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NOVEL HETEROARYL-SUBSTITUTED ARYLAMINOPYRIDINE DERIVATIVES AS MEK INHIBITORS

RELATED APPLICATIONS

This application claims the benefit of U.S. Provisional Application No. 60/793,073, filed on April 19, 2006. The entire teachings of the above application are incorporated herein by reference.

FIELD OF THE INVENTION

The invention relates to novel heteroaryl-substituted arylaminopyridine derivative MEK
5 inhibitors, which are useful in the treatment of hyperproliferative diseases, such as
cancer and inflammation, in mammals. The invention also relates to a method of
treating a mammal suffering from or susceptible to a hyperproliferative disease, such
as cancer and inflammation, comprising administering a therapeutically effective
amount of a heteroaryl-substituted arylaminopyridine derivative according to the
10 invention. Furthermore, the invention relates to pharmaceutical compositions
containing such MEK inhibitors.

BACKGROUND OF THE INVENTION

The Ras/Raf/MEK/MAPK cascade is one of the major pathways transmitting signals
from the cell surface to the nucleus. The Ras family of G-proteins relay signals from
15 activated growth factor receptors to the downstream intracellular Raf family kinases,
which, in turn, trigger the activation of MEK1 and MEK2 extracellular signal-regulated
kinase (ERK1/ERK2) pathway. The MEK family of genes consists of five genes: MEK1,
MEK2, MEK3, MEK4 and MEK5. The structure of MEK consists of an amino-terminal
negative regulatory domain and a carboxy-terminal MAP kinase-binding domain, which
20 is necessary for binding and activation of ERKs. MEK1 is a 393-amino-acid protein with
a molecular weight of 44 kDa (Crews et al., Science 1992, 258, 478-80).

Upstream and downstream signalling of the Ras/Raf/MEK/MAPK cascade involves
multiple pathways, however MEK appears to specifically phosphorylate MAPK. The
role of MAPK in cancer (Reddy et al. Cancer Metastasis Rev. 2004, 22, 395-403), and
25 the dysfunctional activation of signalling components in the MAPK pathway in a high

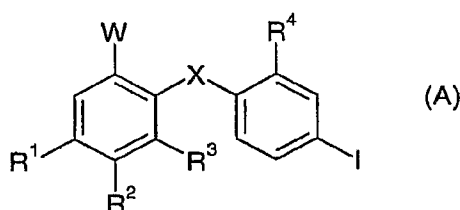
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proportion of tumor types, has lead to an extended interest in MEK as a cancer target and the development of MEK inhibitors (Chang et al., Leukemia 2003, 17,1263-93; Lee et al., Exp. Mol. Med. 2006, 38, 27-35).

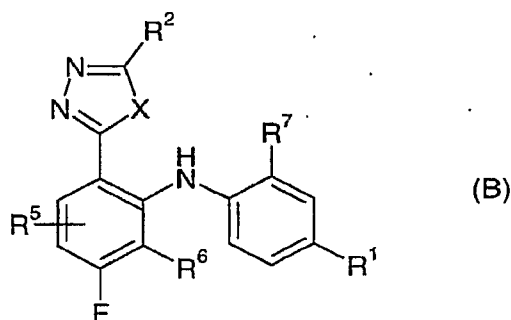
Among the furthest advanced MEK inhibitors is Pfizer's PD-0325901, a diarylamine derived MEK inhibitor, that has entered phase II clinical trials for the potential oral treatment of cancer.

Array Biopharma's ARRY142886, a phenylamino-2-pyridone derived MEK inhibitor, is currently in Phase I clinical trials.

WO 00/42029 (Warner-Lambert Company) further reports about diarylamines (A) that exhibit MEK inhibitory activity and are potentially useful for the treatment of cancer and other proliferative diseases.

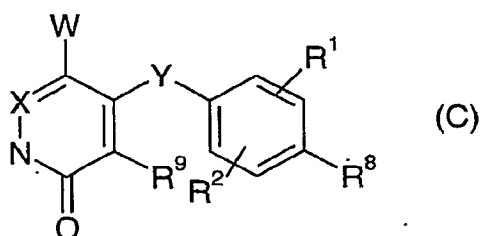


WO 04/056789 (Warner-Lambert Company) reports about oxadiazole- and thiadiazole-phenylamine derivatives (B) as MEK inhibitors for the potential treatment of inflammation and proliferative diseases.

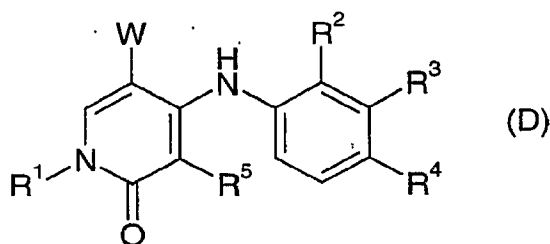


WO 05/051301 (Array Biopharma Inc.) relates to heterocyclic compounds, which are MEK inhibitors and useful for the potential treatment of hyperproliferative diseases.

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WO 05/000818 (Warner-Lambert Company) relates to phenylamino-2-pyridone derivatives as MEK inhibitors that might be useful for the treatment of proliferative diseases.



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SUMMARY OF THE INVENTION

The invention provides in one aspect MEK inhibitors according to Formula (Ia). These compounds are suitable for the treatment of hyperproliferative diseases such as cancer and inflammation.

10 It is another aspect of the present invention to provide MEK inhibitors according to Formula (Ia), which are able to down-regulate, especially inhibit the activity or function of MEK, especially in mammals.

It is another aspect of the present invention to provide a method for treating a mammal suffering from or susceptible to a hyperproliferative disease, comprising administering
15 to the mammal a therapeutically effective amount of a MEK inhibitor according to Formula (Ia). Said diseases include cancer and inflammation

It is another aspect of the present invention to provide a MEK inhibitor according to Formula (Ia) for use as a medicament.

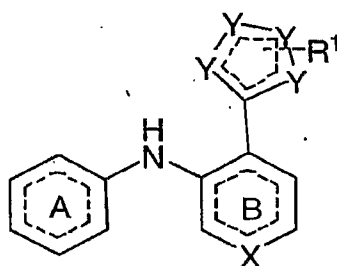
It is another aspect of the present invention to provide a MEK inhibitor according to
20 Formula (Ia) for the preparation of a medicament for the treatment of a hyperproliferative disease.

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It is furthermore an aspect of the present invention to provide a pharmaceutical formulation, which comprises a MEK inhibitor according to Formula (Ia) and a pharmaceutically acceptable carrier.

It is finally another aspect of the present invention to provide a process for making compounds according to Formula (Ia).

MEK inhibitors of the invention have the following general formula:



Formula (Ia)

wherein rings A and B, X, Y and R¹ are as defined in the detailed description.

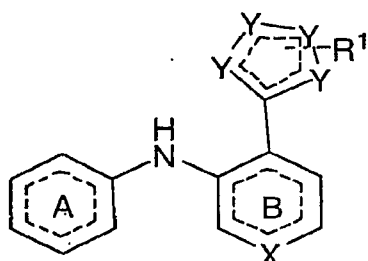
10 DETAILED DESCRIPTION OF THE INVENTION

In view of the foregoing it is one aspect of the present invention to provide novel compounds according to Formula (Ia) that are useful in the treatment of hyperproliferative diseases. Specifically, hyperproliferative diseases related to the hyperactivity of MEK as well as diseases associated to MEK, such as cancer and inflammation, in mammals.

As a result, this invention provides in a first aspect novel compounds as well as pharmaceutically acceptable salts and pharmaceutically active derivatives thereof, that are useful for the treatment of hyperproliferative diseases, such as cancer and inflammation.

20 The compounds are defined by Formula (Ia):

- 5 -



Formula (Ia)

as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof,

wherein:

- 5 rings A and B are optionally and independently substituted at any one or more substitutable ring carbon atoms with halogen, trimethylsilyl, cyano, C₁-C₆ alkyl, C₁-C₆ haloalkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ alkoxy, C₁-C₆ haloalkoxy, or phenyl wherein the phenyl is optionally substituted at any one or more substitutable ring carbon atoms with halogen, trimethylsilyl, cyano, C₁-C₆ alkyl, C₁-C₆ haloalkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, or C₁-C₆ alkoxy, C₁-C₆ haloalkoxy;

X is N or N→O;

Y is independently selected from NR¹, CR¹, S or O, whereby at least one Y is NR¹;

- R¹ is selected from hydrogen, trimethylsilyl, C₁-C₆-alkyl, OR⁶, C(O)OR⁶, NR⁷R⁸, SR⁶, NR⁷S(O)(O)R⁶, NR⁷C(O)R⁶, NR⁷C(O)NR⁷R⁶, NR⁷C(O)OR⁶, NR⁷C(O)C(O)OR⁶, C₁-C₄-alkyl-NR⁷R⁸, C₁-C₄-alkyl-heterocycloalkyl or C₁-C₄ alkyl-(OR⁶)_n;

- R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C₁-C₄ alkyl-C(O)OR⁶, C₁-C₄ alkyl-OR⁶, C₁-C₄-alkyl-NR⁷R⁸, C₁-C₄-alkyl-heterocycloalkyl, C₁-C₄ alkyl-(OR⁶)_n or C₁-C₄ alkyl-C(O)NR⁷R⁸, wherein said alkyl, cycloalkyl, aryl, heteroaryl heterocycloalkyl, and alkyl-heterocycloalkyl is substituted or unsubstituted; or

R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted;

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R' and R'' are independently selected from hydrogen, C₁-C₆-alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted; or

R' and R'' can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

n is 0 to 2.

In preferred embodiments, the variants have the following meanings:

R¹ is selected from OR⁶, NR⁷R⁸, NR⁷C(O)R⁶ or SR⁶;

10 rings A and B are optionally and independently substituted at any one or more substitutable ring carbon atoms with F, Cl, I, Br or phenyl; preferably F or I;

R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR', C₁-C₄ alkyl-(OR')_n, C₁-C₄-alkyl-NR'R'', C₁-C₄ alkyl-C(O)NR'R'' or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl is substituted or unsubstituted; or

15 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; preferably said cyclic ring is morpholinyl, piperidinyl or piperazinyl;

R' and R'' are independently either hydrogen or C₁-C₆-alkyl; or

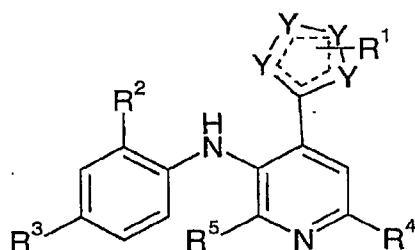
20 R' and R'' can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted.

As set forth above, the variants of each of R¹, R⁶, R⁷, R⁸, R' and R'' are optionally substituted. In this case they are independently substituted with 1 to 5, preferably 1 to 3, more preferably 1 or 2 groups, independently selected from halogen, cyano, nitro, CF₃, CHF₂, CH₂F, OCF₃, OCHF₂, OCH₂F, SCF₃, SCHF₂, SCH₂F, azido, amino, aminosulfonyl, sulfonylamine, sulfanyl, sulfonyl, sulfinyl, sulfonyloxy, acyl, acyloxy, carboxy, alkoxy or hydroxyl, preferably halogen, cyano, nitro, alkoxy, hydroxyl, CF₃, CHF₂, CH₂F, OCF₃, OCHF₂, OCH₂F, SCF₃, SCHF₂, SCH₂F, azido, more preferably

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halogen, cyano, nitro, CF₃, CHF₂, CH₂F, OCF₃, OCHF₂, OCH₂F, SCF₃, SCHF₂, SCH₂F, OH, methoxy, NH₂ or N(methyl)₂.

In one embodiment, the invention provides compounds of Formula (I):



Formula (I)

- 5 as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof,

wherein:

Each Y is independantly selected from NR', CR'' or O, whereby at least one Y is NR';

- 10 R¹ is selected from hydrogen, C₁-C₆-alkyl, OR⁶, NR⁷R⁸, SR⁶ or NR⁷S(O)(O)R',
NR⁷C(O)R⁶; NR⁷C(O)NR⁷R⁸; NR⁷C(O)OR⁶;

R² is halogen;

R³ is selected from Cl, F or I;

R⁴ and R⁵ are independently selected from hydrogen, halogen, C₁-C₆ alkyl or C₁-C₆ alkoxy, wherein said alkyl or alkoxy is substituted or unsubstituted;

- 15 R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, heterocycloalkyl; C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR' or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl, cycloalkyl, aryl, heteroaryl and heterocyclyl is substituted or unsubstituted; or

- 20 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

R' and R'' are independently selected from hydrogen, C₁-C₆ alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted.

In preferred embodiments, the variants have the following meanings:

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R¹ is selected from OR⁶, NR⁷R⁸ or SR⁶;

R² is selected from F, Cl, I or Br; preferably R² is F;

R³ is selected from F, Cl, I or C₁-C₆ alkyl, preferably R³ is I;

5 R⁴ and R⁵ are independently either hydrogen or C₁-C₆ alkyl, preferably R⁴ and R⁵ are hydrogen;

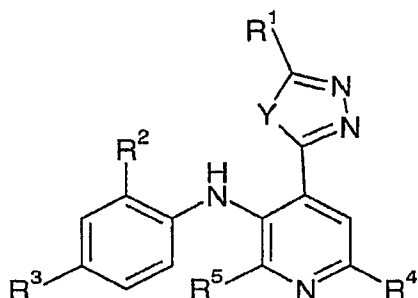
R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR' or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl is substituted or unsubstituted; or

10 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; preferably said cyclic ring is morpholinyl, piperidinyl or piperazinyl;

R' and R'' are independantly either hydrogen or C₁-C₆-alkyl.

15 As set forth above, the variants of each of R⁴, R⁵, R⁶, R⁷, R⁸, R' and R'' are optionally substituted. In this case they are independently substituted with 1 to 5, preferably 1 to 3, more preferably 1 or 2 groups, independently selected from halogen, cyano, nitro, CF₃, CHF₂, CH₂F, OCF₃, OCHF₂, OCH₂F, SCF₃, SCHF₂, SCH₂F, azido, amino, aminosulfonyl, sulfonylamine, sulfanyl, sulfonyl, sulfinyl, sulfonyloxy, acyl, acyloxy, carboxy, alkoxy or hydroxyl, preferably halogen, cyano, nitro, alkoxy, hydroxyl, CF₃,
20 CHF₂, CH₂F, OCF₃, OCHF₂, OCH₂F, SCF₃, SCHF₂, SCH₂F, azido, more preferably halogen, cyano, nitro, CF₃, CHF₂, CH₂F, OCF₃, OCHF₂, OCH₂F, SCF₃, SCHF₂, SCH₂F, OH, methoxy, NH₂ or N(methyl)₂.

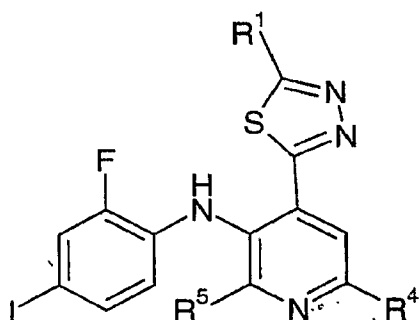
In one embodiment, the invention provides compounds of Formula (II):



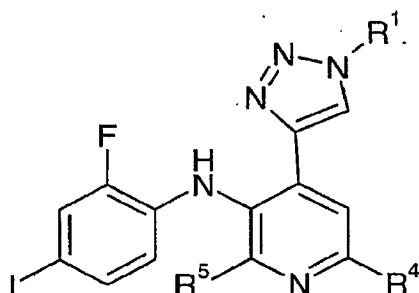
Formula (II) .

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or alternatively compounds of Formula (IIa) or Formula (IIb)



Formula (IIa)



Formula (IIb)

as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active
5 derivatives thereof,

the variables in Formula (II), Formula (IIa) and Formula (IIb) are defined as follows:

Y in Formula (II) is either NR⁷ or O;

R¹ is selected from hydrogen, trimethylsilyl, C₁-C₆-alkyl, OR⁶, C(O)OR⁶, NR⁷R⁸, SR⁶,
NR⁷S(O)(O)R¹, NR⁷C(O)R⁶, NR⁷C(O)NR⁷R⁶, NR⁷C(O)OR⁶, NR⁷C(O)C(O)OR⁶, C₁-
10 C₄-alkyl-NR⁷R⁸, C₁-C₄-alkyl-heterocycloalkyl or C₁-C₄ alkyl-(OR⁶)_n;

R² is selected from halogen, C₁-C₆-alkyl or OR⁶;

R³ is selected from halogen, trimethylsilyl, C₁-C₆-alkyl or OR⁶;

R⁴ and R⁵ are independently selected from hydrogen, halogen, C₁-C₆ alkyl or C₁-C₆
alkoxy, wherein said alkyl or alkoxy is substituted or unsubstituted;

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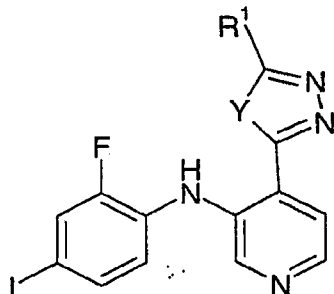
- 5 R^6 , R^7 and R^8 are independently selected from hydrogen, C_1 - C_{10} alkyl, C_3 - C_{10} cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C_1 - C_4 alkyl- $C(O)OR'$, C_1 - C_4 alkyl- OR' , C_1 - C_4 -alkyl- $NR'R''$, C_1 - C_4 -alkyl-heterocycloalkyl, C_1 - C_4 alkyl- $(OR')_n$ or C_1 - C_4 alkyl- $C(O)NR'R''$, wherein said alkyl, cycloalkyl, aryl, heteroaryl, heterocycloalkyl, and alkyl-heterocycloalkyl is substituted or unsubstituted; or
- R^7 and R^8 can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted;
- 10 R' and R'' are independently selected from hydrogen, C_1 - C_6 -alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted; or
- R' and R'' can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and
- n is 0 to 2.
- 15 In preferred embodiments, the variants in Formula (IIa) and Formula (IIb) have the following meanings:
- R^1 is selected from OR^6 , NR^7R^8 or SR^6 ;
- R^2 is selected from F, Cl, I or Br; preferably R^2 is F;
- 20 R^3 is selected from F, Cl or I, preferably R^3 is I;
- R^4 and R^5 are independently either hydrogen or C_1 - C_6 alkyl, preferably R^4 and R^5 are hydrogen;
- R^6 , R^7 and R^8 are independently selected from hydrogen, C_1 - C_{10} alkyl, C_1 - C_4 alkyl- $C(O)OR'$, C_1 - C_4 alkyl- OR' or C_1 - C_4 alkyl- $C(O)NR'R''$, wherein said alkyl is
 25 substituted or unsubstituted; or
- R^7 and R^8 can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; preferably said cyclic ring is morpholinyl, piperidinyl or piperazinyl;
- 30 R' and R'' are independently either hydrogen or C_1 - C_6 -alkyl; or

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R' and R'' can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted.

As set forth above, the variants of each of R⁴, R⁵, R⁶, R⁷, R⁸, R' and R'' are optionally substituted. In this case they are independently substituted with 1 to 5, preferably 1 to 3, more preferably 1 or 2 groups, independently selected from halogen, cyano, nitro, CF₃, CHF₂, CH₂F, OCF₃, OCHF₂, OCH₂F, SCF₃, SCHF₂, SCH₂F, azido, amino, aminosulfonyl, sulfonylamine, sulfanyl, sulfonyl, sulfinyl, sulfonyloxy, acyl, acyloxy, carboxy, alkoxy or hydroxyl, preferably halogen, cyano, nitro, alkoxy, hydroxyl, CF₃, CHF₂, CH₂F, OCF₃, OCHF₂, OCH₂F, SCF₃, SCHF₂, SCH₂F, azido, more preferably halogen, cyano, nitro, CF₃, CHF₂, CH₂F, OCF₃, OCHF₂, OCH₂F, SCF₃, SCHF₂, SCH₂F, OH, methoxy, NH₂ or N(methyl)₂.

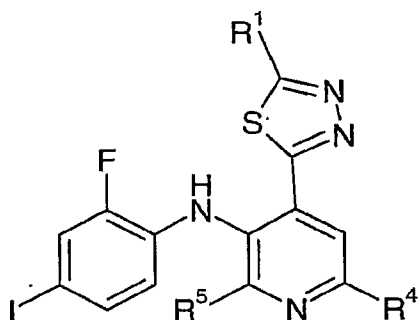
In another embodiment, the invention provides compounds of Formula (III):



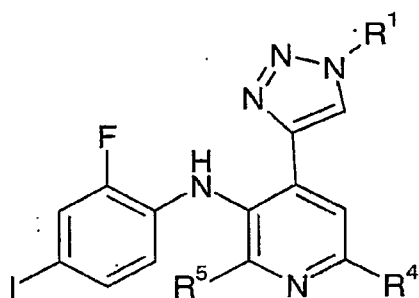
Formula (III)

or alternatively compounds of Formula (IIIa) or Formula (IIIb)

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Formula (IIIa)



Formula (IIIb)

as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof,

5 the variables in Formula (III), Formula (IIIa) and Formula (IIIb) are defined as follows:

Y in Formula (III) is either O or NR';

R¹ is selected from hydrogen, trimethylsilyl, C₁-C₆-alkyl, OR⁶, C(O)OR⁶, NR⁷R⁸, SR⁶, NR⁷S(O)(O)R', NR⁷C(O)R⁶, NR⁷C(O)NR⁷R⁶, NR⁷C(O)OR⁶, NR⁷C(O)C(O)OR⁶, C₁-C₄-alkyl-NR'R'', C₁-C₄-alkyl-heterocycloalkyl or C₁-C₄ alkyl-(OR')_n;

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R⁴ and R⁵ are independently selected from hydrogen, halogen, C₁-C₆ alkyl or C₁-C₆ alkoxy, wherein said alkyl or alkoxy is substituted or unsubstituted;

R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR', C₁-C₄-alkyl-NR'R'', C₁-C₄-alkyl-heterocycloalkyl, C₁-C₄ alkyl-(OR')_n or C₁-C₄

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alkyl-C(O)NR'R'', wherein said alkyl, cycloalkyl, aryl, heteroaryl, heterocycloalkyl, and alkyl-heterocycloalkyl is substituted or unsubstituted; or

5 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

R' and R'' are independently selected from hydrogen, C₁-C₆-alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted; or

10 R' and R'' can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

n is 0 to 2.

In preferred embodiments, the variants in Formula (IIIa) and Formula (IIIb) have the following meanings:

15 R¹ is selected from OR⁶, NR⁷R⁸ or SR⁶;

R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR' or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl is substituted or unsubstituted; or

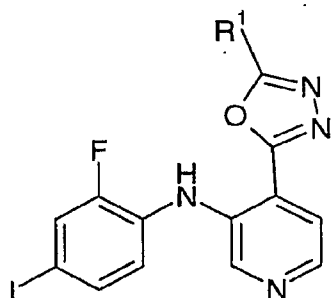
20 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 6 membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; preferably said cyclic ring is morpholinyl, piperidinyl or piperazinyl;

R' and R'' are independently either hydrogen or C₁-C₆-alkyl.

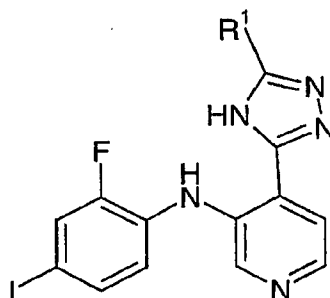
25 As said above, R⁶, R⁷, R⁸, R' and R'' are optionally substituted. In this case they are independently substituted as described above.

In a preferred embodiment, the invention provides compounds of the Formulae (IVa) or (IVb):

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Formula (IVa)



Formula (IVb)

as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof,

wherein

- 5 R¹ is selected from hydrogen, trimethylsilyl, C₁-C₆-alkyl, OR⁶, C(O)OR⁶, NR⁷R⁸, SR⁶, NR⁷S(O)(O)R¹, NR⁷C(O)R⁶, NR⁷C(O)NR⁷R⁶, NR⁷C(O)OR⁶, NR⁷C(O)C(O)OR⁶, C₁-C₄-alkyl-NR⁷R⁸, C₁-C₄-alkyl-heterocycloalkyl or C₁-C₄ alkyl-(OR⁷)_n;

- 10 R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C₁-C₄ alkyl-C(O)OR⁷, C₁-C₄ alkyl-OR⁷, C₁-C₄-alkyl-NR⁷R⁸, C₁-C₄-alkyl-heterocycloalkyl, C₁-C₄ alkyl-(OR⁷)_n or C₁-C₄ alkyl-C(O)NR⁷R⁸, wherein said alkyl, cycloalkyl, aryl, heteroaryl, heterocycloalkyl, and alkyl-heterocycloalkyl is substituted or unsubstituted; or

- 15 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

R⁷ and R⁸ are independently selected from hydrogen, C₁-C₆-alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted; or

- 20 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

n is 0 to 2.

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In preferred embodiments, the variants in Formula (IVa) and Formula (IVb) have the following meanings:

R¹ is selected from OR⁶, NR⁷R⁸ or SR⁶;

5 R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₁-C₄ alkyl-C(O)OR'; C₁-C₄ alkyl-OR', C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl is substituted or unsubstituted; or

10 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; preferably said cyclic ring is morpholinyl, piperidinyl or piperazinyl;

R' and R'' are independently either hydrogen or C₁-C₆-alkyl; or

15 R' and R'' can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted.

As said above R⁶, R⁷, R⁸, R' and R'' are optionally substituted. In this case they are independently substituted as described above.

