

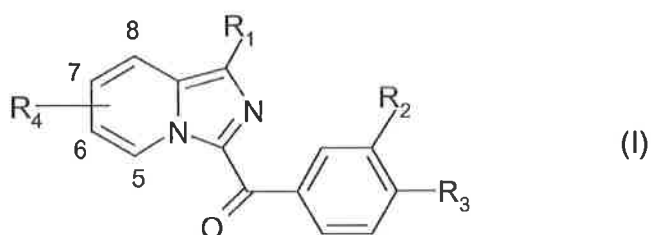
IMIDAZOPYRIDINE DERIVATIVES, PROCESS FOR THE PREPARATION THEREOF AND THERAPEUTIC USE THEREOF

The present invention relates to imidazopyridine derivatives which are inhibitors of
 5 FGFs (Fibroblast Growth Factors), to the process for the preparation thereof and to the therapeutic use thereof.

FGFs are a family of polypeptides synthesized by a large number of cells during embryonic development and by cells of adult tissues in various pathological
 10 conditions.

Indolizine derivatives, which are antagonists of the binding of FGFs to their receptors, are described in international patent applications WO 03/084956 and WO 2005/028476. Imidazo[1,5-a]pyridine derivatives which are FGF antagonists
 15 are described in international patent application WO 2006/097625. Novel imidazopyridine derivatives, which are antagonists of the binding of FGFs to their receptors, have now been identified.

The subject of the present invention is thus compounds, imidazopyridine
 20 derivatives, corresponding to formula (I):



in which:

25

- R₁ represents
 - . a hydrogen or halogen atom,
 - . an alkyl group optionally substituted with -COOR₅,
 - . an alkenyl group optionally substituted with -COOR₅,
 - 30 . a -COOR₅ or -CONR₅R₆ group,

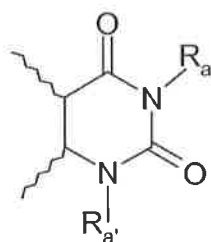
. an $-NR_5COR_6$ or $-NR_5-SO_2R_6$ group,

or

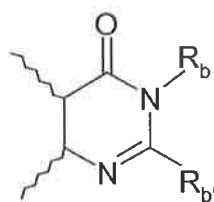
- . an aryl group, in particular phenyl, or a heteroaryl group, said aryl or heteroaryl group being optionally substituted with one or more groups selected from: halogen atoms, alkyl groups, cycloalkyl groups, $-COOR_5$, $-CF_3$, $-OCF_3$, $-CN$, $-C(NH_2)NOH$, $-OR_5$, $-O-Alk-COOR_5$, $-O-Alk-NR_5R_6$, $-O-Alk-NR_7R_8$, $-Alk-OR_5$, $-Alk-COOR_5$, $-CONR_5R_6$, $-CO-NR_5-OR_6$, $-CO-NR_5-SO_2R_7$, $-CONR_5-Alk-NR_5R_6$, $-CONR_5-Alk-NR_7R_8$, $-Alk-NR_5R_6$, $-NR_5R_6$, $-NC(O)N(CH_3)_2$, $-CO-Alk$, $-CO(OAlk)_nOH$, $COO-Alk-NR_5R_6$, $COO-Alk-NR_7R_8$ and 5-membered heteroaryl groups, said heteroaryl groups being optionally substituted with one or more groups selected from halogen atoms and alkyl, $-CF_3$, $-CN$, $-COOR_5$, $-Alk-OR_5$, $-Alk-COOR_5$, $-CONR_5R_6$, $-CONR_7R_8$, $-CO-NR_5-OR_6$, $-CO-NR_5-SO_2R_6$, $-NR_5R_6$ and $-Alk-NR_5R_6$ groups, or with a hydroxyl group or with an oxygen atom,

- n is an integer ranging from 1 to 3,

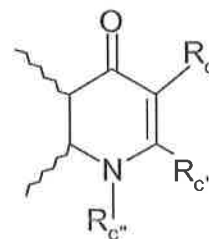
- R_2 and R_3 together form, with the carbon atoms of the phenyl nucleus to which they are attached, a 6-membered nitrogenous heterocycle corresponding to one of formula (A), (B) or (C) below:



(A)



(B)



(C)

20

in which the wavy lines represent the phenyl nucleus to which R_2 and R_3 are attached and:

- . R_a represents a hydrogen atom or an alkyl, haloalkyl, $-Alk-CF_3$, $-Alk-COOR_5$, $-Alk'-COOR_5$, $-Alk-CONR_5R_6$, $-Alk'-CONR_5R_6$, $-Alk-CONR_7R_8$, $-Alk-NR_5R_6$, $-AlkCONR_5-OR_6$, $-Alk-NR_7R_8$, $-Alk-cycloalkyl$, $-Alk-O-R_5$, $-Alk-S-R_5$, $-Alk-CN$, $-OR_5$, $-OAlkCOOR_5$, $-NR_5R_6$, $-NR_5-COOR_6$, $-Alk-aryl$, $-Alk-O-aryl$, $-Alk-O-heteroaryl$, $-Alk-heteroaryl$ or heteroaryl group, where the aryl or heteroaryl group is optionally substituted with one or more halogen atoms and/or alkyl, cycloalkyl, $-CF_3$, $-OCF_3$,

25

-O-R₅ or -S-R₅ groups,

. R_a' represents a hydrogen atom or a linear, branched, cyclic or partially cyclic alkyl group, or an -Alk-OR₅, -Alk-NR₅R₆ or -Alk-NR₇R₈ group, R_a' being optionally substituted with one or more halogen atoms,

5 . R_b represents a hydrogen atom or an alkyl or -Alk-COOR₅ group,

. R_b' represents a hydrogen atom or an alkyl, haloalkyl, cycloalkyl, phenyl or -Alk-COOR₅ group,

. R_c represents a hydrogen atom or an alkyl, -CN, -COOR₅, -CO-NR₅R₆, -CONR₇R₈, -CO-NR₅-Alk-NR₅R₆, -CONR₅-Alk-OR₅, -CONR₅SO₂R₅, -Alk-
10 aryl or -Alk-heteroaryl group, where the aryl or heteroaryl group is optionally substituted with one or more halogen atoms and/or alkyl, cycloalkyl, -CF₃, -OCF₃, -O-alkyl or -S-alkyl groups,

. R_c' represents a hydrogen atom or an alkyl group,

. R_c'' represents a hydrogen atom or an alkyl, alkenyl, haloalkyl, cycloalkyl, -Alk-NR₅R₆, -Alk-NR₇R₈, -Alk-OR₅ or -Alk-SR₅ group,
15

- R₄, located on position 6, 7 or 8 of the imidazopyridine nucleus, represents:

. a hydrogen atom,
20 . a -COOR₅ group,
. a -CO-NR₅-Alk-NR₅R₆ group,
. a -CO-NR₅-Alk-NR₇R₈ group, or
. a -CO-NR₅-Alk-OR₆ group,

- R₅ and R₆, which may be identical or different, represent hydrogen atoms, haloalkyl groups or alkyl groups, cycloalkyl groups or an Ms (mesyl) group,
25

- R₇ and R₈, which may be identical or different, represent hydrogen atoms or alkyl or phenyl groups, or else R₇ and R₈ together form a 3- to 8-membered saturated ring which can optionally contain a heteroatom,

30

- Alk represents a linear or branched alkylene chain, and

- Alk' represents a linear, branched, cyclic or partially cyclic alkylene chain,

these compounds being optionally present in the form of a pharmaceutically acceptable salt thereof.

The compounds of formula (I) may comprise one or more asymmetric carbon atoms. They can therefore exist in the form of enantiomers or of diastereoisomers. These enantiomers and diastereoisomers, and also mixtures thereof, including
5 racemic mixtures, are part of the invention.

The compounds of formula (I) can exist in the form of bases or of acids or can be salified with acids or bases, in particular pharmaceutically acceptable acids or bases. Such addition salts are part of the invention. These salts are
10 advantageously prepared with pharmaceutically acceptable acids or bases, but the salts of other acids or bases that are of use, for example, for purifying or isolating the compounds of formula (I) are also part of the invention.

The compounds of formula (I) can also exist in the form of hydrates or of solvates,
15 namely in the form of associations or combinations with one or more molecules of water or with a solvent. Such hydrates or solvates are also part of the invention.

In the context of the invention, and unless otherwise mentioned in the text, the term:

20 - "alkyl" is intended to mean: a linear or branched, saturated hydrocarbon-based aliphatic group containing from 1 to 6 carbon atoms;

- "cycloalkyl" is intended to mean: a cyclic alkyl group comprising from 3 to 8 ring members, containing between 3 and 6 carbon atoms and optionally comprising one or more heteroatoms, for example 1 or 2 heteroatoms, such as nitrogen and/or
25 oxygen, said cycloalkyl group being optionally substituted with one or more halogen atoms and/or alkyl groups. By way of examples, mention may be made of cyclopropyl, cyclopentyl, piperaziny, pyrrolidiny and piperidiny groups;

- "partially cyclic alkyl group" is intended to mean: an alkyl group of which only a part forms a ring;

30 - "alkylene" is intended to mean: a linear or branched divalent alkyl group containing from 1 to 6 carbon atoms;

- "halogen" is intended to mean: a chlorine, fluorine, bromine or iodine atom;

- "haloalkyl" is intended to mean: an alkyl chain in which all or some of the hydrogen atoms are replaced with halogen atoms, such as fluorine atoms;

- "aryl" is intended to mean: a cyclic aromatic group containing between 5 and 10 carbon atoms, for example a phenyl group;

- "heteroaryl" is intended to mean: a cyclic aromatic group containing between 3 and 10 atoms, including one or more heteroatoms, for example between 1 and 4 heteroatoms, such as nitrogen, oxygen or sulphur, this group comprising one or more, preferably one or two, rings. The heterocycles may comprise several condensed rings. The heteroaryls are optionally substituted with one or more alkyl groups or an oxygen atom. By way of examples, mention may be made of thienyl, pyridinyl, pyrazolyl, imidazolyl, thiazolyl and triazolyl groups; and

- "5-membered heteroaryl" is intended to mean: a heteroaryl group consisting of a 5-membered ring comprising 1 to 4 heteroatoms (such as oxygen and/or nitrogen atoms), optionally substituted with one or more alkyl groups or a hydroxyl group or with an oxygen atom. Mention may, for example, be made of oxadiazolyl and tetrazolyl groups;

- the halogens are preferably selected from F and Cl.

Among the compounds of formula (I) according to the invention, mention may be made of a subgroup of compounds in which R_1 represents:

- . a hydrogen or halogen atom,
- . an alkyl group which is unsubstituted or substituted with $-\text{COOR}_5$,
- . an alkenyl group which is unsubstituted or substituted with $-\text{COOR}_5$,
- . a $-\text{COOR}_5$ group,
- . a $-\text{CONR}_5\text{R}_6$ group,
- . an $-\text{NR}_5-\text{SO}_2\text{R}_6$ group, or
- . a phenyl group optionally substituted with one or two groups selected from:
 - halogen atoms;
 - alkyl groups optionally substituted with $-\text{COOR}_5$;
 - $-\text{CN}$ (cyano), $-\text{C}(\text{NH}_2)\text{NOH}$, $-\text{COOR}_5$, $-\text{CONR}_5\text{R}_6$, $-\text{CO}-\text{NR}_5-\text{OR}_6$, $-\text{CO}-\text{NR}_5-\text{SO}_2\text{R}_6$, $-\text{COAlk}$, $-\text{CO}(\text{OAlk})_n\text{OH}$, $-\text{OR}_5$, $-\text{OCF}_3$, $-\text{O}-\text{Alk}-\text{COOR}_5$, $-\text{Alk}-\text{OR}_5$, $-\text{NR}_5\text{R}_6$ or $-\text{NC}(\text{O})\text{N}(\text{CH}_3)_2$ groups,
 - 5-membered heteroaryls optionally substituted with an alkyl group and/or a hydroxyl group or an oxygen atom,

in which R_5 and R_6 , which may be identical or different, represent hydrogen atoms, or alkyl groups optionally substituted with an $-NR_7R_8$ group,

R_7 represents a hydrogen atom, an alkyl group containing 1 or 2 carbon atoms or a phenyl group, n is an integer ranging from 1 to 3, or

5 . a heteroaryl group which is optionally condensed and/or optionally substituted with one or two groups selected from alkyl groups, OR_5 , $-COOR_5$, $-NR_5R_6$ and cycloalkyl groups, and an oxygen atom, in which R_5 and R_6 , which may be identical or different, represent hydrogen atoms or alkyl groups containing 1 or 2 carbon atoms.

10 Among the compounds of formula (I) according to the invention, mention may be made of another subgroup of compounds in which R_2 and R_3 together form, with the carbon atoms of the phenyl nucleus to which they are attached, a 6-membered nitrogenous heterocycle corresponding to either of formulae (A) and (B) defined above, preferably corresponding to formula (A).

15

Formula (A) or (B) is advantageously such that:

 . R_a represents a hydrogen atom or an alkyl group, optionally substituted with one or more halogens; $-AlkCONR_5R_6$; haloalkyl; $-CH_2-COOR_5$; $-Alk$ -heteroaryl, $-Alk$ -O-phenyl or $-Alk$ -phenyl, where the phenyl group is optionally substituted with
20 one or two alkyl groups and/or OR_5 and/or halogen atoms; $-Alk$ -cycloalkyl,

 . R_a' represents a hydrogen atom or a linear, branched, cyclic or partially cyclic alkyl group, or a $-CH_2-OR_5$ or $-Alk-NR_5R_6$ group,

 . R_b represents a hydrogen atom or an alkyl group,

 . R_b' represents a hydrogen atom or an alkyl, phenyl or $-CH_2-COOR_5$ group,

25 in which the alkyl groups contain 1 to 6 carbon atoms, R_5 being as defined above.

 Among the compounds of formula (I) according to the invention, mention may be made of another subgroup of compounds in which R_4 represents a hydrogen atom
30 or a $-COOH$, $-CO-NH-Alk-NR_7R_8$ or $-CO-NH-Alk-OH$ group, in which Alk , R_7 and R_8 are as defined above, or else an alkyl group, preferably containing 1 to 3 carbon atoms, which is unsubstituted.

 In the compounds of formula (I) according to the invention, R_4 is advantageously

located on position 6 or 7 of the imidazopyridine nucleus.

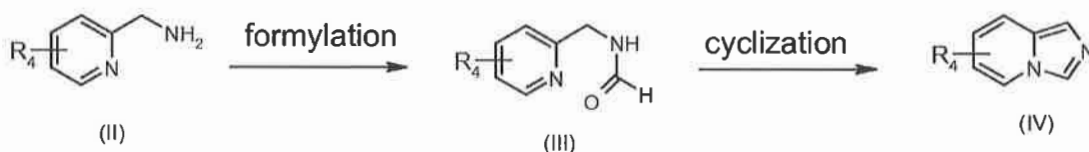
Among the compounds which are subjects of the invention, mention may be made of the following compounds:

- 5 5-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}-2-fluorobenzoic acid
 3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid
 3-{3-[(1-methyl-2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid
10 3-{3-[(3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid
 3-{3-[(3-(4-fluorobenzyl)-1-(methoxymethyl)-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid
15 3-{3-[(3-(4-fluorobenzyl)-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid
 3-{3-[(3-[2-(4-fluorophenyl)ethyl]-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid
 3-{3-[(1-methyl-3-[(5-methylthiophen-2-yl)methyl]-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid
20 3-{3-[(3-[(5-methylthiophen-2-yl)methyl]-2,4-dioxo-1-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid
 3-{3-[(2,4-dioxo-1-propyl-3-(thiophen-2-ylmethyl)-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid
25 3-{3-[(3-[2-(4-fluorophenoxy)ethyl]-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid

30 In the following text, the term "protective group" is intended to mean a group which makes it possible, firstly, to protect a reactive function such as a hydroxyl or an amine during a synthesis and, secondly, to regenerate the intact reactive function at the end of synthesis. Examples of protective groups and also methods of protection and deprotection are given in "Protective Groups in Organic Synthesis", Green *et al.*, 3rd Edition (John Wiley & Sons, Inc., New York).

- In the remainder of the text, the term "leaving group" is intended to mean a group that can be readily cleaved from a molecule by breaking a heterolytic bond, with the departure of a pair of electrons. This group can thus be readily replaced with another group during a substitution reaction, for example. Such leaving groups are,
- 5 for example, halogens or an activated hydroxyl group, such as a mesyl, tosyl, triflate, acetyl, para-nitrophenyl, etc. Examples of leaving groups and also methods for the preparation thereof are given in "Advances in Organic Chemistry", J. March, 3rd Edition, Wiley Interscience, p. 310-316.
- 10 In accordance with the invention, the compounds of general formula (I) can be prepared according to the processes hereinafter.

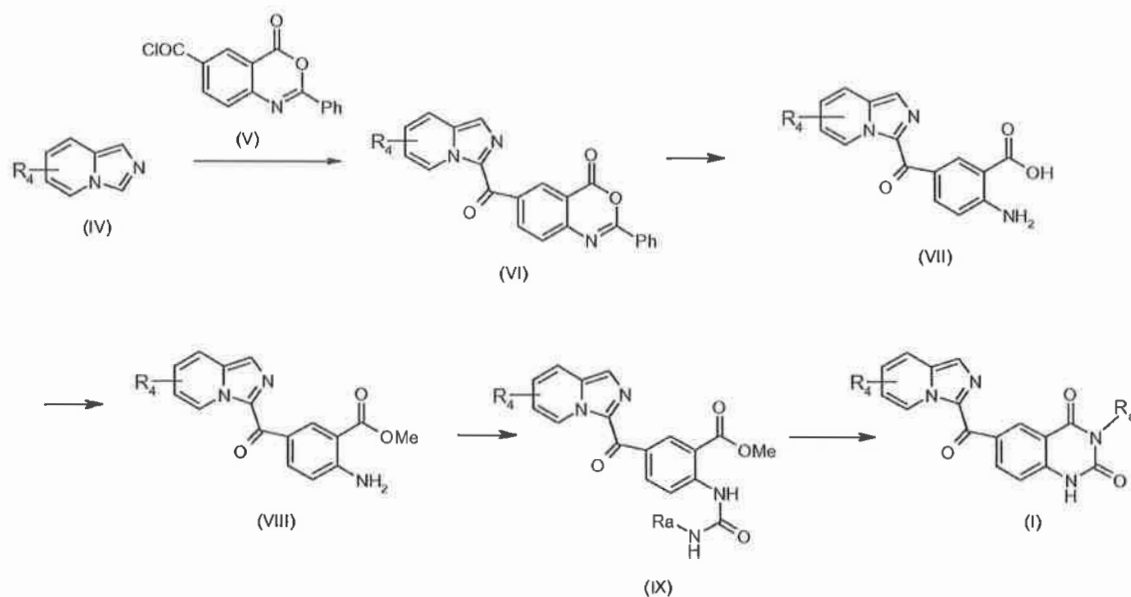
- The compounds of formula (IV) are obtained by methods known in the literature, starting from the suitably substituted corresponding 2-aminomethylpyridines, according to the following reaction scheme, described in *J. Chem. Soc.* (1955),
- 15 2834-2836



- 20 When R_4 represents COOR_5 , the compounds of formula (II) are obtained according to the reaction scheme described in WO 06/097625.

Scheme 1 presents a pathway for obtaining the compounds of formula (I) in which R_2 and R_3 together form a nitrogenous heterocycle of formula (A) as defined above, and in which R_1 and R_a represent hydrogen atoms.

Scheme 1 (Method 1):



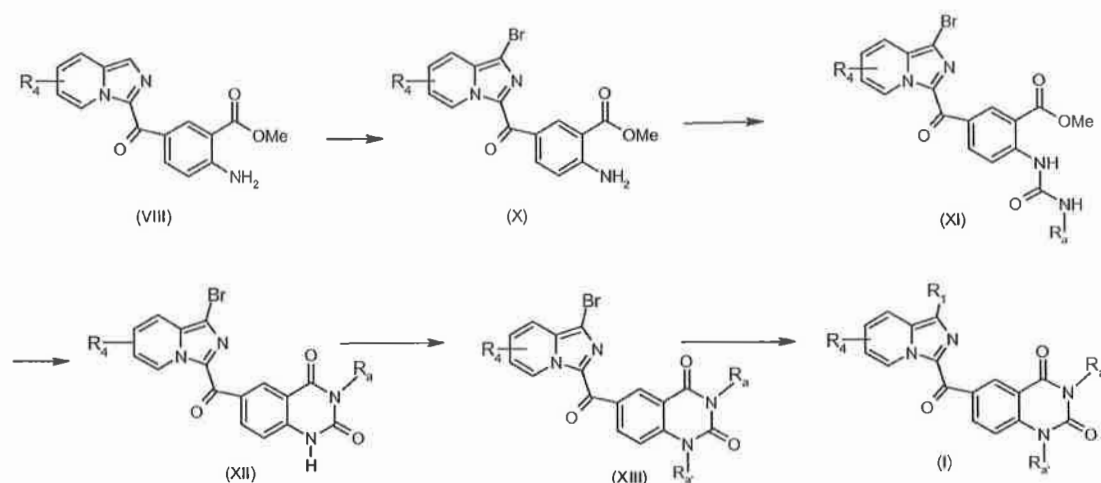
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The compound of formula (IV), in which R_4 is as defined for the compound of formula I, is condensed with the compound of formula V in order to obtain the compound of formula VI. The compound of formula VI is subjected to a basic hydrolysis reaction in order to obtain the compound of formula VII. The esterification of the compound of formula VII produces the compound of formula VIII. By reacting triphosgene, the isocyanate corresponding to the compound of formula VIII is formed, which is condensed with an amine of formula R_aNH_2 in order to obtain the urea of formula IX. The compound of formula IX is subjected to a cyclization reaction in a basic medium in order to obtain the compound of formula I in which R_4 and R_a are as defined above.

