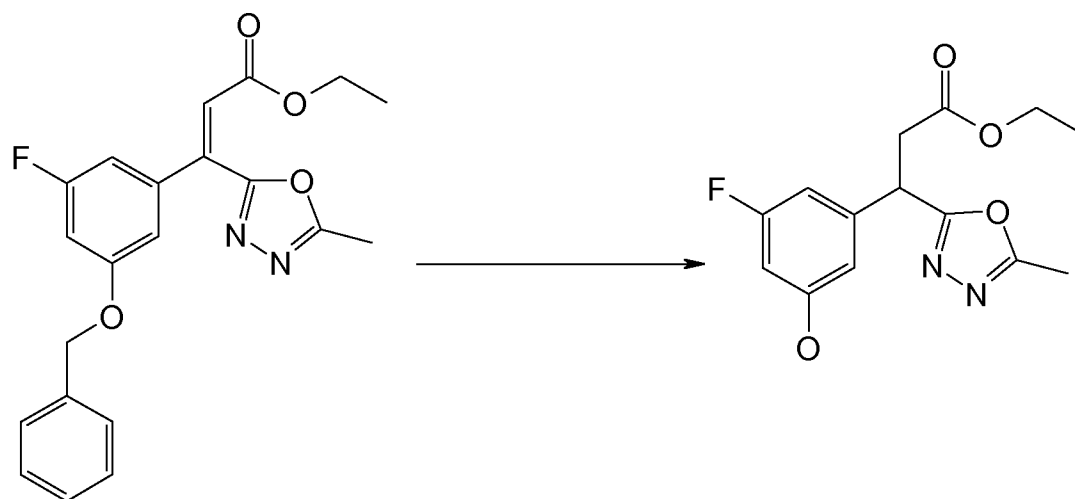


A mixture of starting material (3.0 g, 9.6 mmol) and 2 (3.3g, 9.6 mmol) in benzene (20 mL) was stirred at 100°C for 48h. After cooling, the volatile was removed *in vacuo* and the residue was purified on column chromatography (silica gel, EA/PE =

- 15 1: 8) to give the product (2.8g, 76.0 %) as red oil.

Ethyl 3-(3-fluoro-5-hydroxyphenyl)-3-(5-methyl-1, 3, 4-oxadiazol-2-yl)propanoate

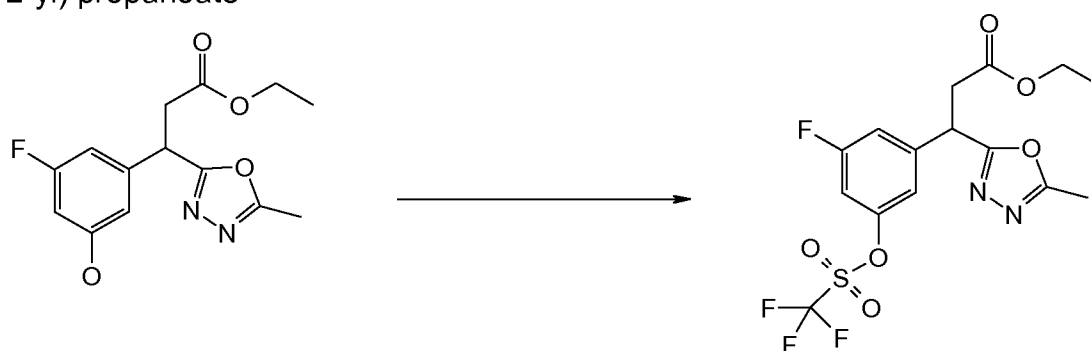
67



To a solution of starting material (5.0 g, 13.1 mmol) in EtOH (100 mL), THF (100 mL), AcOH (2mL) and Pd/C (500 mg) was added. The system was evacuated and then refilled with H₂ for 3 times. The mixture solution was stirred at r.t. overnight under H₂ atmosphere. After the reaction was completed, the mixture was filtrated via on celite pad and the filtrate was evaporated *in vacuo*. The product (4.0g) was obtained with yield 96.3%.

10

Ethyl 3-(3-fluoro-5-(trifluoromethylsulfonyloxy) phenyl)-3-(5-methyl-1, 3, 4-oxadiazol-2-yl) propanoate



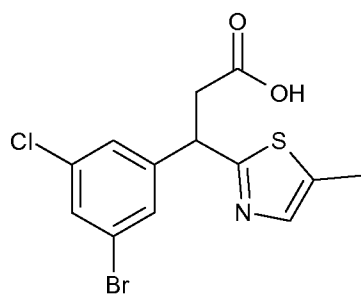
15

A mixture of starting material (2.4 g, 8.16 mmol) and DIPEA (3.2 g, 24.5 mmol) in CH₂Cl₂ (90 mL) was stirred for 0.5h at -78°C. Tf₂O (4.6 g, 16.3 mmol) in CH₂Cl₂ (10 mL) was added dropwise at -78°C. The mixture solution was stirred at -78°C for 1h then allowed warming to r.t. overnight. The volatiles were removed *in vacuo* to give a

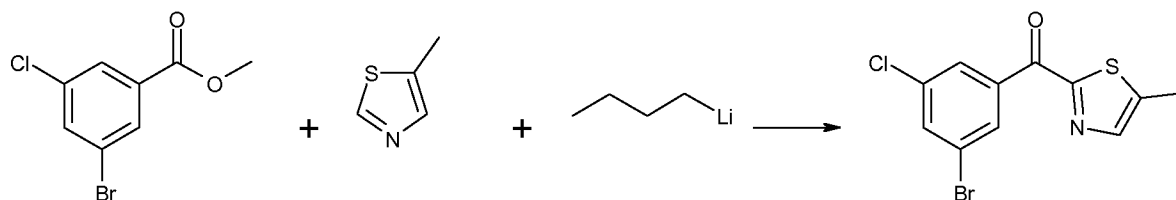
residue which was purified by column chromatography (elution was 12% EA in PE) to give product (2.2g, 63.2 %) as yellow liquid.

5 Synthesis of intermediate A1

3-(3-Bromo-5-chloro-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid

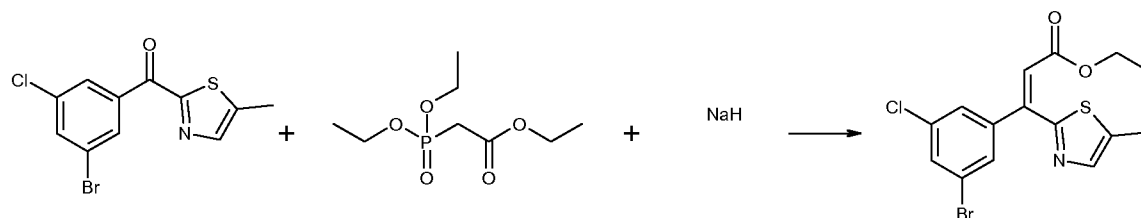


10 (3-Bromo-5-chloro-phenyl)-(5-methyl-thiazol-2-yl)-methanone



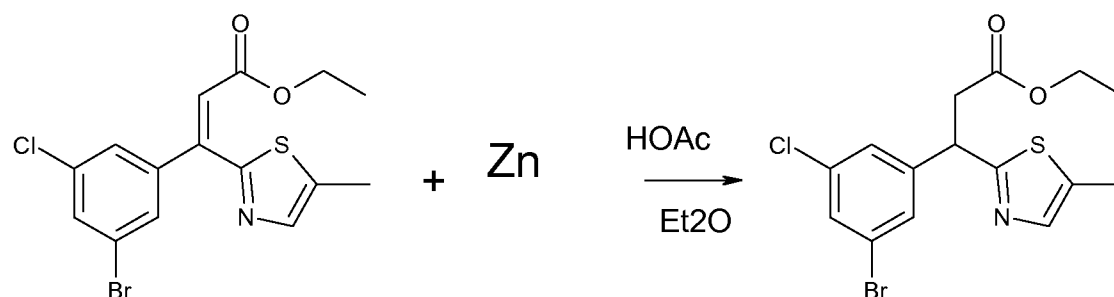
To a stirring mixture of 5-methylthiazole (4.37 g, 44.1 mmol) in dry THF (30L) under nitrogen atmosphere was added n-BuLi (44.1mmol, 17.64 ml solution in hexanes) drop wise over 10 minutes at -78 °C. The mixture was stirred between -78 °C and -60 °C for 30 minutes and then cooled to -78 °C. A solution of 3-bromo-5-chloro-benzoic acid methyl ester (11.0 g, 44.1 mmol) in THF (20 mL) was added drop wise over 10 minutes. The resulting mixture was stirred at -78 °C for 30 minutes and warmed to room temperature with stirring overnight. The mixture was cooled to 0 °C and treated with water (50 mL). The resulting mixture was adjusted to pH around 6 with HCl (1N) and extracted with EtOAc (100 mL x 3). The combined organic layers were dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography eluting with petroleum ether/EtOAc mixtures to afford (3-bromo-5-chloro-phenyl)-(5-methyl-thiazol-2-yl)-methanone (14g, yield = 100%) as a solid.

(Z)-3-(3-Bromo-5-chloro-phenyl)-3-(5-methyl-thiazol-2-yl)-acrylic acid ethyl ester



- 5 To a suspension of NaH (60%, 1.77 g, 44,2 mmol) in dry THF (30 mL) was added ethyl 2-(diethoxyphosphoryl)acetate (10.12 g, 44.2 mmol) at -78°C . After being stirred at -78°C for 1 hour, a solution of 14g (44,2 mmol) (3-Bromo-5-chloro-phenyl)-(5-methyl-thiazol-2-yl)-methanone in THF (10 mL) was added drop wise over 30 min. The mixture was stirred at -78°C for 30 min and then warmed to room temperature with stirring overnight. The mixture was cooled to 0°C and treated with water (50 mL), pH value was adjusted to 6 with HCl (0.5 N) and extracted with EA (50 mL x 3). The combined organic layers were dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was purified by silica gel chromatography eluting with petroleum ether/EtOAc mixtures to afford (Z)-3-(3-Bromo-5-chloro-phenyl)-3-(5-methyl-thiazol-2-yl)-acrylic acid ethyl ester as an gel (18g, yield = 106%) The material is used without further purification in the next step.

3-(3-Bromo-5-chloro-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid ethyl ester

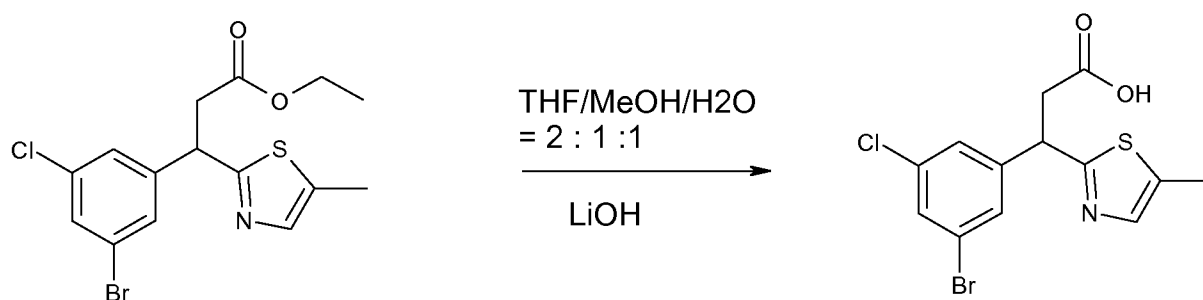


- 20 18g (46,55 mmol) of (Z)-3-(3-Bromo-5-chloro-phenyl)-3-(5-methyl-thiazol-2-yl)-acrylic acid ethyl ester are dissolved in 250 ml Diethylether and 80ml AcOH. 3 Eq (140 mmol) of zinc dust are added over a period of 60 minutes. The resulting mixture is

allowed to stir overnight, diluted with 100ml EtOAc and filtrated over a pad of Celite. The solvent is removed in vacuo, the residue taken up in EtOAc, washed with brine and dried over MgSO₄. After evaporation of the solvent 17g (Yield = 99%) of crude product are obtained, which are used in the last step.

5

3-(3-Bromo-5-chloro-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid



17g (43,73mmol) of 3-(3-Bromo-5-chloro-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid ethyl ester are dissolved in 30ml of a THF/MeOH/H₂O = 2 : 1 : 1 mixture and LiOH (2Eq) is added. After stirring overnight at RT, the solvent is removed in vacuo and the crude material is treated with 30ml of EtOAc and 30ml of water with pH = 4. The organic phase is separated, dried over Na₂SO₄ and the solvent removed in vacuo. The crude material is recrystallized from heptane / EtOAc mixtures delivering 9,2g (Yield = 58%) of 3-(3-Bromo-5-chloro-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid.

20 According to the procedures described above the following additional intermediates can be prepared:

A4: 3-(3-Bromo-5-fluoro-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid

Using methyl-3-bromo-5-fluoro-benzoate as starting material.

25

A6: 3-(3-Bromo-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid

Using methyl-benzoate as starting material.

A7: 3-(3-Bromo-phenyl)-3-(4,5-dimethyl-thiazol-2-yl)-propionic acid

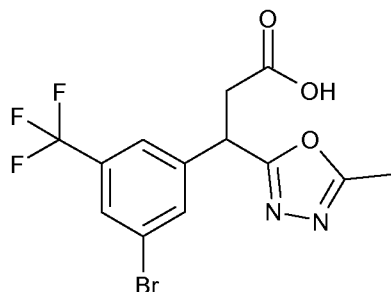
Using methyl-benzoate and 4,5-dimethyl-1,3-thiazole as starting materials.

C2: 3-(3-Bromo-phenyl)-3-thiazol-2-yl-propionic acid

- 5 This material is commercially available from ZereneX (CAS-Number: 1082829-38-6), but can be prepared accordingly starting from methyl-benzoate and 1,3-thaizole as starting materials.