20 Compounds according to Formula (I) include in particular those selected from of the group consisting of:

5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-3H-[1,3,4]oxadiazol-2-one,
[4-(5-Amino-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine,
5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-3H-[1,3,4]oxadiazole-2-thione,
5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-2,4-dihydro-[1,2,4]triazol-3-one,
25 (2-Fluoro-4-iodo-phenyl)-[4-(5-methylsulfonyl-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-amine,
and
(2-Fluoro-4-iodo-phenyl)-[4-(5-morpholin-4-yl-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-amine.

The employed terms have independently the meaning as described below:

30 A "substitutable ring carbon atom" in an aromatic group is a ring carbon bonded to a hydrogen atom. The hydrogen can be optionally replaced with a suitable substituent

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group. Thus, the term "substitutable ring carbon atom" does not include ring carbon atoms which are shared when two aromatic rings are fused. In addition, "substitutable ring atom" does not include ring carbon atoms when the structure depicts that they are already attached to a moiety other than hydrogen. Thus, the carbon atom bonded to R² in Structural Formula (II) is not a "substitutable ring atom" within the meaning of the term, as it is used herein.

"C₁-C₁₀-alkyl" refers to monovalent alkyl groups having 1 to 10 carbon atoms. This term is exemplified by groups such as methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, *tert*-butyl, n-hexyl and the like. By analogy, "C₁-C₁₂-alkyl" refers to monovalent alkyl groups having 1 to 12 carbon atoms, including "C₁-C₆-alkyl" groups and heptyl, octyl, nonyl, decanoyl, undecanoyl and dodecanoyl groups and "C₁-C₁₀-alkyl" refers to monovalent alkyl groups having 1 to 10 carbon atoms, "C₁-C₈-alkyl" refers to monovalent alkyl groups having 1 to 8 carbon atoms and "C₁-C₅-alkyl" refers to monovalent alkyl groups having 1 to 5 carbon atoms.

"Heteroalkyl" refers to C₁-C₁₂-alkyl, preferably C₁-C₆-alkyl, wherein at least one carbon has been replaced by a heteroatom selected from O, N or S, including 2-methoxy ethyl.

"Aryl" refers to an unsaturated aromatic carbocyclic group of from 6 to 14 carbon atoms having a single ring (e.g., phenyl) or multiple condensed rings (e.g., naphthyl). Aryl include phenyl, naphthyl, phenantrenyl and the like.

"C₁-C₆-alkyl aryl" refers to aryl groups having a C₁-C₆-alkyl substituent, including methyl phenyl, ethyl phenyl and the like.

"Aryl C₁-C₆-alkyl" refers to C₁-C₆-alkyl groups having an aryl substituent, including benzyl and the like.

"Heteroaryl" refers to a monocyclic heteroaromatic, or a bicyclic or a tricyclic fused-ring heteroaromatic group. Particular examples of heteroaromatic groups include optionally substituted pyridyl, pyrrolyl, pyrimidinyl, furyl, thienyl, imidazolyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, pyrazolyl, 1,2,3-triazolyl, 1,2,4-triazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, 1,3,4-triazinyl, 1,2,3-triazinyl, benzofuryl, [2,3-dihydro]benzofuryl, isobenzofuryl, benzothienyl, benzotriazolyl, isobenzothienyl, indolyl, isoindolyl, 3H-indolyl, benzimidazolyl, imidazo[1,2-a]pyridyl, benzothiazolyl, benzoxa-zolyl, quinoliziny, quinazoliny, pthalaziny, quinoxaliny,

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cinnolinyl, naphthyridinyl, pyrido[3,4-b]pyridyl, pyrido[3,2-b]pyridyl, pyrido[4,3-b]pyridyl, quinolyl, isoquinolyl, tetrazolyl, 1,2,3,4-tetrahydroquinolyl, 1,2,3,4-tetrahydroisoquinolyl, purinyl, pteridinyl, carbazolyl, xanthenyl or benzoquinolyl.

5 "C₂-C₆-alkenyl" refers to alkenyl groups preferably having from 2 to 6 carbon atoms and having at least 1 or 2 sites of alkenyl unsaturation. Preferable alkenyl groups include ethenyl (-CH=CH₂), n-2-propenyl (allyl, -CH₂CH=CH₂) and the like.

"C₂-C₆-alkynyl" refers to alkynyl groups preferably having from 2 to 6 carbon atoms and having at least 1-2 sites of alkynyl unsaturation, preferred alkynyl groups include ethynyl (-C≡CH), propargyl (-CH₂C≡CH), and the like.

10 "C₃-C₈-cycloalkyl" refers to a saturated carbocyclic group of from 3 to 8 carbon atoms having a single ring (e.g., cyclohexyl) or multiple condensed rings (e.g., norbornyl). C₃-C₈-cycloalkyl include cyclopentyl, cyclohexyl, norbornyl and the like.

"Heterocycloalkyl" refers to a C₃-C₈-cycloalkyl group according to the definition above, in which up to 3 carbon atoms are replaced by heteroatoms chosen from the group
15 consisting of O, S, NR, R being defined as hydrogen or methyl. Heterocycloalkyl include pyrrolidine, piperidine, piperazine, morpholine, tetrahydrofurane and the like.

"Acy" refers to the group -C(O)R where R includes "C₁-C₁₂-alkyl", preferably "C₁-C₆-alkyl", "aryl", "heteroaryl", "C₃-C₈-cycloalkyl", "heterocycloalkyl", "aryl C₁-C₆-alkyl", "heteroaryl C₁-C₆-alkyl", "C₃-C₈-cycloalkyl C₁-C₆-alkyl" or "heterocycloalkyl C₁-C₆-alkyl".

20 "Acyloxy" refers to the group -OC(O)R where R includes H, "C₁-C₆-alkyl", "C₂-C₆-alkenyl", "C₂-C₆-alkynyl", "C₃-C₈-cycloalkyl", "heterocycloalkyl", "aryl", "heteroaryl", "aryl C₁-C₆-alkyl" or "heteroaryl C₁-C₆-alkyl", "aryl C₂-C₆-alkenyl", "heteroaryl C₂-C₆-alkenyl", "aryl C₂-C₆-alkynyl", "heteroaryl C₂-C₆-alkynyl", "cycloalkyl C₁-C₆-alkyl", "heterocycloalkyl C₁-C₆-alkyl".

25 "Alkoxy" refers to the group -O-R where R includes "C₁-C₆-alkyl" or "aryl" or "heteroaryl" or "aryl C₁-C₆-alkyl" or "heteroaryl C₁-C₆-alkyl". Preferred alkoxy groups include for example, methoxy, ethoxy, phenoxy and the like.

"Aminocarbonyl" refers to the group -C(O)NRR' where each R, R' includes independently hydrogen or C₁-C₆-alkyl or aryl or heteroaryl or "aryl C₁-C₆-alkyl" or
30 "heteroaryl C₁-C₆-alkyl", including N-phenyl formamide.

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"Acylamino" refers to the group -NRC(O)R' where each R, R' is independently hydrogen, "C₁-C₆-alkyl", "C₂-C₆-alkenyl", "C₂-C₆-alkynyl", "C₃-C₈-cycloalkyl", "heterocycloalkyl", "aryl", "heteroaryl", "aryl C₁-C₆-alkyl" or "heteroaryl C₁-C₆-alkyl", "aryl C₂-C₆-alkenyl", "heteroaryl C₂-C₆-alkenyl", "aryl C₂-C₆-alkynyl", "heteroaryl C₂-C₆-alkynyl", "cycloalkyl C₁-C₆-alkyl", "heterocycloalkyl C₁-C₆-alkyl".

"Amino" refers to the group -NRR' where each R and R' is independently hydrogen or "C₁-C₆-alkyl" or "aryl" or "heteroaryl" or "C₁-C₆-alkyl aryl" or "C₁-C₆-alkyl heteroaryl", or "cycloalkyl", or "heterocycloalkyl", and where R and R', together with the nitrogen atom to which they are attached, can optionally form a 3-8-membered heterocycloalkyl ring.

"Sulfonylamino" refers to a group $\text{-NRSO}_2\text{-R'}$ where each R, R' includes independently hydrogen, "C₁-C₆-alkyl", "C₂-C₆-alkenyl", "C₂-C₆-alkynyl", "C₃-C₈-cycloalkyl", "heterocycloalkyl", "aryl", "heteroaryl", "aryl C₁-C₆-alkyl" or "heteroaryl C₁-C₆-alkyl", "aryl C₂-C₆-alkenyl", "heteroaryl C₂-C₆-alkenyl", "aryl C₂-C₆-alkynyl", "heteroaryl C₂-C₆-alkynyl", "cycloalkyl C₁-C₆-alkyl", "heterocycloalkyl C₁-C₆-alkyl".

"Aminosulfonyl" refers to a group $\text{-SO}_2\text{-NRR'}$ where each R, R' includes independently hydrogen, "C₁-C₆-alkyl", "C₂-C₆-alkenyl", "C₂-C₆-alkynyl", "C₃-C₈-cycloalkyl", "heterocycloalkyl", "aryl", "heteroaryl", "aryl C₁-C₆-alkyl" or "heteroaryl C₁-C₆-alkyl", "aryl C₂-C₆-alkenyl", "heteroaryl C₂-C₆-alkenyl", "aryl C₂-C₆-alkynyl", "heteroaryl C₂-C₆-alkynyl", "cycloalkyl C₁-C₆-alkyl", "heterocycloalkyl C₁-C₆-alkyl".

"Sulfonyloxy" refers to a group $\text{-OSO}_2\text{-R}$ wherein R is selected from H, "C₁-C₆-alkyl", "C₁-C₆-alkyl" substituted with halogens, e.g., an $\text{-OSO}_2\text{-CF}_3$ group, "C₂-C₆-alkenyl", "C₂-C₆-alkynyl", "C₃-C₈-cycloalkyl", "heterocycloalkyl", "aryl", "heteroaryl", "C₁-C₆-alkyl aryl" or "C₁-C₆-alkyl heteroaryl", "C₂-C₆-alkenyl aryl", "C₂-C₆-alkenyl heteroaryl", "C₂-C₆-alkynyl aryl", "C₂-C₆-alkynylheteroaryl", "C₁-C₆-alkyl cycloalkyl", "C₁-C₆-alkyl heterocycloalkyl".

"Sulfonyl" refers to group $\text{-SO}_2\text{-R}$ wherein R is selected from H, "aryl", "heteroaryl", "C₁-C₆-alkyl", "C₁-C₆-alkyl" substituted with halogens, e.g., an $\text{-SO}_2\text{-CF}_3$ group, "C₂-C₆-alkenyl", "C₂-C₆-alkynyl", "C₃-C₈-cycloalkyl", "heterocycloalkyl", "aryl", "heteroaryl", "C₁-C₆-alkyl aryl" or "C₁-C₆-alkyl heteroaryl", "C₂-C₆-alkenyl aryl", "C₂-C₆-alkenyl heteroaryl", "C₂-C₆-alkynyl aryl", "C₂-C₆-alkynylheteroaryl", "C₁-C₆-alkyl cycloalkyl", "C₁-C₆-alkyl heterocycloalkyl".

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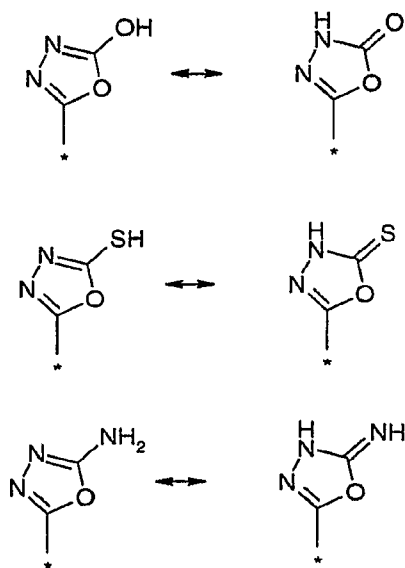
"Sulfinyl" refers to a group $-S(O)-R$ wherein R is selected from H, "C₁-C₆-alkyl", "C₁-C₆-alkyl" substituted with halogens, e.g., an $-SO-CF_3$ group, "C₂-C₆-alkenyl", "C₂-C₆-alkynyl", "C₃-C₈-cycloalkyl", "heterocycloalkyl", "aryl", "heteroaryl", "C₁-C₆-alkyl aryl" or "C₁-C₆-alkyl heteroaryl", "C₂-C₆-alkenyl aryl", "C₂-C₆-alkenyl heteroaryl", "C₂-C₆-alkynyl aryl", "C₂-C₆-alkynylheteroaryl", "C₁-C₆-alkyl cycloalkyl", "C₁-C₆-alkyl heterocycloalkyl".

"Sulfanyl" refers to groups $-S-R$ where R includes H, "C₁-C₆-alkyl", "C₁-C₆-alkyl" optionally substituted with halogens, e.g. a $-S-CF_3$ group, "C₂-C₆-alkenyl", "C₂-C₆-alkynyl", "C₃-C₈-cycloalkyl", "heterocycloalkyl", "aryl", "heteroaryl", "C₁-C₆-alkyl aryl" or "C₁-C₆-alkyl heteroaryl", "C₂-C₆-alkenyl aryl", "C₂-C₆-alkenyl heteroaryl", "C₂-C₆-alkynyl aryl", "C₂-C₆-alkynylheteroaryl", "C₁-C₆-alkyl cycloalkyl", "C₁-C₆-alkyl heterocycloalkyl". Preferred sulfanyl groups include methylsulfanyl, ethylsulfanyl, and the like.

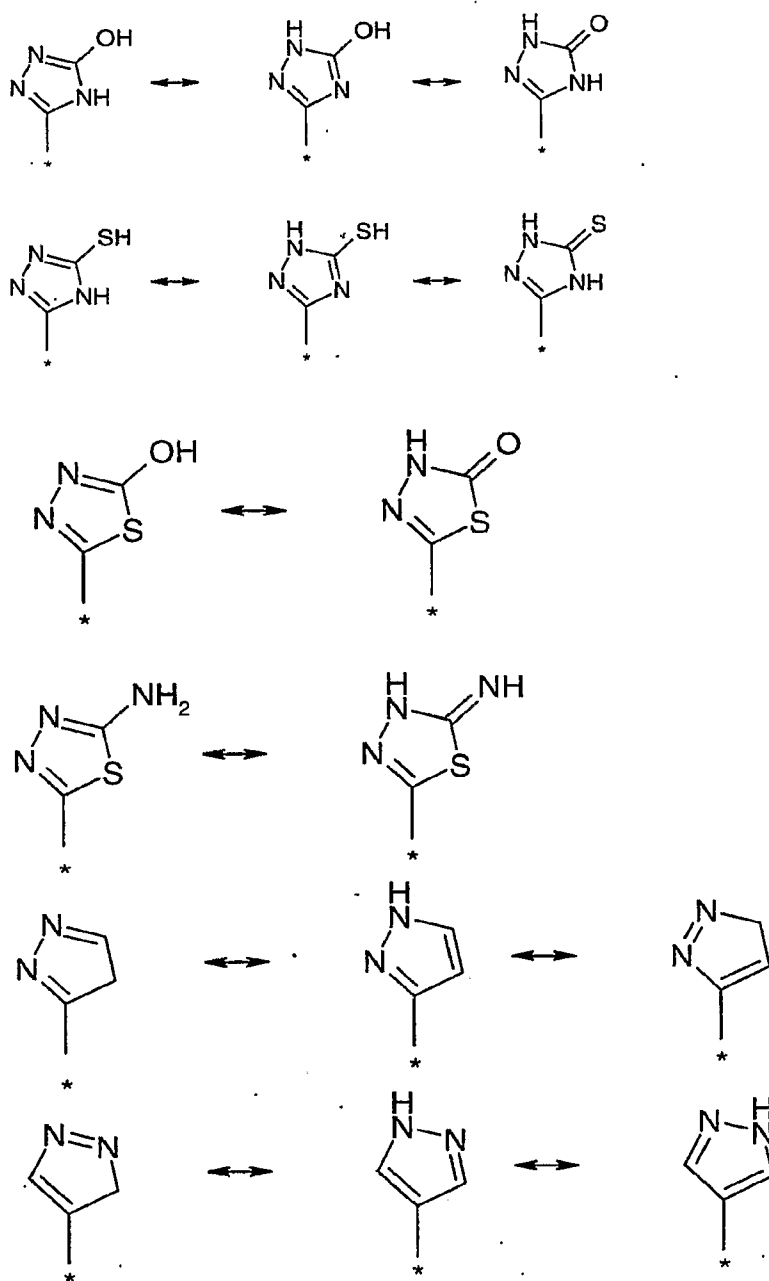
"Halogen" refers to fluoro, chloro, bromo and iodo.

"Cyano" refers to $C\equiv N$.

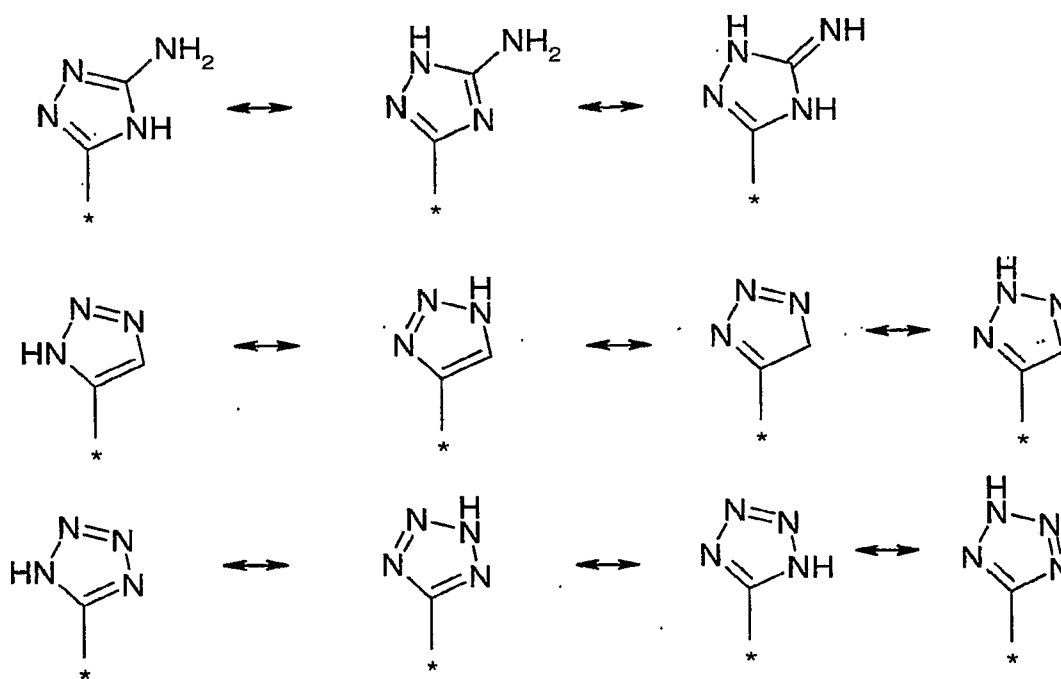
Where tautomerism, like e.g. keto-enol tautomerism, of compounds of the present invention or their prodrugs may occur, the individual forms, like e.g. the keto and enol form, are claimed separately and together as mixtures in any ratio. The same applies for stereoisomers, like e.g. enantiomers, cis/trans isomers, conformers and the like. Tautomers include but are not limited to the following structures:



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Metabolites of compounds of the present invention are also within the scope of the present invention.

In one embodiment the MEK inhibitor, in particular any of the cited MEK inhibitors
 5 inhibits the activity of MEK1 at a concentration of less than 100 μ M. In another
 embodiment the MEK inhibitor inhibits the activity of MEK1 at a concentration of less
 than 10 μ M. In another embodiment the MEK inhibitor inhibits the activity of MEK1 at a
 concentration of less than 1 μ M. In another embodiment the MEK inhibitor inhibits the
 activity of MEK1 at a concentration of less than 0.1 μ M.

10 The invention relates in a second aspect to a method of treating a mammal suffering
 from or susceptible to a hyperproliferative disease comprising administering to the
 mammal a therapeutically effective amount of a compound of Formula (I) or a
 pharmaceutically acceptable salt or a pharmaceutically active derivative thereof.

A hyperproliferative disease may be a cancerous disease including but not limiting to
 15 cancer types such as brain, lung, squamous cell, bladder, gastric, pancreatic, breast,
 head, neck, renal, kidney, ovarian, prostate, colorectal, esophageal, testicular,
 gynecological or thyroid cancer, or a non cancerous hyperproliferative disease such as

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benign hyperplasia of the skin (e.g., psoriasis), restenosis, or prostate (e.g., benign prostatic hypertrophy (BPH)).

In one embodiment the hyperproliferative disease is related to the hyperactivity of MEK as well as diseases associated to MEK in mammals, or diseases mediated by aberrant cell proliferation, such as cancer or inflammation.

A disease related to the "hyperactivity of MEK" refers to a disease, which can be treated by using any compound according to Formula (I) and which encompasses all diseases in which the upregulation and/or activity of MEK needs to be decreased irrespective of the cause of such disease.

10 "Pharmaceutically acceptable salts or complexes" refer to salts or complexes of the below-specified compounds of Formula (I). Examples of such salts include, but are not restricted, to base addition salts formed by reaction of compounds of Formula (I) with organic or inorganic bases such as hydroxide, carbonate or bicarbonate of a metal cation such as those selected in the group consisting of alkali metals (sodium, potassium or lithium), alkaline earth metals (e.g. calcium or magnesium), or with an organic primary, secondary or tertiary alkyl amine. Amine salts derived from methylamine, dimethylamine, trimethylamine, ethylamine, diethylamine, triethylamine, morpholine, N-Me-D-glucamine, N,N'-bis(phenylmethyl)-1,2-ethanediamine, tromethamine, ethanolamine, diethanolamine, ethylenediamine, N-methylmorpholine, procaine, piperidine, piperazine and the like are contemplated being within the scope of the instant invention. Also comprised are salts, which are formed with inorganic acids (e.g. hydrochloric acid, hydrobromic acid, sulfuric acid, phosphoric acid, nitric acid, and the like), as well as salts formed with organic acids such as acetic acid, oxalic acid, tartaric acid, succinic acid, malic acid, fumaric acid, maleic acid, ascorbic acid, benzoic acid, tannic acid, palmoic acid, alginic acid, polyglutamic acid, naphthalene sulfonic acid, methane sulfonic acid, naphthalene disulfonic acid, and poly-galacturonic acid, as well as salts formed with basic amino acids such as Lysine or Arginine.

A "pharmaceutically active derivative" refers to any compound that upon administration to the recipient, is capable of providing directly or indirectly, the activity disclosed herein. The term "indirectly" also encompasses prodrugs which may be converted to the active form of the drug via endogenous enzymes or metabolism. Said prodrug is comprised of the active drug compound itself and a chemical masking group. For

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example, a chemical masking group for alcohol derivatives could be selected from carboxylic acid ester (e.g. acetate, lysine ester) or phosphoric acid esters (e.g. phosphoric acid monoester).

5 In a preferred embodiment the invention relates to method of treating a mammal suffering from or susceptible to a hyperproliferative disease comprising administering to the mammal a therapeutically effective amount of a compound of Formulae (II), (IIIa) or (IIIb) or a pharmaceutically acceptable salt or a pharmaceutically active derivative thereof. In one embodiment said hyperproliferative disease is cancer.

10 The invention also relates to the treatment of a hyperproliferative disease in a mammal that comprises administering to said mammal a therapeutically effective amount of a compound of Formula (I), in combination with an anti-tumor agent selected from the group consisting of mitotic inhibitors, alkylating agents, antimetabolites, intercalating antibiotics, growth factor inhibitors, cell cycle inhibitors, enzyme inhibitors, topoisomerase inhibitors, biological response modifiers, antihormones, angiogenesis
15 inhibitors, and anti-androgens. It is well known in the art, which anti-tumor agent can be selected from combination. In a preferred embodiment, a compound of Formulae (II), (III), (IVa) or (IVb) is administered in combination with an anti-tumor agent as described above.

20 The invention relates in a third aspect to the use of a MEK inhibitor according to Formula (I) or a pharmaceutically acceptable salt or a pharmaceutically active derivative thereof as a medicament.

In a preferred embodiment, the invention relates to the use of a compound according to Formulae (II), (III), (IVa) or (IVb) or a pharmaceutically acceptable salt or a pharmaceutically active derivative thereof as a medicament. In one embodiment the
25 invention relates to the use of a compound selected from the group of

5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-3H-[1,3,4]oxadiazol-2-one,
[4-(5-Amino-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine,
5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-3H-[1,3,4]oxadiazole-2-thione,
5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-2,4-dihydro-[1,2,4]triazol-3-one,
30 (2-Fluoro-4-iodo-phenyl)-[4-(5-methylsulfonyl-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-amine,
and

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(2-Fluoro-4-iodo-phenyl)-[4-(5-morpholin-4-yl-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-amine as a medicament.

The invention relates to all of the compounds disclosed in the examples.