Scheme 2 presents a pathway for obtaining the compounds of formula (I) in which R_2 and R_3 together form a nitrogenous heterocycle of formula (A) as defined above, and in which R_1 represents a group as defined in the general formula, except for a hydrogen atom.

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Scheme 2 (Method 2):

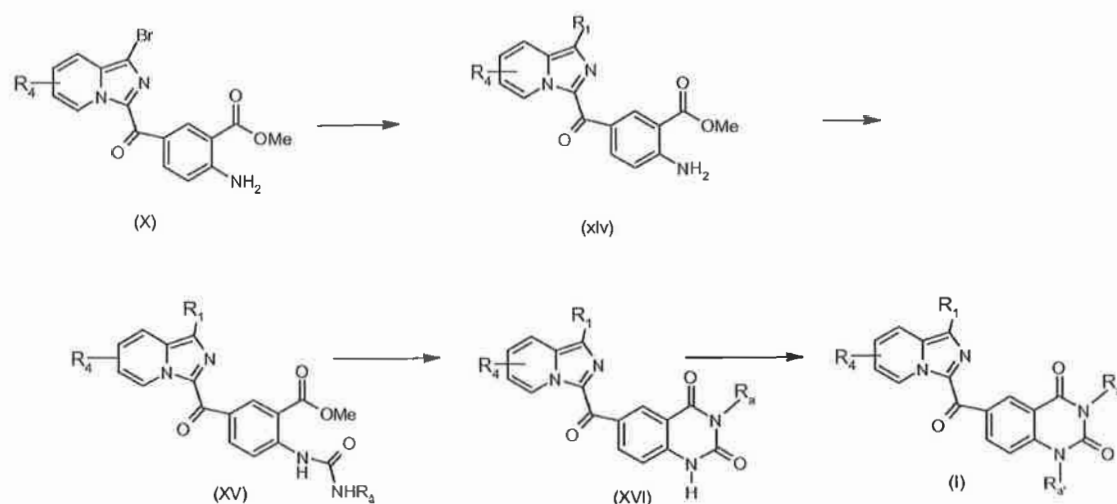


- 5 The compound of formula VIII is subjected to a bromination reaction in order to obtain the compound of formula X. By reacting triphosgene, the isocyanate corresponding to the compound of formula X is formed, which is condensed with an amine of formula R_aNH_2 in order to obtain the urea of formula XI. The compound of formula XI is subjected to a cyclization reaction in a basic medium, in
- 10 order to obtain the compound of formula XII. The compound XII is subjected to an alkylation reaction in the presence of a base and of a halogenated derivative $R_a'X$ in order to obtain the compound of formula XIII. The compound of formula XIII is subjected, in the presence of a palladium catalyst, of a ligand and of a base,
- to a reaction with phenylboronic or heteroarylboronic or phenylboronate ester or
 - 15 heteroarylboronate ester derivatives according to a Suzuki coupling,
 - or else to an imination reaction with benzophenoneimine followed by an acid hydrolysis and by an alkylation reaction with a sulphonyl chloride of formula R_6SO_2Cl ,
 - or else to a cyanation reaction with zinc cyanide, followed by an acid hydrolysis
 - 20 and by an esterification or a peptide coupling with an amine $R_5R_6NH_2$,
- in order to obtain the compound of formula I in which R_1 , R_4 , R_a and R_a' are as defined above.

Scheme 3 presents a pathway for obtaining the compounds of formula (I) in which R_2 and R_3 together form a nitrogenous heterocycle of formula (A) as defined above, and in which R_1 represents a group as defined in the general formula, except for a hydrogen atom, and in which R_4 is as defined above.

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Scheme 3 (Method 3):

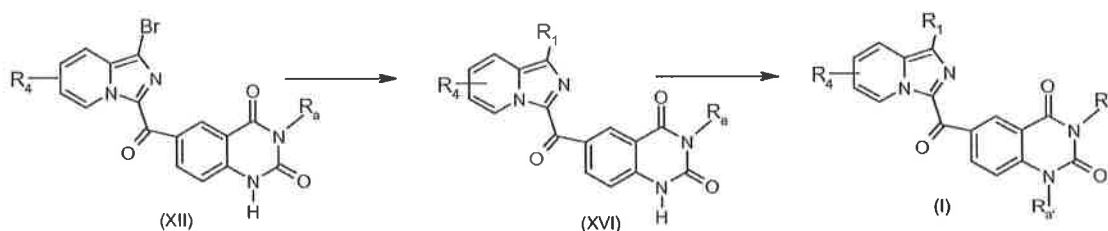


- 10 The compound of formula X is subjected, in the presence of a palladium catalyst, of a ligand and of a base,
- to a reaction with phenylboronic or heteroarylboronic or phenylboronate ester or heteroarylboronate ester derivatives according to a Suzuki coupling,
 - or else to an imination reaction with benzophenoneimine, followed by an acid
- 15 hydrolysis and by an alkylation reaction with a sulphonyl chloride of formula R_6SO_2Cl ,
- or else to a cyanation reaction with zinc cyanide, followed by an acid hydrolysis and by an esterification or a peptide coupling with an amine $R_5R_6NH_2$, R_5 and R_6 being defined above,
- 20 in order to obtain the compound of formula XIV in which R_1 is as defined above. By reacting triphosgene, the isocyanate corresponding to the compound of formula XIV is formed, which is condensed with an amine of formula R_aNH_2 in order to obtain the urea of formula XV. The compound of formula XV is subjected to a cyclization reaction in a basic medium in order to obtain the compound of formula
- 25 XVI. The compound XVI is subjected to an alkylation reaction in the presence of a

base and of a halogenated derivative R_aX in order to obtain the compound of formula I.

Scheme 4 presents a pathway for obtaining the compounds of formula (I) in which R_2 and R_3 together form a nitrogenous heterocycle of formula (A) as defined above, and in which R_1 represents a group as defined in the general formula, except for a hydrogen atom, and in which R_4 is as defined above.

Scheme 4 (Method 4):



The compound of formula XII is subjected, in the presence of a palladium catalyst, of a ligand and of a base,

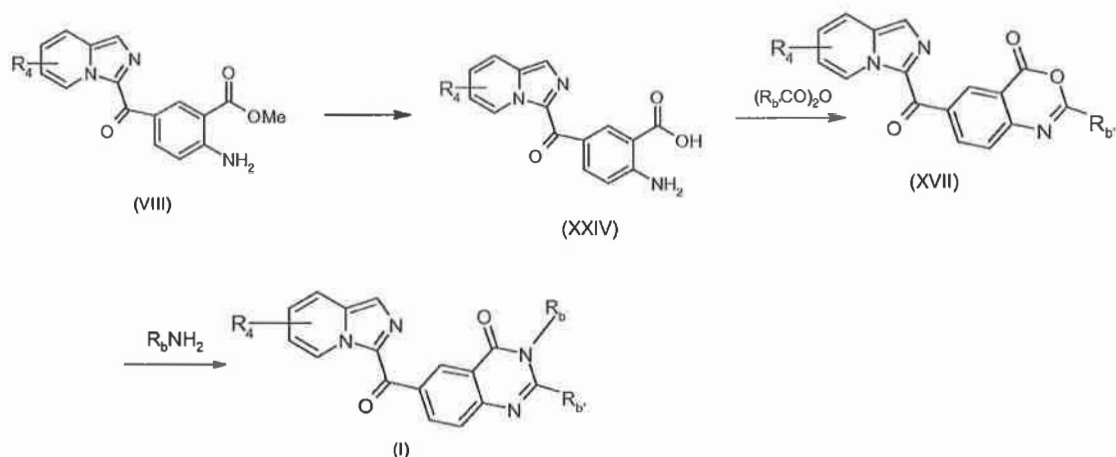
- to a reaction with phenylboronic or heteroarylboronic or phenylboronate ester or heteroarylboronate ester derivatives according to a Suzuki coupling,
- or else to an imination reaction with benzophenoneimine, followed by an acid hydrolysis and by a sulphonylation reaction with a sulphonyl chloride of formula R_6SO_2Cl ,

- or else to a cyanation reaction with zinc cyanide, followed by an acid hydrolysis and by an esterification or a peptide coupling with an amine $R_5R_6NH_2$,
- in order to obtain the compound of formula XVI in which R_1 is as defined above.

The compound XVI is subjected to an alkylation reaction in the presence of a base and of a halogenated derivative R_aX in order to obtain the compound of formula I.

Scheme 5 presents a pathway for obtaining the compounds of formula (I) in which R_2 and R_3 together form a nitrogenous heterocycle of formula (B) as defined above, in which R_1 represents a hydrogen atom and in which R_4 is as defined above.

Scheme 5 (Method 5):

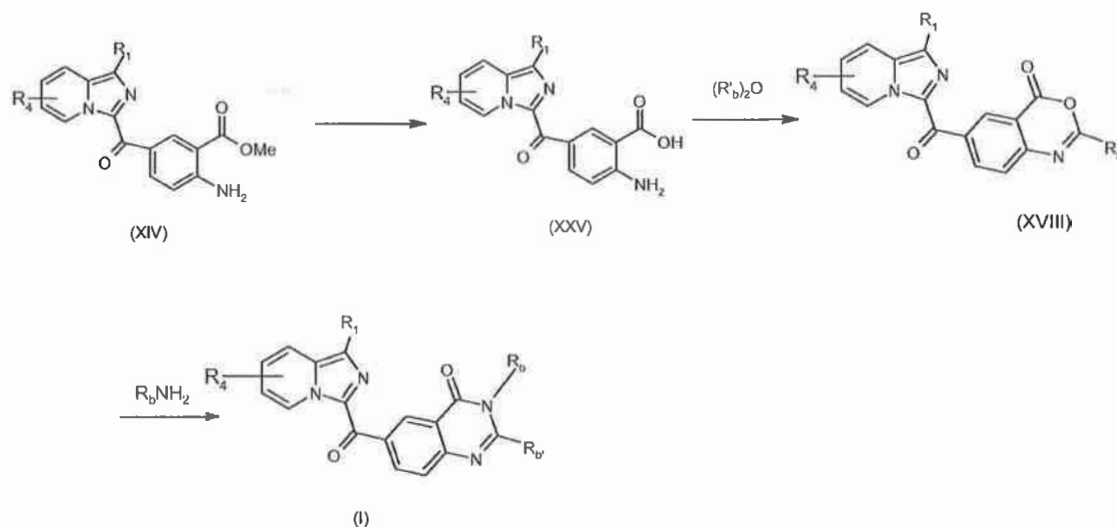


- 5 The compound VIII is subjected to a saponification reaction in order to obtain the compound XXIV. The compound XXIV is subsequently subjected to a condensation reaction with an alkyl or aryl anhydride $(R_bCO)_2O$ in order to obtain the compound of formula XVII. The compound of formula XVII is subjected to a condensation reaction with an amine R_bNH_2 in order to obtain a compound of
- 10 formula I in which R_b and $R_{b'}$ are as defined above.

Scheme 6 presents a pathway for obtaining the compounds of formula (I) in which R_2 and R_3 together form a nitrogenous heterocycle of formula (B) as defined above and in which R_1 is as defined above, except for a hydrogen, and in which R_4 is as

15 defined above.

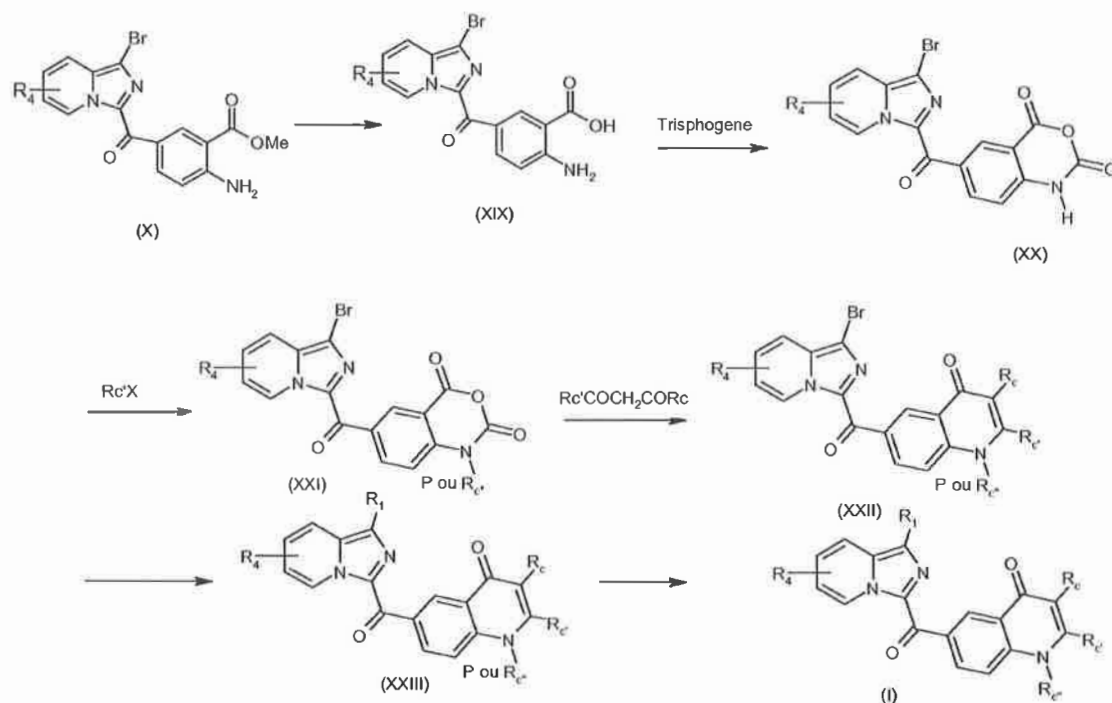
Scheme 6 (Method 6) :



- 5 The compound XIV is subjected to a saponification reaction in order to obtain the compound XXV. The compound XXV is subsequently subjected to a condensation reaction with an alkyl or aryl anhydride $(R'_bCO)_2O$ in order to obtain the compound of formula XVIII. The compound of formula XVIII is subjected to a condensation reaction with an amine R_bNH_2 in order to obtain a compound of formula I in which
- 10 R_b and $R_{b'}$ are as defined above.

Scheme 7 presents a pathway for obtaining the compounds of formula (I) in which R_2 and R_3 together form a nitrogenous heterocycle of formula (C) as defined above and in which $R_{c''}$ and also R_1 and R_4 are as defined above.

Scheme 7 (Method 7):



5

The compound X is subjected to a saponification reaction in order to obtain the compound XIX. The compound XIX is subjected to a condensation reaction in the presence of triphosgene, in order to obtain the compound XX. The compound XX is subjected to an alkylation reaction in the presence of a halogenated derivative $R_c'X$ or of a protective group in order to obtain the compound XXI. The compound XXI is subjected to a condensation reaction with a malonic derivative in order to obtain the compound XXII in which R_c' and R_c are as defined above. The compound XXII is subjected, in the presence of a palladium catalyst, of a ligand and of a base,

- 15 - to a reaction with phenylboronic or heteroarylboronic or phenylboronate ester or heteroarylboronate ester derivatives according to a Suzuki coupling,
- or else to an imination reaction with benzophenoneimine, followed by an acid hydrolysis and by a sulphonylation reaction with a sulphonyl chloride of formula R_6SO_2Cl ,
- 20 - or else to a cyanation reaction with zinc cyanide, followed by an acid hydrolysis and by an esterification or a peptide coupling with an amine $R_5R_6NH_2$,

in order to obtain the compound of formula XXIII in which R_1 is as defined above. The compound XXIII is subjected to a deprotection reaction in order to obtain the compounds of formula I in which R_c is a hydrogen atom.

- 5 In the preceding schemes, the starting compounds and the reactants, when the method for preparing them is not described, are commercially available or described in the literature, or else can be prepared according to methods which are described therein or which are known to those skilled in the art.
- 10 A subject of the invention, according to another of its aspects, is also the compounds of formulae II to XXIII defined above. These compounds are of use as synthesis intermediates for the compounds of formula (I).

The following examples describe the preparation of certain compounds in
15 accordance with the invention. These examples are not limiting and merely illustrate the present invention. The numbers of the compounds exemplified refer back to those given in the table hereinafter, which illustrates the chemical structures and the physical properties of some compounds according to the invention.

20

The reactants and intermediates, when their preparation is not explained, are known in the literature or commercially available. Some intermediates that are of use for preparing the compounds of formula I may also serve as final products of formula (I), as will become apparent in the examples given hereinafter. Similarly,
25 some compounds of formula (I) of the invention can serve as intermediates that are of use for preparing other compounds of formula (I) according to the invention.

By way of example, the derivatives of formula (I) are selected from the following compounds:

- 30 6-(imidazo[1,5-a]pyridin-3-ylcarbonyl)-3-propylquinazoline-2,4(1*H*,3*H*)-dione,
3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid,
3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridine-6-carboxylic acid,

- 3-(3-([3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl)imidazo[1,5-a]pyridin-1-yl)benzoic acid,
 3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzamide,
 5 6-({1-[3-(5-methyl-1,3,4-oxadiazol-2-yl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl-3-propylquinazoline-2,4(1*H*, 3*H*)-dione,
 6-({1-[3-(3-methyl-1,2,4-oxadiazol-5-yl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)-3-propylquinazoline-2,4(1*H*, 3*H*)-dione,
 N-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}methanesulphonamide,
 10 2-morpholin-4-yl-ethyl 3-(3-([3-(4-fluorobenzyl)-1-méthyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl)imidazo[1,5-a]pyridin-1-yl)benzoate,
 N-[2-(dimethylamino)ethyl]-3-(3-([3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl)imidazo[1,5-a]pyridin-1-yl)benzamide,
 15 3-(3-([3-(4-fluorobenzyl)-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl)imidazo[1,5-a]pyridin-1-yl)benzoic acid,
 3-(4-fluorobenzyl)-1-methyl-6-[(1-pyridin-3-yl)imidazo[1,5-a]pyridin-3-yl]carbonyl]quinazoline-2,4(1*H*, 3*H*)-dione,
 3-{3-[(2-methyl-4-oxo-3-propyl-3,4-dihydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid,
 20 a]pyridin-1-yl}benzamide,
 3-{3-[(2-methyl-4-oxo-3-propyl-3,4-dihydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzamide,
 6-(imidazo[1,5-a]pyridin-3-ylcarbonyl)quinazolin-4(3*H*)-one,
 N,N,1,2-tetramethyl-4-oxo-6-[[1-(pyridin-3-yl)imidazo[1,5-a]pyridin-3-yl]carbonyl]-
 25 1,4-dihydroquinoline-3-carboxamide,
 3-[3-({3-[2-(4-fluorophenoxy)ethyl]-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl}carbonyl)imidazo[1,5-a]pyridin-1-yl]benzoic acid.

Abbreviations

- 30 TOTU: O-[(ethoxycarbonyl)cyanomethyleneamino]-N,N,N',N'-tetramethyluronium tetrafluoroborate
 NMP: N-Methylpyrrolidone
 DME: Ethylene glycol dimethyl ether
 DMF: Dimethylformamide
 35 THF: Tetrahydrofuran

Binap: 2,2'-bis(diphenylphosphino)-1,1'-binaphthyl

The NMR analyses were carried out on Bruker Avance 250MHz, 300MHz and 400MHz instruments.

- 5 - The melting points were measured on a Buchi B-450 instrument.
- The mass spectrometry analyses were carried out on a Waters Alliance 2695 (UV: PDA996, MS: LCZ), Alliance 2695 (UV: PDA 996, MS: ZQ (simple Quad) ZQ1), Alliance 2695 (UV: PDA 996, MS: ZQ (simple Quad) ZQ2), Waters UPLC Acquity (UV: Acquity PDA, MS: SQD (simple Quad) SQW), Agilent MSD, Waters
10 ZQ, or Waters SQD instrument.

Example 1: (compound No. 1)

6-(Imidazo[1,5-a]pyridin-3-ylcarbonyl)-3-propylquinazoline-2,4(1*H*,3*H*)-dione

Methyl 2-amino-5-(imidazo[1,5-a]pyridin-3-ylcarbonyl)benzoate

- 15 13.4 ml (96 mmol) of triethylamine are added to 3.5 g (30 mmol) of imidazo[1,5-a]pyridine [described in J. Chem. Soc.; (1955), 2834-2836] in 250 ml of 1,2-dichloroethane, followed, under a nitrogen atmosphere at 0°C, by 13.7 g (48 mmol) of 4-oxo-2-phenyl-4*H*-3,1-benzoxazine-6-carbonyl chloride (described in WO 05/028476). After stirring for 4.5 hours at ambient temperature, the reaction
20 medium is filtered. The residue obtained is washed with 1,2-dichloroethane. After drying overnight at 40°C under reduced pressure, 3 g of a yellow solid are obtained.

- The residue obtained is dissolved in 100 ml of NMP. A solution of 8.4 g (0.15 mol) of KOH in 10 ml of water is added dropwise, under a nitrogen
25 atmosphere, at ambient temperature. The reaction medium is heated at 80°C for 6 hours and then poured, at ambient temperature, into a 1N aqueous solution of hydrochloric acid. The precipitate obtained is filtered off, rinsed with water and then dried at 40°C under reduced pressure overnight. After silica gel column chromatography, elution being carried out with a dichloromethane/methanol/0.1%
30 triethylamine mixture, 5.5 g of a yellow solid are obtained.

- 7 g (0.022 mol) of caesium carbonate and then, dropwise, 1.34 ml (0.022 mol) of methyl iodide are added, under a nitrogen atmosphere at ambient
temperature, to 5.5 g (0.02 mol) of the residue obtained, in 100 ml of DMF. After stirring for 24 hours at ambient temperature, the reaction medium is poured into
35 water. The precipitate obtained is filtered off, rinsed with water and then dried overnight at 40°C under reduced pressure. 5.1 g of a yellow solid are obtained.