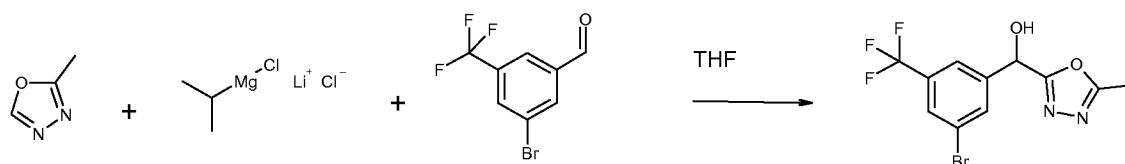
Synthesis of intermediate A2

3-(3-Bromo-5-trifluoromethyl-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid



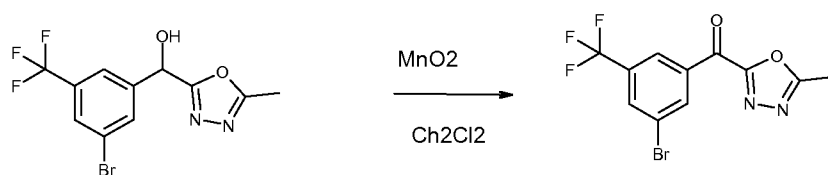
5

(3-Bromo-5-trifluoromethyl-phenyl)-(5-methyl-[1,3,4]oxadiazol-2-yl)-methanol



2,79g (33,2 mmol) of 2-methyl-1,3,4-oxadiazole are dissolved in 45ml THF at -5°C,
 33,2 mmol (25,54ml) of isopropyl magnesium chloride-lithium chloride complex are
 added during 20 minutes while the temperature is kept < 0°C. The mixture is kept at
 0°C for 30 minutes, then 0,8 Eq (6,7g) of 3-Bromo-5-trifluoromethyl-benzaldehyde
 are added and the resulting mixture is allowed to reach RT and stirred for additional
 60 minutes. The reaction is then stopped by the addition of 10ml saturated NH₄Cl
 solution. 50ml MTBE are added, the organic phase is separated, washed with brine
 and dried over MgSO₄. The solvent is removed in vacuo and the crude material (9g)
 is used without further purification in the next step:

(3-Bromo-5-trifluoromethyl-phenyl)-(5-methyl-[1,3,4]oxadiazol-2-yl)-methanone

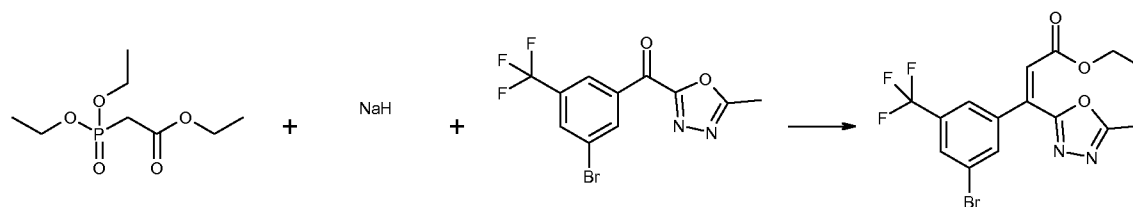


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The crude material from the first step is dissolved in 100ml CH₂Cl₂ and 20g MnO₂ are added. The resulting mixture is stirred for 60 minutes at RT and filtrated. After evaporation of the solvent 9,06g (98%) of crude material is obtained, which is used without further purification in the next step.

5

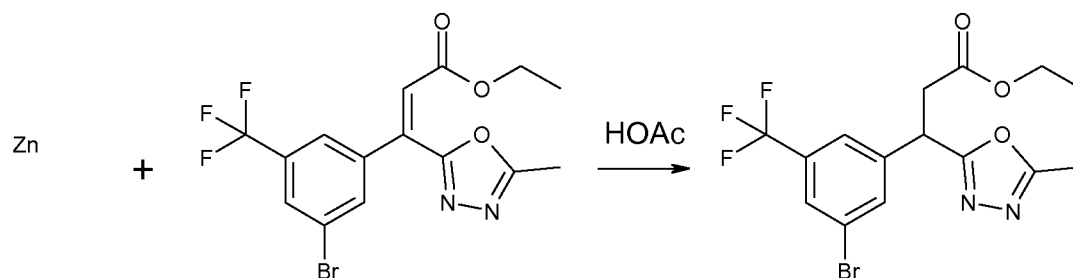
(Z)-3-(3-Bromo-5-trifluoromethyl-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-acrylic acid ethyl ester



To a suspension of NaH (60%, 1.13 g, 28,35 mmol) in dry THF (20 mL) was added ethyl 2-(diethoxyphosphoryl)acetate (6.49 g, 28,35 mmol) at -78 °C. After being stirred at -78 °C for 1 hour, a solution of (3-Bromo-5-trifluoromethyl-phenyl)-(5-methyl-[1,3,4]oxadiazol-2-yl)-methanone (9,047g, 27mmol) in 20ml THF was added drop wise over 30min. The mixture was stirred at -78 °C for 30 min and then warmed to room temperature followed by stirring overnight. The mixture was cooled to 0 °C and treated with water (50 mL), pH value was adjusted to 6 with HCl (0.5 N) and extracted with EtOAc (50 mL x 3). The combined organic layers were dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by silica gel chromatography eluting with a petroleum ether/EtOAc mixture to afford 9g of the desired product (Yield: 90%).

20

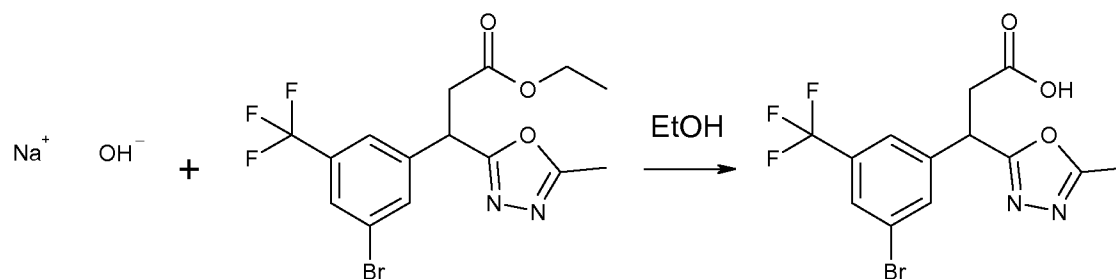
Synthesis of 3-(3-Bromo-5-trifluoromethyl-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid ethyl ester



11.2 g (27,64 mmol) (Z)-3-(3-Bromo-5-trifluoromethyl-phenyl)-3-(5-methyl-

- 5 [1,3,4]oxadiazol-2-yl)-acrylic acid ethyl ester is dissolved in 200 ml acetic acid and 14,1g Zinc dust (215,6 mmol, 7.8 mmol) is added slowly over a period of 30 minutes. The temperature of the reaction mixture is not allowed to reach > 30°C during the addition of the zinc dust. The resulting mixture is allowed to stir overnight. After filtration 200ml EtOAc and 200ml water are added and the organic phase is
- 10 separated and washed with brine. After removal of the solvent the crude material is purified by column chromatography on silica gel using heptane /EtOAc = 3:1 as eluent. 3,97g of 3-(3-Bromo-5-trifluoromethyl-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid ethyl ester (yield = 35%) are isolated.

- 15 Synthesis of 3-(3-Bromo-5-trifluoromethyl-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid



3,97g (9,75mmol) of 3-(3-Bromo-5-trifluoromethyl-phenyl)-3-(5-methyl-

- 20 [1,3,4]oxadiazol-2-yl)-propionic acid ethyl ester are dissolved in 10ml ethanol and 6,34ml of a 2N NaOH solution are added. The mixture is allowed to stir for 4 hours at RT, the ethanol is removed and the residue taken up in 30ml EtOAc and 20ml water.

After phase separation the organic phase is dried over MgSO_4 and the solvent removed in vacuo to deliver 3.3g (Yield = 89 %) of the desired product 3-(3-Bromo-5-trifluoro-methyl-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid.

- 5 According to the procedure described above the following additional intermediates are prepared:

A8: 3-(3-Bromo-5-chloro-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid
Using 3-bromo-5-chloro-benzaldehyde as starting material

10

A9: 3-(3-Bromo-5-fluoro-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid
Using 3-Bromo-5-fluoro-benzaldehyde as starting material

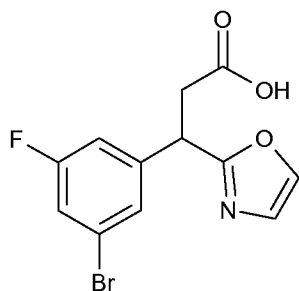
- 15 Using 5-bromo-nicotinaldehyde as starting material

C1: 3-(3-Bromo-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid
This intermediate is commercially available from Aurora Building Blocks (CAS-
Number: 1082916-80-0), but can be obtained using 3-bromo-benzaldehyde

20

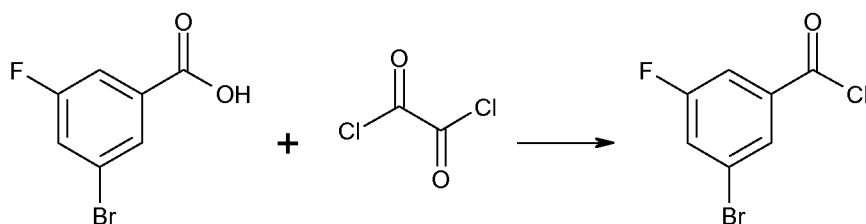
Synthesis of intermediate A3

3-(3-Bromo-5-fluoro-phenyl)-3-oxazol-2-yl-propionic acid



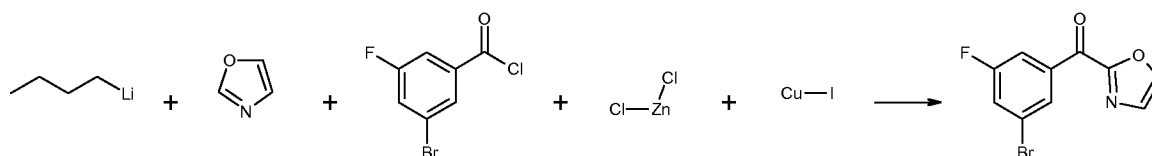
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Synthesis of 3-Bromo-5-fluoro-benzoyl chloride



10g of 3-bromo-5-fluoro-benzoic acid are suspended in 100ml of DCM, 0,5ml DMF and 4.5ml oxalyl chloride (1,15 Eq) are added and the resulting mixture is stirred for 100 minutes at RT. The solvent is removed by vacuo and the benzoic acid chloride is isolated by distillation.

Synthesis of (3-Bromo-5-fluoro-phenyl)-oxazol-2-yl-methanone

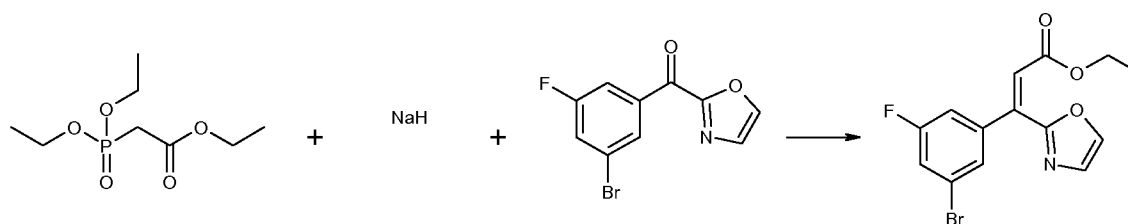


To a stirring mixture of 1,3-oxazole (3,46 g, 50mmol) in dry THF (50mL) was added n-BuLi (1,1 Eq, 20,01 ml solution in hexanes) drop wise over 10 minutes at -78 °C under nitrogen atmosphere. After stirring for 30 minutes at -78°C zinc chloride (0,1mmol, 2,2Eq) as a 2M solution in 5-methyl-tetrahydrofurane was added within 30 minutes and the mixture was allowed to reach -20°C during the addition. Then the mixture was stirred at 0°C for 40 minutes, CuI (8,66g, 45,5mmol) was added and stirring continued for additional 10 minutes. 3-Bromo-5-fluoro-benzoyl chloride (45,5mmol, 8,66g) was dissolved in 50ml of THF and added to the metallated 1,3-oxazole. Stirring was continued until complete conversion was observed by LCMS and 50ml water and 50 ml of a 0,5 M citric acid solution were added. The solid material from the reaction mixture was filtered off and washed with 5-methyl-thf. The organic phase was separated, washed with brine and dried over MgSO₄. 10,8g of crude material (yield = 88%) were obtained and used directly in the next step.

The mixture was stirred between -78 °C and -60 °C for 30minutes and then cooled to -78 °C. A solution of 3-Bromo-5-chloro-benzoic acid methyl ester (11.0 g, 44,1 mmol) in THF (20 mL) was added drop wise over 10 minutes. The resulting mixture was stirred at -78 °C for 30 minutes and warmed to room temperature with stirring

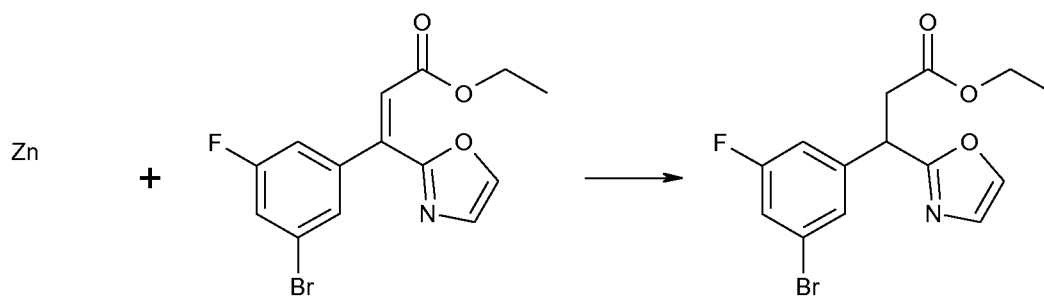
overnight. The mixture was cooled to 0 °C and treated with water (50 mL). The resulting mixture was adjusted to pH around 6 with HCl (1N) and extracted with EtOAc (100 mL x 3). The combined organic layers were dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography eluting with petroleum ether/EtOAc mixtures to afford (3-Bromo-5-chloro-phenyl)-(5-methyl-thiazol-2-yl)-methanone (14g, yield = 100%) as a solid.