- 5 The invention relates in a forth aspect to the use of a MEK inhibitor according to Formula (I) or a pharmaceutically acceptable salt or a pharmaceutically active derivative thereof for the preparation of a medicament for the treatment of a hyperproliferative disease. In one embodiment, said use relates to the preparation of a medicament for the treatment of cancer.
- 10 In a preferred embodiment the invention relates to the use of a compound according to Formulae (II), (III), (IVa) or (IVb) or a pharmaceutically acceptable salt or a pharmaceutically active derivative thereof for the preparation of a medicament for the treatment of a hyperproliferative disease. In one embodiment, said use relates to the preparation of a medicament for the treatment of cancer.
- 15 The invention relates in a fifth aspect to the use of a MEK inhibitor according to Formula (I) or a pharmaceutically acceptable salt or a pharmaceutically active derivative thereof for the treatment of a hyperproliferative disease. In one embodiment, said use relates to the treatment of cancer.

- In a preferred embodiment the invention relates to the use of a compound according to
- 20 Formulae (II), (III), (IVa) or (IVb) or a pharmaceutically acceptable salt or a pharmaceutically active derivative thereof for the treatment of a hyperproliferative disease. In one embodiment, said use relates to the preparation of a medicament for the treatment of cancer.

- Furthermore, the invention provides in a sixth aspect pharmaceutical compositions
- 25 comprising a compound of Formula (I), or a pharmaceutically acceptable salt or a pharmaceutically active derivative thereof, as active ingredient together with a pharmaceutically acceptable carrier.

In a preferred embodiment the invention relates to a pharmaceutical composition comprising a compound of Formulae (II), (III), (IVa) or (IVb), or a pharmaceutically

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acceptable salt or a pharmaceutically active derivative thereof, as active ingredient together with a pharmaceutically acceptable carrier.

"Pharmaceutical composition" means one or more active ingredients, and one or more inert ingredients that make up the carrier, as well as any product which results, directly or indirectly, from combination, complexation or aggregation of any two or more of the ingredients, or from dissociation of one or more of the ingredients, or from other types of reactions or interactions of one or more of the ingredients. Accordingly, the pharmaceutical compositions of the present invention encompass any composition made by admixing a compound of the present invention and a pharmaceutically acceptable carrier.

A pharmaceutical composition of the present invention may additionally comprise one or more other compounds as active ingredients like one or more additional compounds of the present invention, or a prodrug compound or other MEK inhibitors.

The compositions include compositions suitable for oral, rectal, topical, parenteral (including subcutaneous, intramuscular, and intravenous), ocular (ophthalmic), pulmonary (nasal or buccal inhalation), or nasal administration, although the most suitable route in any given case will depend on the nature and severity of the conditions being treated and on the nature of the active ingredient. They may be conveniently presented in unit dosage form and prepared by any of the methods well-known in the art of pharmacy.

In practical use, the compound of Formula (I) can be combined as the active ingredient in admixture with a pharmaceutical carrier according to conventional pharmaceutical compounding techniques. The carrier may take a wide variety of forms depending on the form of preparation desired for administration, e.g., oral or parenteral (including intravenous). In preparing the compositions for oral dosage form, any of the usual pharmaceutical media may be employed, such as, for example, water, glycols, oils, alcohols, flavoring agents, preservatives, coloring agents and the like in the case of oral liquid preparations, such as, for example, suspensions, elixirs and solutions; or carriers such as starches, sugars, microcrystalline cellulose, diluents, granulating agents, lubricants, binders, disintegrating agents and the like in the case of oral solid

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preparations such as, for example, powders, hard and soft capsules and tablets; with the solid oral preparations being preferred over the liquid preparations.

Because of their ease of administration, tablets and capsules represent the most advantageous oral dosage unit form in which case solid pharmaceutical carriers are obviously employed. If desired, tablets may be coated by standard aqueous or nonaqueous techniques. Such compositions and preparations should contain at least 0.1 percent of active compound. The percentage of active compound (a compound of Formulae (I), (II), (III), (IVa) or (IVb) in these compositions may, of course, be varied and may conveniently be between about 2 percent to about 60 percent of the weight of the unit. The amount of active compound in such therapeutically useful compositions is such that an effective dosage will be obtained. The active compounds can also be administered intranasally as, for example, liquid drops or spray.

The tablets, pills, capsules, and the like may also contain a binder such as gum tragacanth, acacia, corn starch or gelatin; excipients such as dicalcium phosphate; a disintegrating agent such as corn starch, potato starch, alginic acid; a lubricant such as magnesium stearate; and a sweetening agent such as sucrose, lactose or saccharin. When a dosage unit form is a capsule, it may contain, in addition to materials of the above type, a liquid carrier such as a fatty oil.

Various other materials may be present as coatings or to modify the physical form of the dosage unit. For instance, tablets may be coated with shellac, sugar or both. A syrup or elixir may contain, in addition to the active ingredient, sucrose as a sweetening agent, methyl and propylparabens as preservatives, a dye and a flavoring such as cherry or orange flavor.

Compounds according to Formulae (I), (II), (III), (IVa) or (IVb) may also be administered parenterally. Solutions or suspensions of these active compounds can be prepared in water suitably mixed with a surfactant such as hydroxy-propylcellulose. Dispersions can also be prepared in glycerol, liquid polyethylene glycols and mixtures thereof in oils. Under ordinary conditions of storage and use, these preparations contain a preservative to prevent the growth of microorganisms.

The pharmaceutical forms suitable for injectable use include sterile aqueous solutions or dispersions and sterile powders for the extemporaneous preparation of sterile

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injectable solutions or dispersions. In all cases, the form must be sterile and must be fluid to the extent that easy syringability exists. It must be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms such as bacteria and fungi. The carrier can be a solvent or dispersion
5 medium containing, for example, water, ethanol, polyol (e.g., glycerol, propylene glycol and liquid polyethylene glycol), suitable mixtures thereof, and vegetable oils.

Any suitable route of administration may be employed for providing a mammal, especially a human, with an effective dose of a compound of the present invention. For example, oral, rectal, topical, parenteral, ocular, pulmonary, nasal, and the like may be
10 employed. Dosage forms include tablets, troches, dispersions, suspensions, solutions, capsules, creams, ointments, aerosols, and the like. Preferably compounds of the present invention are administered orally.

The effective dosage of active ingredient employed may vary depending on the particular compound employed, the mode of administration, the condition being treated
15 and the severity of the condition being treated. Such dosage may be ascertained readily by a person skilled in the art.

When treating cancer or other hyperproliferative diseases for which compounds of Formulae (I), (II), (III), (IVa) or (IVb) are indicated, generally satisfactory results are obtained when the compounds of the present invention are administered at a daily
20 dosage of from about 0.1 milligram to about 100 milligram per kilogram of animal body weight, preferably given as a single daily dose or in divided doses two to six times a day, or in sustained release form. For most large mammals, the total daily dosage is from about 1.0 milligrams to about 1000 milligrams, preferably from about 1 milligram to about 50 milligrams. In the case of a 70 kg adult human, the total daily dose will
25 generally be from about 7 milligrams to about 350 milligrams. This dosage regimen may be adjusted to provide the optimal therapeutic response.

Furthermore, the invention provides in a seventh aspect a method of preparing a compound of Formula (I).

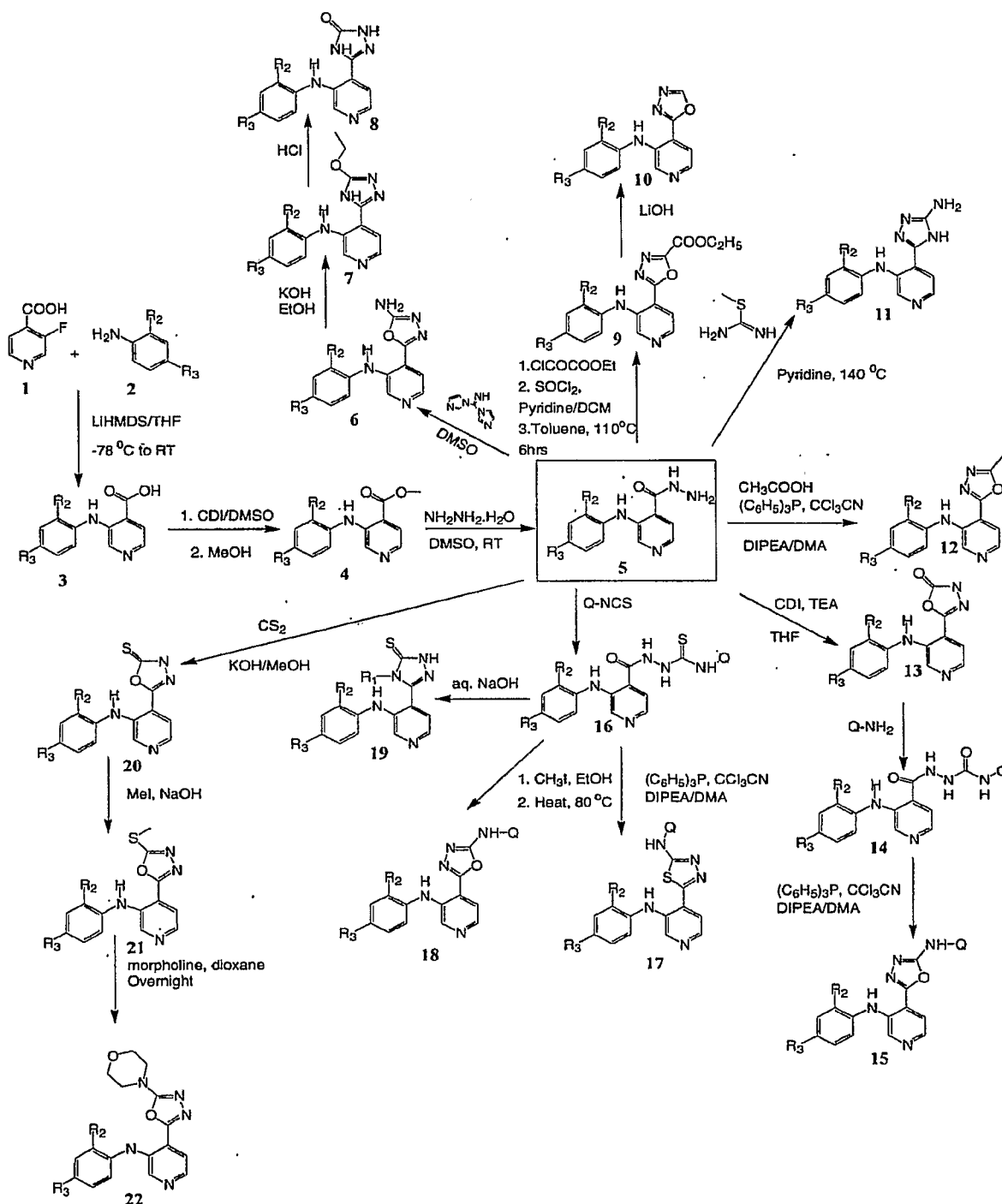
In a preferred embodiment the invention relates to a method of preparing compounds
30 of Formulae (II), (III), (IVa) or (IVb).

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The compounds according to Formula (I) can be prepared according to the procedures of the following Schemes and Examples, using appropriate materials and are further exemplified by the following specific examples. Moreover, by utilizing the procedures described herein, in conjunction with ordinary skills in the art, additional compounds of the present invention claimed herein can be readily prepared. The compounds illustrated in the examples are not, however, to be construed as forming the only genus that is considered as the invention. The examples further illustrate details for the preparation of the compounds according to Formula (I). Those skilled in the art will readily understand that known variations of the conditions and processes of the following preparative procedures can be used to prepare these compounds. The instant compounds are generally isolated in the form of their pharmaceutically acceptable salts, such as those described above. The amine-free bases corresponding to the isolated salts can be generated by neutralization with a suitable base, such as aqueous sodium hydrogencarbonate, sodium carbonate, sodium hydroxide and potassium hydroxide, and extraction of the liberated amine-free base into an organic solvent, followed by evaporation. The amine-free base, isolated in this manner, can be further converted into another pharmaceutically acceptable salt by dissolution in an organic solvent, followed by addition of the appropriate acid and subsequent evaporation, precipitation or crystallization.

Illustrations of the preparation of compounds according to Formula (Ia) are shown in the following schemes. Unless otherwise indicated in the schemes, the variables have the same meaning as described above.

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Scheme 1

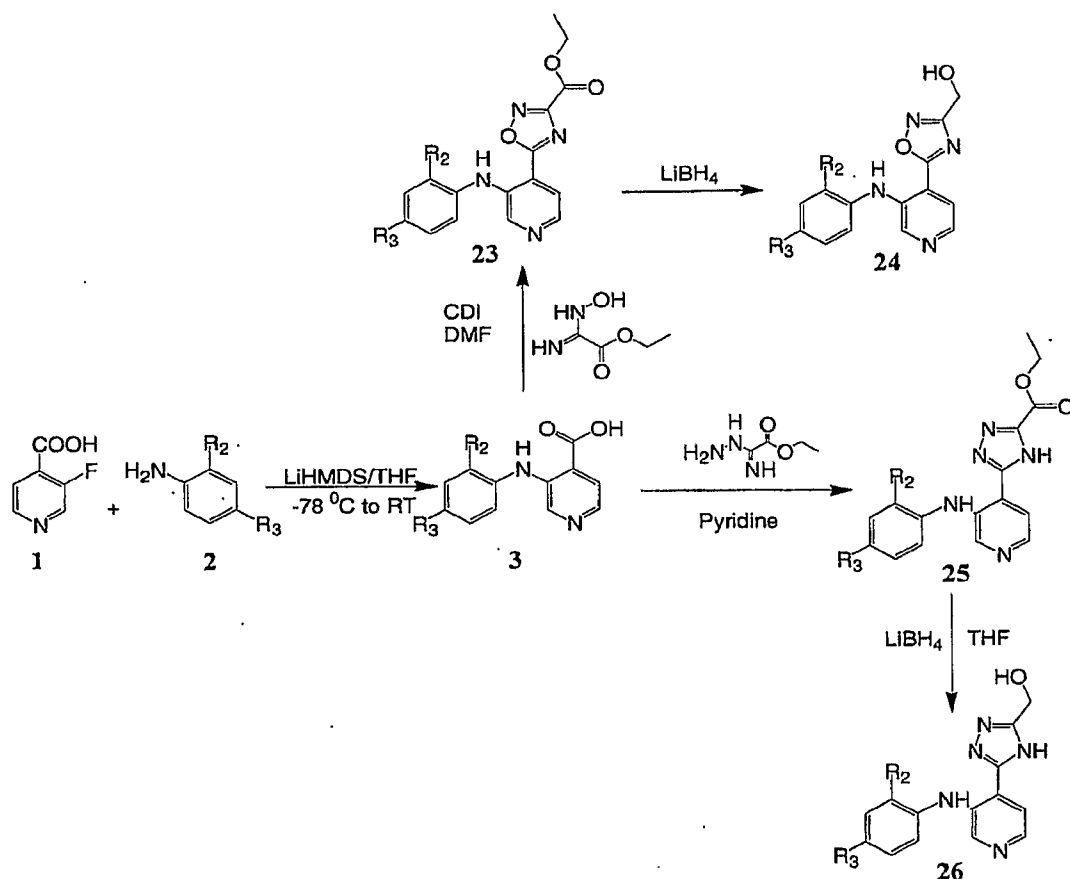
Scheme 1 illustrates the synthesis of a group of example compounds in the present invention. In first step anilines (2) are reacted with 3-fluoro isonicotinic acids (1) in inert

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solvents, preferably THF, by addition of a base, preferably but not limited to LiHMDS. In step 2 the 3-anilino isonicotinic acids (**3**) are transformed to simple esters such as methyl ester (**4**) by employing coupling agents such as PyBOP; CDI etc. and simple alcohols, e.g. methanol. The resulting esters are easily converted to hydrazides (**5**) by heating with the corresponding hydrazines in alcoholic solvents. These hydrazides, as illustrated in Scheme 1, serve as key intermediates for the formation of a great variety of heterocyclic rings. For example, treatment with 1,1-di-1H-imidazol-1-yl-methyleneamine affords [1,3,4]oxadiazol-2-amines such as compound **6** in Scheme 1. These compounds can be rearranged by heating in alcoholic solutions of KOH to 5-alcoxy-[1,2,4]triazoles (**7**) which can be in turn hydrolyzed with acid to the corresponding triazolones (**8**). Hydrazides (**5**) can also react with 3-Chloro-2-oxopropionic acetates and then can be cyclized into [1,3,4]oxadiazole-2-carboxylate derivatives (**9**), which in turn can be decarboxylated to unsubstituted oxadiazoles (**10**) with treatment with lithium or sodium hydroxide or other similar reagents.

15 Coupling/cyclization of **5** with 2-amino-thioacetimidic acid esters affords [1,2,4]triazoles-3-amines (**11**). Condensations of **5** with acids in the presence of triphenylphosphine/trichloroacetonitrile affords 2-alkyl-[1,3,4]oxadiazoles (**12**). Condensation of **5** with carbonyldiimidazole results in [1,3,4]oxadiazol-2-ones (**13**). The latter can be ring-opened with amines to analogs such as **14**, which in turn can be cyclized into *N*-substituted [1,3,4]oxadiazoles-2-amines (**15**). Reaction of **5** with isothiocyanates affords analogs such as **16** which can be cyclized into either substituted [1,3,4]thiazodiazoles-3-amines (**17**), or [1,3,4]oxadiazoles-2-amines (**18**), or [1,2,4]triazole-3-thiones (**19**). Condensation of **5** with carbon disulfide under basic conditions afforded [1,3,4]oxadiazol-2-thiones (**20**), which can be S-alkylated to compounds **21**, which in turn can react with various nucleophiles to afford 2-substituted [1,3,4]oxadiazole analogs, such as **22**.

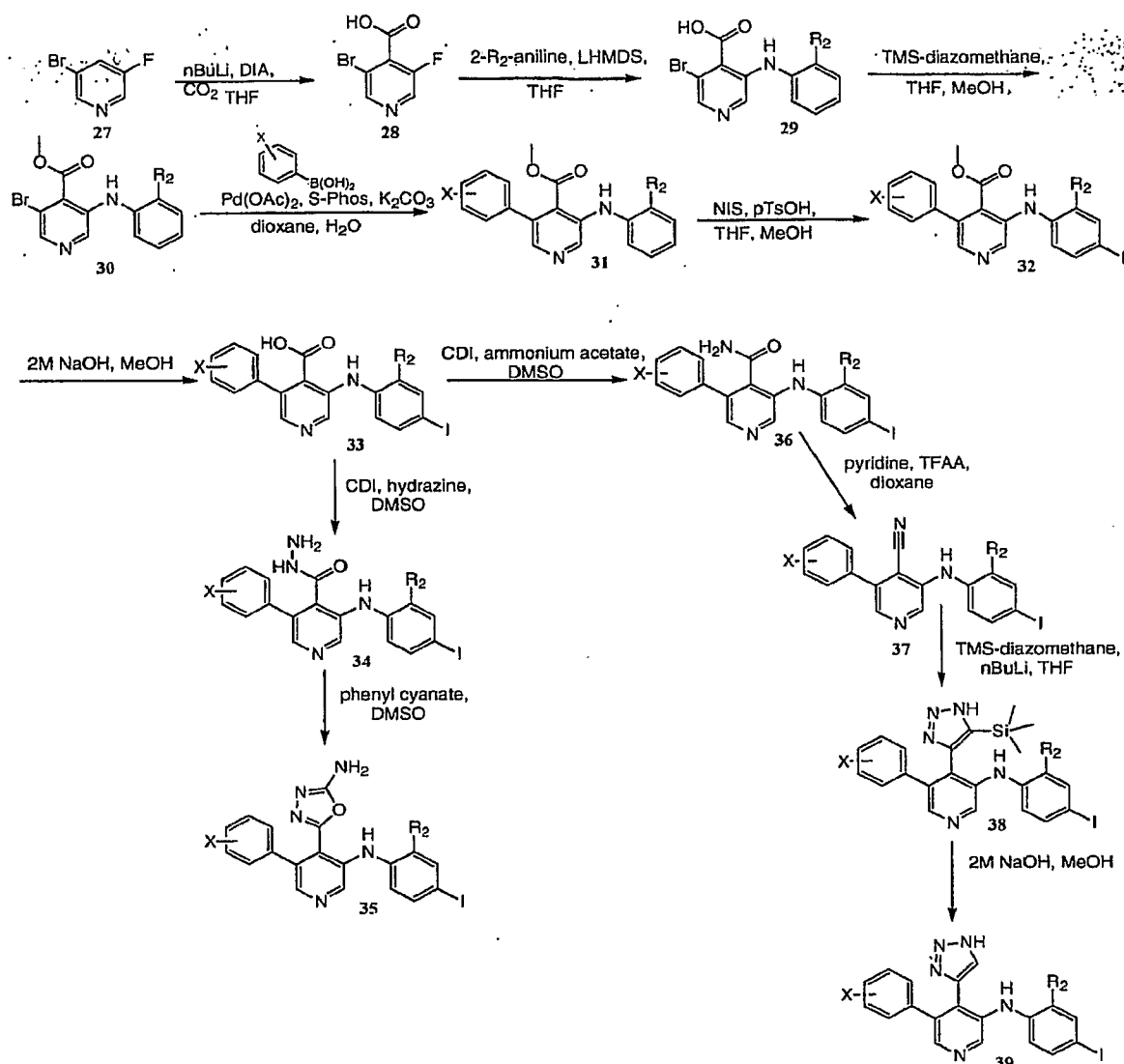
- 31 -



Scheme 2

Scheme 2 illustrates the general synthesis of examples with various heterocyclic systems stemming from intermediate acids **3** instead of hydrazides **5**. Acids **3** can be coupled with hydroxylamino-imino-acetates and eventually cyclize into 1,2,4-oxadiazole-3-carboxylic acid esters (**23**) which in turn can be reduced to the corresponding hydroxymethyl derivatives (**24**). Finally, intermediate **3**, similarly to the reactions that formed **23**, and **24** can react with hydrazine-imino-acetates to form 1,2,4-triazole-3-carboxylic acid esters (**25**) and the corresponding alcohols (**26**) after borohydride reduction.

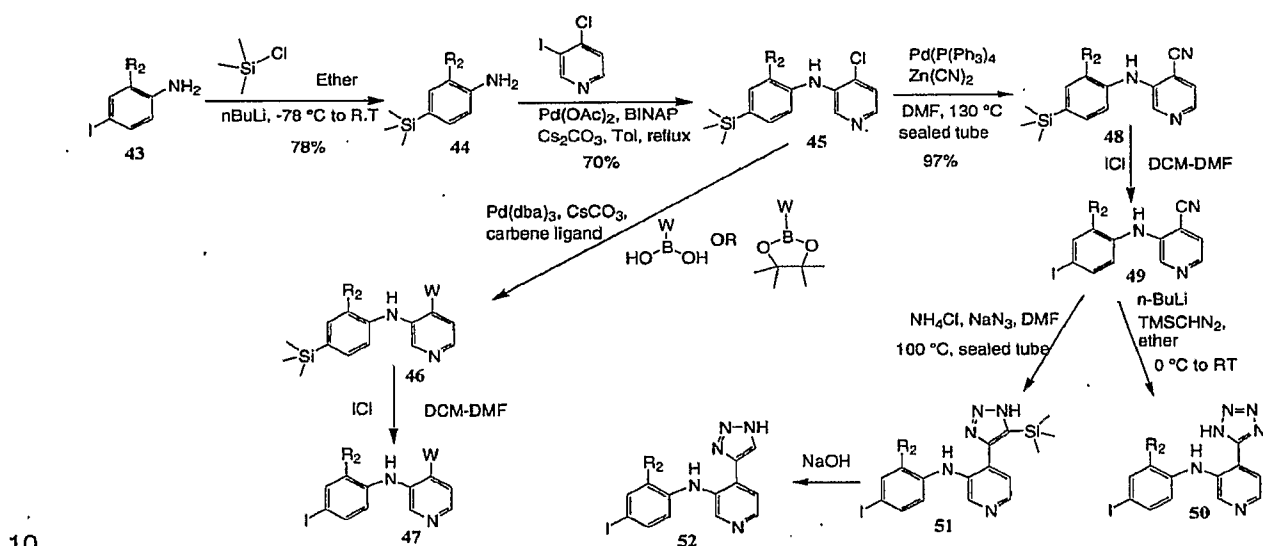
- 32 -

**Scheme 3**

Scheme 3 illustrates the synthesis of examples with substitutions in the 5 position of the pyridine core, and in particularly phenyl substitutions. 3-Bromo-5-Fluoro-pyridine (27) is first 4-lithiated and then treated with CO₂ to form the isonicotinic acid (28), which in turn is coupled with various anilines to afford biaryl systems such as 29. The acid functionality is first protected as simple esters (30) and then reacted under Suzuki or other palladium-catalyzed coupling conditions to afford a variety of 5-substituted analogs, including the ones described with structure 31. Halogenation, such as iodination with NIS under acidic conditions, gives rise to compounds under structure

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32, which are deprotected to the free acids (33). These acids are activated with typical coupling reagents, such as CDI or others, to react with either hydrazines or amines to produce hydrazides (34) or amides (36) respectively. The former are in turn cyclized into systems similar to the ones described in Scheme 1, including aminoxadiazoles such as structure 35. Amides 36 can typically be first dehydrated to the corresponding nitriles (37), which are useful intermediates in the construction of various heterocyclic compounds. The example shown in Scheme 3 is the conversion of nitrile 37 into triazoles 38 and 39 by reaction with diazomethane.