Melting point: 192°C

MH+: 296

Methyl

5-(imidazo[1,5-a]pyridin-3-ylcarbonyl)-2-

5 [(propylcarbamoyl)amino]benzoate

0.35 g (1.2 mmol) of triphosgene is added, at ambient temperature under a nitrogen atmosphere, to a suspension of 0.5 g (1.7 mmol) of methyl 2-amino-5-(imidazo[1,5-a]pyridin-3-ylcarbonyl)benzoate in 20 ml of anhydrous dioxane. After heating for 2 hours at 100°C, 0.28 ml (3.4 mmol) of n-propylamine and then 0.71 ml (5 mmol) of triethylamine are added to the reaction medium at ambient temperature. After stirring for 18 hours at ambient temperature, H₂O is added. The aqueous phase is extracted with dichloromethane. The organic phase is dried over sodium sulphate, filtered, and concentrated under reduced pressure. The yellow solid obtained is purified by silica gel column chromatography, elution being carried out with a dichloromethane/methanol (98/2) mixture. 0.410 g of a yellow solid is obtained.

Melting point: 205°C

MH+: 381

20 6-(Imidazo[1,5-a]pyridin-3-ylcarbonyl)-3-propylquinazoline-2,4(1H,3H)-dione

1.38 ml (1.38 mmol) of a 1N aqueous solution of sodium hydroxide are added, at ambient temperature, to a suspension of 0.436 g (1.15 mmol) of methyl 5-(imidazo[1,5-a]pyridin-3-ylcarbonyl)-2-[(propylcarbamoyl)amino]benzoate in 10 ml of methanol. After refluxing for 2 hours, the methanol is concentrated under reduced pressure. A 1N aqueous solution of hydrochloric acid is added. The precipitate obtained is filtered off, rinsed with water and then dried overnight at 40°C under reduced pressure. 0.27 g of a yellow solid is obtained.

Melting point: 304°C

¹H-NMR (D₆-DMSO, 400 MHz):

0.91 (t, J=7.17Hz, 3H), 1.63 (q, J=7.59Hz, 2H), 3.89 (t, J=7.17Hz, 2H), 7.25-7.37 (m, 2H), 7.39-7.43 (m, 1H), 7.82 (s, 1H), 7.97 (d, J=8.86Hz, 1H), 8.59 (d, J=8.86Hz, 1H), 9.18 (s, 1H), 9.74 (d, J=7.17Hz, 1H), 11.8 (s, 1H).

Example 2: (compound No. 10)

Sodium salt of 3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid

5 **Methyl 2-amino-5-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]benzoate**

0.42 g (2.4 mmol) of N-bromosuccinimide is added, under a nitrogen atmosphere at ambient temperature, to a solution of 0.67 g (2.4 mmol) of methyl 2-amino-5-(imidazo[1,5-a]pyridin-3-ylcarbonyl)benzoate in 20 ml of dichloromethane.
10 After stirring for 2 h 30, water is added. The precipitate formed is filtered off, rinsed with water, and dried overnight at 40°C under reduced pressure. 0.77 g of a yellow solid is obtained.

Melting point: 230°C

MH+: 375, 377

15 **Methyl 2-amino-5-({1-[3-(methoxycarbonyl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)benzoate**

0.248 g (1.38 mmol) of [4-(methoxycarbonyl)phenyl]boronic acid, 0.57 g (2.30 mmol) of potassium carbonate in 2 ml of water, and 0.027 g (0.02 mmol) of tetrakis(triphenylphosphine)palladium are added to a solution of 0.43 g (1.15 mmol) of methyl 2-amino-5-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]benzoate in 10 ml of DME, under an inert argon atmosphere. The reaction medium is heated at 90°C for 2 hours. The reaction medium is acidified with a 1N aqueous solution of hydrochloric acid, and extracted with dichloromethane. The organic phase is
25 washed with water, dried over sodium sulphate, filtered, and concentrated under reduced pressure. The solid obtained is dissolved in 5 ml of DMF. 30 µl (0.5 mmol) of methyl iodide and 0.052 g (0.16 mmol) of caesium carbonate are added. After stirring for 24 hours at ambient temperature, the reaction medium is hydrolysed with water and then extracted with ethyl acetate. The organic phase is dried over
30 sodium sulphate, filtered, and then concentrated under reduced pressure. The solid obtained is taken up in methanol. After filtration and drying overnight at 50°C under reduced pressure, 0.379 g of a yellow powder is obtained.

Melting point: 203°C

MH+: 430

35

Methyl 5-({1-[3-(methoxycarbonyl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)-2-[(propylcarbamoyl)amino]benzoate

0.181 g (0.61 mmol) of triphosgene is added, under an inert atmosphere, to
5 0.75 g (0.87 mmol) of methyl 2-amino-5-({1-[3-(methoxycarbonyl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)benzoate in 10 ml of dioxane. The reaction medium is heated at 100°C for 3 hours. 0.14 ml (1.75 mmol) of propylamine and 0.37 ml (2.62 mmol) of triethylamine are added at ambient temperature. After stirring for 2 hours at ambient temperature, the reaction medium
10 is hydrolysed with water. The medium is filtered, washed with water, and dried under reduced pressure at 50°C overnight. The solid obtained is purified by silica gel column chromatography with a dichloromethane/methanol (95/5) mixture. 0.27 g of a yellow powder is obtained.

Melting point: 212°C

15 MH+: 515

3-{3-[(2,4-Dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid

1.31 ml (1.31 mmol) of a 1N aqueous solution of sodium hydroxide are
20 added to 0.27 g (0.52 mmol) of methyl 5-({1-[3-(methoxycarbonyl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)-2-[(propylcarbamoyl)amino]benzoate in 8 ml of methanol. The reaction medium is heated at 70°C for 5.5 hours. The methanol is concentrated under reduced pressure. The residue is taken up in water. The aqueous phase is acidified with a
25 1N aqueous solution of hydrochloric acid, and then extracted with dichloromethane. The organic phase is dried over sodium sulphate, filtered, and then concentrated under reduced pressure. The solid obtained is taken up in methanol and then filtered, and dried at 50°C under reduced pressure overnight. 0.245 g of a yellow solid is obtained.

30 Melting point: 365°C

MH+: 469

Sodium salt of 3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid

0.51 ml (0.51 mmol) of a 1N aqueous solution of sodium hydroxide is added to 0.245 g (0.52 mmol) of 3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid in 5 ml of methanol. The reaction medium is stirred for 1.5 hours at ambient temperature. After the addition
5 of diisopropyl ether, the precipitate formed is filtered off, rinsed with diisopropyl ether, and dried at 50°C under reduced pressure overnight. 0.242 g of a yellow powder is obtained.

Melting point: 383°C

MH+: 469

10 ¹H NMR (D6-DMSO, 400 MHz):

0.90 (t, J=7.82Hz, 3H), 1.58-1.67 (m, 2H), 3.88 (t, J=7.07Hz, 1H), 7.32-7.35 (m, 2H), 7.45 (t, J=7.82, 1H), 7.53 (t, J=7.82Hz, 1H), 7.88-7.94 (m, 2H), 8.22 (d, J=8.94Hz, 1H), 8.44 (t, J=1.7Hz, 1H), 8.74 (d, J=8.7Hz, 1H), 9.14 (d, J=1.9Hz, 1H), 9.82 (d, J=7Hz, 1H), 11.9 (bs, 1H).

15

Example 3 (compound No. 8)

3-[(2,4-Dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridine-6-carboxylic acid

20 **3-[(4-Amino-3-carboxyphenyl)carbonyl]imidazo[1,5-a]pyridine-6-carboxylic acid**

3.68 ml (0.026 mol) of triethylamine and then, under a nitrogen atmosphere at ambient temperature, 1.5 g (8.5 mmol) of methyl imidazo[1,5-a]pyridine-6-carboxylate [described in WO 06/097625] are added to 4.02 g (0.014 mol) of 4-
25 oxo-2-phenyl-4H-3,1-benzoxazin-6-carbonyl chloride in 60 ml of 1,2-dichloroethane. After stirring for 24 hours at ambient temperature, the reaction medium is filtered, and washed with 1,2-dichloroethane, then with a 1N aqueous solution of hydrochloric acid and then with water. After drying overnight under reduced pressure at 40°C, the product obtained is dissolved in 60 ml of NMP.
30 3.59 g (6.4 mmol) of potassium hydroxide dissolved in 11 ml of water are added. The reaction medium is heated at 100°C for 4 hours and then poured into a 1N aqueous solution of hydrochloric acid. After filtration, the solid obtained is rinsed with water and then dried overnight in an oven under reduced pressure at 40°C. 5.45 g of a yellow solid are obtained.

MH+: 326

Methyl 3-[[4-amino-3-(methoxycarbonyl)phenyl]carbonyl]imidazo[1,5-a]pyridine-6-carboxylate

- 5 9.4 g (2.9 mmol) of caesium carbonate and then 1.8 ml (2.9 mmol) of methyl iodide, at ambient temperature are added, under an inert atmosphere, to 4.2 g (1.3 mmol) of 3-[[4-amino-3-carboxyphenyl]carbonyl]imidazo[1,5-a]pyridine-6-carboxylic acid in 60 ml of DMF. After stirring for 4.5 hours at ambient temperature, the reaction medium is hydrolysed with water. The precipitate obtained is filtered
10 off, rinsed with water, and then dried at 40°C under reduced pressure overnight. The solid obtained is purified by silica gel column chromatography, elution being carried out with dichloromethane. 1.3 g of a yellow solid are obtained.

MH+: 354

- 15 **Methyl 3-[[3-(methoxycarbonyl)-4-[(propylcarbamoyl)amino]imidazo[1,5-a]pyridine-6-carboxylate**

- 0.14 g (0.49 mmol) of triphosgene is added, at ambient temperature under a nitrogen atmosphere, to 0.3 g (0.7 mmol) of methyl 3-[[4-amino-3-(methoxycarbonyl)phenyl]carbonyl]imidazo[1,5-a]pyridine-6-carboxylate in 10 ml of
20 anhydrous dioxane. After heating for 1 h 15 at 100°C, 0.12 ml (1.4 mmol) of n-propylamine and 0.29 ml (2 mmol) of triethylamine are added to the reaction medium at ambient temperature. After stirring for 4 hours at ambient temperature, the reaction medium is hydrolysed with water. The precipitate obtained is filtered off, rinsed with water, and then dried under reduced pressure at 40°C overnight.
25 The solid obtained is triturated from THF and then filtered and dried under reduced pressure at 40°C overnight. 0.21 g of a yellow solid is obtained.

Melting point: 266°C

MH+: 439

- 30 **3-[(2,4-Dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridine-6-carboxylic acid**

1.2 ml (1.2 mmol) of a 1N aqueous solution of sodium hydroxide are added, at ambient temperature, to 0.21 g of methyl 3-[[3-(methoxycarbonyl)-4-[(propylcarbamoyl)amino]imidazo[1,5-a]pyridine-6-carboxylate

in 5 ml of methanol. After refluxing for 4 hours, the reaction medium is acidified with a 1N aqueous solution of hydrochloric acid. The precipitate obtained is filtered off and then rinsed with water and dried under reduced pressure at 40°C overnight. The solid obtained is recrystallized while hot from methanol and then dried under
5 reduced pressure at 40°C overnight. 0.118 g of a yellow solid is obtained.

Melting point: 384°C

MH+: 393

¹H-NMR (D6-DMSO, 400 MHz) :

0.92 (t, J=7.2Hz, 3H), 1.59-1.68 (m, 2H), 3.87-3.94 (m, 2H), 7.33 (d, J=8.2Hz, 1H),
10 7.72 (d, J=9.3Hz, 1H), 7.98 (s, 1H), 8.06 (d, J=9.3Hz, 1H), 8.59 (d, J=8.51Hz, 1H),
9.20(d, J=2.03Hz, 1H), 11.8 (s, 1H), 13.7 (s, 1H).

Example 4: (compound No. 49)

Sodium salt of 3-(3-[[3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-
15 tetrahydroquinazolin-6-yl]carbonyl]imidazo[1,5-a]pyridin-1-yl)benzoic acid

Methyl 2-[[[(4-fluorobenzyl)carbamoyl]amino]-5-({1-[3-(methoxycarbonyl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)benzoate

2.14 g (7.2 mmol) of triphosgene are added, at ambient temperature under
20 an inert atmosphere, to 2.58 g (6 mmol) of methyl 2-amino-5-({1-[3-(methoxycarbonyl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)benzoate in 50 ml of dioxane. After refluxing for 7 hours, 2.25 g (18 mmol) of 4-fluorobenzylamine and 1.82 g (18 mmol) of triethylamine are added at ambient temperature. The reaction medium is refluxed for 3 hours and then concentrated under reduced pressure.
25 The residue is triturated from water. After filtration, the solid is rinsed with methanol and then dried under reduced pressure at 40°C overnight. 3.3 g of a yellow solid are obtained.

MH+: 581

30 **3-(3-[[3-(4-Fluorobenzyl)-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl]imidazo[1,5-a]pyridin-1-yl)benzoic acid**

2.85 ml (0.0285 mol) of a 1N aqueous solution of sodium hydroxide are added to 3.3 g (5.7 mmol) of methyl 2-[[[(4-fluorobenzyl)carbamoyl]amino]-5-({1-[3-(methoxycarbonyl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)benzoate dissolved in

250 ml of methanol. After refluxing for 2 hours, the reaction medium is acidified with 50 ml of a 1N aqueous solution of hydrochloric acid and then diluted with 700 ml of water. The precipitate obtained is filtered off, and dried under reduced pressure at 40°C overnight. 3.01 g of a yellow solid are obtained.

5 MH+: 535

Methyl 3-(3-{[3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoate

2.44 g (7.5 mmol) of caesium carbonate and 1.06 g (7.5 mmol) of methyl iodide are added, under an inert atmosphere, to 1.3 g (2.5 mmol) of 3-(3-{[3-(4-fluorobenzyl)-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoic acid in 50 ml of DMF. The reaction medium is stirred for 3 hours at ambient temperature under a nitrogen atmosphere and then concentrated under reduced pressure. The residue obtained is washed with 200 ml of water and
10 then dried under reduced pressure at 40°C overnight. 1.35 g of a yellow solid are obtained.

MH+: 563

Sodium salt of 3-(3-{[3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoic acid

24 ml (24 mmol) of a 1N aqueous solution of lithium hydroxide are added to 1.3 g (2.4 mmol) of methyl 3-(3-{[3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoate in 120 ml of THF. The reaction medium is refluxed for 5 hours and then acidified at 5°C with
25 45 ml of a 1N aqueous solution of hydrochloric acid and, finally, diluted with 200 ml of water. After filtration, the residue obtained is dried under reduced pressure at 40°C overnight.

0.62 ml (0.62 mmol) of a 1N aqueous solution of sodium hydroxide is added to 0.35 g (0.64 mmol) of the yellow solid obtained, in 20 ml of methanol. After
30 filtration, the residue obtained is dried under reduced pressure at 40°C overnight. 0.38 g of a yellow solid is obtained.

MH+: 549

¹H-NMR (D6-DMSO, 500 MHz):

3.62 (s, 3 H), 5.17 (s, 2 H), 7.11-7.18 (ps t, J = 8.9 Hz, 2 H), 7.35 – 7.40 (ps t, 8.9

Hz, 1 H), 7.42 – 7.48 (m, 3 H), 7.54 – 7.60 (ps t, J = 8.9 Hz, 1 H), 7.70 – 7.74 (ps d, J = 8.9 Hz, 1 H), 7.89 – 7.95 (ps t, J = 8.9 Hz, 2 H), 8.26 – 8.30 (ps d, J = 8.9 Hz, 1 H), 8.44 – 8.48 (m, 1 H), 8.96 – 9.01 (ps d, J = 8.9 Hz, 1 H), 9.22 – 9.24 (m, 1 H), 9.88 – 9.91 (ps d, J = 7.2 Hz, 1 H).

5

Example 5: (compound No. 29)

3-{3-[(2,4-Dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzamide

10.7 mg (0.2 mmol) of ammonium chloride, 5.17 mg (0.4 mmol) of N,N-diisopropylethylamine and 49.2 mg (0.2 mmol) of TOTU are added, at 0°C, under an inert atmosphere, to 46.8 mg (0.1 mmol) of 3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid in 2 ml of DMF. The reaction medium is stirred for 12 hours at ambient temperature and then poured into 30 ml of a saturated solution of sodium hydrogen carbonate. The precipitate obtained is filtered off, washed with water, and dried under reduced pressure at 40°C overnight. 0.042 g of a yellow solid is obtained.

MH+: 468

¹H-NMR (D6-DMSO, 500 MHz):

δ = 0.92 (t, 3 H, J = 7.7 Hz), 1.66 (tq, 2 H, J = 7.7 Hz, 7.3 Hz), 3.94 (t, 2 H, J = 7.3 Hz), 7.34-7.42 (2 m, 2 H), 7.52-7.61 (2m, 2 H), 7.69 (t, 1 H, J = 7.6 Hz), 7.96 (m, 1 H), 8.10 -8.23 (2 m, 2 H), 8.41- 8.46 (m, 2 H), 8.80 (dd, 1 H, J = 8.9 Hz, 2.2 Hz), 9.27 (d, 1 H, 1.9 Hz), 9.88 (d, 1 H, J = 7.1 Hz), 11.83 (s, 1 H).

Example 6: (compound No. 34)

6-({1-[3-(5-Methyl-1,3,4-oxadiazol-2-yl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl-3-propylquinazoline-2,4(1H, 3H)-dione

N'-Acetyl-3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzohydrazide

29.6 mg (0.4 mmol) of acetohydrazide, 98.4 mg (0.3 mmol) of TOTU and 0.104 ml (0.6 mmol) of N,N-diisopropylethylamine are added, under an inert atmosphere, at 0°C, to 93.7 mg (0.2 mmol) of 3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid in 6 ml of DMF. The reaction medium is stirred for 1 hour at 0°C and then for 6 hours at 50°C

and then concentrated under reduced pressure. The residue is taken up in 10 ml of methanol. The precipitate obtained is filtered off, washed with diethyl ether and with pentane, and then dried under reduced pressure at 40°C overnight. 45 mg of a yellow solid are obtained.

5 MH+: 525

6-({1-[3-(5-Methyl-1,3,4-oxadiazol-2-yl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl-3-propylquinazoline-2,4(1H, 3H)-dione

35 mg (0.066 mmol) of N'-acetyl-3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-
10 tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzohydrazide in
1 ml of phosphorus oxychloride are heated at 100°C for 15 minutes. The reaction
medium is concentrated under reduced pressure. The residue obtained is
hydrolysed with water and with a saturated solution of sodium hydrogen carbonate.
The aqueous phase is extracted with dichloromethane. The organic phase is
15 concentrated under reduced pressure. The residue obtained is purified by silica gel
column chromatography, elution being carried out with methanol. 0.025 g of a
yellow solid is obtained.

 MH+: 507

¹H-NMR (D6-DMSO, 500MHz):

20 0.91 (t, J = 7.5 Hz, 3H), 1.65 (qt, J = 7.5 Hz, 7.5 Hz, 2 H), 2.67 (s, 3 H), 3.93 (t, J =
7.5 Hz, 2 H), 7.33 – 7.43 (m, 2 H), 7.58 – 7.64 (m, 1H), 7.77 – 7.84 (m, 1 H), 8.04 –
8.06 (m, 1 H), 8.28 – 8.32 (m, 1H), 8.39 – 8.43 (m, 1 H), 8.59 (s, 1 H), 8.71 – 8.74
(m, 1 H), 9.37 (s, 1 H), 9.86 – 9.90 (s, 1 H), 11.85 (br s, 1H).

25 **Example 7: (compound No. 36)**

6-({1-[3-(3-Methyl-1,2,4-oxadiazol-5-yl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)-3-propylquinazoline-2,4(1H, 3H)-dione

30 **3-{3-[(2,4-Dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}-N-[(1E)-hydroxyethanimidoyl]benzamide**

39 mg (0.24 mmol) of 1,1'-carbonyldiimidazole are added, at ambient temperature under an inert atmosphere, to 94 mg (0.2 mmol) of 3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid in

5 ml of DMF. After stirring for 12 hours at ambient temperature, 22.2 mg (0.3 mmol) of acetamidoxime are added. The reaction medium is stirred for 5 hours at ambient temperature and then concentrated under reduced pressure. The residue is triturated from diethyl ether, filtered, and then dried under reduced pressure at
 5 40°C overnight. 0.101 g of a yellow solid is obtained.

MH+: 525

6-{{1-[3-(3-Methyl-1,2,4-oxadiazol-5-yl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl}-3-propylquinazoline-2,4(1*H*, 3*H*)-dione

10 A solution of 0.1 g (0.19 mmol) of 3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}-N-[(1*E*)-hydroxyethanimidoyl]benzamide in 3 ml of DMF is heated at 120°C for 5 hours. The reaction medium is concentrated under reduced pressure. The residue obtained is taken up in diethyl ether, filtered, and then dried under reduced
 15 pressure at 40°C overnight. 0.083 g of a yellow solid is obtained.

MH+: 507

¹H-NMR (D6-DMSO):

0.91 (t, J = 7.5 Hz, 3H), 1.65 (qt, J = 7.5 Hz, 7.5 Hz, 2 H), 2.47 (s, 3 H), 3.94 (t, J = 7.5 Hz, 2 H), 7.36 – 7.45 (m, 2 H), 7.59 – 7.66 (m, 1H), 7.82 – 7.89 (m, 1H), 8.13 –
 20 8.19 (m, 1H), 8.36 – 8.45 (m, 2H), 8.68 (s, 1H), 8.75 – 8.79 (m, 1H), 9.25. 9.28 (m, 1 H), 9.85 – 9.90 (m, 1 H), 11.85 (br s, 1 H).