Synthesis of (Z)-3-(3-Bromo-5-fluoro-phenyl)-3-oxazol-2-yl-acrylic acid ethyl ester



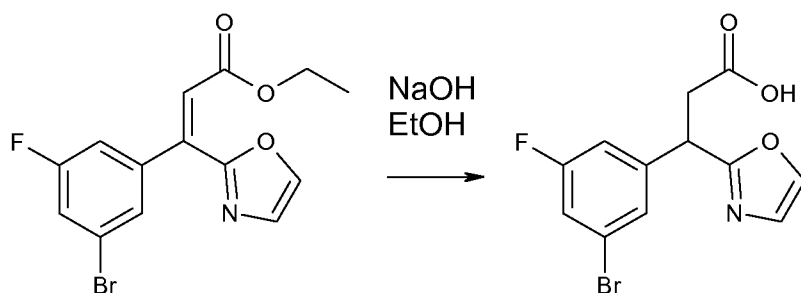
To a suspension of NaH (60%, 1.6 g, 40 mmol) in dry THF (30 mL) was added ethyl 2-(diethoxyphosphoryl)acetate (9.15 g, 40 mmol) at -78°C. After being stirred at -78°C for 1 hour, a solution of (3-Bromo-5-fluoro-phenyl)-oxazol-2-yl-methanone (10.8g, 40mmol) in 20ml THF was added drop wise over 30min. The mixture was stirred at -78 °C for 30 min and then warmed to room temperature followed by stirring overnight. The mixture was cooled to 0 °C and treated with water (50 mL), pH value was adjusted to 6 with HCl (0.5 N) and extracted with EtOAc (50 mL x 3). The combined organic layers were dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by silica gel chromatography eluting with a petroleum ether/EtOAc mixture to afford 9.5g of the desired product (Yield: 61%)

Synthesis of 3-(3-Bromo-5-fluoro-phenyl)-3-oxazol-2-yl-propionic acid ethyl ester



9.5 g (27,93 mmol) (Z)-3-(3-Bromo-5-fluoro-phenyl)-3-oxazol-2-yl-acrylic acid ethyl ester is dissolved in 200 ml acetic acid and 9,5g Zinc dust (11Ea)) is added slowly over a period of 30 minutes. The temperature of the reaction mixture is not allowed to reach > 30°C during the addition of the zinc dust. The resulting mixture is allowed to stir overnight. After filtration 200ml EtOAc and 200ml water are added and the organic phase is separated and washed with brine. After removal of the solvent the crude material is purified by column chromatography on silica gel using heptane /EtOAc = 3:1 as eluent. 9,50g of 3-(3-Bromo-5-fluoro-phenyl)-3-oxazol-2-yl-propionic acid ethyl ester are isolated (yield = 615)

Synthesis of 3-(3-Bromo-5-fluoro-phenyl)-3-oxazol-2-yl-propionic acid



9,50g (27,76mmolmmol) of 3-(3-Bromo-5-fluoro-phenyl)-3-oxazol-2-yl-propionic acid ethyl ester are dissolved in 30ml ethanol and 20,82ml of a 2N NaOH solution are added. The mixture is allowed to stir for 4 hours at RT, the ethanol is removed and the residue taken up in 30ml EtOAc and 20ml water. After phase separation the organic phase is dried over Mg₂SO₄ and the solvent removed in vacuo to deliver 8,35g (Yield = 58 %) of the desired product 3-(3-Bromo-5-fluoro-phenyl)-3-oxazol-2-yl-propionic acid.

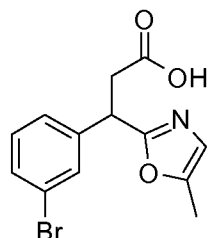
According to this procedure the following additional intermediates are prepared:

A5: 3-(3-Bromo--phenyl)-3-oxazol-2-yl-propionic acid

- 5 Using 3-bromo benzoyl-chloride as starting material

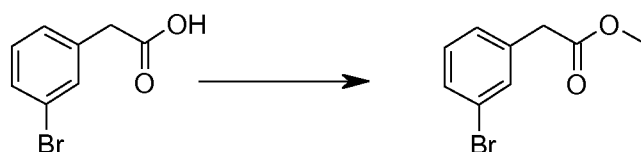
Synthesis of intermediate A12

3-(3-Bromophenyl)-3-(5-methyloxazol-2-yl) propanoic acid



5

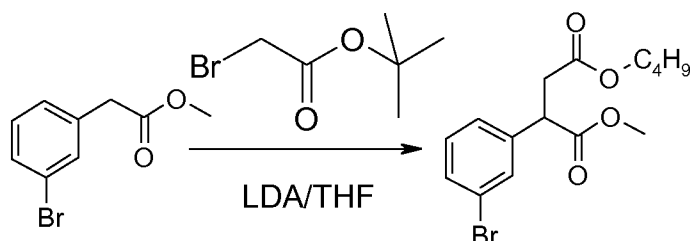
Methyl 2-(3-bromophenyl) acetate



To the solution of the starting material (12.0 g, 55.8 mmol) in MeOH (200 mL) was added SOCl_2 (2 mL). The reaction solution was stirred and heated to reflux overnight. After the reaction was completed, 200 mL of water and 30 mL of NaHCO_3 aq. were added. The mixture was extracted with EA (150 mL x 3). The combined organic layers were washed with brine and dried over Na_2SO_4 . After filtration and evaporation of the solvent, the product (11.0g, 86.1 %) was obtained.

15

4-tert-butyl 1-methyl 2-(3-bromophenyl) succinate

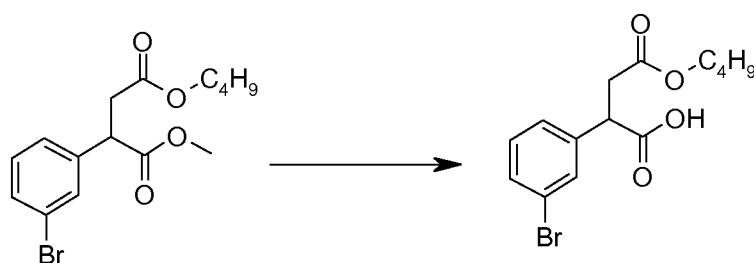


The starting material (11.0 g, 48.0 mmol) in THF (100 mL) was added dropwise to LDA in THF (2M, 24 mL) at -78°C . The resulting mixture was stirred as such for 2h before tert-butyl bromoacetate (10.3 g, 52.8 mmol) in THF (20 mL) was added over 10 min. The reaction mixture was stirred for 2 h at -78°C and allowed to warm slowly to 0°C . The reaction was quenched with saturated aqueous NH_4Cl (50 mL). The

organic phase was extracted with ethyl acetate (100 mL x 3). The organic layers were combined and dried over Na₂SO₄. After filtration and evaporation of the solvent, the resulting residue was purified on column chromatography (silica gel, PE: EA = 20:1) to give the target product (14.4 g, 87.5%) as yellow oil.

5

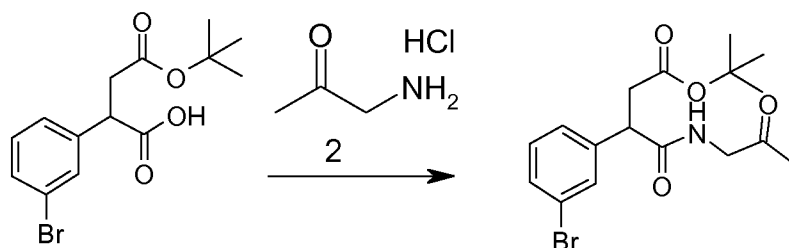
2-(3-Bromophenyl)-4-tert-butoxy-4-oxobutanoic acid



Lithium hydroxide (2.6 g, 109.2 mmol) was added to a solution of the starting material (12.5 g, 36.4 mmol) in MeOH/THF/H₂O (1:1:1, 500 mL). The resulting mixture was stirred at room temperature overnight. After the reaction was completed, the volatile was removed *in vacuo* and aqueous HCl (6N) was added dropwise to adjust the solution to pH = 4. The solution was extracted with EA (150 mL x 3) and the organic layers were combined, dried over Na₂SO₄. After filtration and evaporation of the solvent, the crude product (12.0g, 99%) was obtained as yellow oil.

15

Tert-butyl 3-(3-bromophenyl) -4-oxo-4- (2-oxopropylamino) butanoate

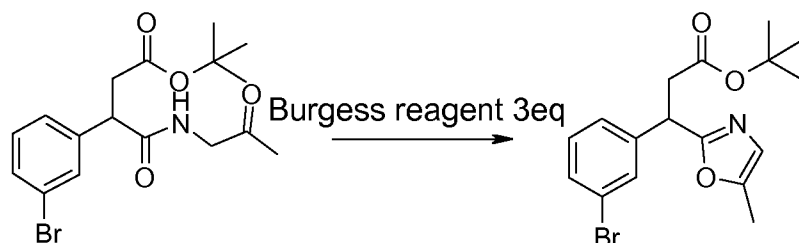


HATU (18.0 g, 47.4 mmol) and DIPEA (10.2 g, 79 mmol) were added sequentially to the starting material (5.2 g, 15.8 mmol) in DMF (80 mL), followed by the addition of 2 (4.3 g, 39.5 mmol). The reaction mixture was stirred at r.t. overnight. The reaction was quenched with water (100 mL) and extracted with ethyl acetate (100 mL x 3). The combined organic extracts were dried over Na₂SO₄. After filtration and evaporation of the solvent, the residue was purified on reverse phase

20

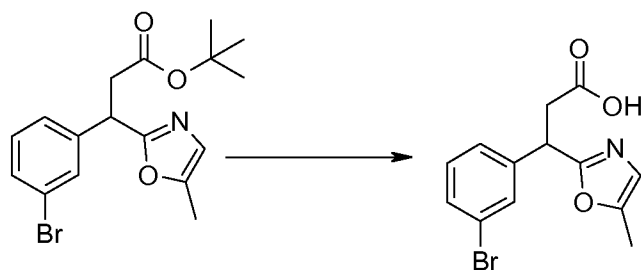
chromatography [A: CH₃CN; B: NH₄HCO₃/H₂O (0.01%), A: 20%-95% in 22minutes, RT=13-14min] to give the product (5.3g, 87.3%) as yellow oil.

5 Tert-butyl 3-(3-bromophenyl)-3-(5-methyloxazol-2-yl) propanoate



To the solution of the starting material (2.1 g, 5.5 mmol) in THF (20 mL) was added Burgess reagent (3.9 g, 16.5 mmol). The mixture solution was stirred and heated to 150 °C under Microwave for 30min. After the reaction completed, the volatile was removed *in vacuo* and the residue was purified by Reverse chromatography [A: CH₃CN; B: NH₄HCO₃.H₂O (0.01%), A: 20%-95% in 15minutes, RT=8.6-9.7min] to give the product (1.85g, 92%) as yellow oil.

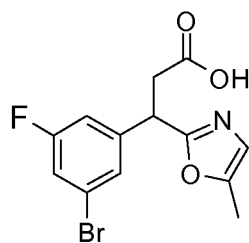
15 3-(3-Bromophenyl)-3-(5-methyloxazol-2-yl) propanoic acid



Lithium hydroxide (1.8 g, 76.5 mmol) was added to a solution of the starting material (5.6 g, 15.3 mmol) in THF/H₂O (1:1, 150 mL). The resulting mixture was stirred at room temperature overnight. After the reaction was completed, the volatile was removed *in vacuo* and aqueous HCl (6N) was added dropwise to adjust the solution to pH = 4. The solution was extracted with EA (150 mL x 3) and the organic layers were combined, dried over Na₂SO₄. After filtration and evaporation of the solvent, the product (4.15 g, 87.5%) was obtained as off-white solid.

Synthesis of intermediate A13:

3-(3-Bromo-5-fluoro-phenyl)-3-(5-methyloxazol-2-yl) propanoic acid



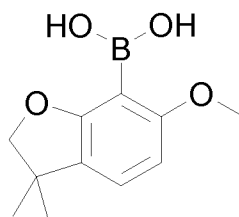
5

This intermediate is synthesized in analogy to intermediate A12 and starting from methyl 2-(3-bromo-5-fluorophenyl) acetate.

10

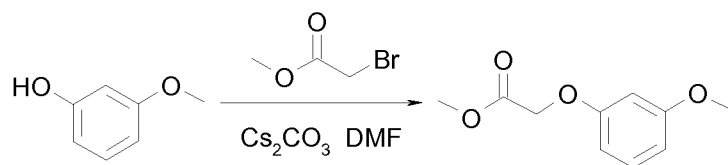
Synthesis of novel boronic acids:

6-methoxy-3,3-dimethyl-2,3-dihydrobenzofuran-7-ylboronic acid



15

Methyl 2-(3-methoxyphenoxy)acetate

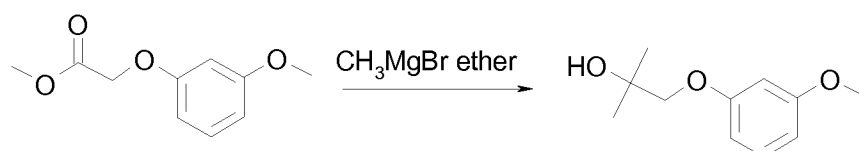


20 To a mixture of 3-methoxyphenol (20 g, 161.3 mmol) and Cs_2CO_3 (52.4 g, 161.3 mmol) in DMF (200 mL), methyl bromoacetate (24.5 g, 161.3 mmol) was added. The reaction mixture was stirred at room temperature under argon overnight. The

inorganic precipitate was filtered off, and the filtrate was concentrated under reduced pressure. The residue was partitioned between water (200 mL) and CH₂Cl₂ (200 mL x 3). The combined organic solution was dried over Na₂SO₄ and evaporated to give (28 g, 90%) product.

5

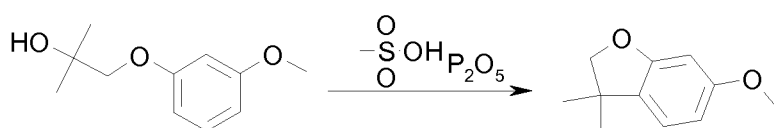
1-(3-methoxyphenoxy)-2-methylpropan-2-ol



10

To a solution of methyl 2-(3-methoxyphenoxy)acetate (28 g, 143 mmol) in ether (200 mL) cooled in an ice water bath was added drop wise a solution of methyl magnesium bromide (143 mmol, 3M in ether) in ether. After 1h the reaction mixture was poured into 500 mL of 2N HCl. The mixture was extracted with EA (200 mL x 3). The combined organic layers were dried over Na₂SO₄. After filtration and evaporation of the solvent, the intermediate (25 g 90%) was obtained.