Scheme 4

Scheme 4 illustrates the general synthesis of additional examples with various types of **W** substituents. According to this scheme, various 4-iodo-anilines (43) are converted to the corresponding 4-trimethylsilyl derivatives (44), which in turn are coupled under palladium-catalyzed conditions with various 4-chloro-pyridines to form key (4-chloro-pyridin-3-yl-phenyl)-amines (45). These intermediates are functionalised using a variety of either Suzuki or other palladium-catalyzed coupling reactions to introduce a great variety of **W** groups in the 4-pyridyl position (46). These compounds are typically converted to their halogenated analogs (47) (e.g.iodo or other) by displacement of the trimethylsilyl group. Alternatively, the 4-chloropyridyl group of 45 can be converted to a 4-cyanopyridyl group by heating in sealed tube in the presence of Pd(PPh₃)₄ and

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$\text{Zn}(\text{CN})_2$. This cyano intermediate (**48**) allows the access of various compounds with certain heterocyclic ring systems in the 4-pyridyl position, such as tertazoles (**50**), triazoles (**51**, **52**), and others.

5 Suitable anilines, pyridines, and other derivatives, reagents, and starting materials are commercially available from Alpha Aesar, Ward Hill, MA, USA; Sigma-Aldrich Milwaukee, WI, USA; or from Acros Organics, Morris Plains, NJ, USA; or can be routinely prepared by procedures described in "March's Advanced Organic Chemistry: Reactions, Mechanisms, and Structure", 5th Edition; John Wiley & Sons.

10 Unless otherwise noted, all non-aqueous reactions were carried out either under an argon or nitrogen atmosphere with commercial dry solvents. Compounds were typically purified using flash column chromatography using Merck silica gel 60 (230-400 mesh). In certain cases reverse phase preparative HPLC was used for purification. The ^1H -NMR spectra were recorded on a Joel ECP-400 (400 MHz for ^1H -NMR) using typically d_6 -dimethylsulfoxide or d_4 -methanol as solvent; chemical shifts are reported in ppm
15 relative to tetramethylsilane.

Analytical LC/MS was performed using the following two methods:

Method A: A Discovery® C^{18} , 5 μm , 3 x 30 mm column was used at a flow rate of 400 $\mu\text{L}/\text{min}$, sample loop 5 μL , mobile phase: (A) water with 0.1% formic acid, mobile phase, (B) methanol with 0.1% formic acid; retention times are given in minutes.

20 Method details: (I) runs on a Quaternary Pump G1311A (Agilent) with UV/Vis diode array detector G1315B (Agilent) and Finnigan LCQ Duo MS detector in ESI + modus with UV-detection at 254 and 280 nm with a gradient of 15-95% (B) in a 3.2 min linear gradient (II) hold for 1.4 min at 95% (B) (III) decrease from 95-15% (B) in a 0.1 min linear gradient (IV) hold for 2.3 min at 15% (B).

25 Method B: A Waters Symmetry® C^{18} , 3.5 μm , 4.6 x 75 mm column at a flow rate of 1 mL/min, sample loop 10 μL , mobile phase (A) is water with 0.05% TFA, mobile phase (B) is ACN with 0.05% TFA; retention times are given in minutes. Methods details: (I) runs on a Binary Pump G1312A (Agilent) with UV/Vis diode array detector G1315B (Agilent) and Applied Biosystems API3000 MS detector in ESI + modus with UV-
30 detection at 254 and 280 nm with a gradient of 20-85% (B) in a 10 min linear gradient (II) hold for 1 min at 85% (B) (III) decrease from 20-85% (B) in a 0.2 min linear gradient (IV) hold for 3.8 min at 20% (B).

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Abbreviations

Designation

CDI	N,N-Carbonyldiimidazole
DCM	Dichloromethane
DIPEA	N-Ethyldiisopropylamine
DMF	N,N-Dimethylformamide
DMSO	Dimethylsulfoxide
EDC	1-(3-Dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride
HPLC	High pressure liquid chromatography
LC/MS	Liquid chromatography/Mass Spectrometry
LiHMDS	Lithium hexamethyldisilazide
MCPBA	3-Chloroperoxybenzoic acid
NT	Not Tested
NMR	Nuclear Magnetic Resonance
PyBOP	Benzotriazole-1-yl-oxy-trispyrrolidinophosphonium hexafluorophosphate
RT	Room Temperature
TEA	Triethylamine
Tert	Tertiary-butyl
TFA	Trifluoroacetic acid
THF	Tetrahydrofuran
TLC	Thin Layer Chromatography

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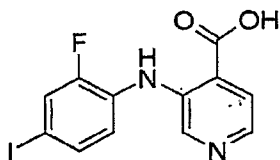
10 WO 05/051301

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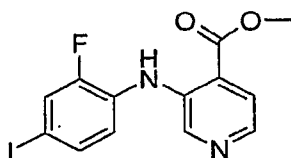
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EXAMPLES

The examples presented below are intended to illustrate particular embodiments of the invention, and are not intended to limit the scope of the specification or the claims in any way. The example and intermediate numbering used in this section are independent of the numbering used in Schemes 1-4.

Intermediate 1: 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid

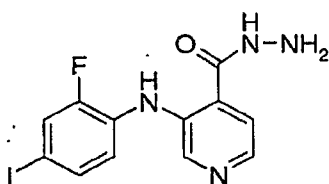
A mixture of 4-iodo-2-fluoroaniline (20.0 g, 84.38 mmol) in anhydrous tetrahydrofuran (80 mL) was cooled to -65°C under an inert atmosphere, prior to slow addition of 1.0 M lithium bis(trimethylsilyl)amide (255 mL, 255 mmol) at a rate that maintained the internal temperature below -55°C . After final addition, the thick slurry was stirred for 30 minutes and then treated with 3-fluoro-isonicotinic acid (8.0 g, 56.69 mmol). The mixture was stirred at room temperature for 4 days and then poured into aqueous 2.0 N sodium hydroxide (1000 mL) and ethyl acetate (250 mL). The layers were separated and the organics were again extracted with aqueous sodium hydroxide (2 x 1000 mL). The pH of the combined aqueous fractions was adjusted to 2 with concentrated hydrochloric acid, which effected precipitation of a solid. The material was filtered, washed with water (300 mL) and dried under high vacuum at 40°C for 18 h to afford 3 (19.05 g, 53.19 mmol, 94%) as a yellow solid. MS [359 (M+1)]. ^1H NMR (400 MHz, DMSO- D_6): 7.3565 (t, $J = 8.8$ Hz, 1H), 7.5509 (d, $J = 8.4$ Hz, 1H), 7.7521 (m, 2H), 8.1175 (d, $J = 4.7$ Hz, 1H), 8.4472 (s, 1H), 9.2184 (s, 1H)

Intermediate 2: 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid methyl ester

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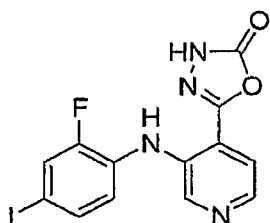
A suspension of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid (12.0 g, 33.51 mmol) and CDI (7.0 g, 43.17 mmol) in anhydrous dimethyl sulfoxide (120 mL) was stirred at room temperature for 16 h, prior to addition of methanol (15.0 g, 468.15 mmol). The dark-yellow solution was stirred at room temperature for 2 h and then slowly poured into water (900 mL). The resultant precipitate was filtered and the cake was washed with water (2 x 300 mL) and dried under high vacuum at 40 °C for 16 h to afford **2** (12.11 g, 32.54 mmol, 97%) as a salmon-colored solid. ¹H-NMR: δ 3.88 (s, 3H); 7.30 (dd, 1H); 7.54 (ddd, 1H); 7.70 (d, 1H); 7.74 (dd, 1H); 8.13 (d, 1H); 8.41 (d, 1H); 8.88 (br s, 1H).

Intermediate 3: 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide



A suspension of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid methyl ester (11.50 g, 30.90 mmol) in isopropanol (200 mL) was treated with hydrazine hydrate (6 mL) and the mixture was heated to gentle reflux for 17 h. The resultant precipitate was filtered, washed with isopropanol (50 mL) and dried under high vacuum for 14 h to give **3** (9.46 g, 25.42 mmol, 82%) as a yellow, crystalline solid. ¹H-NMR: δ 4.65 (br s, 2H); 7.28 (dd, 1H); 7.49 (m, 2H); 7.68 (dd, 1H); 8.14 (d, 1H); 8.52 (s, 1H); 9.34 (br s, 1H); 10.22 (br s, 1H).

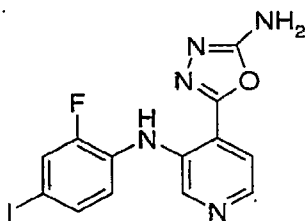
Example 1: 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-3H-[1,3,4]oxadiazol-2-one



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To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (152mg, 0.41mmol) in THF (6mL) contained TEA (60μl, 0.43mmol) were added Di-imidazol-1-yl-methanone (95mg, 0.57mmol) at 0°C. The reaction mixture was warmed to RT and stirred for 5h, additional TEA (40μl) and CDI (50mg) were added, and the reaction mixture was stirred overnight at RT. The precipitate was filtered, the solid washed with MeOH. Yield: yellow solid 123mg. LC-MS (Method A) [5.50min; 399(M+1)].

Example 2: [4-(5-Amino-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine



10

To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (1g, 2.68mmol) in DMSO (6mL) were added 1,1-di-1H-imidazol-1-yl-methyleneamine (864mg, 5.36mmol). The reaction mixture was stirred at RT for 18h. Quenching of the reaction by adding water (60mL). A solid precipitated out. The solid was filtered, washed with water and cold methanol. Yield: white solid 1.0g. LC-MS (Method A) [4.66min; 398(M+1)].

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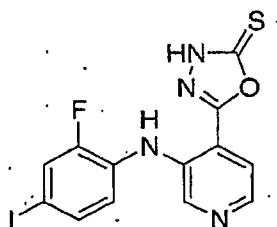
Synthesis of 1,1-di-1H-imidazol-1-yl-methyleneamine

To a solution of 1H-imidazole (3.52g, 51.6mmol), in DCM (250mL) cyanic bromide was added (1.82g, 17.2mmol). The reaction mixture was stirred at 40°C for 30min. Part of the solvent was evaporated, leaving around 30mL, then cooled at 0°C for 1h resulting in a white precipitate which was filtered to afford the expected product 3.3g.

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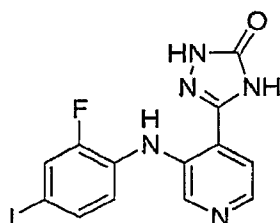
Example 3: 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-3H-[1,3,4]oxadiazole-2-thione

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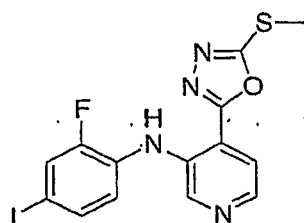
To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (744mg, 2.0mmol) in methanol (15mL) were added dithiomethane (0.22mL, 4.66mmol), followed by potassium hydroxide (2.51mL, 2.13mmol). The reaction mixture was stirred at 60°C for 7h. The solvent was evaporated and the residue was dissolved in ethyl acetate (50mL), washed with Brine (15mLX3), dried over anhydrous MgSO₄, filtered, removed off solvent. Residue was subjected to silica gel column (EtOAc:Hexane=1:1) collected product 210mg. LC-MS (Method A) [5.71min; 415(M+1)]

Example 4 : 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-2,4-dihydro-[1,2,4]triazol-3-one



[4-(5-Ethoxy-4H-[1,2,4]triazol-3-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine (80mg) was added conc. HCl (4mL). The reaction mixture was stirred at 90°C for 4hr, cooled and filtered, the solid was washed with water and methanol. Yield: white solid 30mg. LC-MS (Method A) [5.23min; 398(M+1)]

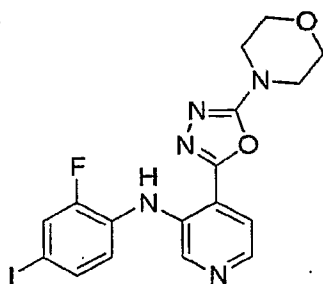
Example 5 : (2-Fluoro-4-iodo-phenyl)-[4-(5-methylsulfanyl-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-amine



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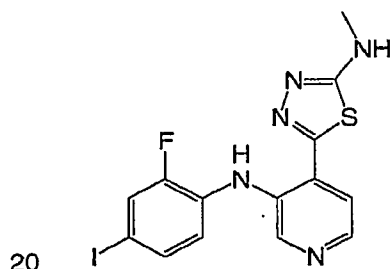
The reaction mixture of 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-3H-[1,3,4]oxadiazole-2-thione (600mg, 1.45mmol), Iodomethane (216mg, 1.52mmol) and 1eq. NaOH (1.45mL) in ethanol (10mL) was stirred for overnight. The solvent was removed, ethyl acetate (50mL) was added to the the residue, which was washed with
5 Brine (20mL x 3). Dried over anhydrous MgSO₄ was added, filtered, and the solution evaporated. Residue was subjected to silica gel column (EtOAc: Hexane=1:1), Yield: 245mg. LC-MS (Method A) [5.72min; 426(M+1)]

10 **Example 6: (2-Fluoro-4-iodo-phenyl)-[4-(5-morpholin-4-yl-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-amine**



To a solution of (2-Fluoro-4-iodo-phenyl)-[4-(5-methylsulfanyl-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-amine (100mg, 0.23mmol) in dioxane (5mL) were added morpholine (101mg, 1.17mmol). The reaction mixture was stirred at 120°C for 8h. The solvent was
15 evaporated, residue was purified with prep plate (CH₂Cl₂:MeOH= 20:1 5% TEA), collected product 24mg. LC-MS (Method A) [5.20min; 468(M+1)]

Example 7: (2-Fluoro-4-iodo-phenyl)-[4-(5-methylamino- [1,3,4] thiadiazol-2-yl)-pyridin-3-yl]-amine

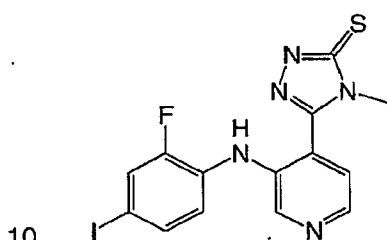


20 To a solution of N'- [3-(2-Fluoro-4-iodo-phenylamino)-pyridine-4-carbonyl] of hydrazinecarbothioic acid N-methylamide (150mg, 0.34 mmol) in DMA (2mL) added

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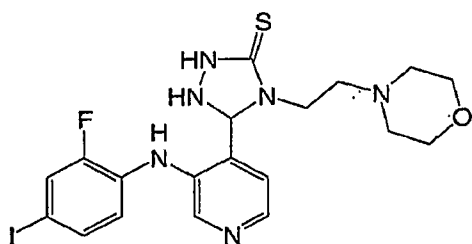
PS-Triphenyl-phosphane (353mg, 1.35 mmol), Trichloro-acetonitrile (97mg, 0.67 mmol), and DIEPA (0.17mL, 1.01mmol). The reaction mixture was stirred at 100°C for 4hrs, and then filtered, and the volatiles evaporated. The crude product was purified by preparative HPLC to afford 8 mg of the desired product. LC/MS (Method A) [5.03min; 428(M+1)].

Example 8: 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-4-methyl-2, 4-dihydro- [1,2,4] triazole-3-thione



N'- [3-(2-Fluoro-4-iodo-phenylamino)-pyridine-4-carbonyl] of hydrazinecarbothioic acid N-methylamide (100mg, 0.22 mmol) was added to 1N of aqueous NaOH solution (1mL) and refluxed for 1h. The pH was adjusted to 7 with 1N HCl. Upon neutralization a white solid precipitated. The desired product (40mg) was isolated from the mixture by filtration LC/MS (Method A) [5.69min; 428(M+1)].

Example 9: 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-4-(2-morpholin-4-yl-ethyl)-[1,2,4] triazolidine-3-thione

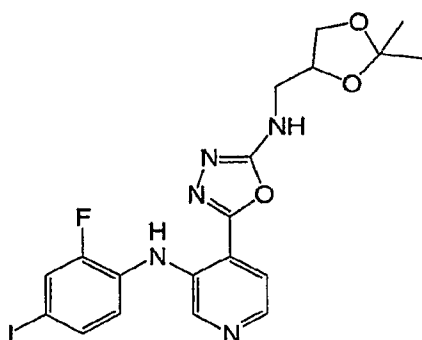


3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid N'- (2-Morpholin-4-yl-ethyl)-thiourea -Hydrazide (100mg, 0.22 mmol) was added to 1N aqueous NaOH solution (1mL), and refluxed for 1h. The pH was adjusted to 7 with 1N HCl. Upon neutralization a white

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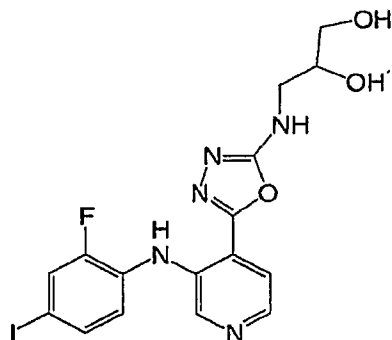
solid precipitated. The desired product (45mg) was isolated from the mixture by filtration. LC/MS (Method A) [5.13min; 527(M+1)]

5 **Example 10: (4-{5-[(2,2-Dimethyl- [1,3] dioxolan-4-ylmethyl)-amino]-[1,3,4] oxadiazol-2-yl}-pyridin-3-yl)-(2-fluoro-4-iodo-phenyl)-amine**



To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic N- (2,2-Dimethyl- [1,3] dioxolan-4-yl)-formamide acidl-hydrazide (189mg, 0.36 mmol) in DMA (2mL) added
 10 PS-Triphenyl-phosphane (375mg, 1.43 mmol,) Trichloro-acetonitrile (104mg, 0.72 mmol) and DIEPA (0.19mL, 1.43mmol). The reaction mixture was stirred at 100°C for 4hrs. After filtration, the solution was directly subjected to preparative HPLC for purification to afford the product (130mg). LC/MS (Method A) [5.25min; 512(M+1)].

15 **Example 11: 3-{5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-propane-1, 2-diol**



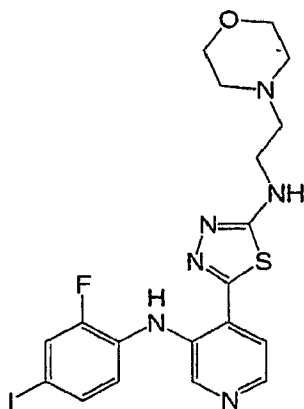
(4-{5-[(2,2-Dimethyl- [1,3] dioxolan-4-ylmethyl)-amino]-[1,3,4] oxadiazol-2-yl}-pyridin-3-yl)-(2-fluoro-4-iodo-phenyl)-amine (100mg, 0.2mmol) was dissolved in dichloromethane
 20 (5mL) and TFA (3mL) was added. The reaction mixture was stirred at RT for 4hrs. The

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solvent was evaporated, and the residue was dissolved in Ethyl acetate, and washed with 1N aqueous NaOH. After solvent evaporation a residue was collected and dissolved in methanol. The product crystallized out the methanolic solution and filtered (63mg). LC/MS (Method A) [6.61min; 472(M+1)].

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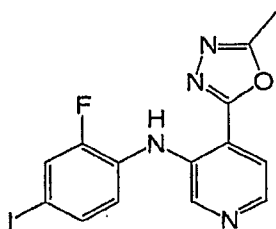
Example 12: (2-Fluoro-4-iodo-phenyl)-[4-[5-(2-morpholin-4-yl-ethylamino)-[1,3,4]thiadiazol-2-yl]-pyridin-3-yl]-amine



To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid N'-(2-Morpholin-4-yl-ethyl)-thiourea-hydrazide (200mg, 0.34 mmol) in DMA (2mL) PS-Triphenyl-phosphane (385mmol, 1.47 mmol,) Trichloro-acetonitrile (106 mg, 0.73 mmol) DIEPA (0.19mL, 0.76mmol) were added. The reaction mixture was stirred at 100°C for 4hrs and then filtered, and the solvent was removed. The crude product was purified by preparative HPLC to afford 51mg of pure product. LC/MS (Method A) [6.06min; 527(M+1)].

15

Example 13: (2-Fluoro-4-iodo-phenyl)-[4-(5-methyl-[1,3,4] oxadiazol-2-yl)-pyridin-3-yl]-amine



A mixture of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid (200mg, 0.54 mmol), acetic acid (32mg, 0.54 mmol), PS-Triphenyl-phosphine (845mg, 3.22 mmol),

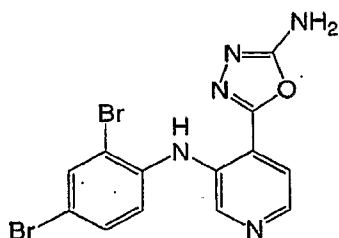
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- 44 -

Trichloro-acetonitrile (155 mg, 1.07 mmol) and DIEPA (0.23mL, 1.34mmol) in acetonitrile was stirred in a sealed tube at 100°C for 6hrs. The mixture was then filtered, and the filtrate was directly subjected to preparative HPLC to afford 55mg of the desired product. LC/MS (Method A) [6.63min; 397(M+1)].

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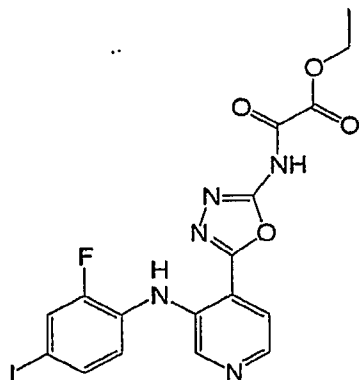
Example 14: [4-(5-Amino- [1,3,4] oxadiazol-2-yl)-pyridin-3-yl]-(2,4-dibromo-phenyl)-amine



10 To a solution of 3-(2,4-Dibromo-phenylamino)-isonicotinic acid hydrazide (200mg, 0.52 mmol) in DMSO (2mL) C- (Di-imidazol-1-yl)-methyleamine (167mg, 1.04 mmol) was added. The reaction mixture was stirred at RT under Argon overnight, then poured into water. A solid precipitated out, which was filtered, and washed with methanol to afford the desired product (100mg). LC/MS (Method A) [5.23min; 412(M+1)].

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Example 15: N- {3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl}-[1,3,4] oxadiazol-2-yl}-oxalamic acid ethyl ester

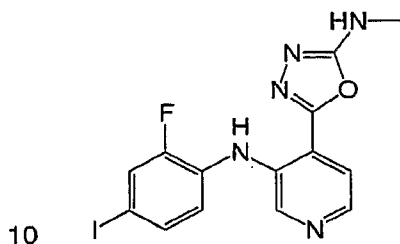


20 To a solution of [4-(5-Amino- [1,3,4] oxadiazol-2-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine (470mg, 1.18mmol, 1.00eq) in DCM (50mL) Ethyl-diisopropyl-amine (0.72mL; 4.14 mmol; 3.5eq), was added and then ethyl chloro (oxo) acetate (0.2mL; 1.78mmol, 1.50eq) was added dropwise at 0°C, under Argon. The reaction mixture

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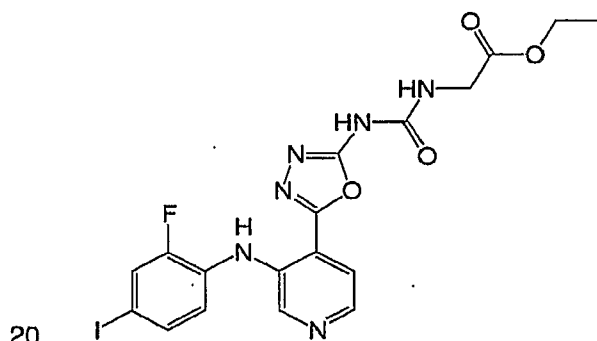
was allowed to warm up to room temperature while stirring overnight. The mixture was then diluted with DMC (50mL), and washed first with 5% aqueous NaHCO₃, and then with brine. The organic layer was dried over anhydrous MgSO₄, filtered, and the solvent was evaporated to afford a residue which was subjected to silica gel column chromatography using DCM:MeOH 50:1 as eluent, to afford 220mg of the desired product. LC/MS (Method A) [5.41min; 498(M+1)]

Example 16: (2-Fluoro-4-iodo-phenyl)-[4-(5-methylamino- [1,3,4] oxadiazol-2-yl)-pyridin-3-yl]-amine



To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid N'-acetyl-hydrazide (400mg; 0.93mmol; 1eq) in DMA (5mL), PS-Triphenyl-phosphine (1.22g; 4.66mmol; 5eq), Trichloro-acetonitrile (269mg; 1.86 mmol; 2eq) and DIEPA (0.49mL; 2.8mmol; 3eq) were added. The reaction mixture was stirred at 100°C for 4hrs. The mixture was filtered and the solution was subjected to preparative HPLC to afford the product (98 mg). LC/MS (Method A) [5.03min; 412(M+1)]

Example 17: (3-[5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-yl]-ureido)-acetic acid ethylester



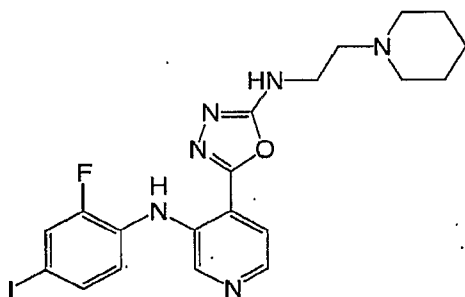
A suspension of [4-(5-Amino- [1,3,4] oxadiazol-2-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine (0.200g, 0.504 mmol) in dry THF (3mL) was treated with ethyl isocyanatoacetate (0.077g, 0.596mmol). The reaction mixture was stirred at RT

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overnight. More isocyanatoacetate (0.083g, 0.643mmol) was added, and the mixture was stirred for another 72hrs. The material was diluted with diethyl ether (2mL), filtered and dried under high vacuum for 2h to afford a yellow solid product (0.144g, 0.274mmol, 54%) LC/MS (Method A) [5.53min; 527(M+1)]

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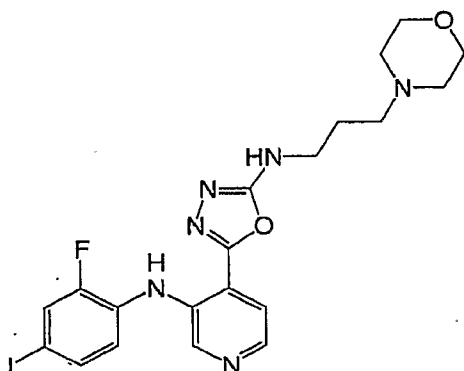
Example 18: (2-Fluoro-4-iodo-phenyl)-{4-[5-(2-piperidin-1-yl-ethylamino)-[1,3,4]oxadiazol-2-yl]-pyridin-3-yl}-amine



To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (300mg; 0.81 mmol; 1eq) in DCM (5mL), 1-(2-Isothiocyanato-ethyl)-piperidine (192mg; 1.13mmol; 1.4eq) was added. The reaction mixture was refluxed for 4hrs under argon. A yellow solid precipitated out and filtered to afford an intermediated that was suspended in ethanol (5mL). Iodomethane (148mg; 1.05mmol; 1.3eq) was added to the reaction mixture and stirred overnight at RT, then heated to 80°C and stirred for another 2hr. The mixture was filtered to give a yellow solid product 40mg. LC/MS (Method A) [3.89min; 509(M+1)]

Example 19: (2-Fluoro-4-iodo-phenyl)-{4-[5-(3-morpholin-4-yl-propylamino)-[1,3,4]oxadiazol-2-yl]-pyridin-3-yl}-amine

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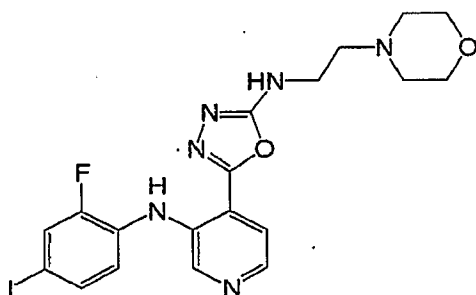


To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (500mg; 1.34 mmol; 1eq) in DCM (5mL) 4-(3-Isothiocyanato-propyl)-morpholine (300mg; 1.61mmol; 1.2eq) was added. The reaction mixture was refluxed for 4hrs under argon.