Example 8: (Compound No. 13)

N-{3-[(2,4-Dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}methanesulphonamide
 25

Methyl 5-[(1-bromoimidazo(1,5-a)pyridin-3-yl)carbonyl]-2-[(propylcarbamoyl)aminobenzoate

0.55 g (0.0019 mol) of triphosgene is added, at ambient temperature under
 30 an inert atmosphere, to 1 g (2.7 mmol) of methyl 2-amino-5-[1-bromo(imidazo[1,5-a]pyridin-3-yl)carbonyl]benzoate in 30 ml of anhydrous dioxane. The reaction medium is heated for 1.5 hours at 100°C. 0.44 ml (5.3 mmol) of n-propylamine and 1.12 ml (8 mmol) of triethylamine are added at ambient temperature. After 2 h 30, the reaction medium is hydrolysed with water. The aqueous phase is extracted with

dichloromethane. The organic phase is dried over sodium sulphate, filtered, and then concentrated under reduced pressure. The solid obtained is triturated from dichloromethane, filtered, and then dried under reduced pressure at 40°C overnight.

- 5 MH+: 459, 461
 Melting point: 236°C

6-[(1-Bromoimidazo(1,5-a)pyridin-3-yl)carbonyl]-3-propylquinazoline-2,4(1H,3H)-dione

- 10 3.14 ml (3.1 mmol) of a 1N aqueous solution of sodium hydroxide are added, at ambient temperature, to 1.2 g (2.6 mmol) of methyl 5-[(1-bromoimidazo(1,5-a)pyridin-3-yl)carbonyl]-2-[(propylcarbamoyl)aminobenzoate in 20 ml of methanol. After refluxing for 3 hours, the reaction medium is hydrolysed with a 1N aqueous solution of hydrochloric acid. The precipitate obtained is filtered off, rinsed with
15 methanol, and then dried under reduced pressure at 40°C overnight. 1.09 g of a yellow solid are obtained.

 MH+: 427, 429
 Melting point: 322°C

- 20 **6-[(1-Aminoimidazo(1,5-a)pyridin-3-yl)carbonyl]-3-propylquinazoline-2,4(1H,3H)-dione**

- 1.45 g (4.7 mmol) of caesium carbonate, 1.13 ml (6.7 mmol) of benzophenoneimine, 0.278 g (0.45 mmol) of binap and 0.204 g (0.22 mmol) of dibenzylideneacetone dipalladium are added, at ambient temperature under an
25 argon atmosphere, to 0.955 g (2 mmol) of 6-[(1-bromoimidazo(1,5-a)pyridin-3-yl)carbonyl]-3-propylquinazoline-2,4(1H,3H)-dione in 20 ml of DMSO. The reaction medium is heated at 110°C for 18 hours. The reaction medium is extracted with ethyl acetate. The organic phase is dried over sodium sulphate, filtered and concentrated under reduced pressure.

- 30 The residue obtained is dissolved in 40 ml of THF. 4.5 ml (9 mmol) of a 2N aqueous solution of hydrochloric acid are added at ambient temperature. After stirring for 4 hours at ambient temperature, the reaction medium is concentrated under reduced pressure. The residue obtained is washed with dichloromethane and with methanol, and then dried under reduced pressure at 40°C overnight.

0.558 g of a red solid is obtained.

MH+: 364

N-{3-[(2,4-Dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}methanesulphonamide

0.1 ml (1.2 mmol) of mesyl chloride is added, at 0°C under an inert atmosphere, to 0.25 g (0.4 mmol) of 6-[(1-aminoimidazo(1,5-a)pyridin-3-yl)carbonyl]-3-propylquinazoline-2,4(1*H*,3*H*)-dione in 5 ml of pyridine. After the addition of methanol, the reaction medium is concentrated under reduced pressure.

The residue is taken up with dichloromethane. The organic phase is washed with a 1N aqueous solution of hydrochloric acid and then with water, dried over sodium sulphate, filtered, and concentrated under reduced pressure. The residue is recrystallized while hot from methanol, purified on a silica gel frit, elution being carried out with DMF. 0.057 g of an orange solid is obtained.

Melting point: 334°C

MH+: 442

¹H-NMR (D6-DMSO, 400 MHz):

0.88 (t, J=7.37Hz, 3H), 1.55-1.65 (m, 2H), 3.29 (s, 3H), 3.87-3.90 (m, 2H), 7.27-7.31 (m, 2H), 7.40-7.44 (m, 1H), 7.92 (d, J=9Hz, 1H), 8.52 (d, J=8.46Hz, 1H), 9.15 (d, J=2.18Hz, 1H), 9.71 (d, J=7.1Hz, 1H), 10.2 (s, 1H), 11.8 (s, 1H).

Example 9: (Compound No. 82)

2-Morpholin-4-yl-ethyl 3-(3-{[3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoate

hydrochloride

0.022 g (0.61 mmol) of 4-(2-chloroethyl)morpholine hydrochloride and 0.189 g (1.37 mmol) of potassium carbonate are added, under an inert atmosphere, to 0.3 g (0.55 mmol) of 3-(3-{[3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoic acid in 8 ml of DMF. After stirring for 18 h at ambient temperature and then for 8 hours at 50°C, the reaction medium is hydrolysed with water, and extracted with ethyl acetate. The organic phase is washed with water, dried over sodium sulphate, filtered, and then concentrated under reduced pressure. The yellow solid obtained is purified by silica gel column chromatography, elution being carried out with a

dichloromethane/methanol (95/5) mixture. 0.61 ml of a 1N aqueous solution of hydrochloric acid is added to 0.334 g of the yellow solid obtained, in 5 ml of methanol. The reaction medium is stirred for 1 hour at ambient temperature. Diethyl ether is added and then the reaction medium is filtered. The precipitate
5 obtained is rinsed with diethyl ether and then dried under reduced pressure at 50°C overnight. 0.298 g of a yellow solid is obtained.

Melting point: 215°C

MH⁺: 662

10 ¹H-NMR (D6-DMSO, 500 MHz):

3.21-3.31 (m, 2H), 3.31 (s, 3H), 3.46-3.54 (m, 2H), 3.6-3.7 (m, 2H), 3.61 (s, 3H),
3.70-3.80 (m, 2H), 3.90-4 (m, 2H), 4.65-4.75 (m, 2H), 5.16 (s, 2H), 7.11-7.16 (m,
2H), 7.37-7.39 (m, 1H), 7.42-7.45 (m, 2H), 7.55-7.58 (m, 1H), 7.67 (d, J=9.28Hz,
1H), 7.73 (t, J=7.69Hz, 1H), 8.07 (d, J=7.69Hz, 1H), 8.29-8.34 (m, 2H), 8.55 (s,
15 1H), 8.82 (d, J=9.01Hz, 1H), 9.27 (d, J=1.85Hz, 1H), 9.83 (d, J=7.16Hz, 1H), 10.9
(s, 1H).

Example 10: (Compound No. 117)

**N-[2-(Dimethylamino)ethyl]-3-(3-[[3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-
20 tetrahydroquinazolin-6-yl]carbonyl]imidazo[1,5-a]pyridin-1-yl]benzamide
hydrochloride**

0.06 ml (0.55 mmol) of N,N-dimethylethylenediamine, 0.134 g (0.41 mmol) of
TOTU and 0.14 ml (0.82 mmol) of diisopropylethylamine are added to 0.15 g
(0.27 mmol) of 3-(3-[[3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-
25 tetrahydroquinazolin-6-yl]carbonyl]imidazo[1,5-a]pyridin-1-yl]benzoic acid in 5 ml of
DMF. The reaction medium is heated at 80°C for 16 hours. The reaction medium is
hydrolysed with water, and extracted with ethyl acetate. The organic phase is
washed with water, dried over sodium sulphate, filtered, and concentrated under
reduced pressure. The yellow solid obtained is purified by silica gel column
30 chromatography, elution being carried out with a dichloromethane/methanol (95/5)
mixture. 0.23 ml of a 1N solution of hydrochloric acid in diethyl ether is added to
0.095 g of the yellow solid obtained. After stirring for 1 hour, diethyl ether is added.
The precipitate obtained is filtered off, rinsed with water, and then dried under
reduced pressure at 50°C overnight. 0.1 g of a yellow solid is obtained.

Melting point: 247°C

MH+: 619

¹H-NMR (D6-DMSO, 400 MHz):

2.50(m, 6H), 2.84 (s, 2H), 3.31 (s, 3H), 3.61(s, 1H), 3.64-6.70 (m, 1H), 5.16 (s, 2H), 7.7.11-7.17 (m, 2H), 7.37-7.46 (m, 3H), 7.55-7.60 (m, 1H), 7.67-7.71 (m, 2H), 7.93 (d, J=8.19Hz, 1H), 8.19 (d, J=7.51Hz, 1H), 8.38-8.43 (m, 2H), 8.87 (d, J=8.88Hz, 1H), 8.92 (t, J=5.12Hz, 1H), 9.27 (d, J=2Hz, 1H), 9.81 (s, 1H), 9.84 (d, J=7.1Hz, 1H).

10 **Example 11: (Compound No. 72)**

Sodium salt of 3-(3-{[3-(4-fluorobenzyl)-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoic acid

15 **Propyl 3-(3-{[3-(4-fluorobenzyl)-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoate**

1.371 g (4.21 mmol) of caesium carbonate and 0.715 g (4.21 mmol) of propyl iodide are added, under an inert atmosphere, to 0.75 g (1.4 mmol) of 3-(3-{[3-(4-fluorobenzyl)-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoic acid in 30 ml of DMF. The reaction medium is stirred for 20 3 hours at ambient temperature under a nitrogen atmosphere, and then concentrated under reduced pressure. The residue obtained is washed with 100 ml of water and then dried under reduced pressure at 40°C overnight. The solid obtained is purified by silica gel column chromatography, elution being carried out with a dichloromethane/methanol (75/1) mixture. 0.55 g of a yellow solid is 25 obtained.

MH+: 619

30 **Sodium salt of 3-(3-{[3-(4-fluorobenzyl)-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoic acid**

8.9 ml (8.9 mmol) of a 1N aqueous solution of lithium hydroxide are added to 0.55 g (0.889 mmol) of propyl 3-(3-{[3-(4-fluorobenzyl)-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoate in 50 ml of THF. The reaction medium is refluxed for 6 hours and then acidified at 5°C with 17 ml of a 1N aqueous solution of hydrochloric acid and, finally, diluted with 100 ml

of water. After filtration, the residue obtained is dried under reduced pressure at 40°C overnight.

0.408 ml (0.408 mmol) of a 1N aqueous solution of sodium hydroxide is added to 0.24 g (0.416 mmol) of the yellow solid obtained, in 20 ml of methanol. After
5 filtration, the residue obtained is dried under reduced pressure at 40°C overnight. 0.24 g of a yellow solid is obtained.

MH⁺: 577

¹H-NMR (D6-DMSO, 500 MHz):

0.97 (t, J = 7.5 Hz, 3H), 1.71 (tq, J₁/J₂ = 7.5 Hz, 2 H), 4.18 (t, J = 7.5 Hz, 2 H), 5.20
10 (s, 2 H), 7.17 (ps t, J = 9.3 Hz, 2 H), 7.37 – 7.41 (m, 1 H), 7.44 – 7.49 (3 m, 3 H), 7.59 (m, 1 H), 7.78 (ps d, J = 8.5 Hz, 1 H), 7.91 (2 m, 2 H), 8.28 (ps d, J = 9.8 Hz, 1 H), 8.45 (m, 1 H), 8.99 – 9.02 (m, 1 H), 9.23 (m, 1 H), 9.90 (ps d, J = 7.5 Hz, 1 H).

Example 12: (Compound No. 113)

15 **3-(4-Fluorobenzyl)-1-methyl-6-[(1-pyridin-3-ylimidazo[1,5-a]pyridin-3-yl)carbonyl]quinazoline-2,4(1H, 3H)-dione**

Methyl 5-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-2-[(4-fluorobenzyl)carbamoyl]amino}benzoate

20 3 g (10.4 mmol) of triphosgene diluted in 40 ml of dioxane are added to 5.57 g (14.9 mmol) of methyl 2-amino-5-[1-bromo(imidazo[1,5-a]pyridin-3-yl)carbonyl]benzoate in 160 ml of dioxane, under an inert atmosphere. The reaction medium is refluxed for 1 hour. 3.7 g (0.030 mol) of 4-fluorobenzylamine and 6.22 ml (0.045 mol) of triethylamine are added at ambient temperature. The
25 reaction medium is stirred for 4 hours at ambient temperature and then hydrolysed with water. The precipitate obtained is filtered off, rinsed with water, and dried under reduced pressure at 50°C overnight. The solid obtained is taken up with methanol, filtered, rinsed with methanol, and dried under reduced pressure overnight.

30 12 g of a yellow solid are obtained (yield=95.5%).

MH⁺: 525, 527

Melting point: 203°C

6-[(1-Bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-3-(4-fluorobenzyl)quinazoline-2,4(1H, 3H)-dione
35

22.33 ml (22.33 mmol) of a 1N aqueous solution of sodium hydroxide are added to 7.8 g (0.0149 mol) of methyl 5-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-2-[(4-fluorobenzyl)carbamoyl]amino]benzoate in 100 ml of methanol. The reaction medium is refluxed for 2.5 hours. After hydrolysis with water, the precipitate obtained is filtered off, rinsed with water, and dried under reduced pressure at 50°C overnight.

The solid obtained is taken up in a 0.1N aqueous solution of hydrochloric acid, filtered, rinsed with water, and dried under reduced pressure at 50°C overnight. 5.4 g of a yellow solid are obtained.

10 Melting point: 325°C
MH+: 494, 496

6-[(1-Bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-3-(4-fluorobenzyl)-1-methylquinazoline-2,4(1H, 3H)-dione

15 1.87 g (5.7 mmol) of caesium carbonate and 0.39 ml (6.2 mmol) of methyl iodide are added, at ambient temperature under an inert atmosphere, to 2.6 g (5.17 mmol) of 6-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-3-(4-fluorobenzyl)quinazoline-2,4(1H,3H)-dione in 50 ml of anhydrous DMF. The reaction medium is stirred for 18 hours at ambient temperature and then filtered.

20 The precipitate is rinsed with water and then dried under reduced pressure at 50°C overnight. 2.54 g of a yellow solid are obtained.

Melting point: 280°C
MH+: 507, 509

25 **3-(4-Fluorobenzyl)-1-methyl-6-[(1-pyridin-3-ylimidazo[1,5-a]pyridin-3-yl)carbonyl]quinazoline-2,4(1H, 3H)-dione**

0.04 g (0.32 mmol) of 3-pyridylboronic acid, 0.2 g (0.81 mmol) of potassium phosphate dihydrate dissolved in 0.29 ml of water, and 6.2 mg (0.01 mmol) of tetrakis(triphenylphosphine)palladium are added to 0.15 g (0.27 mmol) of 6-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-3-(4-fluorobenzyl)-1-methylquinazoline-2,4(1H, 3H)-dione in 3 ml of DMF under an inert argon atmosphere. The reaction medium is microwave-heated at 150°C for 20 minutes. After filtration over talc, the reaction medium is concentrated under reduced pressure. The residue obtained is purified by silica gel column chromatography, elution being carried out with a

dichloromethane/methanol (95/5) mixture. 0.12 g of a yellow solid is obtained.

Melting point: 207°C

MH+: 506

5 **3-(4-Fluorobenzyl)-1-methyl-6-[(1-pyridin-3-ylimidazo[1,5-a]pyridin-3-yl)carbonyl]quinazoline-2,4(1H, 3H)-dione hydrochloride**

0.35 ml (0.35 mmol) of a 1N solution of hydrochloric acid in diethyl ether is added to 0.12 g (0.23 mmol) of 3-(4-fluorobenzyl)-1-methyl-6-[(1-pyridin-3-ylimidazo[1,5-a]pyridin-3-yl)carbonyl]quinazoline-2,4(1H, 3H)-dione in 3 ml of
10 methanol. After stirring for 1 hour at ambient temperature, the reaction medium is filtered. The precipitate obtained is rinsed with diethyl ether, and dried under reduced pressure at 50°C overnight. 0.12 g of a yellow solid is obtained.

MH+: 506

Melting point: 267°C

15 ¹H-NMR (D6-DMSO, 400 MHz):

3.60 (s, 3H), 5.16 (s, 2H), 7.14 (t, J=8.34Hz, 2H), 7.36-7.47 (m, 3H), 7.60 (t, J=7.05Hz, 1H), 7.65 (d, J=8.98Hz, 1H), 7.83 (t, J=7.05Hz, 1H), 8.43 (d, J=8.98Hz, 1H), 8.66-8.75 (m, 2H), 8.83 (d, J=8.98Hz, 1H), 9.30(m, 2H), 9.81 (d, J=7.05Hz, 1H).

20

Example 13: (Compound No. 53)

3-{3-[(2-Methyl-4-oxo-3-propyl-3,4-dihydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid

2-Amino-5-(1-bromoimidazo[1,5-a]pyridin-3-ylcarbonyl)benzoic acid

25 60 ml (60 mmol) of a 1N aqueous solution of sodium hydroxide are added, at ambient temperature, to 3.74 g (10 mmol) of methyl 2-amino-5-[1-bromo(imidazo[1,5-a]pyridin-3-yl)carbonyl]benzoate in 300 ml of methanol and 125 ml of water. The reaction medium is refluxed for 6 hours and then 140 ml of a 1N aqueous solution of hydrochloric acid are added. After concentration of the
30 methanol under reduced pressure, the precipitate obtained is filtered off, washed with water, and then dried under reduced pressure at 40°C for 18 hours. 3.53 g of a yellow solid are obtained.

MH+: 360, 362

2-(Acetylamino)-5-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]benzoic acid

0.92 g (2.56 mmol) of 2-amino-5-(1-bromoimidazo[1,5-a]pyridin-3-ylcarbonyl)benzoic acid in 30 ml of acetic anhydride are refluxed for 5.5 hours. The reaction medium is concentrated under reduced pressure. The residue is taken up in water and then filtered and dried under reduced pressure overnight at 40°C. 1.1 g of a yellow solid are obtained.

MH+: 402, 404

6-[(1-Bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-2-methyl-3-propylquinazolin-4(3H)-one

1.32 g (22.4 mmol) of n-propylamine are added, at 0°C under an inert atmosphere, to 0.9 g (2.2 mmol) of 6-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-2-methyl-4H-3,1-benzoxazin-4-one in 15 ml of glacial acetic acid. The reaction medium is microwave-heated for 45 minutes at 160°C. The reaction medium is concentrated under reduced pressure. The residue obtained is taken up with a saturated aqueous solution of sodium carbonate. The precipitate obtained is filtered off and then dried under reduced pressure at 50°C overnight. 0.67 g of a yellow solid is obtained.

MH+: 425, 427

Methyl 3-{3-[(2-methyl-4-oxo-3-propyl-3,4-dihydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoate

0.35 g (1.95 mmol) of 3-methoxycarbonylphenylboronic acid, 0.689 g (3.24 mmol) of potassium phosphate dissolved in 3 ml of water, and 0.037 g (0.032 mmol) of tetrakis(triphenylphosphine)palladium are added to 0.69 g (1.62 mmol) of 6-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-2-methyl-3-propylquinazolin-4(3H)-one in 15 ml of NMP. The reaction medium is microwave-heated for 15 minutes at 150°C and then concentrated under reduced pressure. After the addition of 100 ml of water, the precipitate is filtered off and then dried under reduced pressure at 50°C overnight. The solid obtained is purified by silica gel column chromatography, elution being carried out with a dichloromethane/methanol (50/1) mixture.

MH+: 481

3-{3-[(2-Methyl-4-oxo-3-propyl-3,4-dihydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid

7.65 ml of a 1N aqueous solution of sodium hydroxide are added to 0.735 g
5 (1.53 mmol) of methyl 3-{3-[(2-methyl-4-oxo-3-propyl-3,4-dihydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoate in 30 ml of THF. The reaction medium is refluxed for 2.5 hours. After acidification with 10 ml of a 1N aqueous solution of hydrochloric acid, the reaction medium is concentrated under reduced pressure. The residue is taken up in 20 ml of water. The precipitate obtained is
10 filtered off, and dried under reduced pressure at 50°C overnight. 0.52 g of a yellow solid is obtained.

MH+: 467

¹H-NMR (D6-DMSO, 500 MHz):

0.97 (t, J = 7.6 Hz, 3 H), 1.69 – 1.76 (m, 2 H), 2.71 (s, 3H), 4.07 – 4.11 (m, 2 H),
15 7.40 – 7.44 (m, 1H), 7.59 – 7.66 (m, 1H), 7.71 – 7.80 (m, 2 H), 8.01 – 8.05 (m, 1 H), 8.28 – 8.39 (2 m, 2 H), 8.55 – 8.58 (m, 1 H), 8.79 – 8.82 (m, 1 H), 9.30 – 9.34 (m, 1 H), 9.88 – 9.22 (m, 1 H), 13.23 (br s, 1H).

Example 14: (Compound No. 55)

20 **3-{3-[(2-Methyl-4-oxo-3-propyl-3,4-dihydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzamide**

0.107 g (2 mmol) of ammonium chloride, 0.328 g (1 mmol) of TOTU and 0.517 g (4 mmol) of N,N-diisopropylethylamine are added, at ambient temperature under an inert atmosphere, to 0.233 g (0.5 mol) of 3-{3-[(2-methyl-4-oxo-3-propyl-
25 3,4-dihydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid in 30 ml of DMF. The reaction medium is stirred for 5 hours at ambient temperature and then concentrated under reduced pressure. 50 ml of a saturated solution of sodium hydrogen carbonate are added to the residue. The precipitate obtained is filtered off, and then dried under reduced pressure at 50°C overnight. 0.230 g of a yellow
30 solid is obtained.