15 6-methoxy-3,3-dimethyl-2,3-dihydrobenzofuran

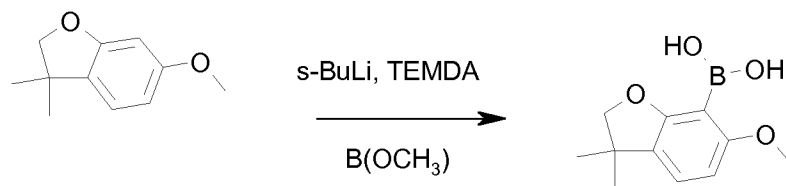


20

To a solution of phosphorous pentoxide (55.1 g, 383 mmol) in methanesulfonic acid (100 mL) was added drop wise 1-(3-methoxyphenoxy)-2-methylpropan-2-ol (25 g, 127.6 mmol) over a 30 min period. The reaction mixture was stirred for 3h at r.t. The reaction mixture was poured into 500 mL of ice water and extracted with ether (200 mL x 3). The combined extracts were dried and evaporated. The residue was purified by silica gel column chromatography eluting with petroleum ether/ ether (9:1) to afford 6-methoxy-3,3-dimethyl-2,3-dihydrobenzofuran (5.5 g, 25%) as a yellow oil.

25

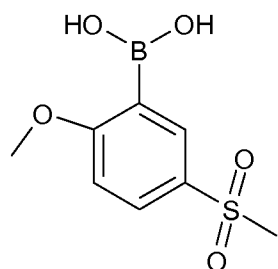
6-methoxy-3,3-dimethyl-2,3-dihydrobenzofuran-7-ylboronic acid



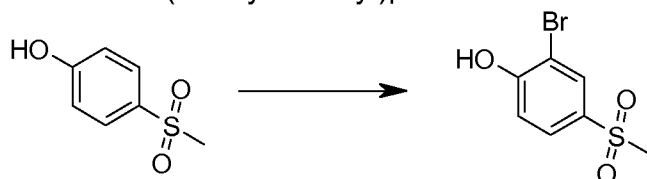
$s\text{-BuLi}$ (35 mL, 1.3M in cyclohexane) was added to TEMDA (6.6 mL) at -78°C drop wise within 30 min, 20 mL of THF was added to keep stirring. After 20min, 6-methoxy-3,3-dimethyl-2,3-dihydrobenzofuran (5.4 g, 30.3 mmol) in THF (30 mL) was added into the lithium solution slowly. The solution was stirred at -78°C for 1.5 h and trimethyl borate (18 mL) was added. The mixture was warmed to room temperature and stirred at room temperature overnight. The solution was acidified to pH= 5-6 and extracted with ethyl acetate. The organic layer was concentrated and the resulted residue was purified on column chromatography (silica gel, dichloromethane/ petroleum ether = 3/1) to give the crude product, which was sonicated with petroleum ether. The white precipitate was filtered and dried to give 6-methoxy-3,3-dimethyl-2,3-dihydrobenzofuran-7-ylboronic acid (3.55 g, 53%) as a white solid.

^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ 7.58 (brs, 2H), 7.07 (d, $J = 8.0\text{Hz}$, 1H), 6.36 (d, $J = 8.4\text{ Hz}$, 1H), 3.68 (s, 3H), 2.87 (s, 2H), 1.38 (s, 6H).

2-(2-methoxy-5-(methylsulfonyl)phenyl)-4,4,5,5-tetramethyl-1,3,2- dioxaborolane



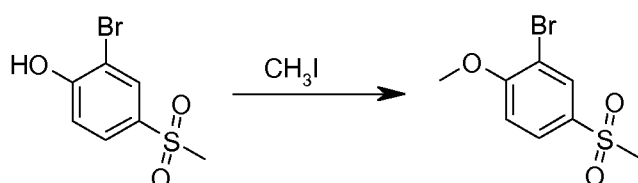
2-bromo-4-(methylsulfonyl)phenol



To a solution of 4-(methylsulfonyl)phenol (5 g, 29.1 mmol) in ether (100 mL) at -15°C was slowly added acetic acid (5 mL). To this cold solution was slowly added Br_2 (5.1

g, 32.0 mmol) and the reaction was stirred at -10°C for 1h and then warmed to r.t. and stirred for further 10h. After the reaction was completed, 100 mL of sat. NaHCO₃ aq. was added and the mixture was extracted with EA (50 mL x 3). The combined organic layers were washed with brine and dried over Na₂SO₄. After filtration and evaporation of the solvent, 2-bromo-4-(methylsulfonyl)phenol (3.2g, 43.6 %) was obtained as a white solid.

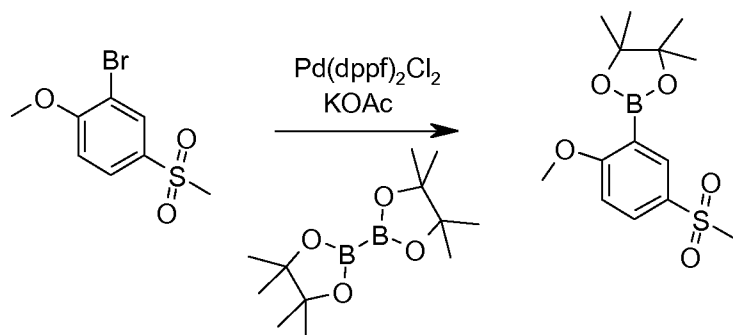
2-bromo-1-methoxy-4-(methylsulfonyl) benzene



To a solution of 2-bromo-4-(methylsulfonyl) phenol (3.2 g, 12.7 mmol) in DMF (35 mL) at 0 °C was added K₂CO₃ (5.3g, 38.1 mmol). The reaction was stirred at 0°C for 0.5h and to this cold solution was slowly added CH₃I (2.7 g, 19.1 mmol) drop wise.

The reaction was stirred at r.t. overnight. After the reaction was completed, 80 mL of ice-water was added. The mixture was extracted with EA (60 mL x 3). The combined organic layers were washed with brine and dried over Na₂SO₄. After filtration and evaporation of the solvent, the obtained residue was purified on column chromatography (silica gel, ethyl acetate / petroleum ether = 1:4) to give 2-bromo-1-methoxy-4-(methylsulfonyl)benzene (2.12 g, 63.0 %) as yellow solid.

2-(2-methoxy-5-(methylsulfonyl)phenyl)-4,4,5,5-tetramethyl-1,3,2- dioxaborolane

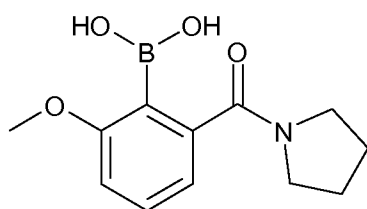


A mixture of 2-bromo-1-methoxy-4-(methylsulfonyl)benzene (2.12 g, 8 mmol), 4,4,4',4',5,5,5',5'-octamethyl-2,2'-bi(1,3,2-dioxaborolane) (2.65 g, 10.4 mmol), potassium acetate (2.4 g, 24 mmol) and [1,1'-Bis(diphenylphosphino)ferrocene] dichloropalladium(II) (600 mg) in 1,4-dioxane (50 mL) were heated to 80°C overnight.

- 5 After cooling, the mixture was filtered. The filtrate was concentrated and the resulted residue was purified on column chromatography (silica gel, dichloromethane/petroleum ether = 3/1) to give 2-(2-methoxy-5-(methylsulfonyl)phenyl)-4,4,5,5-tetramethyl-1,3,2-dioxaborolane (200 mg, 8%) as a white solid.

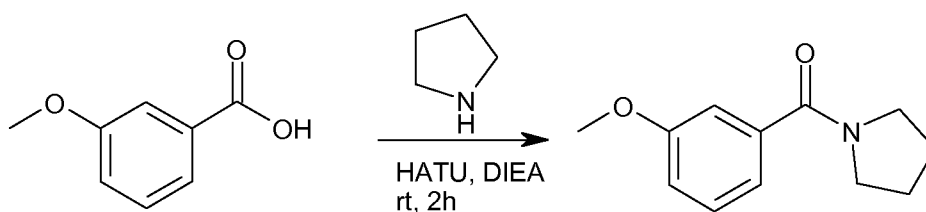
10 ¹H NMR (400 MHz, DMSO-*d*₆) δ 8.01-7.98 (m, 2H), 7.21 (d, *J*=8.8 Hz, 1H), 3.85 (s, 3H), 3.15 (s, 3H), 1.32 (s, 12H).

2-methoxy-6-(pyrrolidine-1-carbonyl)phenylboronic acid



15

(3-methoxyphenyl)(pyrrolidin-1-yl)methanone



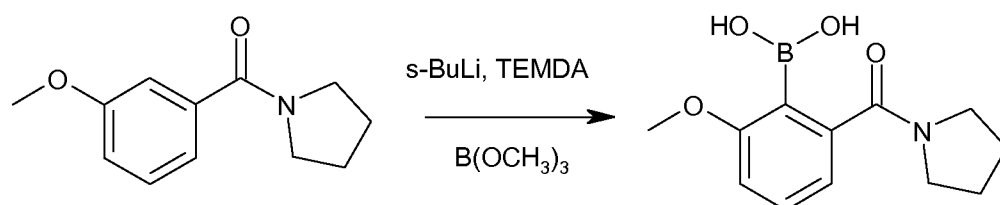
20

A solution of 3-methoxybenzoic acid (13.5 g, 88.8 mmol), pyrrolidine (9 mL, 133 mmol), HATU (33 g, 88.8 mmol) and DIEA (30 mL, 172 mmol) in DMF (100 mL) was stirred at room temperature for 2h. The solution was added water and extracted with ethyl acetate (200 mL x 3). The organic layer was dried and concentrated. The residue was purified by silica gel chromatography (eluted with petroleum ether: ethyl acetate =2:1 to 1:1) to give (3-methoxyphenyl)(pyrrolidin-1-yl)methanone (23 g, 100 %) as a yellow oil.

25

^1H NMR (400 MHz, CDCl_3) δ 7.30-7.26 (m, 1H), 7.05 (m, 2H), 6.94-6.92 (m, 1H), 3.80 (s, 3H), 3.59 (t, $J = 6.8$ Hz, 2H), 3.40 (t, $J = 6.8$ Hz, 2H), 1.94-1.83 (m, 4H).

2-methoxy-6-(pyrrolidine-1-carbonyl)phenylboronic acid

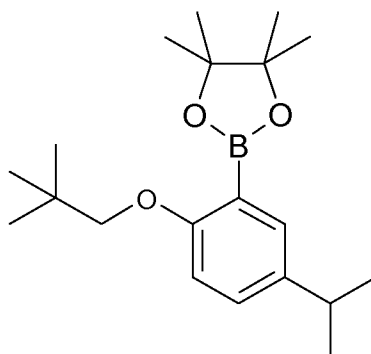


5

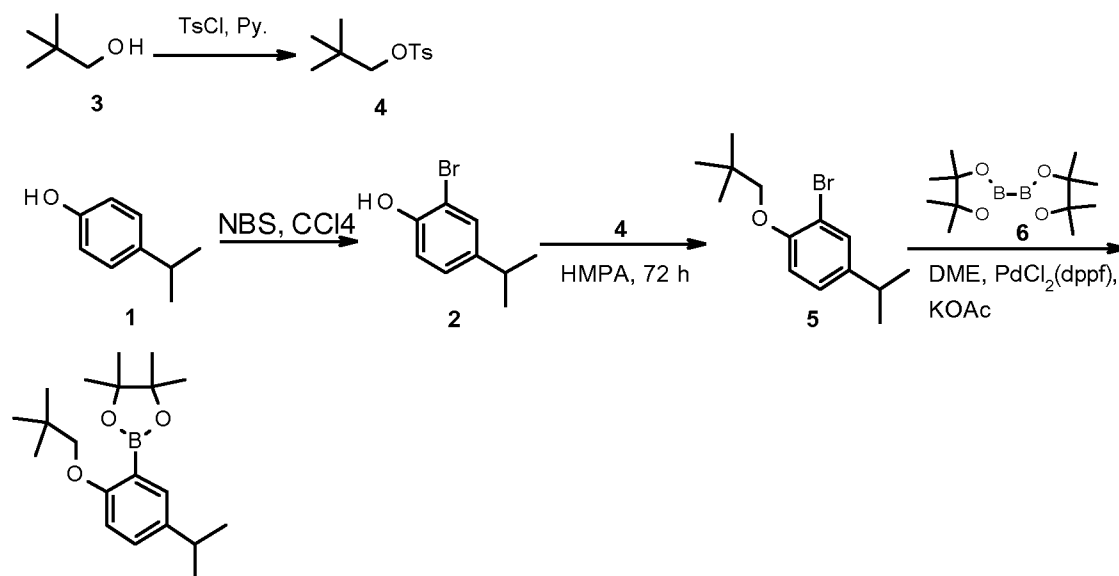
s-BuLi (70 mL, 1.3M in cyclohexane) was added to TEMDA (15.3 g, 131.9 mmol) at -78 °C dropwise within 30 min. After stirring for 20 min, (3-methoxyphenyl)(pyrrolidin-1-yl)methanone (16 g, 78 mmol) in THF (40 mL) was added into the lithium solution slowly. The solution was stirred at -78 °C for 1.5 h and trimethyl borate (36 mL, 322.6 mmol) was added. The mixture was warmed to room temperature and stirred at rt overnight. The solution was acidified to pH= 5-6 and the precipitate were filtered. The filter cake was washed with water and ethyl acetate to give 2-methoxy-6-(pyrrolidine-1-carbonyl)phenylboronic acid (5.5 g, 28%) as a white solid.

^1H NMR (400 MHz, DMSO-d_6) δ 7.49 (d, $J = 8.0$ Hz, 1H), 7.38 (t, $J = 8.0$ Hz, 1H), 7.12 (d, $J = 8.0$ Hz, 1H), 7.01 (brs, 2H), 4.08 (t, $J = 6.0$ Hz, 2H), 3.84-3.74 (m, 5H), 2.11-2.07 (m, 2H), 1.94-1.91 (m, 2H).

2-[2-(2,2-Dimethyl-propoxy)-5-isopropyl-phenyl]-4,4,5,5-tetramethyl-[1,3,2]dioxaborolane



General synthetic scheme:



5 Step 1: Synthesis of 2-Bromo-4-isopropyl-phenol (**2**)

To a solution of 4-isopropylphenol (**1**) (15 g, 110 mmol) in CCl₄ (150 mL) was added NBS (21.6 g, 121 mmol) in several batches at ice/water bath. The mixture was stirred at rt for overnight. DCM (300 ml) was added. The organic layer was washed with water (100 ml x 3), dried over Na₂SO₄ and filtrated, evaporated and purified by silica gel chromatography (from PE to PE:EA = 50:1) to give **2** as a yellow oil (10 g, 42%).