- 5 A yellow solid precipitated out, which was filtered to afford an intermediate that was suspended in ethanol (5mL). Iodomethane (190mg; 1.34mmol; 1.0eq) was then added and the reaction mixture was stirred overnight at RT. It was then heated to 80°C and stirred for another 2hr. Filtration afforded the yellow solid product (128mg). LC/MS (Method A) [3.73min; 525(M+1)]

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Example 20: (2-Fluoro-4-iodo-phenyl)-{4-[5-(2-morpholin-4-yl-ethylamino)-[1,3,4]oxadiazol-2-yl]-pyridin-3-yl}-amine

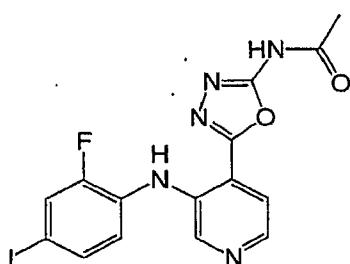


- To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (500mg; 1.34 mmol; 1eq) in DCM (5mL) 4-(3-Isothiocyanato-ethyl)-morpholine (255mg; 1.61mmol; 1.2eq) was added. The reaction mixture was refluxed for 4hrs under argon. A yellow solid precipitated out, which was filtered to afford an intermediate that was suspended in ethanol (5mL). Iodomethane (190mg; 1.34mmol; 1.0eq) was then added and the reaction mixture was stirred overnight at RT. It was then heated to 80°C and

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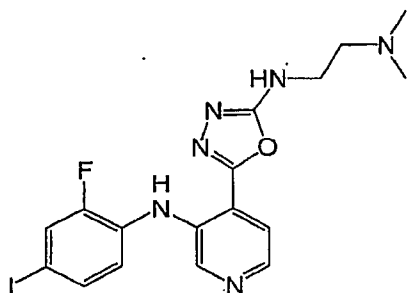
stirred for another 2hr. Filtration afforded the yellow solid product 153mg. LC/MS (Method A) [0.41min; 511(M+1)]

Example 21: N- {5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4]oxadiazol-2-yl}-acetamide



To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (250mg; 0.67 mmol; 1eq) in DCM (5mL) Acetyl isothiocyanate (80mg; 0.8mmol; 1.2eq) was added. The reaction mixture was refluxed for 4hrs under argon. A yellow solid precipitated out, which was filtered to afford an intermediate that was suspended in ethanol (5mL). Iodomethane (95mg; 0.67mmol; 1.0eq) was then added and the reaction mixture was stirred overnight at RT. It was then heated to 80°C and stirred for another 2hr. Filtration afforded the yellow solid product 89mg. LC/MS (Method A) [5.30min; 440(M+1)]

Example 22: N- {5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4]oxadiazol-2-yl}-N', N'-dimethyl-ethane-1, 2-diamine

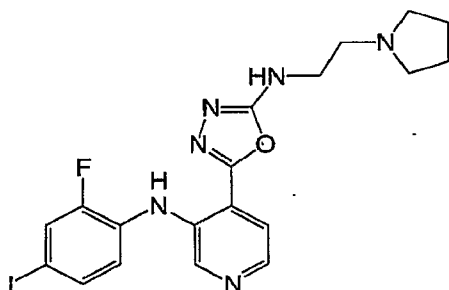


3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid N'- (3-dimethylamino-propionyl)-hydrazide (130mg, 0.27mmol) was dissolved in DMA (2mL), and then PS-Triphenylphosphane (420mg, 1.6 mmol, 6eq), Trichloro-acetonitrile (77mg, 0.53 mmol, 2eq) and

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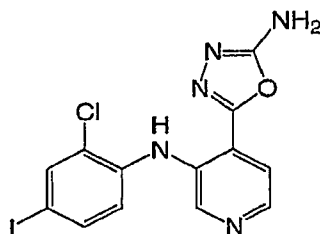
DIEPA (0.19mL, 1.07mmol, 4eq) were added. The reaction mixture was stirred at 100°C overnight, and then filtered. The filtrate was directly subjected to preparative HPLC to afford 32 mg of the desired product. LC/MS (Method A) [3.63min; 467(M+1)]

5 **Example 23: (2-Fluoro-4-iodo-phenyl)-{4-[5-(2-pyrrolidin-1-yl-ethylamino)-[1,3,4]oxadiazol-2-yl]-pyridin-3-yl}-amine**



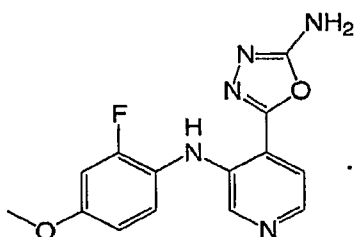
3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid N'-(3-pyrrolidin-1-yl-propionyl)-hydrazide (138mg, 0.27mmol) was dissolved in DMA (2mL), and then PS-Triphenyl-phosphane (424mg, 1.62 mmol, 6eq) Trichloro-acetonitrile (78mg, 0.54 mmol, 2eq) and DIEPA (0.19mL, 1.08mmol, 4eq) were added. The reaction mixture was stirred at 100°C overnight, and then filtered. The filtrate was directly subjected to preparative HPLC to afford 26 mg of the desired product. LC/MS (Method A) [4.20min; 495(M+1)]

15 **Example 24: [4-(5-Amino-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-(2-chloro-4-iodo-phenyl)-amine**



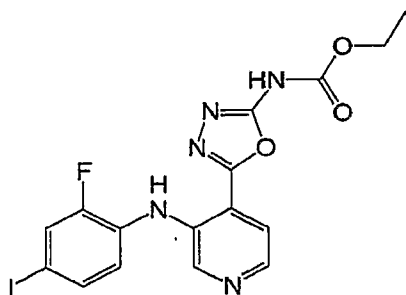
To a solution of 3-(2-Chloro-4-iodo-phenylamino)-isonicotinic acid hydrazide (150mg, 0.39mmol, 1eq), in DMSO (2mL) C-(Di-imidazol-1-yl)-methylethylamine (124mg, 0.77mmol, 2eq) was added. The reaction mixture was stirred at RT under argon overnight and then poured into water. The product was isolated by filtration (135 mg). LC/MS (Method A) [5.22min; 414(M+1)]

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Example 25: [4-(5-Amino- [1,3,4] oxadiazol-2-yl)-pyridin-3-yl]-(2-fluoro-4-methoxy-phenyl)-amine

- 5 To a solution of 3-(2-Fluoro-4-methoxy-phenylamino)-isonicotinic acid hydrazide (150mg, 0.54mmol, 1eq) in DMSO (2mL) C- (Di-imidazol-1-yl)-methylethylamine (175mg, 1.09mmol, 2eq) was added. The reaction mixture was stirred at RT under argon overnight and then poured into water. The product was isolated by filtration (89 mg). LC/MS (Method A) [0.52min; 302(M+1)]

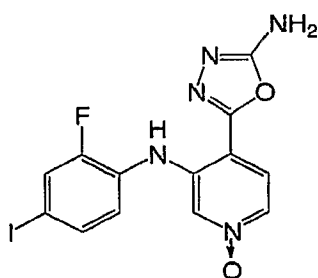
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Example 26: {5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-yl}-carbamic acid ethyl ester

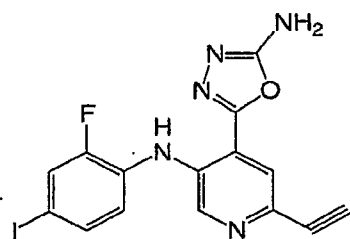
- To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (500mg; 1.34 mmol; 1eq) in DCM (5mL) Isothiocyanatoethylester (2.0g; 5.37mmol; 1.1eq) was added. The reaction mixture was refluxed for 4hrs under argon. A yellow solid precipitated out that was filtered out and suspended in ethanol (20mL). Iodomethane (0.35mL; 5.64mmol; 1.05eq) was added and the reaction mixture was first stirred overnight at RT, and then heated to 80°C while stirring for 2hr. The product was
- 15
- 20 filtered out of the mixture (1.1g).

LC/MS (Method A) [5.33min; 470(M+1)]

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Example 27: [4-(5-Amino- [1,3,4] oxadiazol-2-yl)-1-oxy-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine

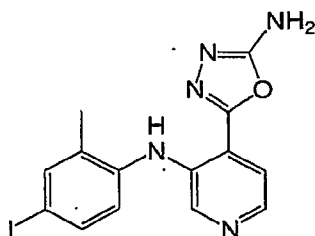
- 5 To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-1-oxy-isonicotinic acid hydrazide (150mg, 0.39mmol, 1eq) in DMSO (2mL) C- (Di-imidazol-1-yl)-methyleamine (124mg, 0.78mmol, 2eq) was added. The reaction mixture was stirred at RT under argon overnight and then poured into water. The product was isolated by filtration (80 mg). LC/MS (Method A) [4.68min; 414(M+1)]

Example 28: [4-(5-Amino- [1,3,4] oxadiazol-2-yl)-6-ethynyl-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine

- To a solution of 2-Ethynyl-5- (2-fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (124mg, 0.31mmol, 1eq) in DMSO (2mL) C- (Di-imidazol-1-yl)-methyleamine (101mg, 0.63mmol, 2eq) was added. The reaction mixture was stirred at RT under argon overnight. and then poured into water. The solid fomed was isolated by filtration and washed with methanol to afford the product (30 mg). LC/MS (Method A) [5.15min; 422(M+1)]

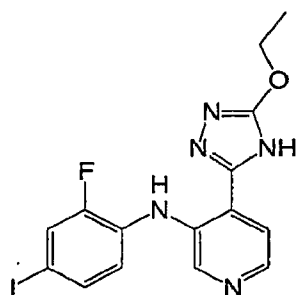
Example 29: [4-(5-Amino- [1,3,4] oxadiazol-2-yl)-pyridin-3-yl]-(4-iodo-2-methyl-phenyl)-amine

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To a solution of 3-(4-iodo-2-methyl-phenylamino)-isonicotinic acid hydrazide (200mg, 0.54mmol, 1eq) in DMSO (2mL) C- (Di-imidazol-1-yl)-methyleamine (175mg, 1.09mmol, 2eq) was added. The reaction mixture was stirred at RT under argon overnight. and then poured into water. The solid fomed was isolated by filtration and washed with methanol to afford the product 1(45 mg). LC/MS (Method A) [4.90min; 394(M+1)]

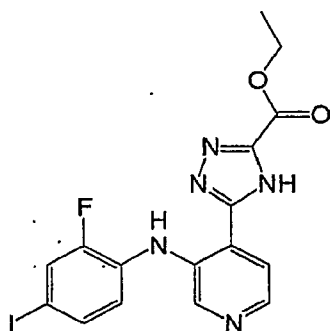
Example 30: [4-(5-Ethoxy-4H- [1,2,4] triazol-3-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine



To a solution of [4-(5-Amino- [1,3,4] oxadiazol-2-yl)-pyridin-3-yl]-(2-floro-4-iodo-phenyl)-amine (150mg, 0.38mmol) in ethanol (10mL) KOH 9212mg, 3.78mmol) was added. The reaction mixture was refluxed for 3hrs. Then, added acetic acid was added to bring the pH to 5. The volatives were stripped and the residue was dissolved in EtOAc, and washed with 5% Na₂CO₃, and brine, and then dried over anhydrous MgSO₄. After filtration and solvent removal the remaining crude product was subjected to silica gel column chromatography (eluent: Hex:ane EtOAc=1:1) to afford the desired product (45mg). [5.08min; 394(M+1)]

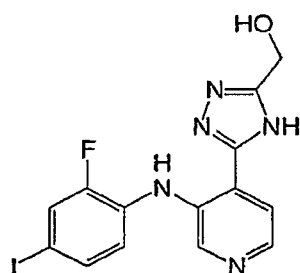
Example 31: 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-4H-[1,2,4] triazole-3-carboxylic acid ethyl ester

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Amino- {[3-(2-fluoro-4-iodo-phenylamino)-pyridine-4-carbonyl]-hydrazono}-acetic acid ethyl ester (1.6g, 3.4mmol) in pyridine (10mL) was stirred at 140°C for 3hrs. Cooled, poured into water, Extracted with EtOAc, Combined the organic layer, and washed with
 5 Brine, dried over MgSO₄, filtered, removed off solvent, got crude product, which was subjected to silica gel column (Hex: EtOAc=1:1). Collected final product 1.02g. [5.23min, 454 (M+1)]

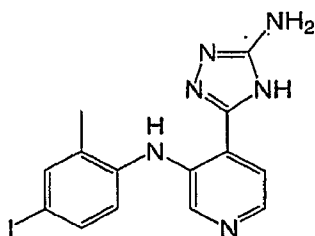
10 **Example 32: {5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-4H-[1,2,4] triazol-3-yl}-methanol**



To a solution of 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-4H-[1,2,4] triazole-3-carboxylic acid ethyl ester (150mg, 0.33mmol, 1eq) in THF (3mL) LiBH₄ (10.8mg, 0.5mmol, 1.5eq) was added. The reaction mixture was stirred overnight at RT. The
 15 solvent was evaporated and the residue was dissolved in EtOAc, washed with brine, and dried over MgSO₄. After filtration and solvent removal, the crude product was purified by preparative HPLC, to afford the desired product (41 mg). [5.57min; 454 (M+1)]

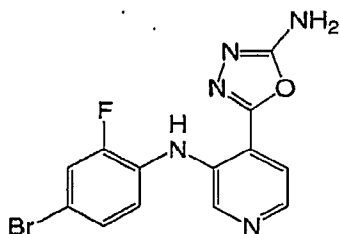
20 **Example 33: [4-(5-Amino-4H-[1,2,4] triazol-3-yl)-pyridin-3-yl]-(4-iodo-2-methyl-phenyl)-amine.**

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A reaction mixture of 3-(4-Iodo-2-methyl-phenylamino)-isonicotinic acid hydrazide (120mg, 0.33mmol, 1eq) and C-Methylsulfanyl-methanediamine hydroiodide (104mg, 0.47mL, 1.45eq) in pyridine (2mL) in sealed tube was heated to 140°C while stirring for 4 hrs. After cooling, it was poured into water, extracted with EtOAc, and the combined organic fractions, dried over anhydrous MgSO₄. After filtration and solvent removal the remaining crude product was subjected to silica gel column chromatography (eluent: Hexane EtOAc=1:1) to afford the desired product (35 mg). [4.26min; 393(M+1)]

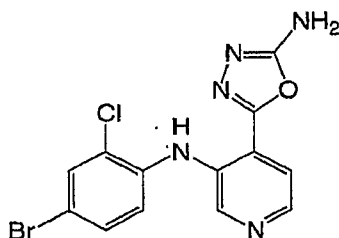
Example 34: [4-(5-Amino- [1,3,4] oxadiazol-2-yl)-pyridin-3-yl]-(4-bromo-2-fluoro-phenyl)-amine



To a solution of 3-(4-Bromo-2-fluoro-phenylamino)-isonicotinic acid hydrazide (200mg, 0.62mmol, 1eq) in DMSO (2mL) C- (Di-imidazol-1-yl)-methylethylamine (198mg, 1.23mmol, 2eq) was added. The reaction mixture was stirred at RT under argon overnight. It was then poured into water and the solid that precipitated out was filtered and washed with methanol to afford the desired product (195 mg). LC/MS (Method A) [0.62min; 351(M+1)]

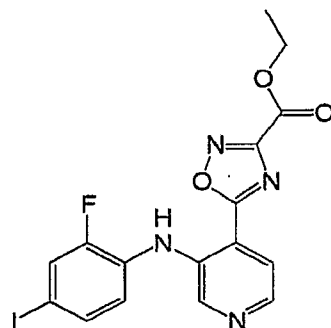
Example 35: [4-(5-Amino- [1,3,4] oxadiazol-2-yl)-pyridin-3-yl]-(4-bromo-2-chloro-phenyl)-amine

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To a solution of 3-(4-bromo-2-chlorophenylamino)-isonicotinic acid hydrazide (200mg, 0.59mmol, 1eq) in DMSO (2mL) C- (Di-imidazol-1-yl)-methylenamine (188mg, 1.17mmol, 2eq) was added. The reaction mixture was stirred at RT under argon overnight and then poured into water. A solid precipitated out that was filtered and washed with methanol to give the desired product (125 mg). LC/MS (Method A) [4.77min; 367(M+1)]

Example 36: 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,2,4] oxadiazole-3-carboxylic acid ethyl ester



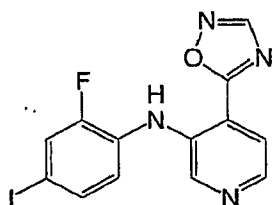
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A reaction mixture of 3-(2-fluoro-4-iodophenylamino)-isonicotinic acid (1000mg, 2.79mmol, 1.00eq), and Di-imidazol-1-yl-methanone ((543mg, 3.35mmol, 1.20eq) in DMF was stirred for 4hrs at RT under argon. Then hydroxyamino-imino-acetic acid ethyl ester (612mg, 4.19mmol, 1.5eq.) was added and the reaction mixture was stirred overnight, and then diluted with water. A yellow precipitate was formed that was filtered to afford the product (785mg). [5.46min; 455 (M+1)]

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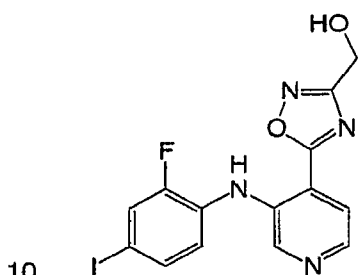
Example 37: (2-Fluoro-4-iodo-phenyl)-(4-[1,2,4] oxadiazol-5-yl-pyridin-3-yl)-amine

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To a solution of 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,2,4] oxadiazole-3-carboxylic acid ethyl ester (1.8g, 3.96 mmol) in THF-H₂O (1:1)(20mL) LiOH (0.19g, 7.93mmL) was added. The reaction mixture was stirred for 2hrs at RT. The pH was
 5 adjusted to 5 with 1N HCl and the mixture was diluted with water. A solid precipitated and was filtered to give the product (1.4g). LC/MS (Method A) [4.39min; 383(M+1)]

Example 38: {5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,2,4] oxadiazol-3-yl}-methanol



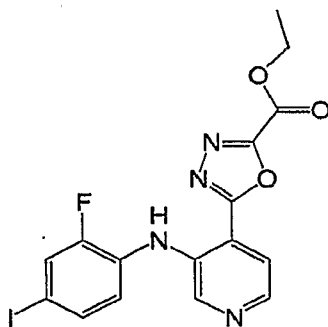
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To a solution of 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,2,4] oxadiazole-3-carboxylic acid ethyl ester (768mg, 1.69mmol, 1eq) in THF (6mL) LiBH₄ (55mg, 2.54mmol, 1.5eq) was added. The reaction mixture was stirred overnight and the solvent was evaporated. The residue was dissolved in EtOAc, washed with brine, and
 15 dried with anhydrous MgSO₄. Filtration and removal of the solvent afforded the crude product which was subjected to silica gel column chromatography (eluent: CH₂Cl₂: MeOH 70:1) to give rise to pure product (497 mg). [5.01min; 413 (M+1)]

Example 39: 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazole-2-carboxylic acid ethyl ester

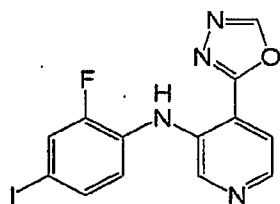
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To a solution of {N'- [3-(2-Fluoro-4-iodo-phenylamino)-pyridine-4-carbonyl]-hydrazino}-oxo-acetic acid ethyl ester (1.27g, 2.69mmol, 1.0eq.) in DCM (20mL) containing pyridine (0.55g, 6.99mmol, 2.66eq.) thionyl dichloride (0.26mL, 3.5mmol; 1.30eq) was added at 0°C. The reaction mixture were stirred at 0°C for 2hrs. The solvent was evaporated, toluene was added and the reaction mixture was stirred at 110°C for 6hrs. Removal of the solvent resulted in a residue that was subjected to silica gel column chromatography (EtOAc: Hex=1:2) to afford the desired product (0.50g). [5.32min; 455 (M+1)]

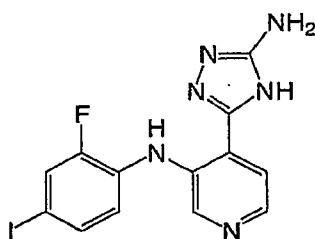
Example 40: (2-Fluoro-4-iodo-phenyl)-(4-[1,3,4] oxadiazol-2-yl-pyridin-3-yl)-amine



To a solution of 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazole-2-carboxylic acid ethyl ester (200mg, 0.44mmol) in THF-H₂O (1:1)(5mL) LiOH (32mg, 1.32mmol) was added. The reaction mixture was stirred for 2hrs and then the pH was adjusted to 5 with 1N HCl. Upon dilution with water a solid precipitated. Filtration afforded the desired product (89mg); LC/MS (Method A) [0.48min; 383(M+1)]

Example 41: [4-(5-Amino-4H- [1,2,4] triazol-3-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine

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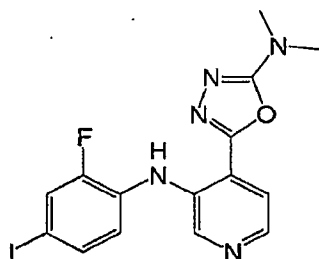


A reaction mixture of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (100mg, 0.27mmol, 1eq) and C-Methylsulfanyl-methanediamine hydroiodide (86mg, 0.39mmol, 1.45eq) in pyridine (2mL) in a sealed tube was heated at 140°C for 4hrs.