MH+: 466

¹H-NMR (D6-DMSO, 500 MHz):

0.98 (t, J = 8 Hz, 3H), 1.74 (m, 2 H), 2.71 (s, 3 H), 4.10 (t, J = 8.1 Hz, 2H), 7.40 – 7.45 (m, 1H), 7.54 – 7.64 (m, 2 H), 7.67 – 7.71 (m, 1 H), 7.75 – 7.80 (m, 1 H), 7.96

– 8.00 (m, 1 H), 8.19 – 8.23 (m, 2 H), 8.42 – 8.48 (m, 2H), 8.82 – 8.85 (m, 1H), 9.39 – 9.41 (m, 1 H), 9.90 – 9.95 (m, 1 H).

Example 15: (Compound No. 3)

5 **6-(Imidazo[1,5-a]pyridin-3-ylcarbonyl)quinazolin-4(3H)-one**

0.36 g (3.6 mmol) of formamidine acetate is added to 0.2 g (0.72 mmol) of 2-amino-5-(imidazo[1,5-a]pyridin-3-ylcarbonyl)benzoic acid (described in WO 06/097625) in 7 ml of ethanol. The reaction medium is microwave-heated at 150°C for 25 minutes. The reaction medium is hydrolysed with a 1N aqueous solution of sodium hydroxide. The aqueous phase is extracted with dichloromethane. The heterogeneous organic phase is filtered. The solid obtained is purified by silica gel column chromatography, elution being carried out with a dichloromethane/methanol (90/10) mixture. 54 mg of a yellow solid are obtained.

MH+: 291

15 Melting point: 289°C

¹H-NMR (D6-DMSO, 400 MHz):

7.29-7.47 (m, 2H), 7.80-7.82 (m, 1H), 7.96 (s, 1H), 8.04-8.07 (m, 1H), 8.23 (s, 1H), 8.67-8.70 (m, 1H), 9.29 (s, 1H), 9.52-9.53 (m, 1H), 12.5 (s, 1H).

20 **Example 16: (Compound No. 177)**

N,N-1,2-Tetramethyl-4-oxo-6-[[1-(pyridin-3-yl)imidazo[1,5-a]pyridin-3-yl]carbonyl]-1,4-dihydroquinoline-3-carboxamide

6-[(1-Bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-2H-3,1-benzoxazine-2,4(1H)-dione

25 1.30 g (4.37 mmol) of triphosgene dissolved in 10 ml of dioxane are added, under an inert atmosphere at ambient temperature, to 1.05 g (2.91 mmol) of 2-amino-5-(1-bromoimidazo[1,5-a]pyridin-3-ylcarbonyl)benzoic acid. The reaction medium is refluxed for 4 hours, and then concentrated under reduced pressure. 100 ml of water are added to the residue. The precipitate obtained is filtered off
30 under reduced pressure at 40°C for 18 hours. 1.1 g of a yellow solid are obtained.

MH+: 386, 388

6-[(1-Bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-1-methyl-2H-3,1-benzoxazine-2,4(1H)-dione

808 mg (5.7 mmol) of methyl iodide and 164 mg (3.42 mmol) of 50% sodium hydride are added, at ambient temperature under an inert atmosphere, to 1.1 g (2.85 mmol) of 6-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-2H-3,1-benzoxazine-2,4(1H)-dione in 20 ml of DMF. After stirring for 3 hours, the reaction medium is poured into 200 ml of ice-cold water. The precipitate is filtered off, rinsed with water, and then dried under reduced pressure for 18 hours at 40°C. 1.13 g of a yellow solid are obtained.

MH+: 402, 403.95

10 **6-[(1-Bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-N,N-1,2-tetramethyl-4-oxo-1,4-dihydroquinoline-3-carboxamide**

1.25 ml of a 2N aqueous solution of sodium hydroxide are added, at ambient temperature, to 333 mg (2.5 mmol) of N,N-dimethylacetamide in 3 ml of DMF. After stirring for 1 hour, 400 mg (1 mmol) of 6-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-1-methyl-2H-3,1-benzoxazine-2,4(1H)-dione dissolved in 25 ml of DMF are added. The reaction medium is heated for 6 hours at 50°C under an inert atmosphere. The reaction medium is concentrated under reduced pressure. The residue obtained is purified by silica gel column chromatography, elution being carried out with a dichloromethane/methanol 20/1 mixture. 220 mg of a yellow oil are obtained.

MH+: 467, 469

N,N-1,2-Tetramethyl-4-oxo-6-[[1-(pyridin-3-yl)imidazo[1,5-a]pyridin-3-yl]carbonyl]-1,4-dihydroquinoline-3-carboxamide

25 115 mg (0.37 mmol) of 3-pyridinylboronic acid, 331 mg (1.56 mmol) of potassium phosphate dissolved in 1 ml of water, and 18 mg (15.6 µmol) of tetrakis(triphenylphosphine)palladium are added, at ambient temperature under an inert atmosphere, to 365 mg (0.78 mmol) of 6-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-N,N-1,2-tetramethyl-4-oxo-1,4-dihydroquinoline-3-carboxamide in 20 ml of N-methylpyrrolidone. The reaction medium is microwave-heated at 150°C for 25 minutes and then concentrated under reduced pressure. The residue obtained is purified by silica gel column chromatography, elution being carried out with a dichloromethane/methanol 9/1 mixture. 0.290 g of a yellow oil is obtained.

MH+: 466

¹H-NMR (D6-DMSO, 500 MHz):

2.44 (s, 3 H), 2.89 (s, 3 H), 3.03 (s, 3 H), 3.89 (s, 3 H), 7.39 – 7.43 (m, 1 H),
7.59 – 7.64 (2 m, 2 H), 8.07 (d, J = 9.7 Hz, 1 H), 8.44 – 8.49 (2 m, 2 H), 8.67 (d, J =
4.8 Hz, 1 H), 8.86 (dd, J = 9.7 Hz and 2.2 Hz, 1 H), 9.32 – 9.34 (m, 1 H), 9.55 (d, J
5 = 2.2 Hz, 1 H), 9.93 (d, J = 7.4 Hz, 1 H).

Example 17: (Compound No. 223)

**Sodium salt of 3-[3-({3-[2-(4-fluorophenoxy)ethyl]-1-propyl-2,4-dioxo-1,2,3,4-
tetrahydroquinazolin-6-yl}carbonyl)imidazo[1,5-a]pyridin-1-yl]benzoic acid**

10

**Methyl 5-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-2-({[2-(4-
fluorophenoxy)ethyl]carbamoyl}amino)benzoate**

4.75 g (16 mmol) of triphosgene are added, at ambient temperature under
an inert atmosphere, to 4.99 g (13.33 mmol) of methyl 2-amino-5-({1-[3-
15 (methoxycarbonyl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)benzoate in 220 ml of
dioxane. After refluxing for 5 hours, 6.21 g (40 mmol) of 2-(4-fluorophenoxy)-1-
ethylamine and 4.05 g (40 mmol) of triethylamine are added at ambient
temperature. The reaction medium is refluxed for 3 hours and then concentrated
under reduced pressure. The residue is triturated from water. After filtration, the
20 solid is rinsed with methanol and then dried under reduced pressure at 40°C
overnight. 6.67 g of a yellow solid are obtained.

MH+: 555

**6-[(1-Bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-3-[2-(4-
25 fluorophenoxy)ethyl] quinazoline-2,4(1H,3H)-dione**

60.1 ml (60.1 mmol) of a 1N aqueous solution of sodium hydroxide are added
to 6.67 g (12 mmol) of methyl 5-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-2-
({[2-(4-fluorophenoxy)ethyl]carbamoyl}amino)benzoate dissolved in 600 ml of
methanol. After refluxing for 2 hours, the reaction medium is acidified with 120 ml
30 of a 1N aqueous solution of hydrochloric acid and then diluted with 2000 ml of
water. The precipitate obtained is filtered off, and dried under reduced pressure at
40°C overnight. 5.83 g of a yellow solid are obtained.

MH+: 523.2, 525.2

6-[(1-Bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-3-[2-(4-fluorophenoxy)ethyl]-1-propylquinazoline-2,4(1H,3H)-dione

7.22 g (22.16 mmol) of caesium carbonate and 5.65 g (33.24 mmol) of propyl iodide are added, under an inert atmosphere, to 5.6 g (11.08 mmol) of 6-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-3-[2-(4-fluorophenoxy)ethyl]quinazoline-2,4(1H,3H)-dione in 300 ml of DMF. The reaction medium is stirred for 12 hours at ambient temperature under a nitrogen atmosphere and then concentrated under reduced pressure. The residue obtained is washed with 700 ml of water and then dried under reduced pressure at 40°C overnight. 5.74 g of a yellow solid are obtained.

MH+: 565, 567

Methyl 3-[3-({3-[2-(4-fluorophenoxy)ethyl]-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl}carbonyl)imidazo[1,5-a]pyridin-1-yl]benzoate

2.178 g (12.1 mmol) of 3-methoxycarbonylphenylboronic acid, 4.279 g (20.16 mmol) of potassium phosphate dissolved in 30 ml of water, and 582.4 g (0.504 mmol) of tetrakis(triphenylphosphine)palladium are added to 5.7 g (10.08 mmol) of 6-[(1-bromoimidazo[1,5-a]pyridin-3-yl)carbonyl]-3-[2-(4-fluorophenoxy)ethyl]-1-propylquinazoline-2,4(1H,3H)-dione in 180 ml of NMP. The reaction medium is microwave-heated for 15 minutes at 120°C and then concentrated under reduced pressure. The solid obtained is purified by silica gel column chromatography, elution being carried out with a dichloromethane/methanol (100/1) mixture. 4.32 g of a yellow solid are obtained.

MH+: 621.3

Sodium salt of 3-[3-({3-[2-(4-fluorophenoxy)ethyl]-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl}carbonyl)imidazo[1,5-a]pyridin-1-yl]benzoic acid

69.6 ml (69.6 mmol) of a 1N aqueous solution of lithium hydroxide are added to 4.32 g (6.96 mmol) of methyl 3-[3-({3-[2-(4-fluorophenoxy)ethyl]-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl}carbonyl)imidazo[1,5-a]pyridin-1-yl]benzoate in 500 ml of THF. The reaction medium is refluxed for 3 hours and then acidified at ambient temperature with 150 ml of a 1N aqueous solution of hydrochloric acid and, finally, diluted with 700 ml of water. After filtration, the

residue obtained is dried under reduced pressure at 40°C overnight.

5.88 ml (5.88 mmol) of a 1N aqueous solution of sodium hydroxide are added to 4.11 g (6 mmol) of the yellow solid obtained, in 100 ml of methanol. After filtration, the residue obtained is dried under reduced pressure at 40°C overnight. 3.46 g of a yellow solid are obtained.

MH⁺: 607.3

Mp: 190-205°C (decomposition)

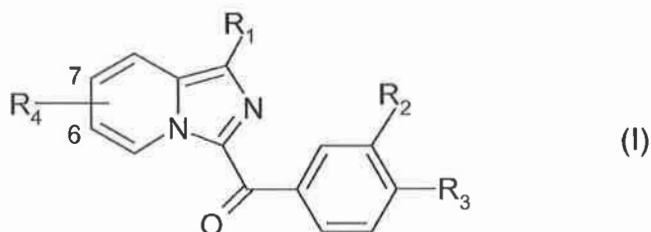
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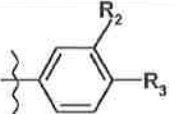
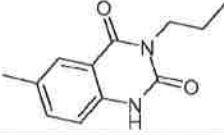
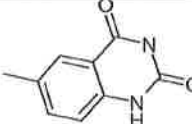
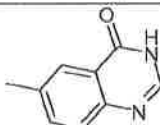
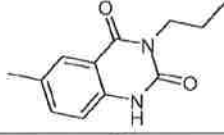
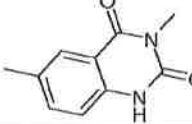
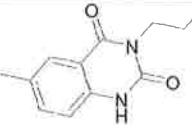
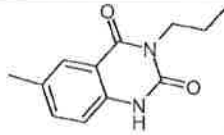
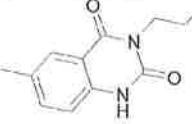
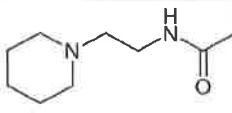
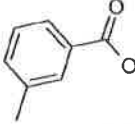
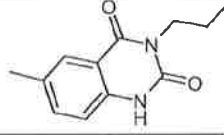
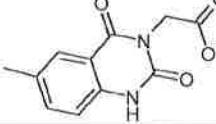
0.98 (t, J=7.7Hz, 3H), 1.71 (tq, J₁=J₂=7.7Hz, 2H), 4.17 (t, J=7.7Hz, 2H), 4.24 (t, J=6.6Hz, 2H), 4.39 (t, J=6.6Hz, 2H), 6.97-7.00 (2m, 2H), 7.10-7.16 (2m, 2H), 7.38-7.41 (m, 1H), 7.47- 7.52 (m, 1H), 7.57-7.61 (m, 1H), 7.75-7.79 (m, 1H), 7.94-7.98 (2m, 2H), 8.26-8.30 (m, 1H), 8.49-8.52 (m, 1H), 8.97-9.02 (m, 1H), 9.26-9.28 (m, 1H), 9.89-9.93(m, 1H).

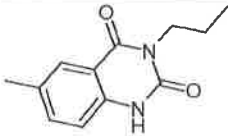
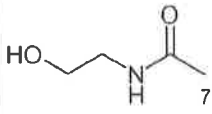
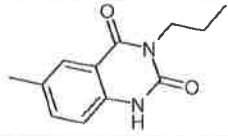
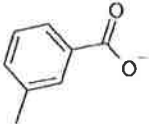
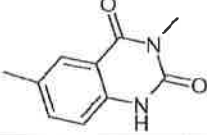
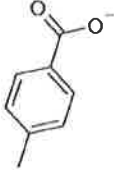
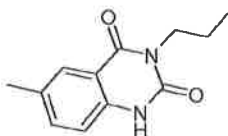
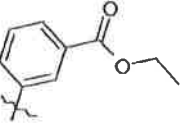
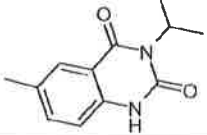
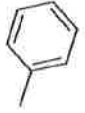
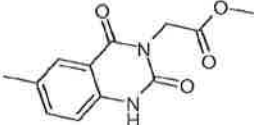
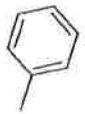
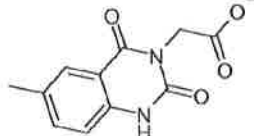
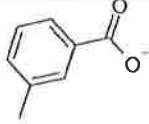
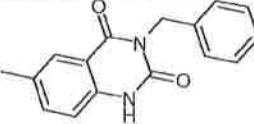
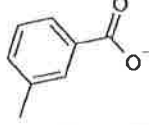
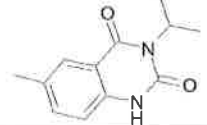
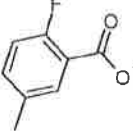
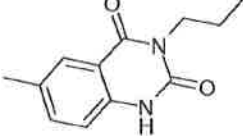
15 The table which follows illustrates the chemical structures and the physical properties of some compounds according to the invention. In this table:

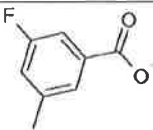
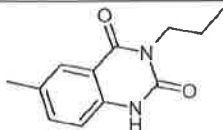
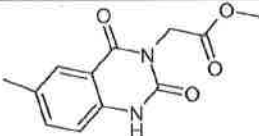
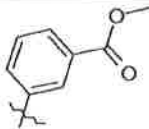
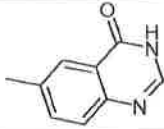
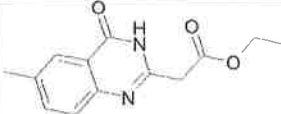
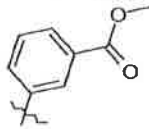
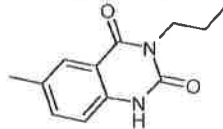
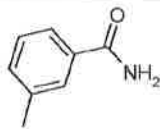
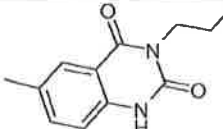
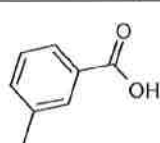
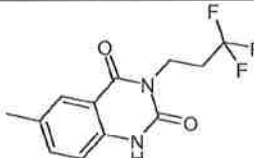
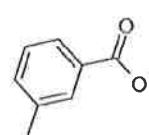
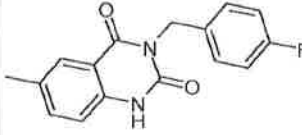
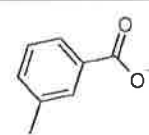
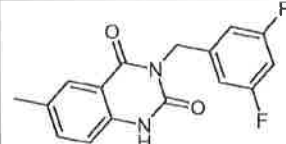
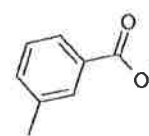
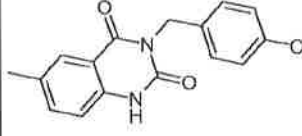
- Me and Et represent, respectively, methyl and ethyl groups;
- the wavy lines indicate the bond attached to the rest of the molecule;
- "Mp" represents the melting point of the compound, expressed in degrees Celsius;
- "M+H⁺" represents the mass of the compound, obtained by LC-MS (Liquid Chromatography - Mass Spectroscopy).

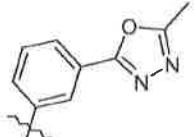
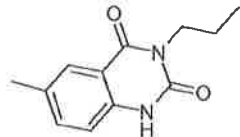
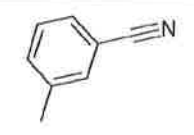
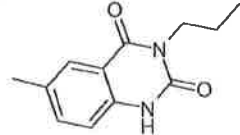
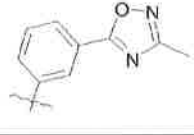
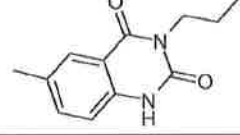
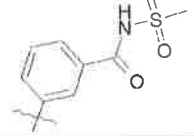
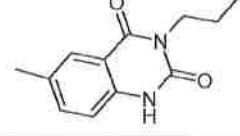
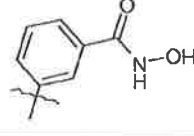
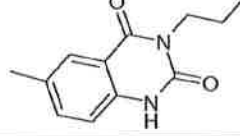
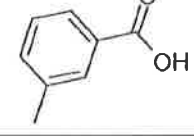
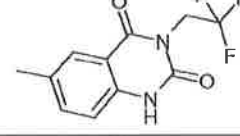
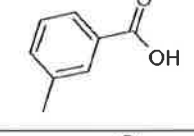
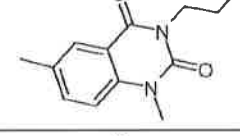
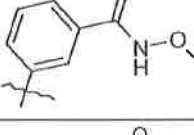
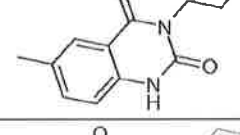
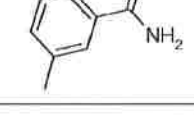
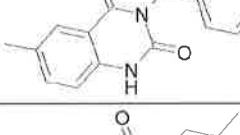
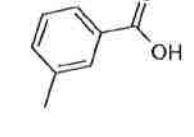
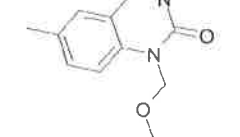
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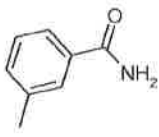
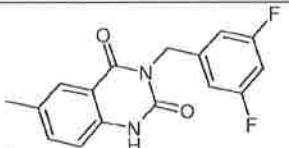
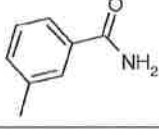
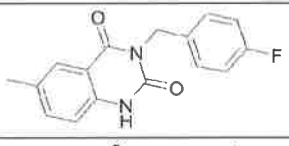
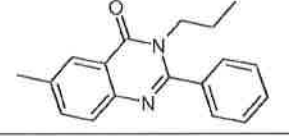
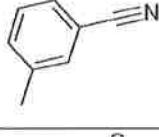
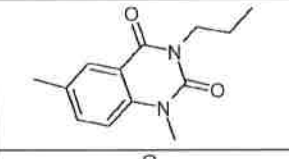
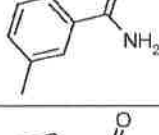
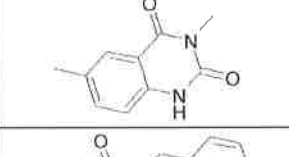
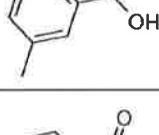
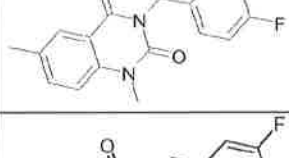
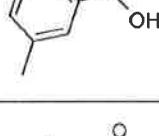
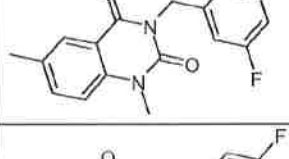
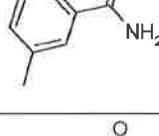
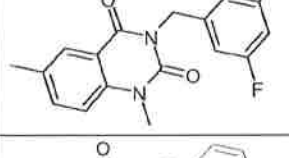
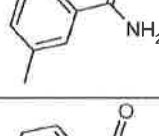
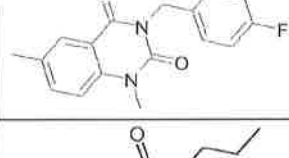
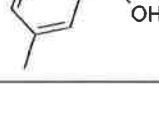
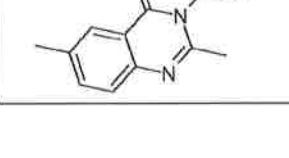