Step 2: Synthesis of Toluene-4-sulfonic acid 2,2-dimethyl-propyl ester (**4**)

To a stirred solution of **3** (10.0 g, 114 mmol) in pyridine (70 mL) at ice/water bath was added a solution of TsCl (32.5 g, 170 mmol) in pyridine (50 ml). The mixture was stirred at rt for overnight. The mixture was concentrated and water (200 ml) was added. Then it was extracted with EA (200 ml x 3). The combined organic layer was washed with HCl (1N, 150 ml), NaHCO₃ (aq, 150 ml), water (150 ml) and brine (150 ml), dried over MgSO₄, filtered, concentrated and purified by silica gel chromatography (from PE to PE:EA = 50:1) to afford **4** as a white solid (16.0 g, 58%).

Step 3: 2-Bromo-1-(2,2-dimethyl-propoxy)-4-isopropyl-benzene (**5**)

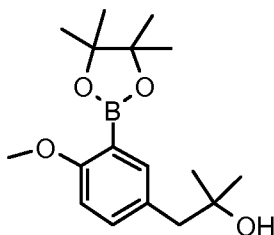
To a suspension of **2** (10.0 g, 45.6 mmol) and **4** (15.7 g, 65.1 mmol) in HMPA was added KOH (3.9 g, 70 mmol). The mixture was stirred vigorously at 100 °C for 72 h. The mixture was poured into water (300 ml). Then it was extracted with EA (200 ml x 3). The combined organic layer was washed with HCl (1N, 200 ml), NaHCO₃ (aq, 200 ml), water (200 ml) and brine (200 ml), dried over MgSO₄, filtered, concentrated and purified by silica gel chromatography (from PE to PE:EA = 50:1) to afford **5** as a light yellow oil (7.8 g, 59%).

Step 4: Synthesis of 2-[2-(2,2-Dimethyl-propoxy)-5-isopropyl-phenyl]-4,4,5,5-tetramethyl-[1,3,2]dioxaborolane

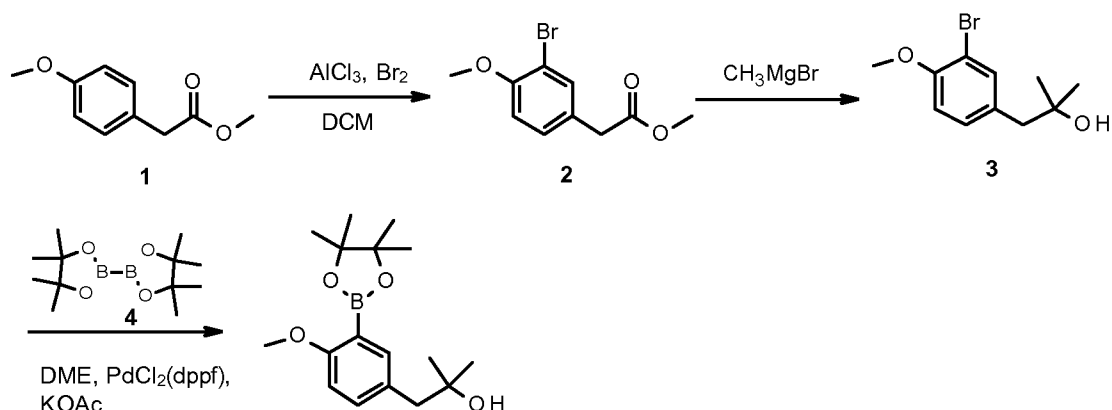
To a solution of **5** (7.8 g, 27.3 mmol) in DME (100 mL), **6** (8.6 g, 33.6 mmol), KOAc (8.3 g, 84.2 mmol) and PdCl₂(dppf) (2.3 g, 2.8 mmol) was added. The mixture was purged with N₂ and stirred at 100 °C under N₂ atmosphere for 16 h. The mixture was filtered, concentrated and purified by silica gel chromatography (from PE to PE:EA = 100:1) to afford 7.5 g light yellow solid. PE (8 ml) was added. The mixture was filtered to afford 2-[2-(2,2-Dimethyl-propoxy)-5-isopropyl-phenyl]-4,4,5,5-tetramethyl-[1,3,2]dioxaborolane as a white solid (3.1 g, 36%).

¹HNMR (CDCl₃): 1.03 (s, 9H), 1.21 (d, 6H), 1.34 (s, 12H), 2.85 (m, 1H), 6.73 (d, 1H), 7.22 (m, 1H), 7.47 (d, 1H)

1-[4-Methoxy-3-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenyl]-2-methyl-propan-2-ol



General synthetic scheme:



Step 1: Synthesis of (3-Bromo-4-methoxy-phenyl)-acetic acid methyl ester (**2**)

To a solution of **1** (9.7 g, 54 mmol) and AlCl_3 (7 g, 54 mmol) in DCM (100 mL) was added Br_2 (9.4 g, 59.4 mmol) slowly at ice/water bath. The mixture was stirred at 0°C for 20 min and then poured into ice-water (100 mL). The product was extracted with DCM. The organic layer was washed with saturated $\text{Na}_2\text{S}_2\text{O}_3$, brine, dried over Na_2SO_4 and filtrated. Evaporation of solvent to give the product as a colorless oil (12.1 g, 80%).

Step 2: Synthesis of 1-(3-Bromo-4-methoxy-phenyl)-2-methyl-propan-2-ol (**3**)

To a stirred solution of **2** (12.1 g, 47 mmol) in THF (100 mL) at ice/water bath was added CH_3MgBr (3M solution in diethyl, 47 mL, 141 mmol) and the mixture was stirred at rt for 3 hours. The mixture was then re-cooled to 0°C and slowly quenched with saturated aqueous ammonium chloride solution. The organic solution was washed with brine, dried over sodium sulfate and concentrated. The crude product was purified by silica gel chromatography (from PE: EA = 20:1 to PE: EA = 10:1) to give **3** as a yellow oil (6 g, 96%).

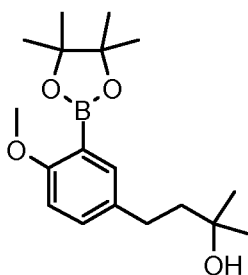
Step 3: Synthesis of 1-[4-Methoxy-3-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenyl]-2-methyl-propan-2-ol

To a solution of **3** (6.0 g, 23.2 mmol) in DME (100 mL, degassed by sparging with N_2), **5** (7.5 g, 30 mmol), KOAc (6.7 g, 69 mmol) and $\text{PdCl}_2(\text{dppf})$ (1.9 g, 2.3 mmol) was added. The mixture was stirred at 110°C under N_2 atmosphere for 16 h. The mixture was filtered, concentrated and purified by silica gel chromatography (from PE: EA =

10:1 to PE: EA = 4:1) to afford 1-[4-Methoxy-3-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenyl]-2-methyl-propan-2-ol as a white solid (3.5 g, 50%).
¹H-NMR (CDCl₃): 1,22 (s, 6H); 1,35 (s, 12H), 2,71 (s, 2H), 3,82 (s, 3H), 6,81 (d, 1H); 7,25 (m, 1H), 7,49 (d, 1H)

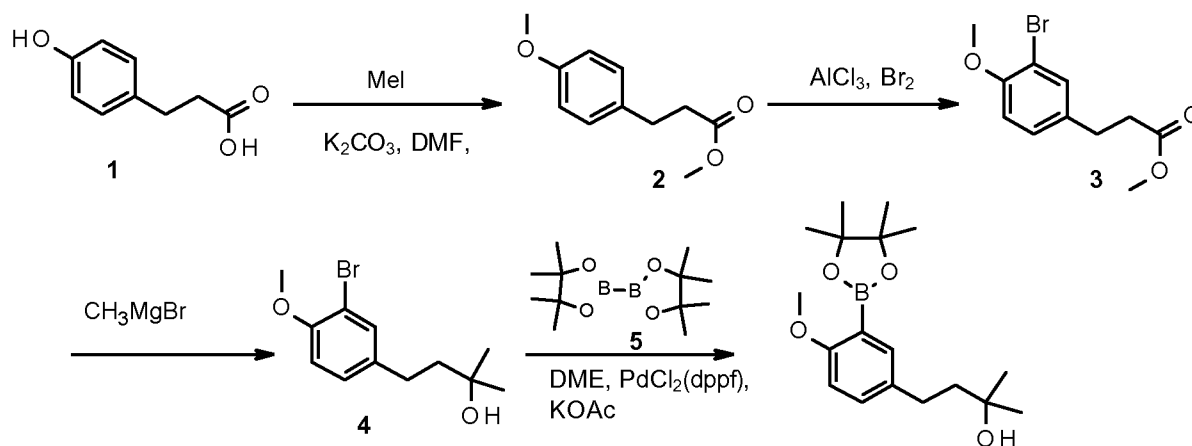
5

4-[4-Methoxy-3-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenyl]-2-methyl-butan-2-ol



10

General synthetic scheme:



Step 1: Synthesis of 3-(4-Methoxy-phenyl)-propionic acid methyl ester (**2**)

15 To a suspension of **1** (10.0 g, 60 mmol) and K₂CO₃ (21.0 g, 150 mmol) in DMF was added CH₃I (26 g, 180 mmol). The mixture was stirred at rt for overnight. EA (500 ml) was added. The mixture was washed with water (150 ml x 2) and brine (100 ml). The organic layer was dried over Na₂SO₄, filtered and concentrated to afford **2** as a light yellow oil (12.0 g).

20

Step 2: Synthesis of 3-(3-Bromo-4-methoxy-phenyl)-propionic acid methyl ester (**3**)

To a solution of **2** (12.0 g, 61.9 mmol) in DCM (150 ml) was added AlCl₃ (8.3 g, 61.9 mmol) at ice/salt bath. Then Br₂ (9.9 g, 61.9 mmol) was added dropwise. The mixture was stirred at 0 °C for 20 minutes. The reaction mixture was poured into
5 ice/water (200 ml) and extracted with DCM (150 ml x 3). The combined organic layer was washed with Na₂S₂O₃ (aq, 150 ml) and brine (150 ml), dried over Na₂SO₄, filtered and concentrated to afford **3** as a light yellow oil (16.8 g).

Step 3: Synthesis of 4-(3-Bromo-4-methoxy-phenyl)-2-methyl-butan-2-ol (**4**)

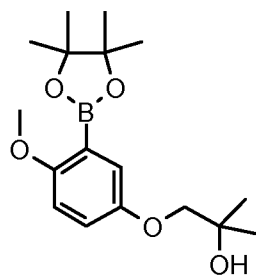
10 To a solution of **3** (16.8 g, 61.5 mmol) in THF (100 ml) at ice/water bath was added CH₃MgBr (3M, 37 ml) dropwise. Then it was stirred at rt for 3 h. NH₄Cl (aq, 150 ml) was added to quench the reaction. EA (300 ml) was added and the aqueous layer was separated. The organic layer was washed with brine (150 ml), dried over Na₂SO₄, filtered, concentrated and purified by silica gel chromatography (from PE:EA = 20:1
15 to PE:EA = 8:1) to afford **4** as a colorless oil (10.0 g).

Step 4: Synthesis of 4-[4-Methoxy-3-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenyl]-2-methyl-butan-2-ol

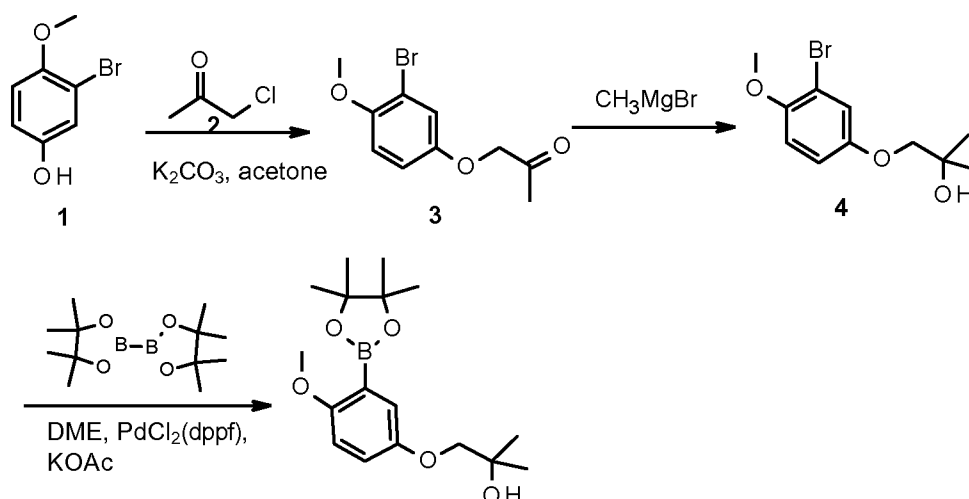
To a solution of **4** (10.0 g, 36.8 mmol) in DME (100 mL), **5** (11.2 g, 44.1 mmol),
20 KOAc (10.8 g, 110 mmol) and PdCl₂(dppf) (3.1 g, 3.7 mmol) was added. The mixture was purged with N₂ and stirred at 100 °C under N₂ atmosphere for 16 h. The mixture was filtered, concentrated and purified by silica gel chromatography (from PE: EA = 50:1 to PE:EA = 5:1) to afford 8.7 g colorless oil. PE/EA (100:1, 40ml) was added and placed at rt for 2 days. The mixture was filtered to afford **00238** as a white
25 solid (3.8 g).

¹H NMR (CDCl₃): 1,28 (s, 6H); 1,36 (s, 12H), 1,76 (m, 2H), 2,64 (m, 2H), 3,81 (s, 3H); 6,79 (d, 1H), 7,23 (m, 1H), 7,50 (d, 1H)

30 1-[4-Methoxy-3-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenoxy]-2-methyl-propan-2-ol



General synthetic scheme:



Step 1: Synthesis of 1-(3-Bromo-4-methoxy-phenoxy)-propan-2-one (**3**)

- 5 To a suspension of **1** (8.0 g, 40 mmol) and K_2CO_3 (6.0 g, 44 mmol) in acetone was added **2** (4.1 g, 44 mmol). The mixture was stirred at 60°C for 8 hours maintaining gentle reflux. The mixture was poured into water, extracted with EA. The organic layer was washed with brine, dried over Na_2SO_4 , filtered and concentrated to give **3** as oil (12.8 g, 80%).