- 5 After cooling the mixture was poured into water, extracted with EtOAc, and the combined organic fractions were dried over anhydrous MgSO_4 . After filtration and solvent evaporation the resulting residue was subjected to silica gel column chromatography (EtOAc:Hex=1:1) to give the desired product (46mg). [4.99min; 397(M+1)]

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Example 42: [4-(5-Dimethylamino- [1,3,4] oxadiazol-2-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine

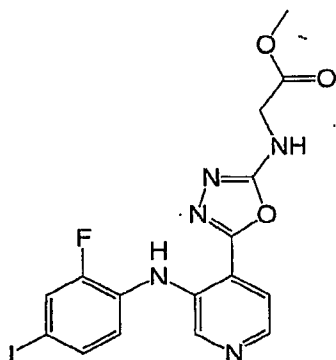


- To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid N'-formamid-
 15 hydrazide (100mg, 0.23mmol, 1eq) in acetonitrile, PS-Triphenyl-phosphane (236mg, 0.9 mmol, 4eq), Trichloro-acetonitrile (0.09mL, 0.9mmol, 4eq) and DIEPA (0.16mL, 0.9mL, 4eq) were added. The reaction mixture was stirred at 100C for 4hrs. After filtration the resulting solution was directly subjected to preparative HPLC to afford the desired product (45 mg). LC/MS (Method A) [4.70min; 426(M+1)]

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Example 43: {5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-acetic acid methyl ester

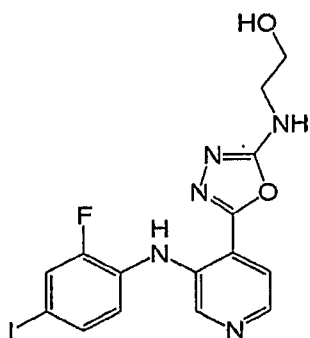
- 59 -



To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (1.50g 4.03mmol; 1eq) in DCM (20mL) Isothiocyanatoethylester (0.58g; 4.43mmol; 1.1eq) was added. The reaction mixture was refluxed for 4hrs under argon. A yellow solid precipitated out and filtered to afford 2.05g of an intermediate. 1.0g (1.99mmol, 1eq) of this intermediate was suspended in ethanol (20mL) and iodomethane (0.13mL; 2.09mmol; 1.05eq) was added. The reaction mixture was stirred overnight at RT, and filtered, to isolate a solid. 100mg (0.19mmol) of that solid was suspended in ethanol (3mL) and heated at 80°C for 2hr. Filtration resulted in the desired product (20mg).

LC/MS (Method A) [5.09min; 470(M+1)]

Example 44: 2-({5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-ethanol)

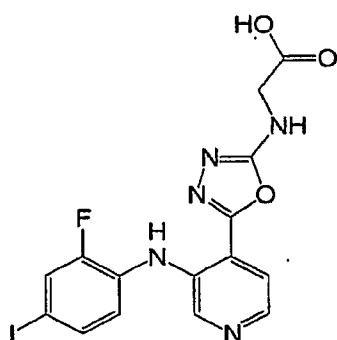


To a solution of {5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-acetic acid methyl ester (400mg, 0.85mmol, 1eq) in THF (5mL) LiBH₄ (37mg, 1.7mmol, 2eq) was added. The reaction mixture was stirred at RT for 20hrs and then quenched with methanol and then added water. The solvent was partially removed

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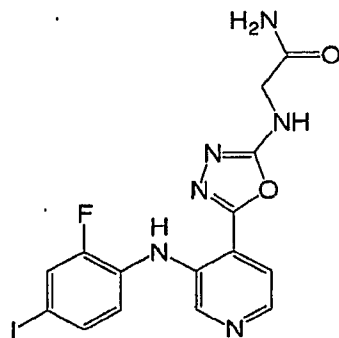
and EtOAc was added. A solid precipitated out that was filtered to give the desired product (190mg). LC/MS (Method A) [4.48min; 442(M+1)]

Example 45: {5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-acetic acid



To a solution of {5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4]oxadiazol-2-ylamino}-acetic acid methyl ester (262mg, 0.56mmol, 1eq) in THF-H₂O (1:1) (2mL) LiOH (27mg, 1.12mmol, 2eq) was added. The reaction mixture was stirred at RT for 2hrs. The solvent was removed patially and more water was added. The mixture was washed with ether and the pH of the aqueous layer was adjusted to 3 with 1N HCl. A solid precipitated and filtered to afford the desired product (190mg). LC/MS (Method A) [4.83min; 456(M+1)]

Example 46: 2-{5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-acetamide



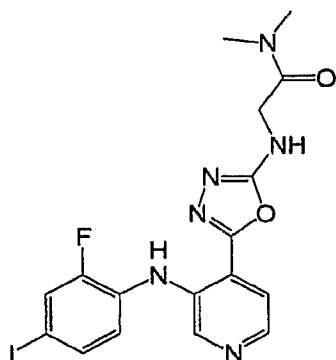
To a solution of {5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-acetic acid (75mg, 0.16mmol, 1eq) in DMF CDI (35mg, 0.21mmol, 1.3eq) was

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added. The reaction mixture was stirred at RT under argon for 3hrs and then ammoniumacetate (30mg, 0.41mmol, 2.5eq) was added. The mixture was stirred for another 2hrs. The resulting mixture was directly purified by preparative HPLC to afford the desired product (35mg). LC/MS (Method A) [4.57min; 455(M+1)]

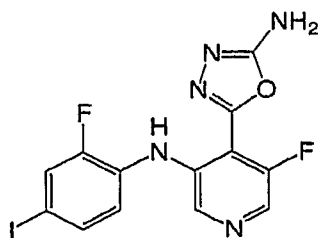
5

Example 47: 2-{5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-N, N-dimethyl-acetamide



To a solution of {5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-acetic acid (75mg, 0.16mmol, 1eq) in DMF CDI (35mg, 0.21mmol, 1.3eq) was added. The reaction mixture was stirred at RT under argon for 3hrs. Then diisopropanlethylamine (352mg, 0.41mmol, 2.5eq), and dimethylamine hydrichloride (40mg, 0.41mmol, 2.5eq) was added and the mixture was stirred for another 2hrs. It was then directly purified by reverse phase preparative HPLC to give the desired product (35mg). LC/MS (Method A) [4.87min; 483 (M+1)]

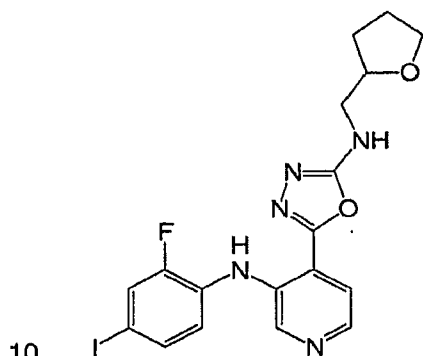
Example 48: [4-(5-Amino- [1,3,4] oxadiazol-2-yl)-5-fluoro-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine



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To a solution of 3-Fluoro-5- (2-fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (100mg, 0.26mmol, 1eq) in DMSO (2mL) C- (Di-imidazol-1-yl)-methylethylamine (123mg, 0.77mmol, 3eq) was added. The reaction mixture was stirred at RT under argon overnight and then poured into water. A solid precipitated out that was filtered out and washed with methanol to afford the desired product (65 mg). LC/MS (Method A) [5.19min; 416(M+1)]

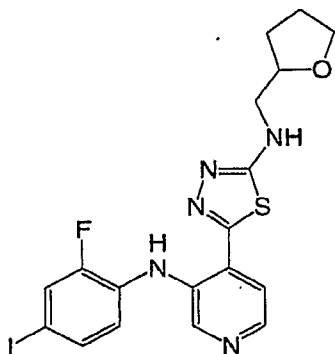
Example 49: (2-Fluoro-4-iodo-phenyl)-(4-{5-[(tetrahydro-furan-2-ylmethyl)-amino]-[1,3,4] oxadiazol-2-yl}-pyridin-3-yl)-amine



A reaction mixture of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid N'- (Tetrahydro-furan-2-ylmethyl)-thiourea-hydrazide (200mg, 0.39mmol, 1eq), methyl iodide (0.03mL, 0.5mmol, 1.3eq) and DIEPA (0.2mL, 0.58mmol, 1.5eq) were stirred at RT overnight. It was then heated at 85°C for 2hrs. As the solvent was partially removed a solid precipitated out. It was filtered to produce the desired product (105mg). LC/MS (Method A) [5.27min; 482(M+1)]

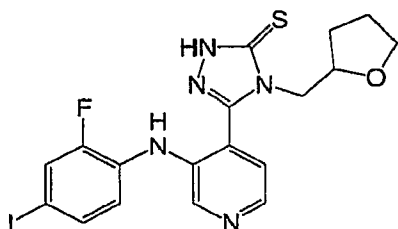
Example 50: (2-Fluoro-4-iodo-phenyl)-(4-{5-[(tetrahydro-furan-2-ylmethyl)-amino]-[1,3,4] thiadiazol-2-yl}-pyridin-3-yl)-amine

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To a solution of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid, N'- (Tetrahydro-furan-2-ylmethyl)-thiourea-hydrazide (260mg, 0.50 mmol) in DMA (2mL), PS-Triphenyl-phosphane (794mg, 3.03 mmol,) Trichloro-acetonitrile (146mg, 1.01 mmol) and DIEPA (0.26mL, 1.51mmol) were added. The reaction mixture was stirred at 80°C for 4hrs and then filtered, and the solvent was removed. The residue was washed with methanol to afford the product (97mg). LC/MS (Method A) [5.31min; 498(M+1)]

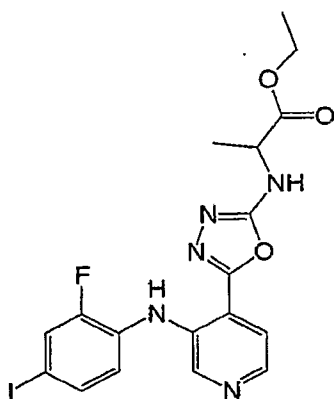
Example 51: 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-4-(tetrahydro-furan-3-ylmethyl)-2,4-dihydro- [1,2,4] triazole-3-thione



3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid N'- (Tetrahydro-furan-2-ylmethyl)-thiourea-hydrazide (100mg, 0.19mmol) was treated with 1N NaOH (1mL). The reaction mixture was heated to 110°C in sealed tube for 2hrs. The mixture was then neutralized with 1N HCl, filtered, and the resulting solid was washed with methanol to give the desired product (17mg). LC/MS (Method A) [5.01min; 498(M+1)]

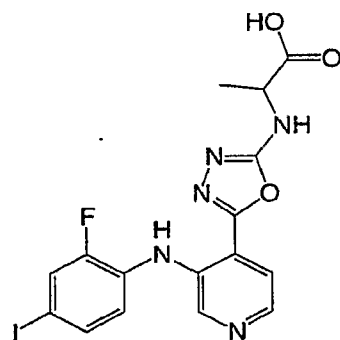
Example 52: 2-[5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-propionic acid ethyl ester

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A mixture of 3-(2-Fluoro-4-iodo-phenylamino)-isonicotinic acid N'-thiocarbamic acid ethyl ester (600mg, 1.13mmol, 1eq), methyl iodide (0.09mL, 1.47mmol, 1.3eq), and DIEPA (0.3mL, 1.69mmol, 1.5eq) were stirred at RT overnight. Then it was heated at 85°C for 2hrs. The solvent was partially removed resulting in the precipitation of a solid that was filtered to afford the product (75mg). LC/MS (Method A) [5.22min; 498(M+1)]

Example 53: 2-{5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-propionic acid



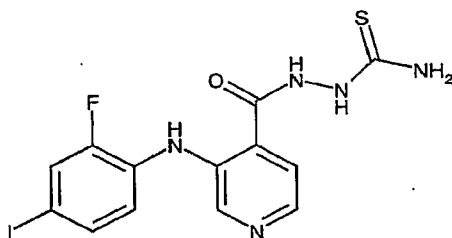
10

A mixture of 2-{5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-[1,3,4] oxadiazol-2-ylamino}-propionic acid ethyl ester (45 mg, 0.09mmol, 1.0eq) and LiOH (5mg, 0.18mmol, 2eq) in THF-H₂O (1:1) (0.5mL) was stirred at RT for 2hr. The pH was adjusted to pH 5 with 1N HCl and the mixture was diluted with EtOAc, washed with brine, and dried over anhydrous MgSO₄. Filtration and solvent removal resulted in a residue that was washed with ether to give rise to the desired product (41mg). LC/MS (Method A) [5.04min; 470(M+1)]

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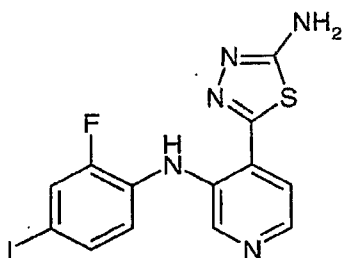
- 65 -

Intermediate 4: 2-{3-(2-fluoro-4-iodophenylamino)isonicotinoyl}hydrazinecarbothioamide



- 5 To a solution of 3-[(2-fluoro-4-iodophenyl)amino]isonicotinic acid (360 mg; 1.01 mmol) was added 1,1'-carbonylbis(1H-imidazole) (196 mg; 1.21mmol). The mixture was stirred overnight. The dark yellow colored solution was treated with thiosemicarbazide (110 mg; 1.21 mmol). The reaction mixture was stirred at room temperature overnight. Water (10 mL) was added, and extracted with ethyl acetate. Concentrated
10 and purified on Flashmaster to obtain 2-{3-[(2-fluoro-4-iodophenyl)amino]isonicotinoyl}hydrazine carbothioamide. (300 mg, 69%) MS m/z: 432 M+1

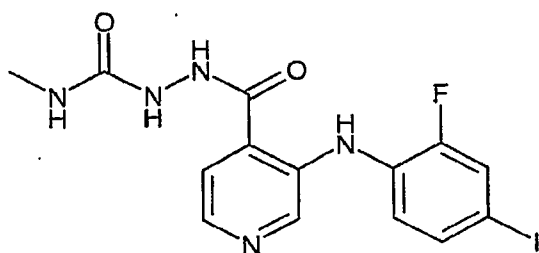
Example 54: [4-(5-Amino-[1,3,4]thiadiazol-2-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine



- 15 To a solution of 2-{3-[(2-fluoro-4-iodophenyl)amino]isonicotinoyl} hydrazine carbothioamide (220 mg; 0.51 mmol) in DMF (5.00 mL) were added resin bound triphenylphosphine (401 mg; 1.53 mmol), carbon tetrabromide tetrabromomethane (508 mg; 1.53 mmol), and triethyl amine (180 mg; 1.53 mmol). The mixture was
20 heated at 50 °C for 5h. Filtered and purified on Agilent Prep. HPLC system to get 10 mg of [4-(5-Amino-[1,3,4]thiadiazol-2-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine. MS m/z: 414 M+1

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Intermediate 5: 2-(3-(2-fluoro-4-iodophenylamino)isonicotinoyl)-N-methylhydrazinecarboxamide

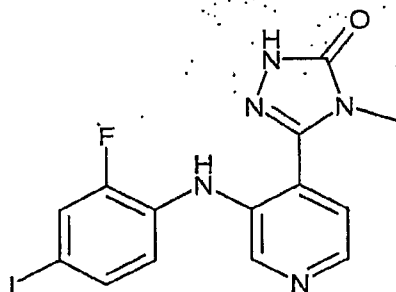


5

To a suspension of 3-[(2-fluoro-4-iodophenyl)amino]isonicotinohydrazide (200 mg; 0.54 mmol) in methanol (10 mL) was added (methylimino)(oxo)methane (61 mg; 1.07 mmol) (methyl isocyanate). After few minutes the reaction mixture became homogeneous, and then a yellow colored solid, 2-{3-[(2-fluoro-4-iodophenyl)amino]isonicotinoyl}-N-methylhydrazinecarboxamide (50 mg) was separated out. The filtrate upon concentration gave 158 mg which was also pure. (208 mg, 90%) MS m/z: 429 M+1

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Example 55: 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-4-methyl-2,4-dihydro-[1,2,4]triazol-3-one



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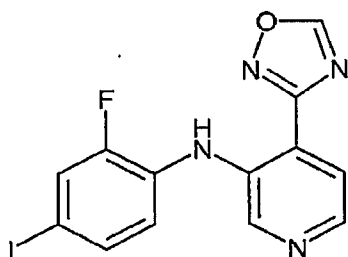
To a suspension of 2-{3-[(2-fluoro-4-iodophenyl)amino]isonicotinoyl}-N-methylhydrazinecarboxamide (165 mg; 0.38 mmol) in water (3 mL) was added 1N NaOH (3 mL). The mixture became homogeneous. The reaction mixture was heated at 120 °C for 3 days. The progress of the reaction was monitored by LCMS.

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Concentration of the solvent and repeated purifications gave 10 mg of 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-4-methyl-2,4-dihydro-[1,2,4]triazol-3-one

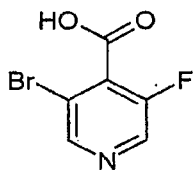
Mass: MS m/z: 412 M+1

5 **Example 56: (2-Fluoro-4-iodo-phenyl)-(4-[1,2,4]oxadiazol-3-yl-pyridin-3-yl)-amine**



To a solution of ethyl 3-{3-[(2-fluoro-4-iodophenyl)amino]pyridin-4-yl}-1,2,4-oxadiazole-5-carboxylate (1.80 g; 3.96 mmol; 1.00 eq.) in THF-H₂O (1:1) was added lithium hydroxide (0.19 g; 7.93 mmol). Stirred for 2h at room temperature. After work up, 1.42g
 10 of (2-Fluoro-4-iodo-phenyl)-(4-[1,2,4] oxadiazol-3-yl-pyridin-3-yl)-amine was obtained in 92% yield. MS m/z: 383 M+1

Intermediate 6: 3-Bromo-5-fluoro-isonicotinic acid

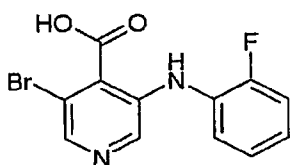


15 n-Butyllithium (12.5 mL, 2.5 M in THF, 31.3 mmol) was added slowly to a solution of diisopropylamine (4.4 mL, 31.3 mmol) in THF (200 mL) at 0 °C, stirred for 15 minutes, and then cooled to -78 °C. 3-Bromo-5-fluoro pyridine (5.0 g, 28.4 mmol) was added as a solid, and the resulting solution was stirred for 10 minutes at -78 °C. Gaseous carbon dioxide was bubbled into the solution via a cannula for 30 minutes, and the
 20 resulting solution was stirred for 2 h warming from -78 °C to room temperature. The reaction solution was concentrated via rotary evaporation, diluted with 2M NaOH, and washed with EtOAc. The aqueous solution was acidified with 1 M HCl, and extracted

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with EtOAc. The organic extracts were washed with brine, dried over MgSO_4 , filtered, and concentrated to afford the desired product (4.75 g, 76 %) as a solid.

Intermediate 7: 3-Bromo-5-(2-fluoro-phenylamino)-isonicotinic acid



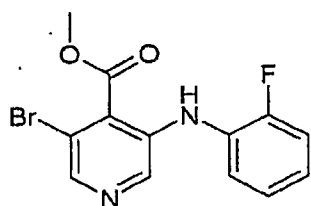
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Lithium bis(trimethylsilyl)amide (54.6 mL, 1.0 M in THF, 54.6 mmol) was added to a solution of 2-fluoro aniline (3.5 mL, 36.4 mmol) in THF (100 mL) at -78°C . The resulting solution was for 1 h at -78°C . 3-Bromo-5-fluoro-isonicotinic acid (**1**) (4.0 g, 18.2 mmol) was added as a solid, and the reaction solution was stirred for 48 h at room temperature. The reaction solution was concentrated via rotary evaporation; diluted with satd. NaHCO_3 , and washed with EtOAc. The aqueous solution was acidified with concentrated HCl. The resulting precipitate was filtered, washed with H_2O , and dried under vacuum to afford the desired product (4.2 g, 74%) as a yellow solid.

10

Intermediate 8: 3-Bromo-5-(2-fluoro-phenylamino)-isonicotinic acid

3-Bromo-5-(2-fluoro-phenylamino)-isonicotinic acid methyl ester

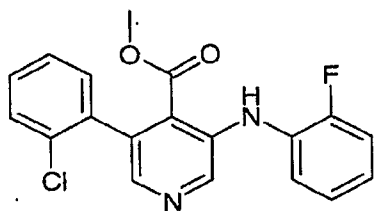


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(Trimethylsilyl)diazomethane (8.1 mL, 2M in THF, 16.2 mmol) was added to a solution of 3-bromo-5-(2-fluoro-phenylamino)-isonicotinic acid (4.2 g, 13.5 mmol) in THF / MeOH (72 mL, 9 / 1, v / v) and stirred for 2 h at room temperature. The reaction solution was quenched with acetic acid, and then concentrated. The crude product was purified via Biotage eluting with a gradient of 0 to 50 % EtOAc in hexanes to afford the desired product (4.1 g, 94%) as a yellow solid.

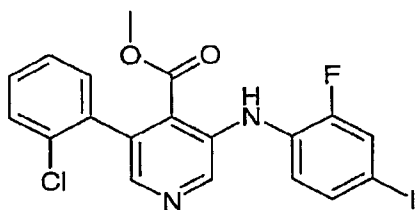
- 69 -

Intermediate 9: 3-(2-Chloro-phenyl)-5-(2-fluoro-phenylamino)-isonicotinic acid methyl ester



3-Bromo-5-(2-fluoro-phenylamino)-isonicotinic acid methyl ester (250 mg, 0.77 mmol),
5 2-chlorophenylboronic acid (0.18 g, 1.2 mmol), Pd(OAc)₂ (10 mg, 0.025 mmol), S-Phos
(20 mg, 0.05 mmol), and K₂CO₃ (0.32 g, 2.3 mmol) were suspended in dioxane / H₂O
(1.65 mL, 9 / 1, v / v) and stirred overnight at 100 °C. The reaction solution was diluted
with EtOAc, and filtered through an Extrelut column. The column was washed with
EtOAc, and the filtrate was concentrated. The crude product was purified via Biotage
10 eluting with a gradient of 0 to 50 % EtOAc in hexanes to afford the desired product
(220 mg, 80 %) as a yellow oil. LC-MS (M+H = 357, obsd. = 357).

Intermediate 10: 3-(2-Chloro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid methyl ester

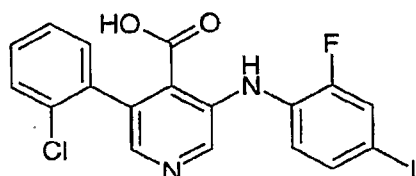


15

N-iodosuccinimide (236 mg, 1.05 mmol) and *p*-toluenesulfonic acid monohydrate (400
mg, 2.1 mmol) were added to a solution of 3-(2-chloro-phenyl)-5-(2-fluoro-
phenylamino)-isonicotinic acid methyl ester (300 mg, 0.84 mmol), and stirred overnight
at 60 °C. The reaction mixture was quenched with Na₂S₂O₃ (1 mL, 10 % aqueous
20 solution), diluted with CH₂Cl₂, and filtered through an Extrelut column. The column was
washed with CH₂Cl₂, and the filtrate was concentrated. The crude product was purified
via Biotage eluting with a gradient of 0 to 55 % EtOAc in hexanes to afford the desired
product (343 mg, 85 %). LC-MS (M+H = 483, obsd. = 483).

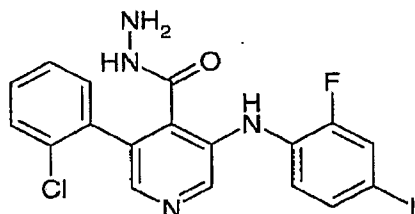
- 70 -

Intermediate 11: 3-(2-Chloro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid



NaOH (3.6 mL, 2M solution in H₂O, 7.2 mmol) was added to a solution of 3-(2-chloro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid methyl ester (343 mg, 0.71 mmol) in MeOH (3 mL) and stirred overnight at 60 °C. The MeOH was removed and the aqueous solution was acidified with concentrated HCl. The resulting precipitate was filtered, washed with H₂O, and dried under vacuum to afford the desired product (253 mg, 76 %). LC-MS (M+H = 469, obsd. = 469).

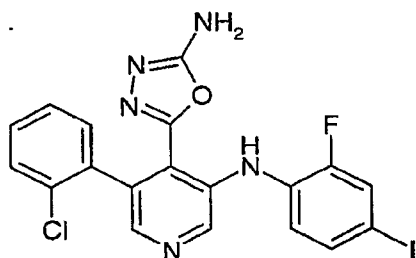
Intermediate 12: 3-(2-Chloro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide



1,1'-carbonyldiimidazole (21 mg, 0.13 mmol) was added to a solution of 3-(2-chloro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid from above (40 mg, 0.09 mmol) in DMSO (1 mL), and stirred overnight at 60 °C. Hydrazine hydrate (13 mg, 0.27 mmol) was added, and the reaction solution was stirred for an additional 4 h at 60 °C. The reaction solution was quenched with H₂O, diluted with EtOAc, and filtered through an Extrelut column. The column was washed with EtOAc, and the filtrate was concentrated. The crude product was purified via Biotage eluting with a gradient of 0 to 70 % EtOAc in hexanes to afford the desired product (17 mg, 41 %). LC-MS (M+H = 483, obsd. = 438).

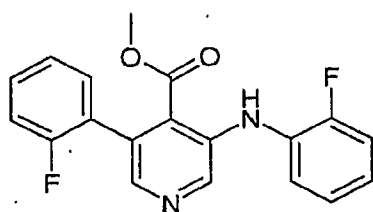
- 71 -

Example 57: [4-(5-Amino-[1,3,4]oxadiazol-2-yl)-5-(2-chloro-phenyl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine



Phenyl cyanate (5 mg, 0.04 mmol) was added to a solution of 3-(2-chloro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (**7**) (17 mg, 0.04 mmol) in DMSO (1 mL), and stirred overnight at room temperature. The reaction solution was quenched with H₂O. The resulting precipitate was filtered, washed with H₂O, and dried under vacuum to afford the desired product (5 mg, 28 %). LC-MS (M+H = 508, obsd. = 508). ¹H NMR : δ 8.92 (s, 1H), 8.47 (d, 1H), 8.03 (d, 1H), 7.74 (dd, 1H), 7.55 (m, 2H), 7.43 (m, 3H), 7.24 (t, 1H), 7.11 (s, 2H).