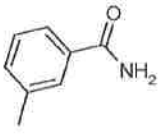
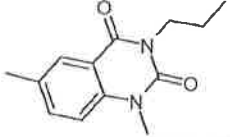
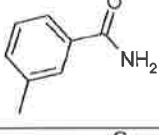
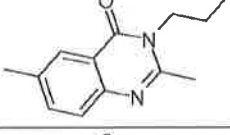
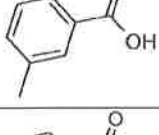
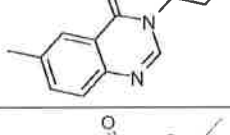
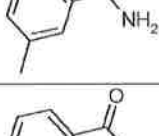
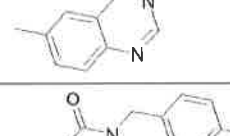
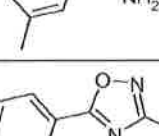
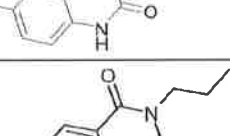
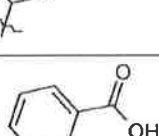
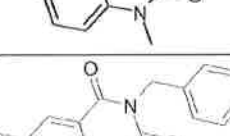
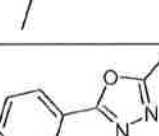
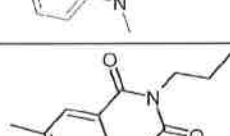
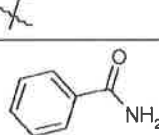
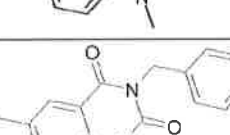
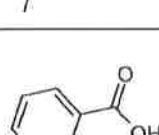
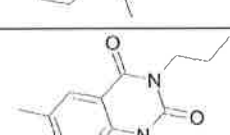
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2	H		H	/	341	307
3	H		H	/	289	291
4	H		7-COOH	/	380	393
5	H		7-COOH	/	404	365
7	-CO-NH ₂		H	/	/	392
8	H		6-COOH	/	384	393
9	H			HCl	234	503
10			H	Na	383	469
11	H		H	Na	398	365

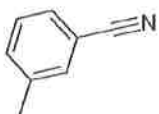
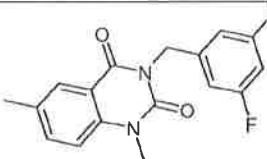
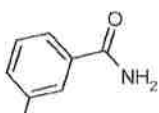
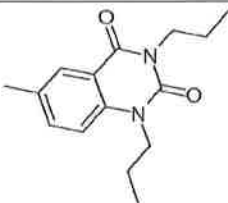
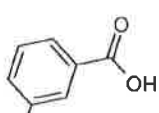
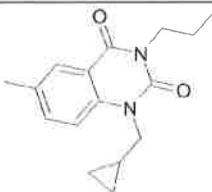
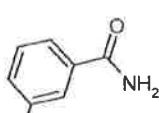
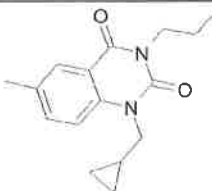
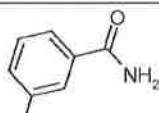
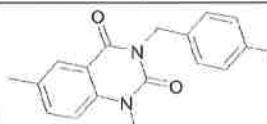
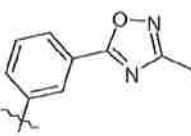
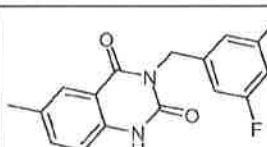
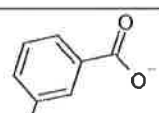
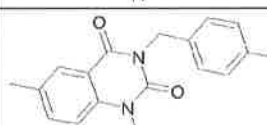
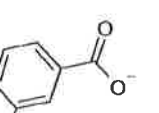
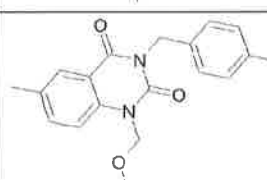
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13	-NH-SO ₂ Me		H	/	334	442
14			H	Na	>41	441
15			H	Na	>410	469
16			H	/	257	497
17			H	/	361	455
18			H	Na	345	440
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20			H	Na	363	469
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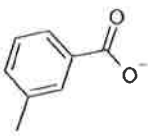
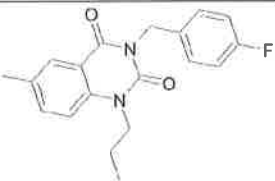
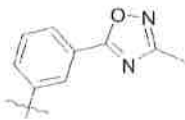
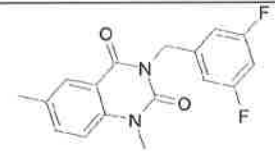
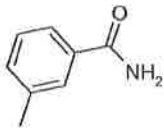
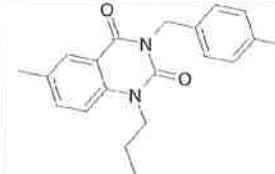
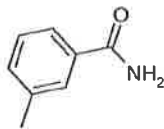
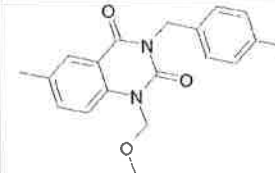
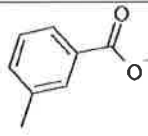
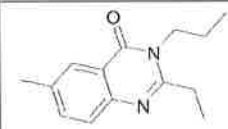
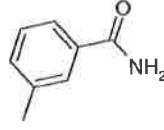
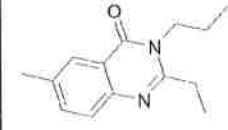
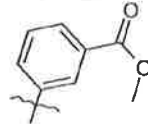
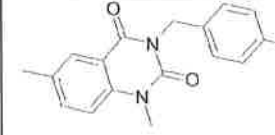
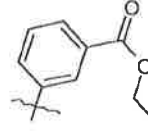
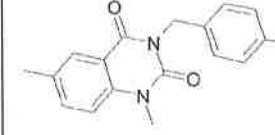
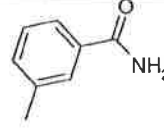
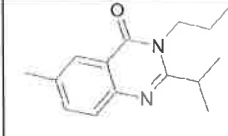
22			H	Na	390	487
23	H		H	/	308	379
24			H	/	318	425
26	H		H	/	263	377
28			H	/	271	483
29			H	/	345-346	468
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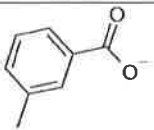
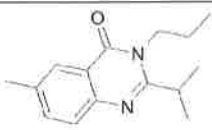
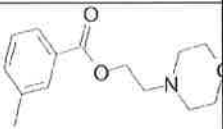
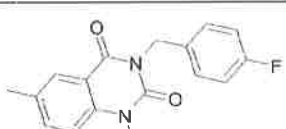
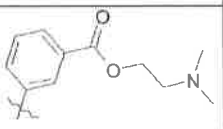
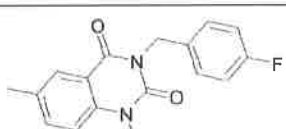
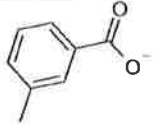
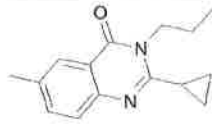
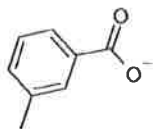
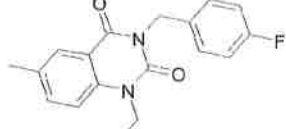
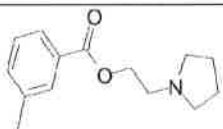
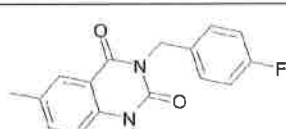
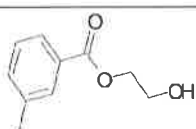
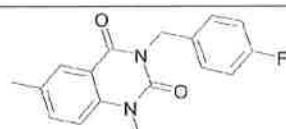
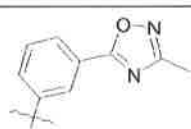
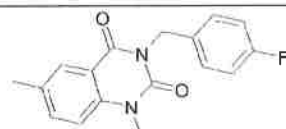
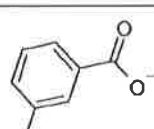
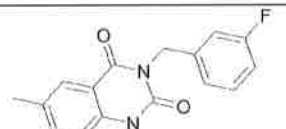
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36			H	/	276-277	507
37			H	/	250-260	546
38			H	/	269	484
39			H	/	387	509
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42			H	/	341	516
43			H	/	268	513

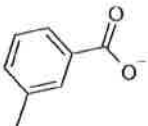
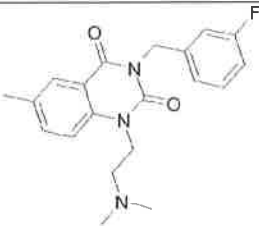
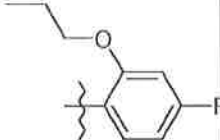
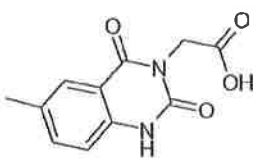
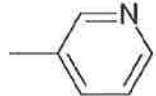
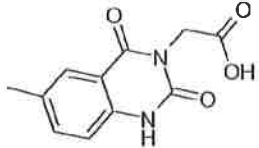
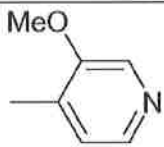
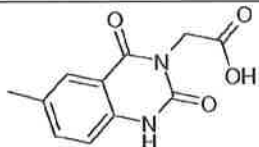
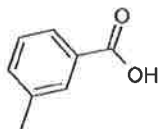
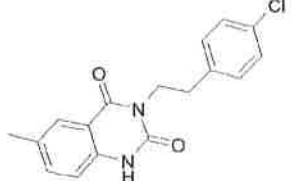
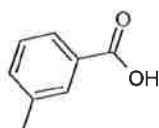
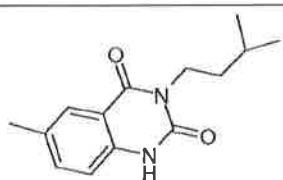
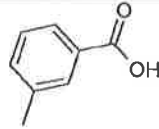
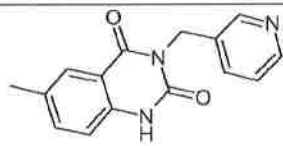
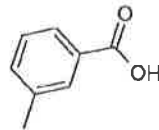
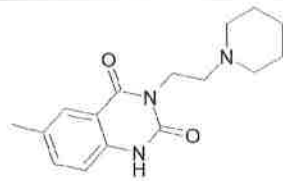
44			H	/	340	552
45			H	/	319	534
46	H		H	/	/	409
47			H	/	230	464
48			H	/	274	440
49			H	Na	182	549
50			H	/	300-301	567
51			H	/	290	566
52			H	/	305	548
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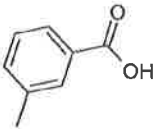
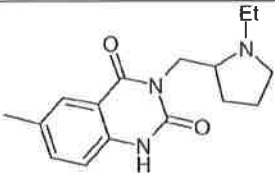
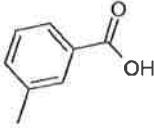
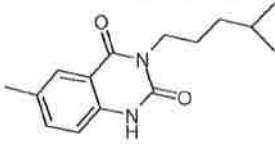
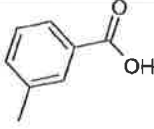
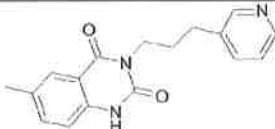
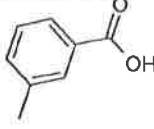
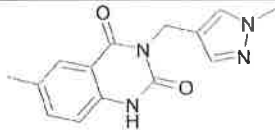
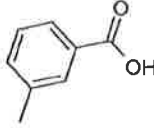
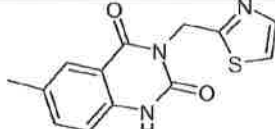
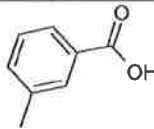
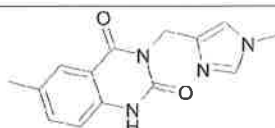
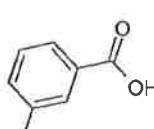
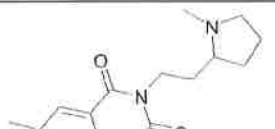
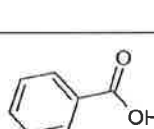
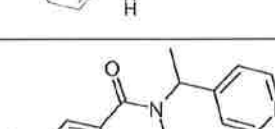
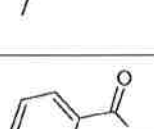
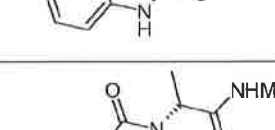
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55			H	/	238	466
56			H	/	311-312	453
57			H	/	251	452
58			H	/	338	550
59			H	/	241	521
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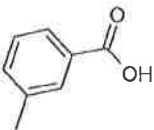
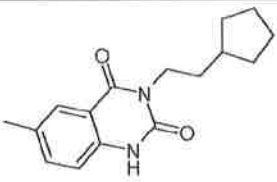
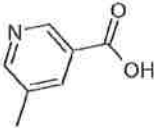
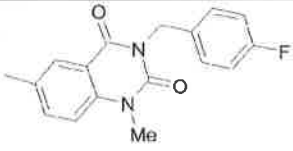
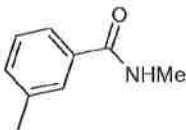
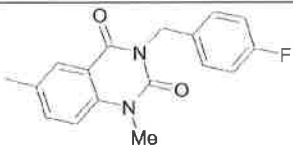
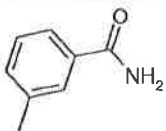
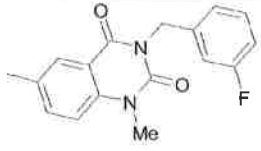
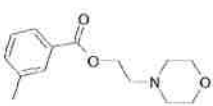
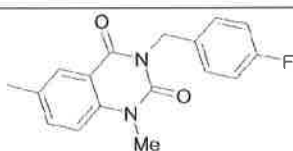
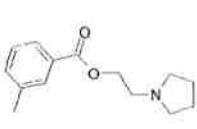
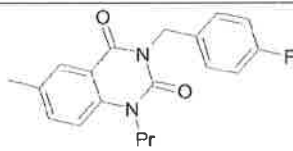
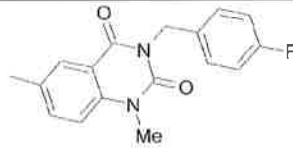
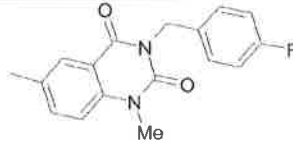
64			H	/	285	548
65			H	/	231	510
66			H	/	271	523
67			H	/	254	522
68			H	/	310	564
69			H	/	319	591
70			H	Na	255	564
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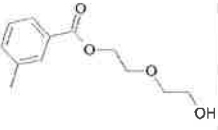
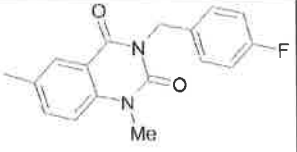
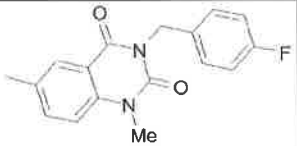
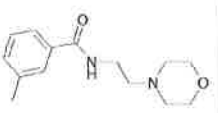
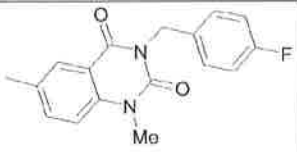
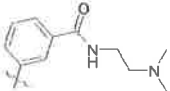
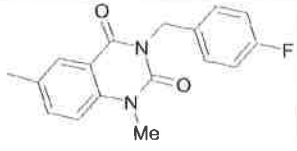
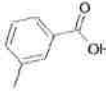
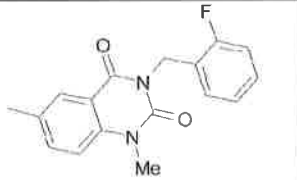
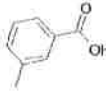
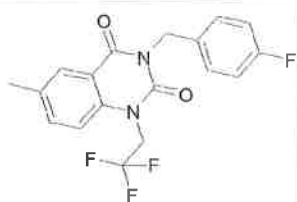
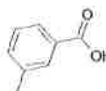
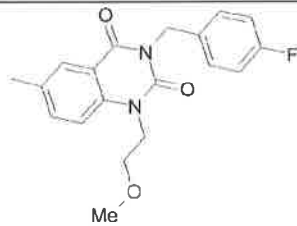
72			H	Na	239-245	577
73			H	/	258	605
74			H	/	250-252	576
75			H	/	297-298	578
76			H	/	150-153	481
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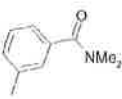
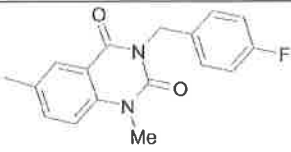
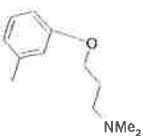
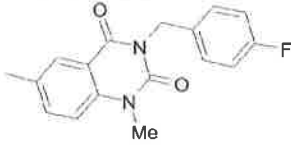
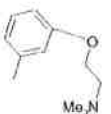
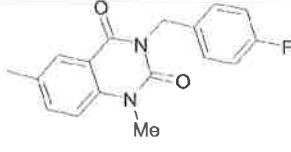
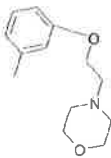
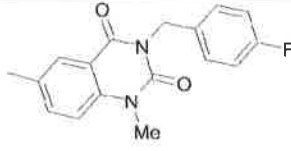
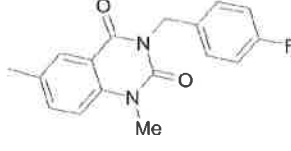
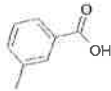
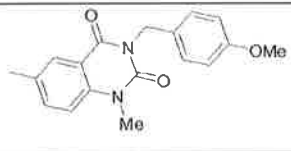
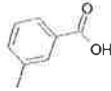
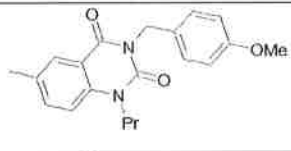
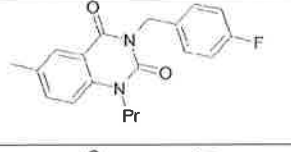
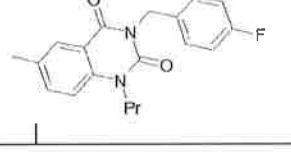
81			H	Na	226-230	494
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83			H	/	238	620
84			H	Na	235-238	493
85			H	Na	244-246	563
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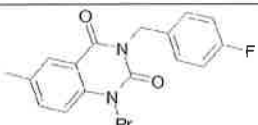
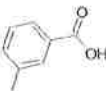
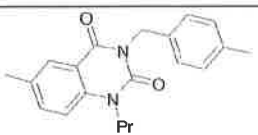
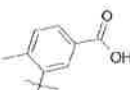
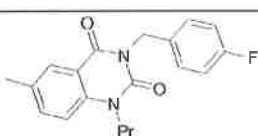
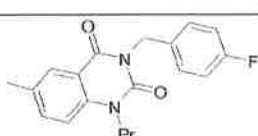
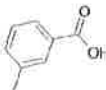
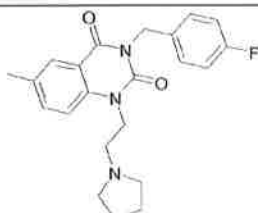
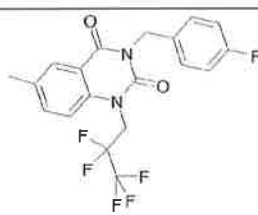
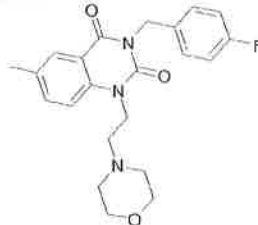
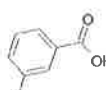
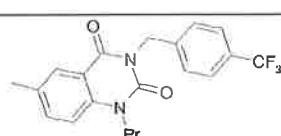
90			H	Na	222-225	606
91			H	/	/	517
92			H	/	/	442
93			H	/	/	472
94			H	HCl	/	602
95			H	/	/	497
95a			H	HCl	/	554
96			H	HCl	/	575

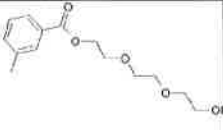
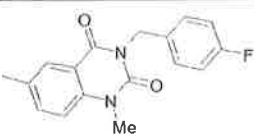
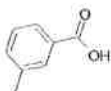
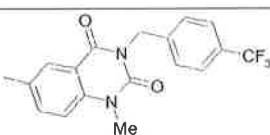
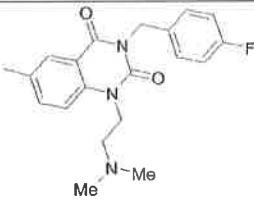
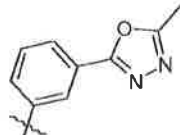
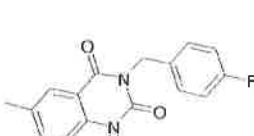
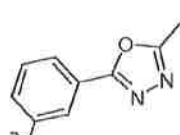
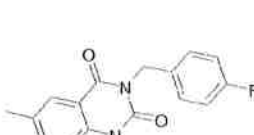
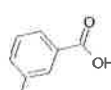
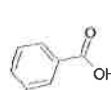
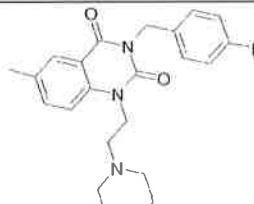
97			H	HCl	/	575
98			H	/	/	511
99			H	HCl	/	583
100			H	/	/	521
101			H	/	/	524
102			H	/	/	521
103			H	/	/	538
104			H	HCl	/	568
105			H	/	/	512