10

Step 2: Synthesis of 1-(3-Bromo-4-methoxy-phenoxy)-2-methyl-propan-2-ol (**4**)

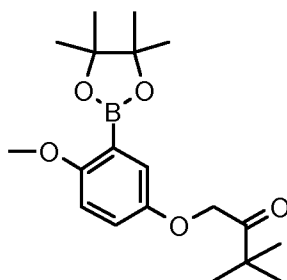
- To a stirred solution of **3** (10.3 g, 40 mmol) in THF (120 mL) at ice/water bath was added CH_3MgBr (3M solution in diethyl, 20 mL, 60 mmol) and the mixture was stirred at room temperature for 3 hours. The mixture was then re-cooled to 0°C and slowly quenched with saturated aqueous ammonium chloride solution. The organic solution was washed with brine, dried over sodium sulfate and concentrated. The crude product was purified by silica gel chromatography (from PE: EA = 10:1 to PE: EA = 8:1) to give **4** as yellow oil (7.2 g, 92%)
- 15

Step 3: Synthesis of 1-[4-Methoxy-3-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenoxy]-2-methyl-propan-2-ol

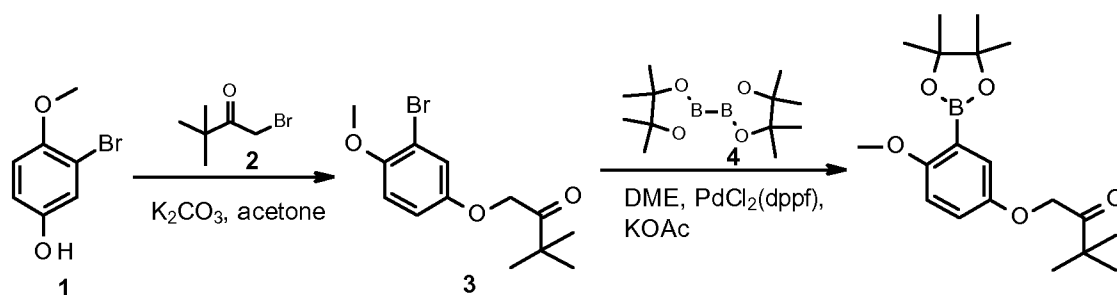
To a solution of **3** (7.2 g, 26.2 mmol) in DME (100 mL, degassed by sparging with N₂), **5** (8.0 g, 31.4 mmol), KOAc (7.7 g, 78.6 mmol) and PdCl₂(dppf) (2.2 g, 2.6 mmol) was added. The mixture was stirred at 110°C under N₂ atmosphere for 16 h. The mixture was filtered, concentrated and purified by silica gel chromatography (from PE: EA = 7:1 to PE: EA = 4:1) to afford **00239** as a white solid (3.1 g, 37%).

¹H-NMR: 1,33 (s, 6H), 1,35 (s, 12H); 3,77 (s, 2H); 3,79 (s, 3H); 6,80 (d, 1H); 6,95 (m, 1H); 7,24 (d, 1H)

1-[4-Methoxy-3-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenoxy]-3,3-dimethyl-butan-2-one



General synthetic scheme:



Step 1: Synthesis of 1-(3-Bromo-4-methoxy-phenoxy)-3,3-dimethyl-butan-2-one (**3**)

To a suspension of **1** (5.0 g, 25 mmol) and K₂CO₃ (3.8 g, 27.5 mmol) in acetone (50 mL) was added **2** (4.9 g, 27.5 mmol). The mixture was stirred at 60°C for 8 hours while maintaining gentle reflux. The mixture was poured into water, extracted

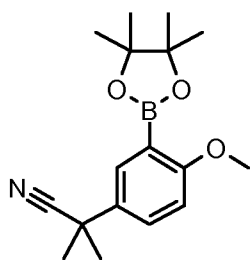
with EA. The organic layer was washed with brine, dried over Na_2SO_4 , filtered and concentrated gave **3** as a colorless oil (7.4 g, 90%).

Step 2: Synthesis of 1-[4-Methoxy-3-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenoxy]-3,3-dimethyl-butan-2-one

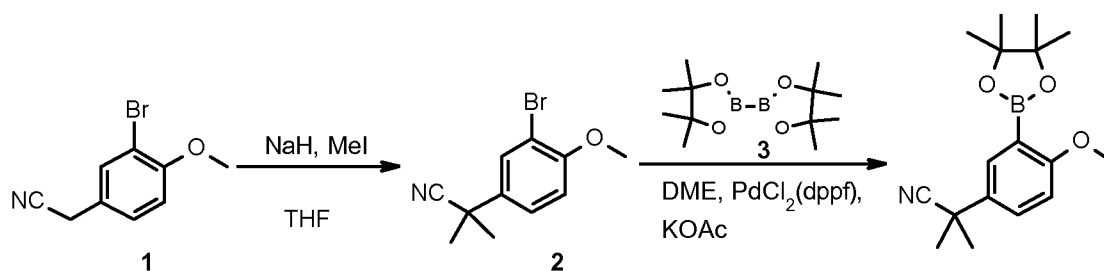
To a solution of **3** (6.7 g, 22.3 mmol) in DME (100 mL, degassed by sparging with N_2), **4** (6.3 g, 24.6 mmol), KOAc (6.5 g, 66.9 mmol) and $\text{PdCl}_2(\text{dppf})$ (1.9 g, 2.23 mmol) was added. The mixture was stirred at 110°C under N_2 atmosphere for 2 hours. The mixture was filtered, concentrated and purified by silica gel chromatography (from PE: EA = 30:1 to PE: EA = 6:1) to afford a white solid (4.0 g, 51%).

$^1\text{H-NMR}$ (CDCl_3): 1,25 (s, 9H); 1,34 (s, 12H), 3,78 (s, 3H), 4,82 (s, 2H); 6,79 (d, 1H); 6,97 (m, 1H); 7,18 (d, 1H).

2-[4-Methoxy-3-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenyl]-2-methyl-propionitrile



General synthetic scheme:



Step1: Synthesis of 2-(3-Bromo-4-methoxy-phenyl)-2-methyl-propionitrile (**2**)

NaH (3.7 g, 93 mmol) was added into THF (100 mL) slowly at 0°C , followed by **1** (7 g, 31 mmol). The solution was stirred at 0°C for 1 hour and then MeI (26.4 g, 186 mmol) was added. This reaction mixture was stirred at rt for overnight. Water (10 mL)

was added and the mixture was extracted with EA. The organic layer was washed with brine, dried over EA and concentrated to give **2** as a colourless oil (9 g, 85%).

Step 2: Synthesis of 2-[4-Methoxy-3-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenyl]-2-methyl-propionitrile

To a solution of **2** (7.8 g, 31 mmol) in DME (100 mL, degassed by sparging with N₂), **3** (9.4 g, 37.2 mmol), KOAc (9.1 g, 93 mmol) and PdCl₂(dppf) (2.6 g, 3.1 mmol) was added. The mixture was stirred at 110°C under N₂ atmosphere for 2 hours. The mixture was filtered, concentrated and purified by silica gel chromatography to afford a white solid (4.1 g, 44%).

¹H-NMR (CDCl₃): 1,36 (s, 12H), 1,72 (s, 6H), 3,84 (s, 3H), 6,85 (d, 1H); 7,51 (m, 1H); 7,68 (d, 1H).

Analogously as described in the synthesis examples, the example compounds of the formula I listed in Table 1 were prepared.

Table 1. Example compounds of the formula I

No.	CHEMICAL_NAME	St. mat	Suzuki reaction	CATH A IC50 [μM]	Obs. Mass	Rt [min]	LC/MS
1	3-(3'-tert-Butyl-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	1,89	366.14	4.88	LC3
2	3-(3'-Methyl-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	0,784	322.15	1.81	LC1
3	3-(2'-Chloro-3'-methyl-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	0,797	356.2	1.86	LC1

4	3-(3'-Chloro-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	0,967	342.09	1.83	LC1
5	3-(3'-Chloro-2'-methyl-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	1,668	356.16	1.9	LC1
6	3-(4'-Chloro-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	1,749	342.12	1.84	LC1
7	3-(2'-Chloro-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	2,040	342.14	1.78	LC1
8	3-(2',3'-Dimethyl-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	2,483	336.16	1.86	LC1
9	3-Biphenyl-3-yl-3-thiazol-2-yl-propionic acid	C2	A	2,586	310.21	1.74	LC1
10	3-(2'-Chloro-3'-fluoro-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	2,720	360.14	1.79	LC1
11	3-(2'-Methyl-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	3,539	322.14	1.8	LC1
12	3-(2'-Fluoro-5'-trifluoromethyl-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	3,619	394.2	1.85	LC1
13	3-(4'-Fluoro-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	4,304	326.09	1.76	LC1
14	3-(2',3'-Dichloro-biphenyl-3-yl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	C1	A	0,634	375.08	1.76	LC1
15	3-(5-Methyl-[1,3,4]oxadiazol-2-yl)-3-(3-pyridin-4-yl-phenyl)-propionic acid	C1	A	23,2	310.16	0.77	LC1
16	3-(2'-Fluoro-3'-methoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	0,588	358.17	1.71	LC1
17	3-[3-(1-Methyl-1H-indazol-3-yl)-phenyl]-3-thiazol-2-yl-propionic acid	C2	A	0,769	364.19	1.69	LC1

18	3-[3-(2,3-Dihydro-benzo[1,4]dioxin-6-yl)-phenyl]-3-thiazol-2-yl-propionic acid	C2	A	0,987	368.18	1.7	LC1
19	3-(2'-Fluoro-5'-propoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	1,11	386.21	1.9	LC1
20	3-(3'-Methoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	1,179	340.2	1.74	LC1
21	3-(2',5'-Difluoro-4'-methoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	1,32	376.14	1.75	LC1
22	3-[3-(6-Amino-2-fluoropyridin-3-yl)-phenyl]-3-thiazol-2-yl-propionic acid	C2	A	1,39	344.18	1.44	LC1
23	3-(3'-Ethoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	1,499	354.2	1.81	LC1
24	3-(4'-Methoxy-3',5'-dimethyl-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	1,651	368.2	1.84	LC1
25	3-(3'-Propoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	1,67	368.22	1.9	LC1
26	3-(3'-Cyanomethoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	1,94	365.18	1.67	LC1
27	3-Thiazol-2-yl-3-(3'-trifluoromethoxy-biphenyl-3-yl)-propionic acid	C2	A	2,25	394.14	1.89	LC1
28	3-(4'-Hydroxy-3'-methoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	2,28	356.17	1.54	LC1
29	3-(4'-Cyclopropylmethoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	2,29	380.2	1.87	LC1
30	3-(5'-Fluoro-3'-isobutoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	2,31	400.23	2	LC1
31	3-(4'-Cyanomethoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	2,42	365.15	1.65	LC1

32	3-(4'-Benzyloxy-3'-fluoro-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	2,426	434.2	1.94	LC1
33	3-Thiazol-2-yl-3-[3'-(2,2,2-trifluoro-ethoxy)-biphenyl-3-yl]-propionic acid	C2	A	2,46	408.16	1.84	LC1
34	3-(3'-Benzyloxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	2,772	416.24	1.93	LC1
35	3-(4'-Isobutoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	2,810	382.21	1.97	LC1
36	3-(2'-Formyl-5'-methoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	3,202	368.18	1.65	LC1
37	3-(3'-Benzyloxy-4'-chloro-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	C2	A	7,87	450.16	1.99	LC1
38	3-[3-(6-Methoxy-pyridin-3-yl)-phenyl]-3-thiazol-2-yl-propionic acid	C2	A	8,35	341.18	1.62	LC1
39	3-[3-(2-Chloro-pyridin-3-yl)-phenyl]-3-thiazol-2-yl-propionic acid	C2	A	11,60	345.11	1.55	LC1
40	3-[3-(3-Chloro-pyridin-4-yl)-phenyl]-3-thiazol-2-yl-propionic acid	C2	A	19,44	345.12	1.54	LC1
41	3-(3-Pyridin-3-yl-phenyl)-3-thiazol-2-yl-propionic acid	C2	A	21,19	311.19	1.1	LC1
42	3-[3-(6-Cyano-pyridin-3-yl)-phenyl]-3-thiazol-2-yl-propionic acid	C2	A	22,58	336.16	1.53	LC1
43	3-[3-(2-Fluoro-pyridin-4-yl)-phenyl]-3-thiazol-2-yl-propionic acid	C2	A	23,6	329.15	1.56	LC1
44	3-[3-(3-Fluoro-pyridin-4-yl)-phenyl]-3-thiazol-2-yl-propionic acid	C2	A	24,8	329.14	1.46	LC1
45	3-(3-Pyridin-4-yl-phenyl)-3-thiazol-2-yl-propionic acid	C2	A		311.19	1	LC1

46	3-{3-[5-(Morpholine-4-carbonyl)-pyridin-3-yl]-phenyl}-3-thiazol-2-yl-propionic acid	C2	A		424.17	1.32	LC1
47	3-(5'-Chloro-2'-fluoro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A6	A	0,228	376.17	1.88	LC1
48	3-(2'-Fluoro-3'-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A6	A	0,397	372.21	1.77	LC1
49	3-(2',5'-Difluoro-4'-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A6	A	0,466	390.25	1.8	LC1
50	3-(2'-Fluoro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A6	A	0,631	342.21	1.8	LC1
51	3-(2'-Fluoro-4'-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A6	A	0,989	372.23	1.8	LC1
52	3-(2'-Acetyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A6	A	2,303	366.23	1.69	LC1
53	3-(2'-Carbamoyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A6	A	5,384	367.23	1.45	LC1
54	3-[3-(3-Fluoro-pyridin-4-yl)-phenyl]-3-(5-methyl-thiazol-2-yl)-propionic acid	A6	A	7,06	343.19	1.55	LC1
55	3-(5-Methyl-thiazol-2-yl)-3-(3-pyridin-3-yl-phenyl)-propionic acid	A6	A	7,535	325.21	1.25	LC1
56	3-(5-Methyl-thiazol-2-yl)-3-{3-[5-(morpholine-4-carbonyl)-pyridin-3-yl]-phenyl}-propionic acid	A6	A	13,59	436.26	1.41	LC1
57	3-(5-Methyl-thiazol-2-yl)-3-(3-pyridin-4-yl-phenyl)-propionic acid	A6	A	13,74	325.2	1.13	LC1
58	3-(3'-Chloro-2'-fluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	0,954	406.12	1.87	LC1