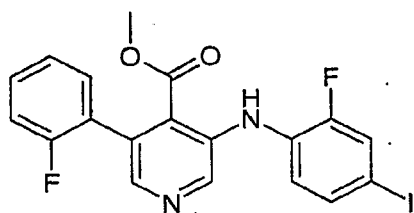
Intermediate 13: 3-(2-Fluoro-phenyl)-5-(2-fluoro-phenylamino)-isonicotinic acid methyl ester



3-Bromo-5-(2-fluoro-phenylamino)-isonicotinic acid methyl ester (1.25 g, 3.84 mmol), 2-fluorophenylboronic acid (0.81 g, 5.77 mmol), Pd(OAc)₂ (30 mg, 0.12 mmol), S-Phos (90 mg, 0.24 mmol), and K₂CO₃ (1.6 g, 11.5 mmol) were suspended in dioxane / H₂O (9 mL, 9 / 1, v / v) and stirred overnight at 100 °C. The reaction solution was diluted with EtOAc, and filtered through an Extrelut column. The column was washed with EtOAc, and the filtrate was concentrated. The crude product was purified via Biotage eluting with a gradient of 0 to 50 % EtOAc in hexanes to afford the desired product (680 mg, 52 %) as a yellow solid. LC-MS (M+H = 341, obsd. = 341).

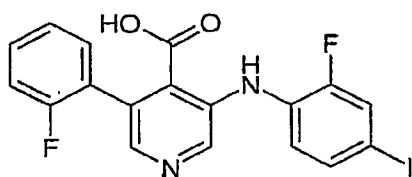
- 72 -

Intermediate 14: 3-(2-Fluoro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid methyl ester



5 N-iodosuccinimide (400 mg, 1.76 mmol) and *p*-toluenesulfonic acid monohydrate (670 mg, 3.53 mmol) were added to a solution of 3-(2-fluoro-phenyl)-5-(2-fluoro-phenylamino)-isonicotinic acid methyl ester (480 mg, 1.41 mmol), and stirred overnight at 60 °C. The reaction mixture was quenched with Na₂S₂O₃ (1 mL, 10 % aqueous solution), diluted with CH₂Cl₂, and filtered through an Extrelut column. The column was washed with CH₂Cl₂, and the filtrate was concentrated. The crude product was purified
10 via Biotage eluting with a gradient of 0 to 55 % EtOAc in hexanes to afford the desired product (516 mg, 78 %). LC-MS (M+H = 467, obsd. = 467).

Intermediate 15 : 3-(2-Fluoro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid

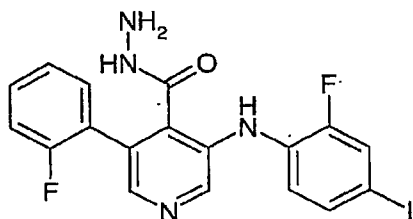


15

NaOH (3.2 mL, 2M solution in H₂O, 6.4 mmol) was added to a solution of 3-(2-fluoro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid methyl ester (300 mg, 0.64 mmol) in MeOH (3 mL) and stirred overnight at 60 °C. The MeOH was removed and the aqueous solution was acidified with concentrated HCl. The resulting precipitate
20 was filtered, washed with H₂O, and dried under vacuum to afford the desired product (215 mg, 74 %). LC-MS (M+H = 453, obsd. = 453).

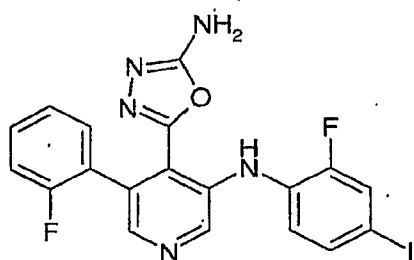
Intermediate 16: 3-(2-Fluoro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide

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1,1'-carbonyldiimidazole (21 mg, 0.13 mmol) was added to a solution of 3-(2-fluoro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid (40 mg, 0.09 mmol) in DMSO (1 mL), and stirred overnight at 60 °C. Hydrazine hydrate (13 mg, 0.27 mmol) was added, and the reaction solution was stirred for an additional 4 h at 60 °C. The reaction solution was quenched with H₂O, diluted with EtOAc, and filtered through an Extrelut column. The column was washed with EtOAc, and the filtrate was concentrated. The crude product was purified via Biotage eluting with a gradient of 0 to 70 % EtOAc in hexanes to afford the desired product (22 mg, 53 %). LC-MS (M+H = 467, obsd. = 467).

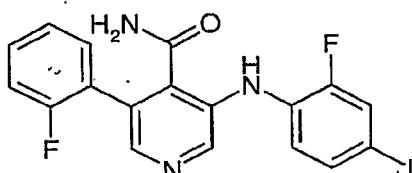
Example 58: [4-(5-Amino-[1,3,4]oxadiazol-2-yl)-5-(2-fluoro-phenyl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine



Phenyl cyanate (6 mg, 0.05 mmol) was added to a solution of 3-(2-fluoro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid hydrazide (22 mg, 0.05 mmol) in DMSO (1 mL), and stirred overnight at room temperature. The reaction solution was quenched with H₂O. The resulting precipitate was filtered, washed with H₂O, and dried under vacuum to afford the desired product (15 mg, 65 %). LC-MS (M+H = 492, obsd. = 492). ¹H NMR : δ 8.78 (s, 1H), 8.43 (d, 1H), 8.11 (d, 1H), 7.73 (dd, 1H), 7.44 (m, 3H), 7.30 (m, 3H), 7.14 (s, 2H).

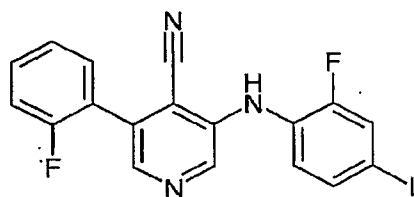
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Intermediate 17: 3-(2-Fluoro-4-iodo-phenylamino)-5-(2-fluoro-phenyl)-isonicotinamide



1,1'-carbonyldiimidazole (70 mg, 0.43 mmol) was added to a solution of 3-(2-fluoro-phenyl)-5-(2-fluoro-4-iodo-phenylamino)-isonicotinic acid (130 mg, 0.29 mmol) in DMSO (1 mL), and stirred overnight at 60 °C. Ammonium acetate (89 mg, 1.15 mmol) was added, and the reaction solution was stirred for an additional 4 h at 60 °C. The reaction solution was quenched with H₂O, diluted with EtOAc, and filtered through an Extrelut column. The column was washed with EtOAc, and the filtrate was concentrated. The crude product was purified via Biotage eluting with a gradient of 0 to 70 % EtOAc in hexanes to afford the desired product (78 mg, 60 %). LC-MS (M+H = 452, obsd. = 452).

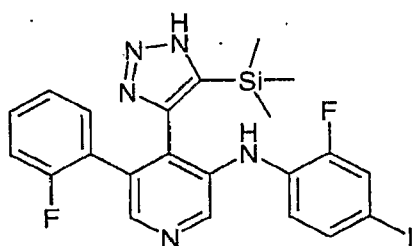
Intermediate 18 : 3-(2-Fluoro-4-iodo-phenylamino)-5-(2-fluoro-phenyl)-isonicotinonitrile



Pyridine (28 µL, 0.35 mmol) and trifluoroacetic anhydride (27 µL, 0.19 mmol) were added to a solution of 3-(2-fluoro-4-iodo-phenylamino)-5-(2-fluoro-phenyl)-isonicotinamide (78 mg, 0.17 mmol) in dioxane (2 mL) at 0 °C. The resulting solution was stirred for 3 h at room temperature. The reaction was quenched with 2M NaOH and stirred for 10 minutes. The reaction solution was diluted with EtOAc and satd. NH₄Cl, and filtered through an Extrelut column. The column was washed with EtOAc, and the filtrate was concentrated to provide a precipitate. The precipitate was washed with MeOH and dried under vacuum to afford the desired product (60 mg, 80 %). LC-MS (M+H = 434, obsd. = 434).

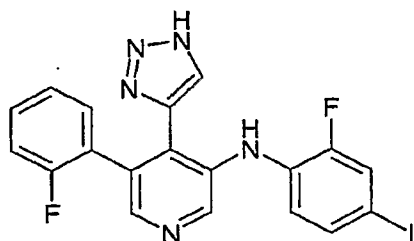
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Example 59: (2-Fluoro-4-iodo-phenyl)-[5-(2-fluoro-phenyl)-4-(5-trimethylsilyl-1H-[1,2,3]triazol-4-yl)-pyridin-3-yl]-amine



- 5 n-Butyllithium (28 μ L, 2.5 M in THF, 0.07 mmol) was added to a solution of (trimethylsilyl)diazomethane (35 μ L, 2.0 M in THF, 0.07 mmol) in THF (1 mL) at 0 °C, and stirred for 20 minutes. 3-(2-Fluoro-4-iodo-phenylamino)-5-(2-fluoro-phenyl)-isonicotinonitrile (20 mg, 0.05 mmol) was added, and the resulting solution was stirred overnight at room temperature. The reaction solution was diluted with EtOAc and satd.
- 10 NH_4Cl , and filtered through an Extrelut column. The column was washed with EtOAc, and the filtrate was concentrated. The crude product was purified via Biotage eluting with a gradient of 0 to 75 % EtOAc in hexanes to afford the desired product (10 mg, 40 %). LC-MS ($M+H$ = 548, obsd. = 548).

15 **Example 60: (2-Fluoro-4-iodo-phenyl)-[5-(2-fluoro-phenyl)-4-(1H-[1,2,3]triazol-4-yl)-pyridin-3-yl]-amine**

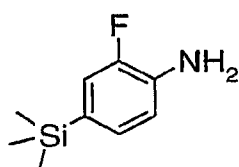


- NaOH (50 μ L, 2.0 M in H_2O , 0.09 mmol) was added to a solution of (2-fluoro-4-iodo-phenyl)-[5-(2-fluoro-phenyl)-4-(5-trimethylsilyl-1H-[1,2,3]triazol-4-yl)-pyridin-3-yl]-amine (10 mg, 0.02 mmol) in MeOH (1 mL), and stirred overnight at 50 °C. The reaction was quenched with satd. NH_4Cl , diluted with EtOAc, and filtered through an Extrelut column. The column was washed with EtOAc, and the filtrate was
- 20

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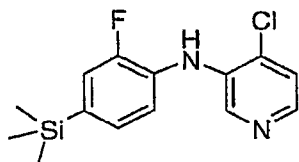
concentrated. The crude product was purified via Biotage eluting with a gradient of 0 to 75 % EtOAc in hexanes to afford the desired product (6 mg, 69 %) as a yellow solid. LC-MS ($M+H = 476$, obsd. = 476). 1H NMR : δ 8.51 (s, 1H), 8.19 (s, 1H), 7.62 (dd, 1H), 7.44 (m, 1H), 7.38 (m, 2H), 7.28 (m, 2H), 7.19 (t, 1H), 6.99 (s, 1H).

5

Intermediate 19: 2-fluoro-4-(trimethylsilyl)aniline

To a stirring solution of 2-fluoro-4-iodoaniline (5 g, 21.1 mmol) in dry ether (20 mL) at $-78^\circ C$, Butyl lithium (2.5 M in Hexane, 25 mL) was added dropwise under nitrogen. The mixture was stirred at $-78^\circ C$ for one hour. Liquid TMSCl (13.8 mL, 105.5 mmol) and DMAP was added into the mixture. The reaction was allowed to warm to amber temperature over 5 h. The reaction was then quenched with slow addition of Et_3N (4 mL) and MeOH (8 mL). The mixture was diluted with ether and washed with water, brine and dry with Na_2SO_4 . The pure product was obtained by vacuum distillation (1-2 mmHg) at $100^\circ C$ (3.0 g, 78% yield). LC/MS [Method A: rt: 5.75 min; m/z: 184 ($M+1$)].

15

Intermediate 20: 4-chloro-N-[2-fluoro-4-(trimethylsilyl)phenyl]pyridin-3-amine

To a solution of 4-chloro-3-iodopyridine (1.37 g, 5.73 mmol) in dry toluene 25 mL, was added sequentially $Pd(OAc)_2$ (122.5 mg, 0.55 mmol), rac-BINAP (0.34 g, 0.55 mmol), Cs_2CO_3 (8.9 g, 27.3 mmol), and 2-fluoro-4-(trimethylsilyl)aniline (1 g, 5.45 mmol). The mixture was degassed with nitrogen twice. The mixture was then refluxed at $130^\circ C$ under nitrogen for 3 days. The mixture was filtered and the filtrate was diluted with EtOAc, washed with H_2O , brine, dried over anhydrous Na_2SO_4 . The organic phase

25

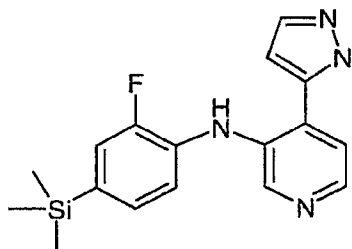
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was concentrated and the resulting residue was purified by flash column with 20% EtOAc-Hexane (0.3% Et₃N in Hexane) to afford desired adduct (1.14 g, 70.1%). LC/MS [Method A: rt: 6.14 min; m/z: 295 (M+1)].

5 *General procedure for Suzuki Coupling:*

(Ding, S.; Gray, N. S.; Wu, X.; Ding, Q.; Schultz, P. G. *J. Am. Chem. Soc.* 2002, 124, 1594-1596.)

Intermediate 21: (2-Fluoro-4-trimethylsilanyl-phenyl)-[4-(2H-pyrazol-3-yl)-pyridin-3-yl]-amine



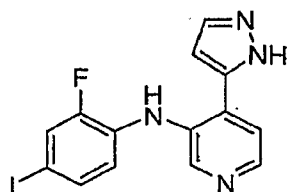
10

4-chloro-N-[2-fluoro-4-(trimethylsilyl)phenyl]pyridin-3-amine (70mg, 0.24 mmol), (0.95 g, 6.8 mmol), Tris(dibenzylideneacetone)diPalladium(0) (21.7 mg, 0.02 mmol), 1,3-dimesityl-1H-imidazol-3-ium chloride (16.2 mg, 0.05 mmol) and cesium carbonate (2.76 g, 8.5 mmol) were dispensed into a seal tube equipped with a stirring bar. The sealed tube was degassed by nitrogen before addition of anhydrous dioxane (14 mL). The mixture was heated at 130 °C over night. The mixture was then co-rotavapped with silica gel and applied to chromatography in silica gel. The desired product was obtained in pure form (63.2 mg, 63%). MS: m/z: 327 (M+1).

15

20 *General procedure for iodination*

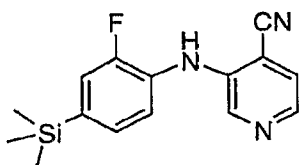
Example 61: (2-Fluoro-4-iodo-phenyl)-[4-(2H-pyrazol-3-yl)-pyridin-3-yl]-amine



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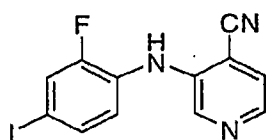
To a solution of (2-Fluoro-4-(trimethylsilyl)phenyl)-[4-(2H-pyrazol-3-yl)-pyridin-3-yl]-amine (45 mg, 0.14 mmol) in DCM (0.5 mL) was added ICI (2M, 0.21 mL, 0.41 mmol) dropwise. The mixture was stirred at RT overnight. The mixture was then diluted with DCM and washed with Na₂S₂O₃ (10%), water, brine and dried over Na₂SO₄. The organic phase was concentrated and the resulting residue was applied to chromatography in silica gel to afford product in 39% yield (52.4 mg). LC/MS [Method A: rt: 5.43 min; m/z: 381 (M+1)].

Intermediate 22: 3-[[2-fluoro-4-(trimethylsilyl)phenyl]amino]isonicotinonitrile



4-chloro-N-[2-fluoro-4-(trimethylsilyl)phenyl]pyridin-3-amine (122 mg, 0.41 mmol), Zn(CN)₂ (49 mg, 0.41 mmol) and Pd(Ph₃)₄ (48 mg, 0.04 mmol) were dispensed into a seal tube equipped with a stirring bar. The seal tube was degassed by vacuum and recharging with nitrogen three times before addition of anhydrous DMF (2 mL). The mixture was heated at 130 °C for 12 h. (Alterman, M; Hallberg, A. *J. Org. Chem.* **2000**, **65**, 7984-7989) The mixture was diluted with EtOAc and washed with water and dried over solid Na₂SO₄. The organic phase was concentrated. The resulting residue was applied to chromatography in silica gel to give the desired product in 97% yield (118 mg). LC/MS [Method B: rt: 6.71 min; m/z: 286 (M+1)].

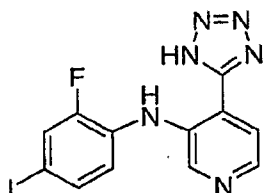
Intermediate 23: 3-[[2-fluoro-4-iodophenyl]amino]isonicotinonitrile



The title compound was prepared from 3-[[2-fluoro-4-(trimethylsilyl)phenyl]amino]isonicotinonitrile according to the general procedure for iodination mentioned above. LC/MS [Method A: rt: 5.00 min; m/z: 340 (M+1)].

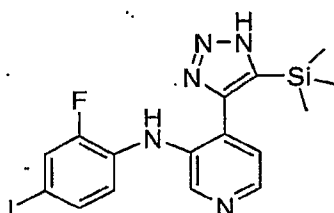
Example 62: N-(2-fluoro-4-iodophenyl)-4-(1H-tetrazol-5-yl)pyridin-3-amine

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3-[(2-fluoro-4-iodophenyl)amino]isonicotinonitrile (100 mg, 0.29 mmol), ammonium chloride (28 mg, 0.52 mmol), and sodium azide (35 mg, 0.54 mmol) were dispensed into a seal tube equipped with a stirring bar. The seal tube was degassed by vacuum and recharging with nitrogen three times before addition of anhydrous DMF (3 mL). The mixture was heated at 100 °C for three days. The mixture was diluted with saturated NaHCO₃ and washed with EtOAc. The aqueous phase was acidified with concentrated HCl to pH < 1, and washed with EtOAc. The aqueous layer was dried to give the desired product. LC/MS [Method B: rt: 6.008 min; m/z: 383 (M+1)].

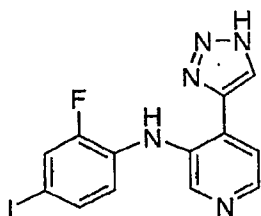
Example 63: N-(2-fluoro-4-iodophenyl)-4-[5-(trimethylsilyl)-1H-1,2,3-triazol-4-yl]pyridin-3-amine



To a solution of TMSdiazomethane (2 M, 90 µL, 0.18 mmol) in ether (2 mL) at 0 °C, was added butyl lithium (2.5 M, 72 µL, 0.18 mmol) dropwise. The mixture was stirred for 20 min at 0 °C. 3-[(2-fluoro-4-iodophenyl)amino]isonicotinonitrile (50 mg, 0.15 mmol) was then added into the solution. The reaction was allowed to warm to room temperature and proceed for three days. The mixture was diluted with ether and washed with saturated ammonium chloride solution, water, and dried over solid Na₂SO₄. The organic phase was concentrated, and the resulting residue was applied to chromatography in silica gel to give the desired product in 40% yield (27 mg). LC/MS [Method A: rt: 0.40 min; m/z: 454 (M+1)].

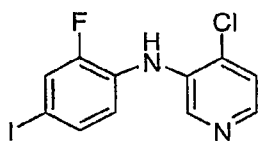
Example 64: N-(2-fluoro-4-iodophenyl)-4-(1H-1,2,3-triazol-4-yl)pyridin-3-amine

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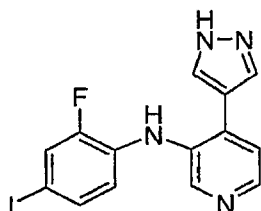
N-(2-fluoro-4-iodophenyl)-4-[5-(trimethylsilyl)-1H-1,2,3-triazol-4-yl]pyridin-3-amine (20 mg, 0.044 mmol) was added into a solution of NaOH (3.5 mg, 0.088 mmol) in THF (0.5 mL) for 5 days. The mixture was diluted with MeOH and neutralized to pH7 with acidic resin (Dowex). After filtration and the filtrate was concentrated to give pure product (11.5 mg, 69%). LC/MS [Method A: rt: 0.40 min; m/z: 381 (M+1)].

Intermediate 24: 4-chloro-N-(2-fluoro-4-iodophenyl)pyridin-3-amine



The title compound was prepared from 4-chloro-N-[2-fluoro-4-(trimethylsilyl)phenyl]pyridin-3-amine according the general procedure for iodination mentioned above. LC/MS [Method B: rt: 6.28 min; m/z: 349 (M+1)].

Example 65: (2-Fluoro-4-iodo-phenyl)-[4-(1H-pyrazol-4-yl)-pyridin-3-yl]-amine

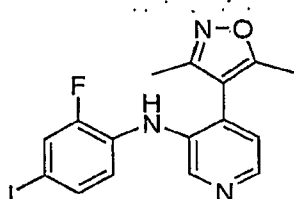


The title compound phenol was prepared from 4-chloro-N-[2-fluoro-4-(trimethylsilyl)phenyl]pyridin-3-amine according to the Suzuki Coupling conditions mentioned above. The resulting trimethylsilylated compound was then subjected to the

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iodination condition (see general procedure above) to afford the title compound.
LC/MS [Method A: rt: 4.48 min; m/z: 381(M+1)].

Example 66: 4-(3,5-Dimethyl-isoxazol-4-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine



The title compound was prepared according the same procedure above. LC/MS
[Method A: rt: 0.51 min; m/z: 410 (M+1)].

Functional Assays

Assay 1: MEK-1 enzyme assay (LANCE-HTRF)

Inhibition of human MEK1 kinase activity was monitored with a homogenous, fluorescence-based assay. The assay uses time resolved fluorescence resonance energy transfer to probe for phosphorylation of ERK1 by MEK1. The detailed procedure is described in the following:

Constitutively active MEK-1EE (150 ng/well) and ERK-2K52A (110 ng/well) were incubated in black, 384 well Optiplates (Perkin Elmer) for 1.5 hrs at room temperature, in the presence of increasing concentrations of inhibitors. The total reaction volume was 50 μ l. The inhibitors were serial diluted in kinase buffer (9 μ M ATP, 50 mM Tris-HCl pH 8.0, 10 mM MgCl₂, 1 mM DTT and 100 μ M Na₃VO₄), to a final concentration of 1% DMSO. At the end of the incubation, the enzyme reaction was quenched by adding 50 μ l of assay buffer containing 50 mM Tris-HCl, 50 mM EDTA, 0.1% BSA and a mixture of 1.25 μ g/mL europium labeled anti-phosphotyrosine mAb (Perkin Elmer) and 5 μ g/mL of allophycocyanin labeled anti-GST antibody (CIS-US). Plates were agitated on a plate shaker for 30 minutes and the phosphorylation of the ERK-2K52A substrate was measured by homogeneous time resolved fluorescence (HTRF) at 340 nm excitation and 665 nm (Europium) / 615 nm (APC) emission filters on the VICTOR V fluorescence plate reader. Phosphorylation is expressed as a ratio of 665 nm signal x

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10,000 / 615 nM signal. To assess the inhibitory potential of the compounds, IC₅₀-values were determined. Results are shown in Table 1, column 2, where “+++” indicates IC₅₀ values below or equal 3μM, “++” indicates IC₅₀ values below or equal 25μM and above 3μM, and “+” indicates IC₅₀ values above 25μM.

5 **Assay 2: Tumor cell proliferation assays (ATP Lite)**

Murine colon C26 (1500 cells/well), human melanoma A375 (2000 cells/well), or human pancreatic MiaPaCa-2 cells (2000 cells/well) were plated in white, 96 well plates (Corning) in growth media (DMEM with 10% fetal bovine serum, 2mM glutamine for C26 and MiaPaCa-2, and RPMI with 10% fetal bovine serum, 2mM glutamine for A375) and cultured overnight at 37°C in a humidified, 5% CO₂ incubator. Inhibitors were serially diluted in 100% DMSO and subsequently added to the cells, in growth media, to reach a final concentration of 0.25% DMSO. The cells were incubated for 4 days in the presence of the test compounds and the proliferation was quantitated using the ATPlite cell proliferation kit (Packard). Results of the cell proliferation assay are shown in Table 1, columns 3-5, where “+++” indicates IC₅₀ values below or equal to 3μM, “++” indicates IC₅₀ values below or equal to 25μM and above 3μM, and “+” indicates IC₅₀ values above 25μM.