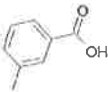
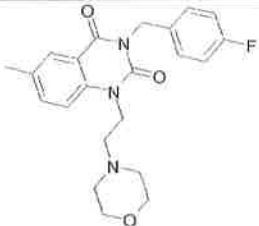
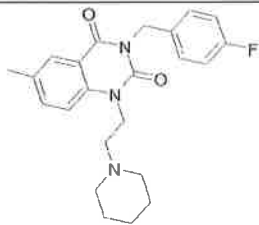
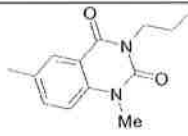
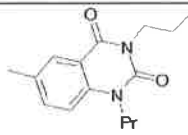
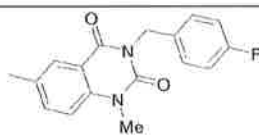
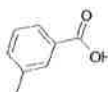
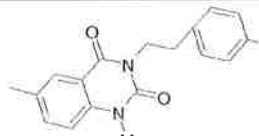
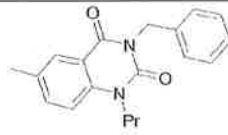
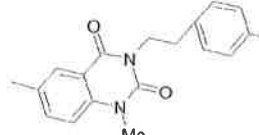
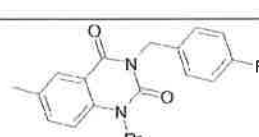
106			H	/	/	523
107			H	HCl	/	550
108			H	/	318	562
109			H	/	/	548
110			H	HCl	177	690
111			H	HCl	187	674
112			H	/	224	505
113			H	HCl	267	506

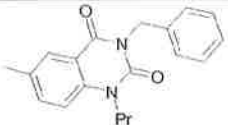
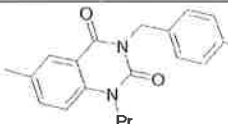
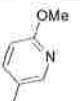
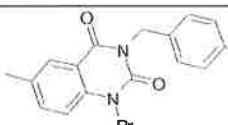
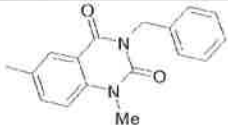
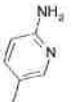
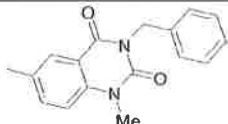
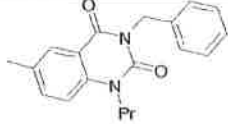
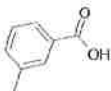
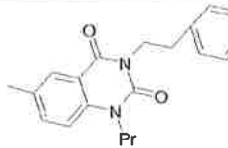
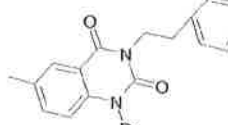
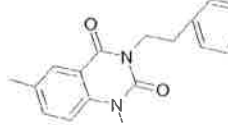
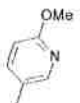
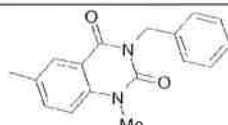
114			H	/	106	637
115			H	/	/	530
116			H	HCl	214	661
117			H	HCl	247	619
118			H	Na	/	598
119			H	Na	/	617
120			H	Na	/	593

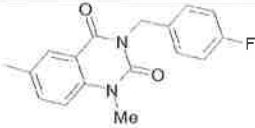
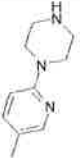
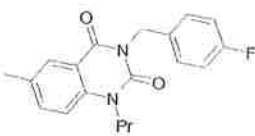
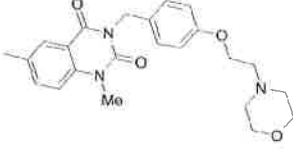
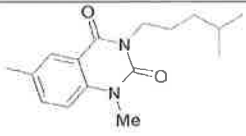
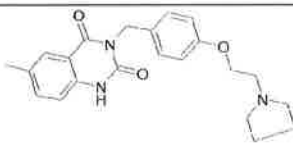
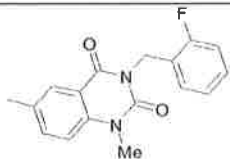
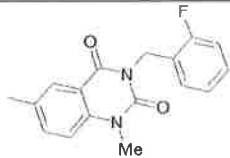
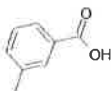
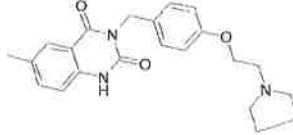
121a			H	/	259	576
121			H	HCl	247	606
122			H	HCl	287	592
123			H	HCl	274	634
124			H	/	216	521
125			H	Na	258	561
126			H	Na	240	589
127			H	HCl	275	534
128			H	HCl	273	534

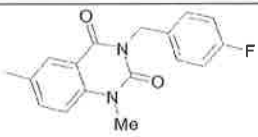
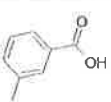
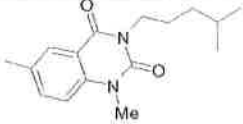
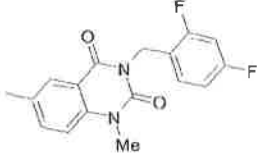
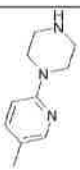
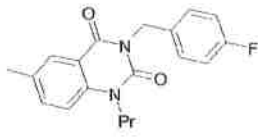
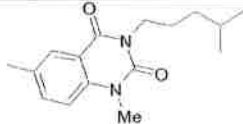
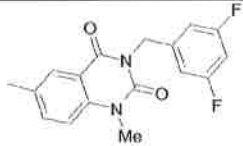
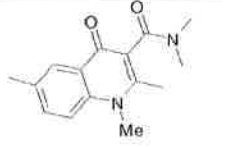
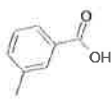
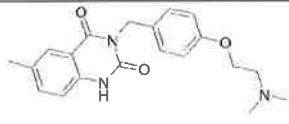
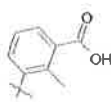
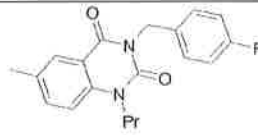
129			H	/	223	550
130			H	Na	246	573
131			H	Na	256	591
132			H	HCl	291	550
133			H	Na	/	632
134			H	/	/	624
135			H	/	/	605
136			H	Na	236	627

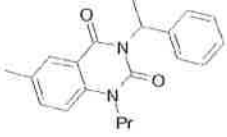
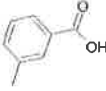
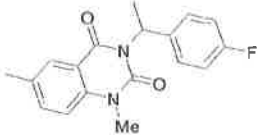
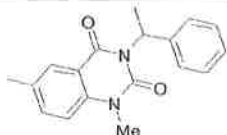
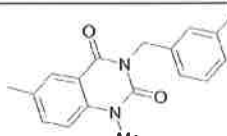
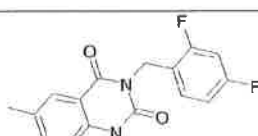
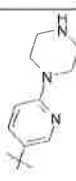
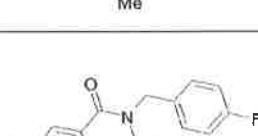
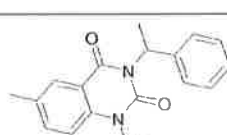
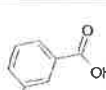
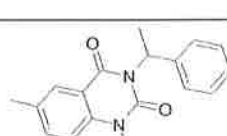
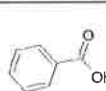
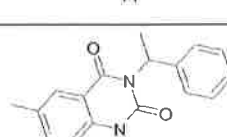
137			H	/	112	681
138			H	Na	237	599
139			H	HCl	226	563
140			H	/	/	573
141			H	/	/	587
142			H	Na	/	667
143			H	Na	/	646

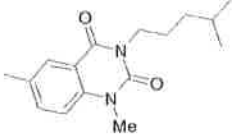
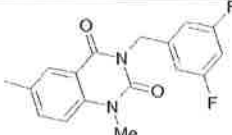
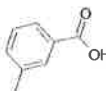
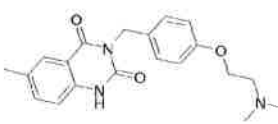
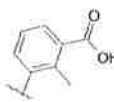
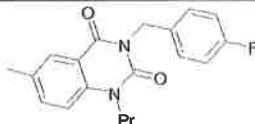
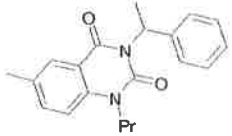
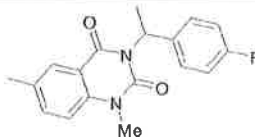
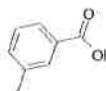
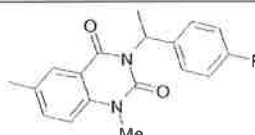
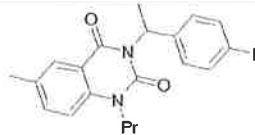
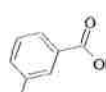
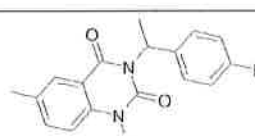
144			H	Na	/	648
145			H	/	/	603
146			H	HCl	250	440
147			H	HCl	278	468
148			H	/	294	522
149			H	Na	256	563
150			H	HCl	225	516
151			H	/	257	536
152			H	/	259	549

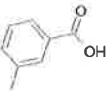
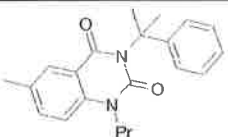
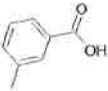
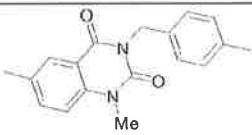
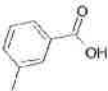
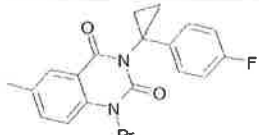
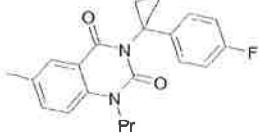
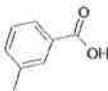
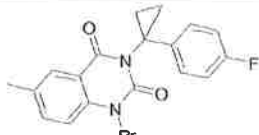
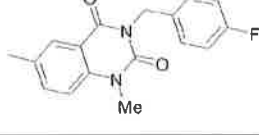
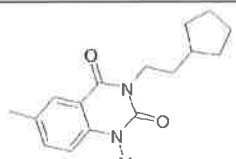
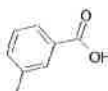
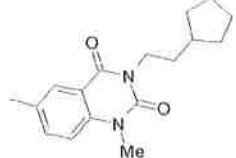
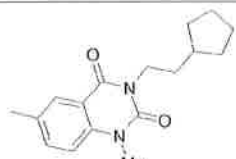
153			H	/	128	532
154			H	/	233	537
155			H	HCl	128	564
156			H	/	/	504
157			H	/	278	521
158	Me		H	/	/	471
159			H	Na	221	591
160			H	HCl	525	548
161			H	/	250	564
162			H	HCl	230	536

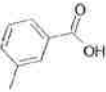
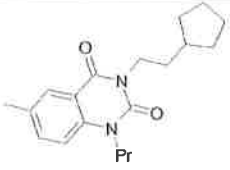
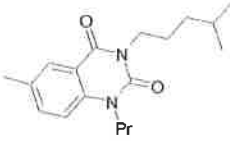
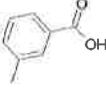
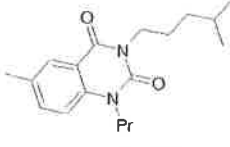
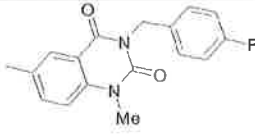
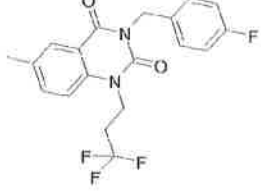
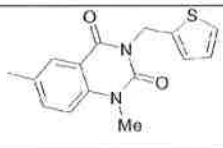
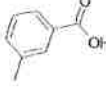
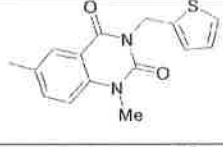
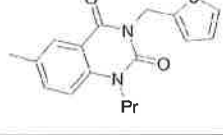
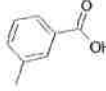
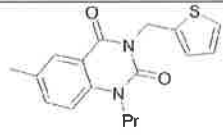
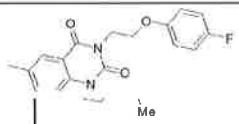
163			H	/	194	509
164			H	/	160	618
165			H	/	/	617
166			H	/	/	482
167			H	/	/	587
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169			H	/	/	522
170			H	/	/	630

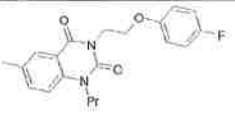
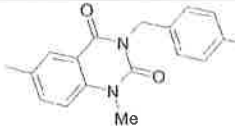
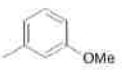
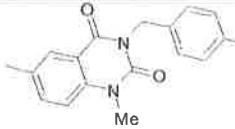
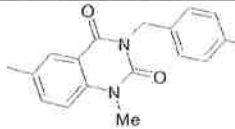
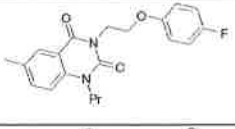
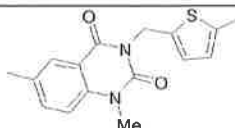
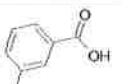
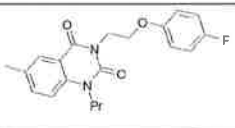
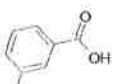
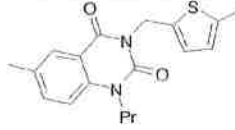
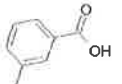
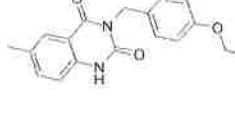
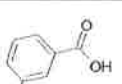
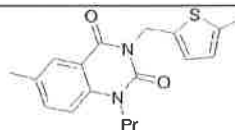
171	Me		H	/	/	443
172			H	Na	/	525
173			H	/	/	524
174			H	/	261	590
175			H	/	/	498
176			H	/	/	524
177			H	/	/	466
178			H	Na	/	604
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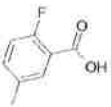
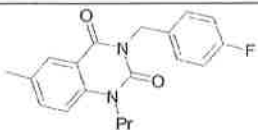
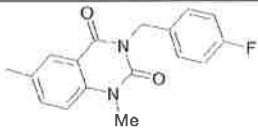
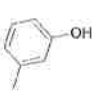
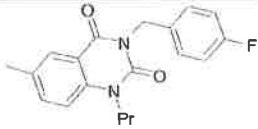
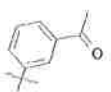
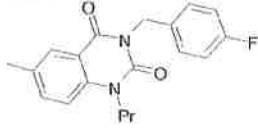
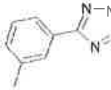
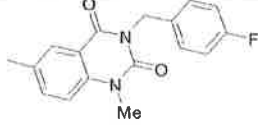
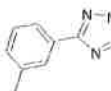
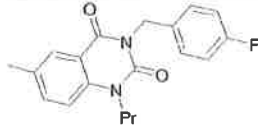
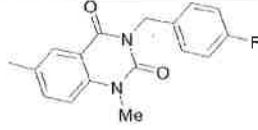
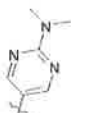
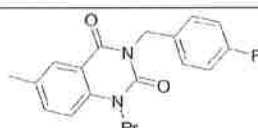
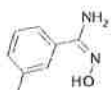
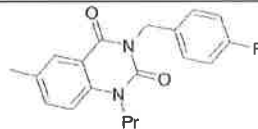
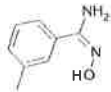
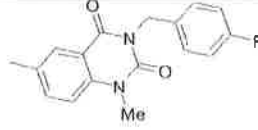
180			H	HCl	229	530
181			H	Na	270	563
182			H	HCl	261	502
183			H	/	/	506
184			H	/	/	524
185			H	/	261	590
186			H	HCl	261	502
187			H	Na	238	573
188			H	Na	274	545

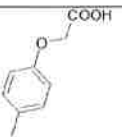
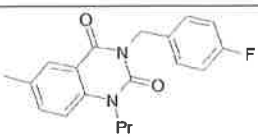
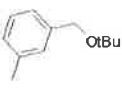
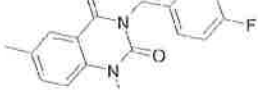
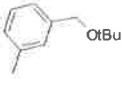
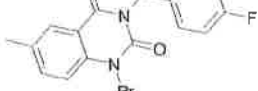
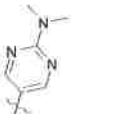
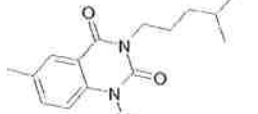
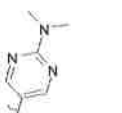
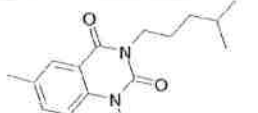
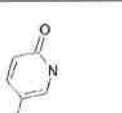
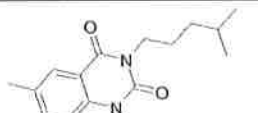
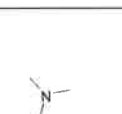
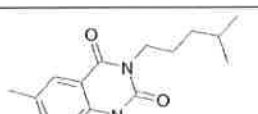
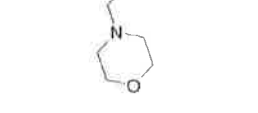
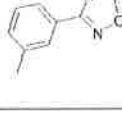
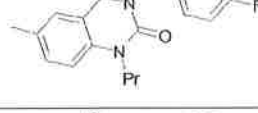
189			H	/	/	498
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191			H	Na	/	604
192			H	Na	236	591
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194			H	HCl	198	520
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196			H	HCl	218	548
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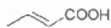
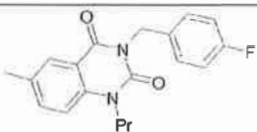
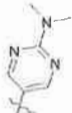
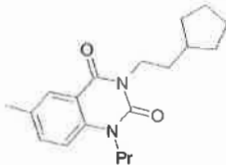
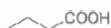
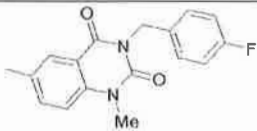
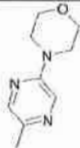
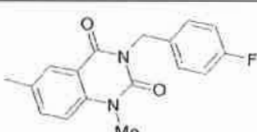
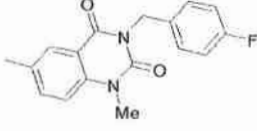
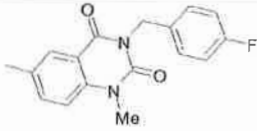
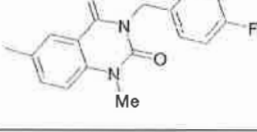
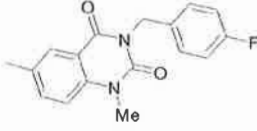
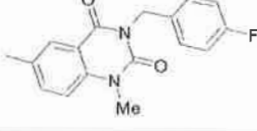
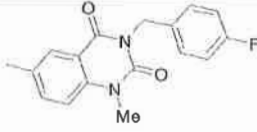
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199			H	Na	257	545
200			H	Na	285	575
201			H	HCl	264	560
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203			H	/	/	509
204			H	/	/	494
205			H	Na	/	537
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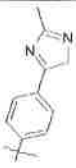
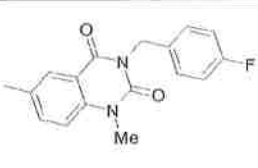
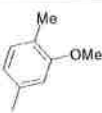
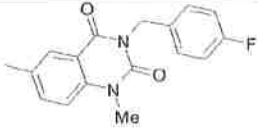
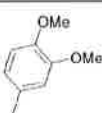
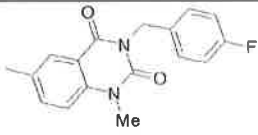
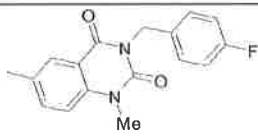
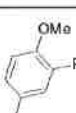
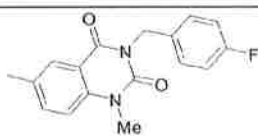
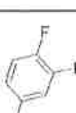
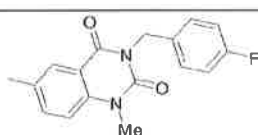
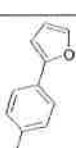
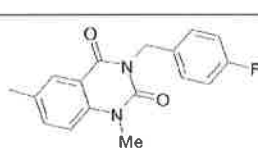
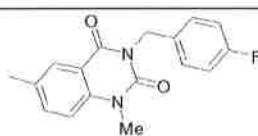
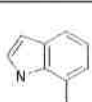
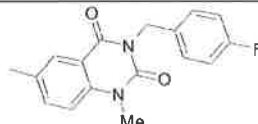
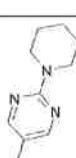
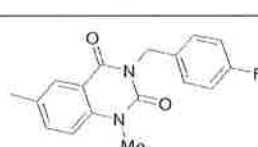
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208			H	/	/	510
209			H	Na	/	553
210			H	HCl	195	506
211			H	/	240	588
212			H	/	/	494
213			H	Na	/	537
214			H	/	/	522
215			H	Na	/	565
216			H	/	/	536