59	3-(2'-Fluoro-4-methoxy-4'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	0,97	386.18	1.86	LC1
60	3-(2',5'-Difluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	0,975	390.15	1.81	LC1
61	3-(5'-Fluoro-4-methoxy-2'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	1,2	386.15	1.86	LC1
62	3-(5'-Chloro-2'-fluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	1,23	406.1	1.88	LC1
63	3-(2'-Chloro-5'-fluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	1,25	406.12	1.85	LC1
64	3-(3'-Chloro-4-methoxy-2'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	1,31	402.15	1.94	LC1
65	3-(4-Methoxy-4'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	1,59	368.17	1.86	LC1
66	3-(2',3'-Difluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	1,77	390.14	1.81	LC1
67	3-(3'-Fluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	1,96	372.14	1.81	LC1
68	3-(4-Methoxy-2',3'-dimethyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	2,3	382.19	1.91	LC1
69	3-(2'-Fluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	2,34	372.16	1.79	LC1
70	3-(2'-Fluoro-4-methoxy-3'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	2,42	386.15	1.86	LC1

71	3-(2',4'-Difluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	2,65	390.13	1.82	LC1
72	3-(4-Methoxy-2',4'-dimethyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	3,09	382.21	1.92	LC1
73	3-(4'-Chloro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	3,2	388.12	1.88	LC1
74	3-(2'-Fluoro-4,4'-dimethoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	3,27	402.18	1.8	LC1
75	3-(4-Methoxy-2'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	3,28	368.17	1.85	LC1
76	3-(4'-Chloro-2'-fluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	3,57	406.11	1.9	LC1
77	3-(4'-Fluoro-4-methoxy-2'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	3,73	386.16	1.86	LC1
78	3-(3'-Chloro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	3,74	388.1	1.88	LC1
79	3-(2'-Chloro-3'-fluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	4,01	406.1	1.84	LC1
80	3-(4-Methoxy-4'-trifluoromethyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	4,38	422.15	1.9	LC1
81	3-(4'-Fluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	4,39	372.16	1.8	LC1
82	3-(4,3'-Dimethoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	5,56	384.17	1.78	LC1

83	3-(2'-Fluoro-4-methoxy-5'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	5,68	386.18	1.86	LC1
84	3-(2'-Chloro-4'-fluoro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	6,36	406.13	1.86	LC1
85	3-(2'-Chloro-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	6,69	388.12	1.84	LC1
86	3-(4-Methoxy-4'-trifluoromethoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	6,73	438.15	1.93	LC1
87	3-(4-Methoxy-3'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	7,12	368.16	1.86	LC1
88	3-(4-Methoxy-3'-trifluoromethyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	7,77	422.15	1.9	LC1
89	3-(4,4'-Dimethoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	7,81	384.16	1.77	LC1
90	3-(4'-Cyano-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	8,8	379.15	1.71	LC1
91	3-(3'-Cyano-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	9,26	379.16	1.72	LC1
92	3-(4-Methoxy-2'-trifluoromethyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	10,2	422.16	1.86	LC1
93	3-(4-Methoxy-3'-trifluoromethoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	18,1	438.15	1.92	LC1
94	3-(2'-Carbamoyl-4-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B	19,3	397.16	1.46	LC1

95	3-(2-Methoxy-5-pyridin-4-yl-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B		355.15	1.16	LC1
96	3-(2-Methoxy-5-pyridin-3-yl-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B		355.13	1.24	LC1
97	3-[5-(3-Fluoro-pyridin-4-yl)-2-methoxy-phenyl]-3-(5-methyl-thiazol-2-yl)-propionic acid	P3	B		373.13	1.53	LC1
98	3-(5-Chloro-2',5'-difluoro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,136	394.14	1.91	LC1
99	3-(5-Chloro-2'-fluoro-3'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,184	390.18	1.97	LC1
100	3-(5,3'-Dichloro-2'-fluoro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,197	410.11	1.97	LC1
101	3-(5-Chloro-2'-trifluoromethoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,221	442.15	1.96	LC1
102	3-(5-Chloro-2'-fluoro-5'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,232	390.16	1.96	LC1
103	3-(2'-Acetyl-5-chloro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,25	400.19	1.8	LC1
104	3-(5-Chloro-2'-fluoro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,328	376.16	1.9	LC1
105	3-(5-Chloro-2',6'-difluoro-4'-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,335	424.17	1.91	LC1
106	3-(5-Chloro-2',3'-difluoro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,354	394.14	1.91	LC1
107	3-(2'-Carbamoyl-5-chloro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,404	401.17	1.58	LC1

108	3-(5-Chloro-3'-trifluoromethoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,406	442.15	2.01	LC1
109	3-(3-Chloro-5-pyridin-4-yl-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid; compound with trifluoro-acetic acid	A1	A	0,459	359.14	1.36	LC1
110	3-(5-Chloro-2'-fluoro-4'-methoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,563	406.18	1.9	LC1
111	3-(5-Chloro-2'-cyano-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	0,567	383.16	1.79	LC1
112	3-(3-Chloro-5-pyridin-3-yl-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	1,06	359.15	1.48	LC1
113	3-(5-Chloro-2'-dimethylcarbamoyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	1,44	429.23	1.69	LC1
114	3-[3-Chloro-5-(3-fluoro-pyridin-4-yl)-phenyl]-3-(5-methyl-thiazol-2-yl)-propionic acid	A1	A	1,56	377.14	1.69	LC1
115	3-(5,2'-Difluoro-5'-methyl-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A3	A	0,437	374.19	1.9	LC1
116	3-(5,2'-Difluoro-3'-methyl-biphenyl-3-yl)-3-oxazol-2-yl-propionic acid	A3	A	0,488	344.15	1.8	LC1
117	3-(5-Methyl-thiazol-2-yl)-3-(5,2',5'-trifluoro-biphenyl-3-yl)-propionic acid	A3	A	0,649	378.14	1.84	LC1
118	3-(2'-Acetyl-5-fluoro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A3	A	0,707	384.18	1.73	LC1
119	3-(5,2'-Difluoro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A3	A	0,78	360.15	1.83	LC1

120	3-(5,2'-Difluoro-5'-methyl-biphenyl-3-yl)-3-oxazol-2-yl-propionic acid	A3	A	0,903	344.14	1.79	LC1
121	3-(5-Fluoro-3'-trifluoromethoxy-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A3	A	0,975	426.2	1.95	LC1
122	3-(5,2'-Difluoro-4'-methyl-biphenyl-3-yl)-3-oxazol-2-yl-propionic acid	A3	A	1,16	344.14	1.8	LC1
123	3-Oxazol-2-yl-3-(5,2',6'-trifluoro-4'-methoxy-biphenyl-3-yl)-propionic acid	A3	A	1,22	378.13	1.74	LC1
124	3-(2'-Carbamoyl-5-fluoro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A3	A	1,23	385.19	1.5	LC1
125	3-(3-Fluoro-5-pyridin-3-yl-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid; compound with trifluoro-acetic acid	A3	A	1,33	343.16	1.38	LC1
126	3-Oxazol-2-yl-3-(5,2',3'-trifluoro-biphenyl-3-yl)-propionic acid	A3	A	1,45	348.12	1.74	LC1
127	3-Oxazol-2-yl-3-(5,2',5'-trifluoro-biphenyl-3-yl)-propionic acid	A3	A	1,49	348.11	1.73	LC1
128	3-(2'-Cyano-5-fluoro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A3	A	1,69	367.16	1.72	LC1
129	3-(2'-Dimethylcarbamoyl-5-fluoro-biphenyl-3-yl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A3	A	1,7	413.24	1.63	LC1
130	3-(5,2'-Difluoro-biphenyl-3-yl)-3-oxazol-2-yl-propionic acid	A3	A	1,72	330.13	1.71	LC1
131	3-(2'-Carbamoyl-5-fluoro-biphenyl-3-yl)-3-oxazol-2-yl-propionic acid	A3	A	1,77	355.18	1.35	LC1
132	3-(5,2'-Difluoro-4'-methoxy-biphenyl-3-yl)-3-oxazol-2-yl-propionic acid	A3	A	2,02	360.13	1.73	LC1

133	3-(2'-Acetyl-5-fluoro-biphenyl-3-yl)-3-oxazol-2-yl-propionic acid	A3	A	2,03	354.18	1.6	LC1
134	3-(2'-Cyano-5-fluoro-biphenyl-3-yl)-3-oxazol-2-yl-propionic acid	A3	A	2,03	337.13	1.6	LC1
135	3-(5-Fluoro-3'-trifluoromethoxy-biphenyl-3-yl)-3-oxazol-2-yl-propionic acid	A3	A	2,28	396.16	1.86	LC1
136	3-(3-Fluoro-5-pyridin-4-yl-phenyl)-3-(5-methyl-thiazol-2-yl)-propionic acid	A3	A	8,65	343.17	1.25	LC1
137	3-Oxazol-2-yl-3-(5,2',6'-trifluoro-biphenyl-3-yl)-propionic acid	A3	A		348.18	1.7	LC1
138	3-(3-Fluoro-5-pyridin-4-yl-phenyl)-3-oxazol-2-yl-propionic acid	A3	A		313.17	0.99	LC1
139	3-(3-Fluoro-5-pyridin-3-yl-phenyl)-3-oxazol-2-yl-propionic acid	A3	A		313.18	1.13	LC1
140	3-[3-Fluoro-5-(3-fluoro-pyridin-4-yl)-phenyl]-3-oxazol-2-yl-propionic acid	A3	A		331.14	1.45	LC1
141	3-(2'-Dimethylcarbamoyl-5-fluoro-biphenyl-3-yl)-3-oxazol-2-yl-propionic acid	A3	A		383.15	1.49	LC1
142	3-(2'-Acetyl-biphenyl-3-yl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	C1	A	1,72	351.21	1.5	LC1
143	3-(2'-Cyano-biphenyl-3-yl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	C1	A	4,74	334.18	1.5	LC1
144	3-(5-Methyl-[1,3,4]oxadiazol-2-yl)-3-(3-pyridin-3-yl-phenyl)-propionic acid	C1	A	6,71	310.17	0.88	LC1

145	3-(5-Methyl-[1,3,4]oxadiazol-2-yl)-3-(3-pyridin-4-yl-phenyl)-propionic acid	C1	A	20,6	310.17	0.74	LC1
146	3-(2'-Carbamoyl-5-chloro-biphenyl-3-yl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A8	A		386.17	3.16	LC3
147	3-(2'-Carbamoyl-5-fluoro-biphenyl-3-yl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A9	A		370.18	2.93	LC3
148	3-(2'-Cyano-5-fluoro-biphenyl-3-yl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A9	A		352.18	3.73	LC3
149	3-(3-Fluoro-5-pyridin-4-yl-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A9	A		328.14	2.07	LC3
150	3-(3-Chloro-5-pyridin-4-yl-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid; compound with trifluoro-acetic acid	A8	A		344.13	2.32	LC3
151	3-(3-Fluoro-5-pyridin-3-yl-phenyl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A9	A		328.14	2.3	LC3
152	3-(2'-Dimethylcarbamoyl-5-fluoro-biphenyl-3-yl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A9	A		398.26	3.22	LC3
153	3-[5-(2,6-Difluoro-phenyl)-pyridin-3-yl]-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A10	A		346.12	1.33	LC1

154	3-[3,3']Bipyridinyl-5-yl-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A10	A		311.11	0.52	LC1
155	3-[5-(2-Carbamoyl-phenyl)-pyridin-3-yl]-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A10	A		353.16	0.8	LC1
156	3-[5-(2,6-Dimethoxy-phenyl)-pyridin-3-yl]-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A10	A		370.19	1.21	LC1
157	3-[5-(2-Cyano-phenyl)-pyridin-3-yl]-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A10	A		335.12	1.23	LC1
158	3-[5-(5-Chloro-2-methoxy-phenyl)-pyridin-3-yl]-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A10	A		374.11	1.44	LC1
159	3-[5-(5-Fluoro-2-methoxy-phenyl)-pyridin-3-yl]-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A10	A		358.13	1.32	LC1
160	3-[5-(2-Acetyl-phenyl)-pyridin-3-yl]-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A10	A		352.15	1.14	LC1
161	3-(5-Methyl-[1,3,4]oxadiazol-2-yl)-3-(3-pyridin-4-yl-5-trifluoromethyl-phenyl)-propionic acid	A1	A		378.16	1.17	LC1
162	3-(5-Methyl-[1,3,4]oxadiazol-2-yl)-3-(3-pyridin-3-yl-5-trifluoromethyl-phenyl)-propionic acid	A1	A		378.14	1.3	LC1
163	3-(2'-Carbamoyl-5-trifluoromethyl-biphenyl-3-yl)-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	A1	A		420.16	1.42	LC1

164	3-Biphenyl-3-yl-3-(5-methyl-[1,3,4]oxadiazol-2-yl)-propionic acid	C1	A	0,935	309.13	1.77	LC1
165	3-(4-Methoxy-biphenyl-3-yl)-3-thiazol-2-yl-propionic acid	P4	B	4,942	340.09	4.4	LC3
166	3-[3-[3-(2-amino-2-oxo-ethyl)phenyl]-5-chloro-phenyl]-3-(5-methylthiazol-2-yl)propanoic acid	A1	A	0,092 ₁	415.03	1.6	LC1
167	3-[3-[3-(2-amino-2-oxo-ethyl)phenyl]-5-fluoro-phenyl]-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoic acid	A9	A	0,318	384.13	1.33	LC1
168	3-[3-fluoro-5-(2-fluorophenyl)phenyl]-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoic acid	A9	A	0,327	345.1	1.63	LC1
169	3-[3-(2,5-difluorophenyl)-5-fluoro-phenyl]-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoic acid	A9	A	0,332	363.09	1.64	LC1
170	3-(3-fluoro-5-phenyl-phenyl)-3-(5-methyl-oxazol-2-yl)propanoic acid	A13	A	0,507	326.1	1.75	LC1
171	3-(3-fluoro-5-phenyl-phenyl)-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoic acid	A9	A	0,508	327.1	1.63	LC1
172	3-[3-[3-(2-amino-2-oxo-ethyl)phenyl]-5-fluoro-phenyl]-3-oxazol-2-yl-propanoic acid	A3	A	0,552	369.08	1.4	LC1

173	3-[3-fluoro-5-(2-fluorophenyl)phenyl]-3-(5-methyloxazol-2-yl)propanoic acid	A13	A	0,651	344.08	1.75	LC1
174	3-[3-(2,5-dimethylphenyl)-5-fluoro-phenyl]-3-(5-methyloxazol-2-yl)propanoic acid	A13	A	0,886	354.15	1.88	LC1
175	3-[3-(2,5-difluorophenyl)-5-fluoro-phenyl]-3-(5-methyloxazol-2-yl)propanoic acid	A13	A	0,980	362.08	1.76	LC1
176	3-[4-(2,5-difluorophenyl)-2-pyridyl]-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoic acid	P5	B	9,01	346.09	1.43	LC1
177	3-[5-[3-(2-amino-2-oxo-ethyl)phenyl]-3-pyridyl]-3-(5-methyl-1,3,4-oxadiazol-2-yl)propanoic acid	A10	A	24	367.09	0.87	LC1

(1) Mass spectroscopic characterization; observed mass number of the ion $[(M+H)^+]$, unless specified otherwise

(2) Cathepsin A inhibitory activity determined in the pharmacological test "Cathepsin

5 A inhibitory activity" described below.

(3) St. mat is the starting material.