Table 1: Results of MEK enzyme assay and tumor cell proliferation assays

Example	MEK1	C26	A375	MiaPaCa2
Example 1	+++	++	+++	++
Example 2	+++	+++	+++	+++
Example 3	+++	++	+++	+++
Example 4	+++	+++	+++	+++
Example 5	+++	+++	+++	+++
Example 6	+++	+++	+++	+++
Example 7	+++	+++	+++	+++
Example 8	+++	+++	+++	+++
Example 9	++	+++	+++	+++

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Example 10	+++	+++	+++	+++
Example 11	+++	+++	+++	+++
Example 12	+++	+++	+++	+++
Example 13	+++	+++	+++	+++
Example 14	+++	+++	+++	+++
Example 15	+++	+++	+++	+++
Example 16	+++	+++	+++	+++
Example 17	+++	+	++	++
Example 18	+++	+++	+++	+++
Example 19	+++	+++	+++	+++
Example 20	+++	+++	+++	+++
Example 21	+++	+++	+++	+++
Example 22	+++	+++	+++	+++
Example 23	+++	+++	+++	+++
Example 24	+++	+++	+++	+++
Example 25	+++	+++	+++	+++
Example 26	+++	+++	+++	+++
Example 27	+++	+++	+++	+++
Example 28	+++	NT	NT	NT
Example 29	+++	+++	+++	+++
Example 30	+++	NT	NT	NT
Example 31	+++	++	+++	++
Example 32	+++	+++	+++	+++
Example 33	+++	+++	+++	+++
Example 34	+++	+++	+++	+++

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Example 35	+++	+++	+++	+++
Example 36	+++	NT	NT	NT
Example 37	+++	+++	+++	+++
Example 38	+++	+++	+++	++
Example 39	+++	+	+++	+
Example 40	+++	+++	+++	+++
Example 41	+++	+++	+++	+++
Example 42	+++	+++	+++	+++
Example 43	+++	+++	+++	++
Example 44	+++	+++	+++	+++
Example 45	+++	++	+++	++
Example 46	+++	+++	+++	+++
Example 47	+++	+++	+++	+++
Example 48	+++	+++	+++	+++
Example 49	+++	+++	+++	+++
Example 50	+++	+++	+++	+++
Example 51	+++	++	+++	++
Example 52	+++	+++	+++	+++
Example 53	+++	+++	+++	++
Example 54	+++	+++	+++	+++
Example 55	+++	+++	+++	+++
Example 56	+++	+	++	++
Example 57	+++	+++	+++	+++
Example 58	+++	+++	+++	+++
Example 60	+++	+++	+++	+++

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Example 61	+++	++	+++	+++
Example 62	+++	+	+	+
Example 63	+++	+++	+++	+++
Example 64	+++	+++	+++	+++
Example 65	+++	+++	+++	+++
Example 66	+++	NT	NT	NT
Example 67	++	NT	NT	NT
Example 68	+++	NT	NT	NT
Example 69	+++	NT	NT	NT
Example 70	+	+	++	+

In vivo efficacy evaluation: Tumor growth inhibition in A375 Xenograft studies

(Malignant melanoma model)

Male nude (*nu/nu*) mice (100 total, 24-26 g body wt) were purchased from Charles
 5 River Laboratories. Animals were injected subcutaneously above the right foreleg with
 5 x 10⁶ A375 cells in 100 μ l of PBS. Tumors were measured with calipers 4, 7, 11 and
 14 days after cells were implanted. Tumor length (*l*) and width (*w*) were measured and
 tumor volume was calculated with the equation $l \cdot w^2 / 2$. At 16 days after cancer cells
 were implanted, tumors had reached an average volume of 93 mm³, at which time the
 10 animals were sorted into 6 groups so that each group had the same mean tumor
 volume. The vehicle used for dosing the compound in suspension was 10% Tween 80
 in water.

Treatments began on the afternoon of the 16th day after cancer cell injections, which
 was designated as Day 0. Tumor volume and body weight was measured for each
 15 animal 3 times per week until Day 14. On Day 14, all animals were euthanized, tumors
 removed and weighted.

The compound of Example 2 produced tumor regression at a dose of 10mg/kg and
 significant tumor growth inhibition at a dose of 0.5 mg/kg.

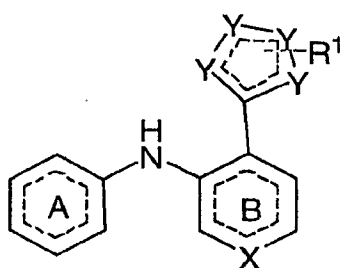
- 86 -

The entire teachings of all references cited herein are hereby incorporated by reference.

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CLAIMS

1. A compound of Formula (Ia),



Formula (Ia)

- 5 as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof, wherein:

10 rings A and B are optionally and independently substituted at any one or more substitutable ring carbon atoms with halogen, trimethylsilyl, cyano, C₁-C₆ alkyl, C₁-C₆ haloalkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ alkoxy, C₁-C₆ haloalkoxy, or phenyl wherein the phenyl is optionally substituted at any one or more substitutable ring carbon atoms with halogen, trimethylsilyl, cyano, C₁-C₆ alkyl, C₁-C₆ haloalkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, or C₁-C₆ alkoxy, C₁-C₆ haloalkoxy;

X is N or N→O;

Y is independently selected from NR', CR', S or O, whereby at least one Y is NR';

- 15 R¹ is selected from hydrogen, trimethylsilyl, C₁-C₆-alkyl, OR⁶, C(O)OR⁶, NR⁷R⁸, SR⁶, NR⁷S(O)(O)R', NR⁷C(O)R⁶, NR⁷C(O)NR⁷R⁶, NR⁷C(O)OR⁶, NR⁷C(O)C(O)OR⁶, C₁-C₄-alkyl-NR'R'', C₁-C₄-alkyl-heterocycloalkyl or C₁-C₄ alkyl-(OR')_n;

- 20 R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR', C₁-C₄-alkyl-NR'R'', C₁-C₄-alkyl-heterocycloalkyl, C₁-C₄ alkyl-(OR')_n or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl, cycloalkyl, aryl, heteroaryl, heterocycloalkyl, and alkyl-heterocycloalkyl is substituted or unsubstituted; or

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R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted;

5 R' and R'' are independently selected from hydrogen, C₁-C₆-alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted; or

R' and R'' can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

n is 0 to 2.

10 2. The compound according to claim 1, wherein

rings A and B are optionally and independently substituted at any one or more substitutable ring carbon atoms with halogen, C₁-C₆ alkyl or phenyl wherein the phenyl is optionally substituted at any one or more substitutable ring carbon atoms with halogen;

15 R¹ is selected from hydrogen, trimethylsilyl, OR⁶, NR⁷R⁸, NR⁷C(O)R⁶, NR⁷C(O)OR⁶, or C₁-C₄ alkyl-(OR')_n;

R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR', C₁-C₄-alkyl-NR'R'', C₁-C₄-alkyl-heterocycloalkyl, or C₁-C₄ alkyl-C(O)NR'R''; or

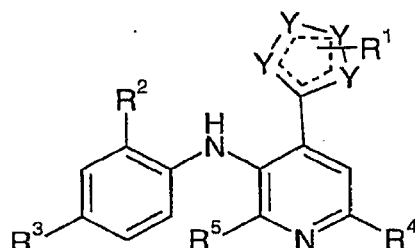
20 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom; and

R' and R'' are either hydrogen or C₁-C₆-alkyl; or

25 R' and R'' can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom.

3. A compound of Formula (I),

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Formula (I)

as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof, wherein:

Y is independantly selected from NR', CR'' or O, whereby at least one Y is NR';

5 R¹ is selected from hydrogen, C₁-C₆-alkyl, OR⁶, NR⁷R⁸, SR⁶, NR⁷S(O)(O)R', NR⁷C(O)R⁶, NR⁷C(O)NR⁷R⁶, or NR⁷C(O)OR⁶;

R² is halogen;

R³ is selected from Cl, F or I;

10 R⁴ and R⁵ are independently selected from hydrogen, halogen, C₁-C₆ alkyl or C₁-C₆ alkoxy, wherein said alkyl or alkoxy is substituted or unsubstituted;

R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR' or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl, cycloalkyl, aryl, heteroaryl and heterocyclyl substituted or unsubstituted; or

15 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

R' and R'' are independently selected from hydrogen, C₁-C₆-alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted.

20 4. The compound according to claim 3, wherein

R¹ is selected from OR⁶, NR⁷R⁸ or SR⁶;

R² is selected from F, Cl, I or Br;

R³ is selected from F, Cl or I;

R⁴ and R⁵ are independently either hydrogen or C₁-C₆ alkyl;

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R^6 , R^7 and R^8 are independently selected from hydrogen, C_1 - C_{10} alkyl, C_1 - C_4 alkyl- $C(O)OR'$, C_1 - C_4 alkyl- OR' or C_1 - C_4 alkyl- $C(O)NR'R''$, wherein said alkyl is substituted or unsubstituted; or

R^7 and R^8 can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; and

R' and R'' are either hydrogen or C_1 - C_6 -alkyl.

5. The compound according to claim 4, wherein

R^2 is F;

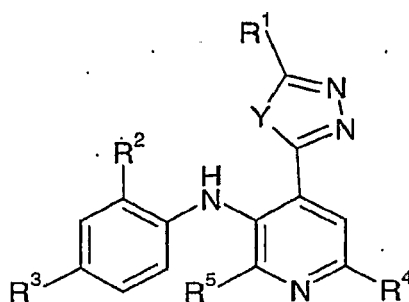
10 R^3 is I;

R^4 and R^5 are hydrogen;

R^6 , R^7 and R^8 are independently selected from hydrogen, C_1 - C_{10} alkyl, C_1 - C_4 alkyl- $C(O)OR'$, C_1 - C_4 alkyl- OR' or C_1 - C_4 alkyl- $C(O)NR'R''$, wherein said alkyl is substituted or unsubstituted; or

15 R^7 and R^8 form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted;

6. A compound of Formula (II),



Formula (II)

20 as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof, wherein, wherein

Y is either NR' or O ;

R^1 is selected from hydrogen, C_1 - C_6 -alkyl, OR^6 , NR^7R^8 , SR^6 , $NR^7S(O)(O)R'$, $NR^7C(O)R^6$, $NR^7C(O)NR^7R^6$, or $NR^7C(O)OR^6$;

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R² is halogen;

R³ is selected from Cl, F or I;

R⁴ and R⁵ are independently selected from hydrogen, halogen, C₁-C₆ alkyl or C₁-C₆ alkoxy, wherein said alkyl or alkoxy is substituted or unsubstituted;

5 R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR' or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl, cycloalkyl, aryl, heteroaryl and heterocyclyl substituted or unsubstituted; or

10 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

R' and R'' are independently selected from hydrogen, C₁-C₆-alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted.

7. A compound according to claim 6, wherein

15 R¹ is selected from OR⁶, NR⁷R⁸ or SR⁶;

R² is selected from F, Cl, I or Br;

R³ is selected from F, Cl or I;

R⁴ and R⁵ are independently either hydrogen or C₁-C₆ alkyl;

20 R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR' or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl is substituted or unsubstituted; or

R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; and

25 R' and R'' are either hydrogen or C₁-C₆-alkyl.

8. The compound according claim 7, wherein

R² is F;

R³ is I;

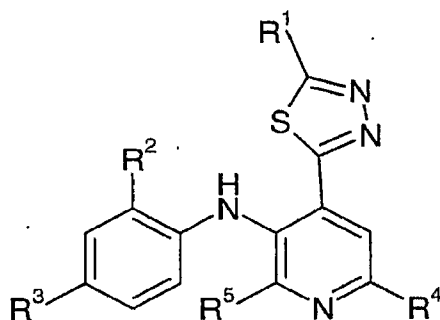
- 92 -

R^4 and R^5 are hydrogen;

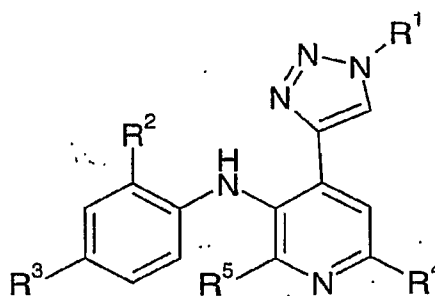
R^6 , R^7 and R^8 are independently selected from hydrogen, C_1 - C_{10} alkyl, C_1 - C_4 alkyl- $C(O)OR'$, C_1 - C_4 alkyl- OR' or C_1 - C_4 alkyl- $C(O)NR'R''$, wherein said alkyl is substituted or unsubstituted; or

5 R^7 and R^8 form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted.

9. A compound of Formula (IIa) or Formula (IIb),



Formula (IIa)



Formula (IIb)

10 as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof, wherein

R^1 is selected from hydrogen, trimethylsilyl, C_1 - C_6 -alkyl, OR^6 , $C(O)OR^6$, NR^7R^8 , SR^6 , $NR^7S(O)(O)R'$, $NR^7C(O)R^6$, $NR^7C(O)NR^7R^6$, $NR^7C(O)OR^6$, $NR^7C(O)C(O)OR^6$, C_1 - C_4 -alkyl- $NR'R''$, C_1 - C_4 -alkyl-heterocycloalkyl or C_1 - C_4 alkyl- $(OR')_n$;

15 R^2 is selected from halogen, C_1 - C_6 -alkyl or OR^6 ;

R^3 is halogen, trimethylsilyl, C_1 - C_6 -alkyl or OR^6 ;

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R⁴ and R⁵ are independently selected from hydrogen, halogen, C₁-C₆ alkyl or C₁-C₆ alkoxy, wherein said alkyl or alkoxy is substituted or unsubstituted;

R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR', C₁-C₄-alkyl-NR'R'', C₁-C₄-alkyl-heterocycloalkyl, C₁-C₄ alkyl-(OR')_n or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl, cycloalkyl, aryl, heteroaryl, heterocycloalkyl, and alkyl-heterocycloalkyl is substituted or unsubstituted; or

R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted;

R' and R'' are independently selected from hydrogen, C₁-C₆-alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted; or

R' and R'' can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

n is 0 to 2.

10. The compound according to claim 9, wherein

R¹ is selected from hydrogen, trimethylsilyl, OR⁶, NR⁷R⁸, NR⁷C(O)R⁶, NR⁷C(O)OR⁶, or C₁-C₄ alkyl-(OR')_n;

R² is selected from F or Cl;

R³ is I;

R⁴ and R⁵ are hydrogen;

R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR', C₁-C₄-alkyl-NR'R'', C₁-C₄-alkyl-heterocycloalkyl, or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl and alkyl-heterocycloalkyl is substituted or unsubstituted; or

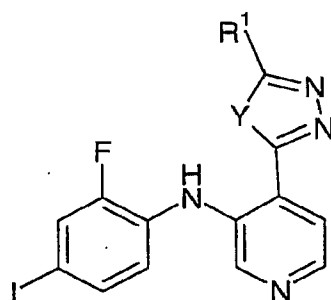
R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; and

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R' and R'' are either hydrogen or C₁-C₆-alkyl; or

R' and R'' can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted.

5 11. A compound of Formula (III)



Formula (III)

as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof, wherein, wherein

10 Y is either O, or NR⁵;

R¹ is selected from hydrogen, C₁-C₆-alkyl, OR⁶, NR⁷R⁸ or SR⁶ or NR⁷S(O)(O)R⁷;

R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, or heterocycloalkyl, C₁-C₄ alkyl-C(O)OR⁷, C₁-C₄ alkyl-OR⁷, C₁-C₄ alkyl-C(O)NR⁷R⁸, wherein said alkyl, cycloalkyl, aryl, heteroaryl and heterocycloalkyl substituted or unsubstituted; or

R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10 membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

R' and R'' are independently selected from hydrogen, C₁-C₆-alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted.

20

12. A compound according to claim 11, wherein

R¹ is selected from OR⁶, NR⁷R⁸ or SR⁶;

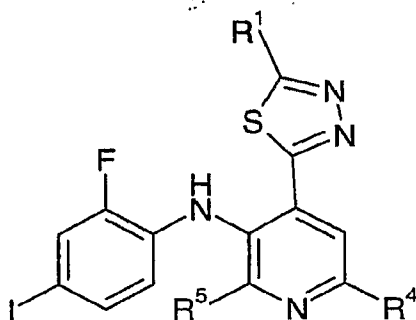
- 95 -

R^6 , R^7 and R^8 are independently selected from hydrogen, C_1 - C_{10} alkyl, C_1 - C_4 alkyl- $C(O)OR'$, C_1 - C_4 alkyl- OR' or C_1 - C_4 alkyl- $C(O)NR'R''$, wherein said alkyl is substituted or unsubstituted; or

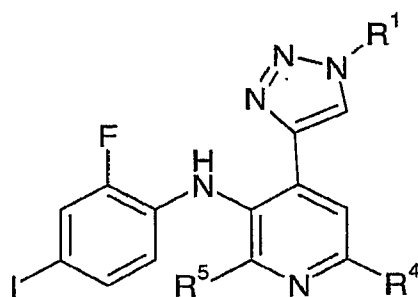
5 R^7 and R^8 can be taken together with the atom to which they are attached to form a 6 membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; and

R' and R'' are independantly either hydrogen or C_1 - C_6 -alkyl.

13. A compound of Formula (IIIa) or Formula (IIIb)



Formula (IIIa)



Formula (IIIb)

10 as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof, wherein, wherein

R^1 is selected from hydrogen, trimethylsilyl, C_1 - C_6 -alkyl, OR^6 , $C(O)OR^6$, NR^7R^8 , SR^6 , $NR^7S(O)(O)R'$, $NR^7C(O)R^6$, $NR^7C(O)NR^7R^6$, $NR^7C(O)OR^6$,
15 $NR^7C(O)C(O)OR^6$, C_1 - C_4 -alkyl- $NR'R''$, C_1 - C_4 -alkyl-heterocycloalkyl or C_1 - C_4 alkyl- $(OR')_n$;

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R⁴ and R⁵ are independently selected from hydrogen, halogen, C₁-C₆ alkyl or C₁-C₆ alkoxy, wherein said alkyl or alkoxy is substituted or unsubstituted;

R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR', C₁-C₄-alkyl-NR'R'', C₁-C₄-alkyl-heterocycloalkyl, C₁-C₄ alkyl-(OR')_n or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl, cycloalkyl, aryl, heteroaryl, heterocycloalkyl, and alkyl-heterocycloalkyl is substituted or unsubstituted; or

R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted;

R' and R'' are independently selected from hydrogen, C₁-C₆-alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted; or

R' and R'' can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

n is 0 to 2.

14. The compound according to claim 13, wherein

R¹ is selected from hydrogen, trimethylsilyl, OR⁶, NR⁷R⁸, NR⁷C(O)R⁶, NR⁷C(O)OR⁶, or C₁-C₄ alkyl-(OR')_n;

R⁴ and R⁵ are hydrogen;

R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR', C₁-C₄-alkyl-NR'R'', C₁-C₄-alkyl-heterocycloalkyl, or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl and alkyl-heterocycloalkyl is substituted or unsubstituted; or

R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; and

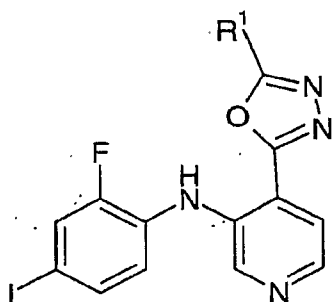
R' and R'' are either hydrogen or C₁-C₆-alkyl; or

- 97 -

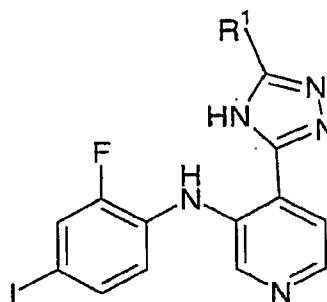
R' and R'' can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted.

- 5 15. The compound according to claim 13, wherein
- R¹ is selected from hydrogen, C₁-C₆-alkyl, OR⁶, NR⁷R⁸ or SR⁶ or NR⁷S(O)(O)R';
- R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, or heterocycloalkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR', C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl, cycloalkyl, aryl, heteroaryl and heterocyclyl substituted or unsubstituted; or
- 10 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10 membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and
- R' and R'' are independently selected from hydrogen, C₁-C₆-alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted.
- 15
16. The compound according to claim 15, wherein
- R¹ is selected from OR⁶, NR⁷R⁸ or SR⁶;
- R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR' or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl is substituted or unsubstituted; or
- 20 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 6 membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; and
- 25 R' and R'' are independantly either hydrogen or C₁-C₆-alkyl.
17. A compound of Formulae (IVa) or (IVb)

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Formula (IVa)



Formula (IVb)

as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof, wherein

R¹ is selected from hydrogen, C₁-C₆-alkyl, OR⁶, NR⁷R⁸ or SR⁶ or NR⁷S(O)(O)R¹;

5 R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₃-C₁₀ cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR' or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl, cycloalkyl, aryl, heteroaryl and heterocyclyl substituted or unsubstituted; or

10 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

R' and R'' are independently selected from hydrogen, C₁-C₆-alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted.

18. A compound according to claim 17, wherein

15 R¹ is selected from OR⁶, NR⁷R⁸ or SR⁶;

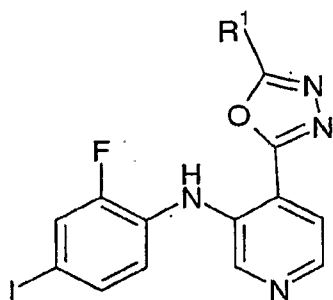
R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR' or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl is substituted or unsubstituted; or

20 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; and

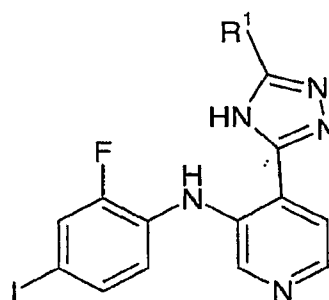
R' and R'' are independently either hydrogen or C₁-C₆-alkyl.

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19. A compound of Formulae (IVa) or (IVb)



Formula (IVa)



Formula (IVb)

as well as tautomers, pharmaceutically acceptable salts and pharmaceutically active derivatives thereof, wherein

5 R^1 is selected from hydrogen, trimethylsilyl, C_1 - C_6 -alkyl, OR^6 , $C(O)OR^6$, NR^7R^8 , SR^6 , $NR^7S(O)(O)R'$, $NR^7C(O)R^6$, $NR^7C(O)NR^7R^6$, $NR^7C(O)OR^6$, $NR^7C(O)C(O)OR^6$, C_1 - C_4 -alkyl- $NR'R''$, C_1 - C_4 -alkyl-heterocycloalkyl or C_1 - C_4 alkyl- $(OR')_n$;

10 R^6 , R^7 and R^8 are independently selected from hydrogen, C_1 - C_{10} alkyl, C_3 - C_{10} cycloalkyl, aryl, heteroaryl, heterocycloalkyl, C_1 - C_4 alkyl- $C(O)OR'$, C_1 - C_4 alkyl- OR' , C_1 - C_4 -alkyl- $NR'R''$, C_1 - C_4 -alkyl-heterocycloalkyl, C_1 - C_4 alkyl- $(OR')_n$ or C_1 - C_4 alkyl- $C(O)NR'R''$, wherein said alkyl, cycloalkyl, aryl, heteroaryl, heterocycloalkyl, and alkyl-heterocycloalkyl is substituted or unsubstituted; or

15 R^7 and R^8 can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted;

R' and R'' are independently selected from hydrogen, C_1 - C_6 -alkyl or aryl, wherein said alkyl or aryl is substituted or unsubstituted; or

20 R' and R'' can be taken together with the atom to which they are attached to form a 4 to 10-membered heteroaryl or heterocyclic ring, each of which is substituted or unsubstituted; and

n is 0 to 2.

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20. The compound according to claim 19, wherein

R¹ is selected from hydrogen, trimethylsilyl, OR⁶, NR⁷R⁸, NR⁷C(O)R⁶, NR⁷C(O)OR⁶, or C₁-C₄ alkyl-(OR')_n;

5 R⁶, R⁷ and R⁸ are independently selected from hydrogen, C₁-C₁₀ alkyl, C₁-C₄ alkyl-C(O)OR', C₁-C₄ alkyl-OR', C₁-C₄-alkyl-NR'R'', C₁-C₄-alkyl-heterocycloalkyl, or C₁-C₄ alkyl-C(O)NR'R'', wherein said alkyl and alkyl-heterocycloalkyl is substituted or unsubstituted; or

10 R⁷ and R⁸ can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted; and

R' and R'' are either hydrogen or C₁-C₆-alkyl; or

R' and R'' can be taken together with the atom to which they are attached to form a 6-membered heterocyclic ring with 1 or 2 N atoms and optionally an O atom, which is substituted or unsubstituted.

15

21. A compound according to claim 1, wherein the compound is selected from the group consisting of:

5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-3H-[1,3,4]oxadiazol-2-one,
[4-(5-Amino-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-(2-fluoro-4-iodo-phenyl)-amine,
20 5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-3H-[1,3,4]oxadiazole-2-thione,
5-[3-(2-Fluoro-4-iodo-phenylamino)-pyridin-4-yl]-2,4-dihydro-[1,2,4]triazol-3-one,
(2-Fluoro-4-iodo-phenyl)-[4-(5-methylsulfanyl-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-amine, and
(2-Fluoro-4-iodo-phenyl)-[4-(5-morpholin-4-yl-[1,3,4]oxadiazol-2-yl)-pyridin-3-yl]-amine.
25

22. A method of treating a mammal suffering from or susceptible to a hyperproliferative disease comprising administering to said mammal a therapeutically effective amount of a compound corresponding to claim 1.

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23. The method according to claim 22, wherein said hyperproliferative disease is selected from cancer or inflammation.
24. The method according to claim 22 or 23, wherein said hyperproliferative disease is related to the hyperactivity of MEK or a disease associated to MEK in mammals.
25. A pharmaceutical composition, which comprises a compound of claim 1 and a pharmaceutically acceptable carrier.
26. A compound according to claim 1 for use as a medicament.
27. A use of a compound according to claim 1 for the preparation of a medicament for the treatment of a hyperproliferative disease.
28. The use according to claim 27, wherein said hyperproliferative disease is cancer or inflammation.
29. The use according to claim 27, wherein said hyperproliferative disease is related to the hyperactivity of MEK as well as a disease associated to MEK in mammals.