Pharmacological tests

10 a) Cathepsin A inhibitory activity

Recombinant human cathepsin A (residues 29-480, with a C-terminal 10-His tag; R&D Systems, # 1049-SE) was proteolytically activated with recombinant human

cathepsin L (R&D Systems, # 952-CY). Briefly, cathepsin A was incubated at 10 $\mu\text{g/ml}$ with cathepsin L at 1 $\mu\text{g/ml}$ in activation buffer (25 mM 2-(morpholin-4-yl)-ethanesulfonic acid (MES), pH 6.0, containing 5 mM dithiothreitol (DTT)) for 15 min at 37 °C. Cathepsin L activity was then stopped by the addition of the cysteine
5 protease inhibitor E-64 (N-(trans-epoxysuccinyl)-L-leucine-4-guanidinobutylamide; Sigma-Aldrich, # E3132; dissolved in activation buffer/DMSO) to a final concentration of 10 μM .

The activated cathepsin A was diluted in assay buffer (25 mM MES, pH 5.5,
10 containing 5 mM DTT) and mixed with the test compound (dissolved in assay buffer containing (v/v) 3 % DMSO) or, in the control experiments, with the vehicle in a multiple assay plate. After incubation for 15 min at room temperature, as substrate then bradykinin carrying an N-terminal [®]Bodipy FL (4,4-difluoro-5,7-dimethyl-4-bora-3a,4a-diaza-s-indacene-3-propionyl) label (JPT Peptide Technologies GmbH;
15 dissolved in assay buffer) was added to the mixture. The final concentration of cathepsin A was 833 ng/ml and the final concentration of labeled bradykinin 2 μM . After incubation for 15 min at room temperature the reaction was stopped by the addition of stop buffer (130 mM 2-(4-(2-hydroxy-ethyl)-piperazin-1-yl)-ethanesulfonic acid, pH 7.4, containing (v/v) 0.013 % [®]Triton X-100, 0.13 % Coating Reagent 3
20 (Caliper Life Sciences), 6.5 % DMSO and 20 μM ebelactone B (Sigma, # E0886)).

Uncleaved substrate and product were then separated by a microfluidic capillary electrophoresis on a LabChip[®] 3000 Drug Discovery System (12-Sipper-Chip; Caliper Life Sciences) and quantified by determination of the respective peak areas.

25 Substrate turnover was calculated by dividing product peak area by the sum of substrate and product peak areas, and the enzyme activity and the inhibitory effect of the test compound thus quantified. From the percentage of inhibition of cathepsin A activity observed with the test compound at several concentrations, the inhibitory concentration IC_{50} , i.e. the concentration which effects 50 % inhibition of enzyme
30 activity was, calculated. IC_{50} values of various example compounds are given in Table 1, wherein "A" means an IC_{50} value of less than 0.1 μM , "B" means an IC_{50}

value between 0.1 μ M and 1 μ M, and "C" means an IC_{50} value between 1 μ M and 30 μ M.

B) In vivo antihypertrophic and renoprotective activity

5

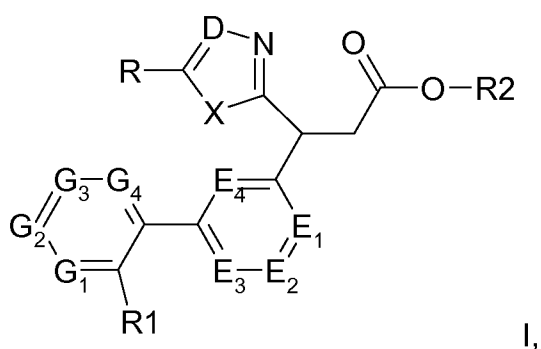
The in vivo pharmacological activity of the compounds of the invention can be investigated, for example, in the model of DOCA-salt sensitive rats with unilateral nephrectomy. Briefly, in this model unilateral nephrectomy of the left kidney (UNX) is performed on Sprague Dawley rats of 150 g to 200 g of body weight. After the operation as well as at the beginning of each of the following weeks 30 mg/kg of body weight of DOCA (desoxycorticosterone acetate) are administered to the rats by subcutaneous injection. The nephrectomized rats treated with DOCA are supplied with drinking water containing 1 % of sodium chloride (UNX/DOCA rats). The UNX/DOCA rats develop high blood pressure, endothelial dysfunction, myocardial hypertrophy and fibrosis as well as renal dysfunction. In the test group (UNX/DOCA Test) and the placebo group (UNX/DOCA Placebo), which consist of randomized UNX/DOCA rats, the rats are treated orally by gavage in two part administrations at 6 a.m. and 6 p.m. with the daily dose of the test compound (for example 10 mg/kg of body weight dissolved in vehicle) or with vehicle only, respectively. In a control group (control), which consists of animals which have not been subjected to UNX and DOCA administration, the animals receive normal drinking water and are treated with vehicle only. After five weeks of treatment, systolic blood pressure (SBP) and heart rate (HR) are measured non-invasively via the tail cuff method. For determination of albuminuria and creatinine, 24 h urine is collected on metabolic cages. Endothelial function is assessed in excised rings of the thoracic aorta as described previously (W. Linz et al., JRAAS (Journal of the renin-angiotensin-aldosterone system) 7 (2006), 155-161). As a measure of myocardial hypertrophy and fibrosis, heart weight, left ventricular weight and the relation of hydroxyproline and proline are determined in excised hearts.

30

DE2013/051 WOPCT

Claims

- 5 1. A compound of the formula I, in any of its stereoisomeric forms or a mixture of stereoisomeric forms in any ratio, or a physiologically acceptable salt thereof, or a physiologically acceptable solvate of any of them,



10

wherein the meanings are:

X is S or O;

D is N or $-\text{C}(\text{R}_3)=$;

- 15 R is H, $(\text{C}_1\text{-C}_6)$ -alkyl, $(\text{C}_3\text{-C}_8)$ -cycloalkyl;

R₁ is H, F, Cl, CF₃, CN, CO-H, CO- $(\text{C}_1\text{-C}_6)$ -alkyl, CO-NR₂₀R₂₁; wherein R₂₀ and R₂₁ are independently from each other H or $(\text{C}_1\text{-C}_6)$ -alkyl;

R₃ is H, methyl or ethyl;

R₂ is Hydrogen or $(\text{C}_1\text{-C}_6)$ -alkyl;

20

E₁ is N or $-\text{C}(\text{R}_4)=$;

E₂ is N or $-\text{C}(\text{R}_5)=$;

E₃ is N or $-\text{C}(\text{R}_6)=$;

E₄ is N or $-\text{C}(\text{R}_7)=$; wherein none or one of E₁, E₂, E₃ or E₄ is N;

25

R₄ is H or O- $(\text{C}_1\text{-C}_6)$ -alkyl;

R5 is H, F, Cl, CF₃, (C₁-C₆)-alkyl, (C₁-C₆)-cycloalkyl;

R6 is H;

R7 is H;

- 5 G₁ is N or -C(R8)=;
G₂ is N or -C(R9)=;
G₃ is N or -C(R10)=;
G₄ is N or -C(R11)=; wherein none or one of G₁, G₂, G₃ or G₄ is N;
or G₃ and G₄ are -C(R10)= and -C(R11)=, wherein R10 and R11 form a 4 to 7
10 membered saturated carbocycle- or heterocycle with one or two oxygen atoms;
which is optionally mono- or disubstituted by (C₁-C₃)-alkyl;

R8 is H, F, Cl, (C₁-C₆)-alkyl, O-(C₁-C₆)-alkyl, CF₃, -CH₂CO-NH₂ or OCF₃;

- 15 R9 is H, F, Cl, OH, O-(C₁-C₆)-alkyl, CH₂OH, CO-NH₂, (C₁-C₆)-alkyl, O-(C₁-C₆)-alkyl,
CF₃ or OCF₃;

R10 is H, F, Cl, OH, (C₁-C₆)-alkyl, CH₂OH, CO-O-(C₁-C₆)-alkyl, SO₂-(C₁-C₆)-alkyl,
CN, O-(C₁-C₆)-alkyl, CF₃ or OCF₃;

- 20 R11 is H, F, Cl, OH, O-(C₁-C₆)-alkyl, CH₂OH, CO-(C₁-C₆)-methyl, CO-N(R20R21),
CO-O-(C₁-C₆)-alkyl, CN, (C₁-C₆)-alkyl, O-(C₁-C₆)-alkyl or OCF₃; wherein R20 and
R21 are independently from each other H or (C₁-C₃)-alkyl or form together with the
nitrogen to which they are attached a 5 or 6 membered saturated ring.

- 25 2. A compound of the formula I, in any of its stereoisomeric forms or a mixture of
stereoisomeric forms in any ratio, or a physiologically acceptable salt thereof, or a
physiologically acceptable solvate of any of them, as claimed in claim 1, wherein
R2 is Hydrogen.

- 30 3. A compound of the formula I, in any of its stereoisomeric forms or a mixture of
stereoisomeric forms in any ratio, or a physiologically acceptable salt thereof, or a
physiologically acceptable solvate of any of them, as claimed in any one or more of
claims 1 and 2, wherein

R is Hydrogen, methyl or ethyl;

R1 is H, F, Cl, CF₃, CN, CO-H, CO-methyl, CO-NH₂, CO-N(CH₃)₂.

R2 is Hydrogen;

R4 is H or O-methyl;

5 R5 is H F, Cl, CF₃, (C₁-C₆)-alkyl, cyclopropyl;

R8 is H, F, Cl, methyl, O-methyl, CF₃ or OCF₃;

R9 is H, F, Cl, OH, O-propyl, CH₂OH, CO-NH₂, methyl, O-methyl, CF₃ or OCF₃;

R10 is H, F, Cl, OH, i-propyl, t-butyl, CH₂OH, CO-O-methyl, SO₂-methyl, CN, methyl, O-methyl, CF₃ or OCF₃;

10 R11 is H, F, Cl, OH, O-methyl, O-i-propyl, CH₂OH, CO-methyl, CO-N(methyl)₂, CO-pyrrolidin, CO-O-methyl, CN, methyl or OCF₃.

4. A compound of the formula I, in any of its stereoisomeric forms or a mixture of

stereoisomeric forms in any ratio, or a physiologically acceptable salt thereof, or a

15 physiologically acceptable solvate of any of them, as claimed in any one or more of claims 1 to 3, wherein

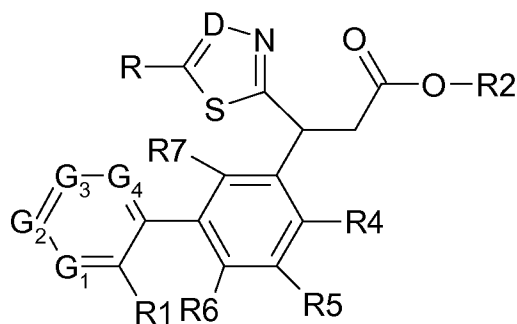
G₃ and G₄ are -C(R10)= and -C(R11)=, wherein R10 and R11 form a 5 or 6 membered saturated carbocycle or heterocycle with one or two oxygen atoms; which is optionally mono- or disubstituted by halogen and or (C₁-C₃)-alkyl.

20

5. A compound of the formula I-1 or I-11, in any of its stereoisomeric forms or a

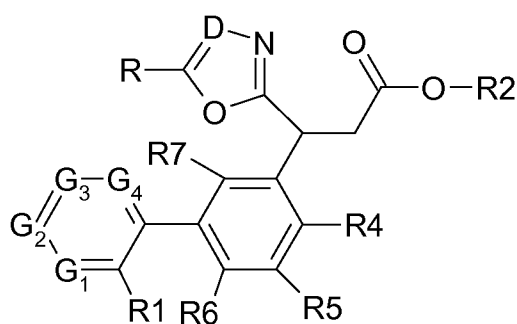
mixture of stereoisomeric forms in any ratio, or a physiologically acceptable salt

thereof, or a physiologically acceptable solvate of any of them, wherein



25

I-1,



I-11

wherein the groups D, R, R1, R2, R4, R5, R6, R7, R8, R9, R10, R11 are defined as in the compounds of formula I in claims 1 to 4, wherein

G₁ is—C(R8)=;

G₂ is—C(R9)=;

5 G₃ is—C(R10)=;

G₄ —C(R11)=.

6. A compound of the formula I as claimed in any one or more of claims 1 to 5 or a physiologically acceptable salt thereof or a physiologically acceptable solvate of any
10 of them for use as a pharmaceutical.

7. A pharmaceutical composition, which comprises at least one compound of the formula I as claimed in any one or more of claims 1 to 5 or a physiologically acceptable salt thereof or a physiologically acceptable solvate of any of them and a
15 pharmaceutically acceptable carrier.

8. Use of a compound of the formula I as claimed in any one or more of claims 1 to 5 or a physiologically acceptable salt thereof or a physiologically acceptable solvate of any of them for the manufacture of a medicament for the treatment of heart failure,
20 congestive heart failure, cardiomyopathy, myocardial infarction, left ventricular dysfunction, cardiac hypertrophy, valvular heart diseases, hypertension, atherosclerosis, peripheral arterial occlusive disease, restenosis, vasvular permeability disorders, treatment of edema, thrombosis, rheumatoid arthritis, osteoarthritis, renal failure, cystic fibrosis, chronic bronchitis, chronic obstructive
25 pulmonary disease, asthma, immunological diseases, diabetic complications, fibrotic diseases, pain, ischemia or reperfusion damage or neurodegenerative diseases, or for cardioprotection or renoprotection or as a diuretic (stand-alone treatment or in combination with established diuretics).

30 9. Use of a compound of the formula I as claimed in any one or more of claims 1 to 5 or a physiologically acceptable salt thereof or a physiologically acceptable solvate of

any of them for the manufacture of a medicament for the treatment of atrial fibrillation.