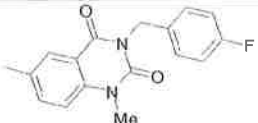
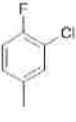
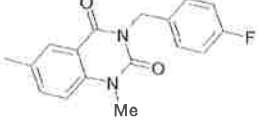
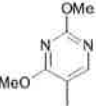
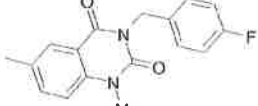
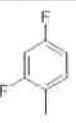
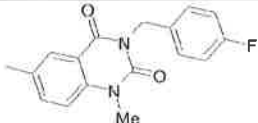
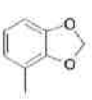
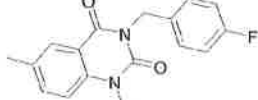
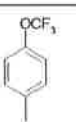
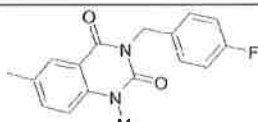
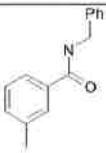
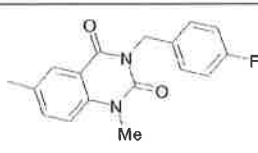
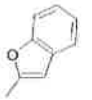
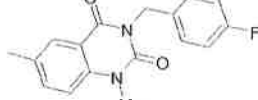
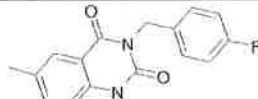
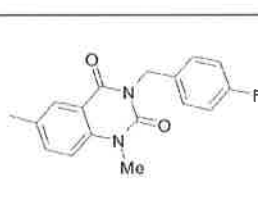
217			H	/	/	564
218			H	/	243	511
219			H	/	255	535
220	nBu		H	/	/	485
221			H	/	/	580
222			H	/	/	508
223			H	Na	/	607
224			H	/	/	536
225			H	Na	/	645
226			H	Na	/	579

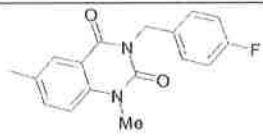
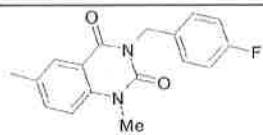
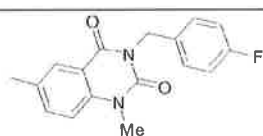
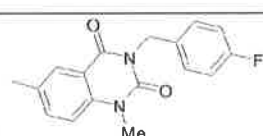
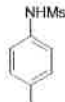
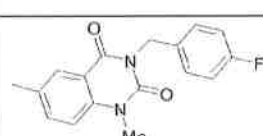
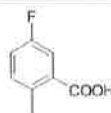
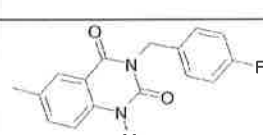
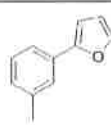
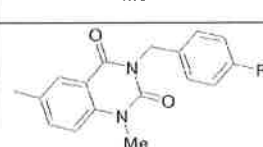
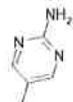
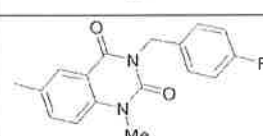
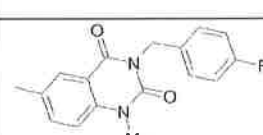
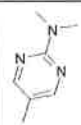
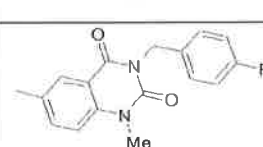
227			H	Na	313	595
228	H		H	/	/	429
229			H	/	146	549
230			H	/	227	575
231			H	/	/	573
232			H	/	/	601
233			H	HCl	290	556
234			H	/	/	578
235			H	/	/	591
236			H	/	/	563

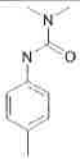
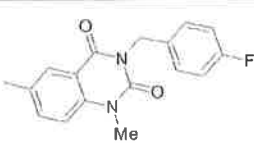
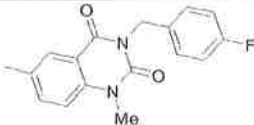
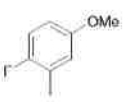
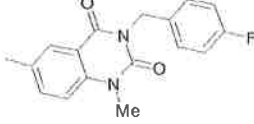
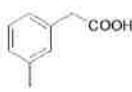
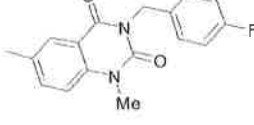
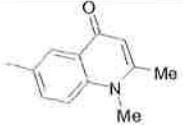
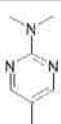
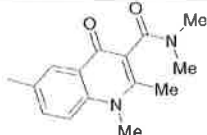
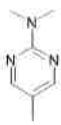
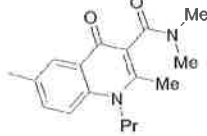
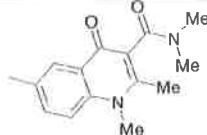
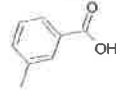
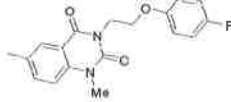
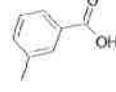
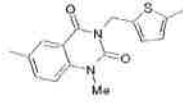
237			H	Na	217	607
238			H	/	182	591
239			H	/	108	619
240			H	/	/	526
241			H	/	/	554
242			H	/	285	550
243			H	/	/	625
244			H	/	/	617
245			H	/	/	589

246			H	Na	342	527
247			H	/	/	566
248			H	Na	349	501
249			H	HCl	/	592
250			H	HCl	/	495
251			H	HCl	/	551
252			H	HCl	/	520
253			H	HCl	/	495
254			H	HCl	/	561
255			H	HCl	/	589

256			H	HCl	/	602
257			H	HCl	/	549
258			H	HCl	/	565
259			H	HCl	/	539
260			H	HCl	/	553
261			H	HCl	/	541
262			H	HCl	/	571
263			H	HCl	/	545
264			H	HCl	/	544
265			H	HCl	/	590

266			H	HCl	/	591
267			H	HCl	/	557
268			H	HCl	/	567
269			H	HCl	/	541
270			H	HCl	/	549
271			H	HCl	/	589
272			H	HCl	/	638
273			H	HCl	/	545
274			H	HCl	/	556
275			H	HCl	/	632

276			H	HCl	/	509
277			H	HCl	/	555
278			H	HCl	/	556
279			H	HCl	/	573
280			H	HCl	/	598
281			H	HCl	/	567
282			H	HCl	/	571
283			H	HCl	/	522
284			H	HCl	/	522
285			H	HCl	/	550

286			H	HCl	/	522
287			H	HCl	/	563
288			H	HCl	/	553
289			H	HCl	/	563
290			H	/	/	395
291			H	/	/	510
292			H	/	/	538
293			H	/	/	466
294			H	Na	/	579
295			H	Na	/	551

The compounds according to the invention were the subject of pharmacological assays for determining their FGF-inhibiting effect.

Example 18: FGF-2-induced *in vitro* angiogenesis of HUVEC cells

5 In order to demonstrate the ability of the FGF-R antagonists of the present invention to inhibit FGF-induced angiogenesis, *in vitro* angiogenesis experiments were carried out with human endothelial cells of HUVEC type stimulated with FGF-2 or b-FGF.

To do this, matrices composed of matrigel (growth factor reduced matrigel, 10 Becton Dickinson 356230) and collagen (rat tail collagen type I, Becton Dickinson 354236) are deposited, at a rate of 160 μ l, into each chamberslide well (Biocoat Cellware collagen, Type I, 8-well cultureslides: Becton Dickinson 354630), or 60 μ l per well of 96-well plates (Biocoat collagen I cellware, Becton Dickinson 354407). The matrix is prepared by mixing 1/3 of matrigel, 1 mg/ml final concentration of 15 collagen, 0.1N NaOH (0.026x the volume of collagen in μ l) and 1x PBS, and the volume is then adjusted with water. The gels are kept at 37°C for 1 hour so as to allow them to polymerize. Next, the human vein endothelial cells (HUVECs ref: C-12200 – Promocell) were seeded at 15×10^3 or 6×10^3 cells/well in 400 or 120 μ l (for the 8-well or 96-well plates respectively) of EBM medium (Clonetics C3121) + 20 2% FBS + 10 μ g/ml hEGF. They were stimulated with 1 or 3 ng/ml of FGF-2 (R&D systems, 133-FB-025; Invitrogen, PHG0026) for 24 h at 37°C in the presence of 5% CO₂. After 24 hours, the length of the network of microtubules formed was measured using a computer-assisted image analysis system (Imagenia Biocom, Courtaboeuf, France) and the total length of the pseudotubules in each well was 25 determined. The average total length of the microcapillary network was calculated in μ m for each condition corresponding to the average of 6 replicates.

Stimulation with FGF2 makes it possible to induce the formation of new tubules. An FGF-R antagonist is considered to be active in this test as long as it is capable of partially inhibiting this angiogenesis at a dose less than or equal to 30 300 nM.

Example of screening for FGF-R antagonists

In this experiment, the molecules are evaluated at 3 and 30 nM on induction of the angiogenesis of HUVEC human cells by FGF-2. Compounds No. 71, 72

(example 11) and 68 are declared active since they exhibit an inhibitory activity of pseudotubule formation which is greater than or equal to 20% at a dose less than or equal to 300 nM.

- 5 Table 1: *In vitro* angiogenesis of HUVEC cells stimulated with FGF-2 and effect of FGF-R antagonists (inhibition of angiogenesis as a percentage of the control)

Compounds No.	3 nM	30 nM
71	-1	41
72	36	24
68	37	52

Example 19: FGF-2-induced *in vitro* proliferation of HUVEC cells

- 10 In order to demonstrate the ability of the FGF-R antagonists of the present invention to inhibit FGF-induced cell proliferation, *in vitro* proliferation experiments were carried out with human endothelial cells of HUVEC type stimulated with FGF-2 or b-FGF.

- To do this, HUVEC human vein endothelial cells (promocell, C-12200) are
 15 seeded at a rate of 5000 cells per well of a 96-well plate (Biocoat collagen I cellware, Becton Dickinson 354650) in 100 μ l of RPMI 1640 deprivation medium (Invitrogen, 31872-025) supplemented with 0.5% or 1% FCS, 2 mM glutamine, 1x sodium pyruvate (Invitrogen, 11360-039) and 1x NEAA (Invitrogen, 11140-035), overnight at 37°C in the presence of 5% CO₂. The following morning, the medium
 20 is suctioned-off and replaced with 50 μ l of deprivation medium containing the antagonists at a 2x concentration, to which are added 50 μ l of FGF-2 (R&D systems, 133-FB-025; Invitrogen, PHG0026) at 0.2 ng/ml (i.e. 2x). After 48 or 72 h, 100 μ l of Cell Titer-GLO™ Luminescent Cell Viability Assay (Promega, G7571) are added for 10 min in order to measure, by means of a luminometer, the amount of
 25 ATP present in the cells and which is in relation to the number of cells per well corresponding to the cell proliferation.

The antagonists of the present invention are considered to be active as long as they are capable of inhibiting FGF-2-induced proliferation of HUVEC cells at a dose less than or equal to 300 nM.

Example of HUVEC cell proliferation induced by FGF-2 and inhibited by FGF-R antagonists

Compounds No. 49 (example 4), 71 and 72 (example 11) are capable of inhibiting the FGF-2-induced cell proliferation since, in their presence, a reduction
5 in proliferation of greater than or equal to 20% is observed for doses less than or equal to 300 nM.

Table 2: Cell proliferation of HUVEC cells stimulated with FGF-2 and effect of FGF-R antagonists (inhibition of proliferation as percentage of the control)

Compounds No.	30 nM	300 nM
49	9	29
71	35	34
72	38	55

10

More generally, all the compounds according to the invention are active, at the dose of 300 nM, in *in vitro* angiogenesis of HUVEC cells induced by FGF-2 or in *in vitro* proliferation of HUVEC cells induced by FGF-2.

15 **Example 20: Model of inflammatory angiogenesis in mice**

Angiogenesis is required for the development of chronic inflammatory diseases such as rheumatoid arthritis. The formation of new vessels allows not only the perfusion of pathological tissues, but also the transport of cytokines responsible for establishing the chronicity of the disease.

20 The model described by Colville-Nash *et al.*, in 1995, makes it possible to study pharmacological agents capable of modulating the occurrence of angiogenesis in an inflammatory context. The model is developed on OF1 female mice (Charles River Laboratories) weighing approximately 25 g, and by groups of 12. The animals are anaesthetized with sodium pentobarbital (60 mg/kg; Sanofi
25 Nutrition Santé animale)) intraperitoneally. An air pocket is created on the back of the mouse by subcutaneous injection of 3 ml of air. After they have awoken, the animals receive a treatment generally by gavage, and receive an injection of 0.5 ml of Freund's adjuvant (Sigma) with 0.1% of croton oil (Sigma) in the pocket. Seven days later, the mice are again anaesthetized and placed on a hot plate at 40°C.
30 One ml of carmine red (Aldrich Chemicals, 5% in 10% of gelatin) is injected into the

tail vein. The animals are then placed at 4°C for 2-3 hours. The skins are then taken and dried for 24 h in an oven at 56°C. The dry tissues are weighed and placed in 1.8 ml of digestion solution (2 mM dithiothreitol, 20 mM Na₂HPO₄, 1 mM EDTA, 12 U/ml papain) for 24 h. The dye is then dissolved in 0.2 ml of 5M NaOH.

- 5 The skins are centrifuged at 2000 rpm for 10 min at ambient temperature. The supernatants are filtered through 0.2 µm cellulose acetate membranes. The filtrates are read in a spectrophotometer at 492 nm against a carmine red calibration range. Two parameters are studied: the dry weight of the granuloma and the amount of dye after digestion of the tissues. The results are expressed as
- 10 mean values (± sem). The differences between the groups are tested with an ANOVA followed by a Dunnett's test, of which the reference group is the "solvent control" group.

The FGF-R antagonists are evaluated between 1 and 50 mg/kg using methylcellulose/tween (0.6% v/v) as vehicle or any other vehicle which allows the

15 active ingredient to be solubilized. The molecules are administered daily, orally (one or two times a day) by gavage. The antagonists of the present invention are considered to be active as long as they enable a significant reduction in the angiogenic parameter, i.e. a reduction in the amount of carmine red dye in the skins of the animals tested.

20

Example of evaluation of FGF-R antagonists in the model of inflammatory angiogenesis in mice. Compounds No. 49 (example 4) and 72 (example 11) at 10 mg/kg, after one week of daily treatment, significantly reduce the two

25 parameters measured: the weight of the granuloma (dry weight of the skin) corresponding to the inflammation part of the model, and the dye content corresponding to the angiogenesis.

Table 3: Effect of the FGF-R antagonists, in a model of inflammatory angiogenesis, on the dry weight of the skins or on their content of carmine red dye.

30

Model of inflammatory angiogenesis	% inhibition of the inflammatory parameter (mass of the granuloma)	% inhibition of the angiogenic parameter (dye content)
Compound No. 49 (example 4); 10 mg/kg	40	46
Compound No. 223 (example 17); 10 mg/kg	25	43
Compound No. 72 (example 11); 10 mg/kg	38	43
Compound No. 71; 30 mg/kg	28	40
Compound No. 149; 10 mg/kg	14	30
Compound No. 215; 10 mg/kg	21	39
Compound No. 10 (example 2); 30 mg/kg	19	19

Example 21: 4T1 orthotopic mammary carcinoma model in mice

In order to evaluate the effect of the FGF-R antagonists in a murine tumour model, 4T1 mouse mammary carcinoma cells are injected into the mammary gland.

- 5 The cells proliferate until the formation of a tumour after infiltration of the cells of the tumour microenvironment.

The 4T1 cells are cultured in RPMI 1640 medium containing 10% FCS and 1% glutamine, supplemented with 1 mg/ml of geneticin. On the day of the injection into the mouse, the 4T1 cell concentration is adjusted to 2×10^6 cells/ml in PBS in order to inject 1×10^5 cells in 50 μ l.

Mice (Balb/c, female, Charles River, approximately 8 $\pm$ 2 weeks old) are anaesthetized by intraperitoneal injection of a mixture of 5% Rompun (xylazine), 10% Imalgene (ketamine) and 85% NaCl, in a proportion of 10 ml/kg. The injection zone (top-right nipple) is disinfected with hexomedine. After having vortexed the cells, 50 μ l are removed in a syringe and injected into the nipple with a 26G needle. The day of injection corresponds to D1. There are 15 mice in each group of mice (10 mice will be devoted to the ELISA assays and 5 mice to the histology). The FGF-R antagonists are evaluated at between 1 and 50 mg/kg in methylcellulose/tween (0.6% v/v) or any other vehicle which makes it possible to solubilize the active ingredient. The molecules are administered daily, orally (one

or two times a day) by gavage, this taking place from D5 to D21, which is the day before the samples are taken. From D5, the tumours are measured as soon as possible, every two days, or even every day at the end of the experiment, using a caliper (sliding caliper). It is done in the following way: the longest length (L) and the perpendicular to the centre (l) are measured in mm. The volume in mm³ is then defined by means of the mathematical formula which determines the volume of an ellipsoid: $(l^2 \times L) \times 0.52$. On the day the samples are taken, generally D22, the mice are sacrificed by means of an excess of sodium pentobarbital after having measured the volume of the tumours. The tumours are then cleared, photographed and weighed. The lungs are also removed and the metastases are counted after boin staining.

The antagonists of the present invention are considered to be active as long as they allow a significant reduction in the volume of the tumour and/or any number of lung metastases.

Example of 4T1 mammary carcinoma in mice

The compounds considered to be active in the inflammatory angiogenesis model are evaluated in the 4T1 mammary carcinoma model in mice at between 1 and 50 mg/kg and showed a reduction in tumour volume of up to 49% and a decrease in the number of lung metastases of up to 33%.

It therefore appears that the compounds of formula (I) according to the present invention, by virtue of their FGF antagonist effect, reduce *in vitro* and *in vivo*, angiogenesis, tumour growth and metastasization.

Generally, FGFs and their receptors play an important role, by means of autocrine, paracrine or juxtacrine secretions, in phenomena where there is dysregulation of the stimulation of cancer cell growth. Furthermore, FGFs and their receptors affect tumour angiogenesis which plays a predominant role both on tumour growth and also on metastasization phenomena.

Angiogenesis is a process in which new capillary vessels are generated from pre-existing vessels or by mobilization and differentiation of bone marrow cells. Thus, both uncontrolled proliferation of endothelial cells and mobilization of angioblasts from the bone marrow are observed in tumour neovascularization processes. It has been shown, *in vitro* and *in vivo*, that several growth factors stimulate endothelial

proliferation, and in particular FGF-1 or a-FGF and FGF-2 or b-FGF. These two factors induce proliferation, migration and protease production by endothelial cells in culture and neovascularization *in vivo*. a-FGF and b-FGF interact with endothelial cells by means of two classes of receptors, high-affinity receptor

5 tyrosine kinases (FGF-Rs) and low-affinity receptors of heparin sulphate proteoglycan type (HSPGs) located at the surface of cells and in extracellular matrices. Although the paracrine role of these two factors on endothelial cells is widely described, these FGFs could also intervene on the cells through an

10 autocrine process. Thus, FGFs and their receptors represent very relevant targets for therapies aimed at inhibiting angiogenesis processes (Keshet E, Ben-Sasson SA., *J. Clin. Invest.*, (1999), Vol. 501, pp. 104-1497; Presta M, Rusnati M, Dell'Era P, Tanghetti E, Urbinati C, Giuliani R *et al*, *New York: Plenum Publishers*, (2000), pp. 7-34, Billottet C, Janji B, Thiery J.P., Jouanneau J, *Oncogene*, (2002) Vol. 21, pp. 8128-8139).

15 Moreover, systematic studies aimed at determining the expression due to FGFs and their receptors (FGF-Rs) of various types of tumour cells demonstrate that a cell response to these two factors is functional in a large majority of human tumour lines studied. These results support the hypothesis that an FGF receptor

20 antagonist could also inhibit tumour cell proliferation (Chandler LA, Sosnowski BA, Greenlees L, Aukerman SL, Baird A, Pierce GF., *Int.J.Cancer*, (1999), Vol. 58, pp. 81-451).

FGFs play an important role in the growth and maintenance of prostate cells. It has

25 been shown, both in animal models and in humans, that an impairment in the cell response to these factors plays an essential role in the progression of prostate cancer. Specifically, in these pathological conditions, both an increase in the production of a-FGF, b-FGF, FGF-6, FGF-8 etc., by the fibroblasts, stromal cells, residual basal cells and endothelial cells present in the tumour and an increase in

30 the expression of FGF receptors and ligands by the tumour cells are recorded. Thus, a paracrine stimulation of prostate cancer cells takes place, and this process appears to be a major component of this pathological condition. A compound which has an FGF receptor antagonist activity, such as the compounds of the present invention, may represent a therapy of choice in these pathological conditions (Giri

35 D, Ropiquet F., *Clin.Cancer Res.*, (1999), Vol. 71, pp. 5-1063; Doll JA, Reiher FK, Crawford SE, Pins MR, Campbell SC, Bouck NP., *Prostate*, (2001), Vol. 305, pp. 49-293) (Sahadevan *et al.*, 2007) (Kwabi-Addo *et al.*, 2004).

Several studies show the presence of FGFs and of their receptors, FGF-Rs, both in human breast tumour lines (in particular MCF7) and in tumour biopsies. These factors appear to be responsible, in this pathological condition, for the appearance of the very aggressive phenotype and induce a strong metastasization. Thus, a compound which has FGF-R receptor antagonist activity, such as the compounds of formula I, may represent a therapy of choice in these pathological conditions (Vercoutter-Edouart A-S, Czeszak X, Crépin M, Lemoine J, Boilly B, Le Bourhis X *et al.*, *Exp. Cell Res.*, (2001), Vol. 262, pp. 59-68) (Schwertfeger, 2009).

Cancerous melanomas are tumours which induce metastases with a high frequency and which are very resistant to the various chemotherapy treatments. The angiogenesis processes play a predominant role in the progression of a cancerous melanoma. Furthermore, it has been shown that the probability of the occurrence of metastases increases very greatly with the increase in the vascularization of the primary tumour. Melanoma cells produce and secrete various angiogenic factors, including a-FGF and b-FGF. Moreover, it has been shown that inhibition of the cellular effect of these two factors by means of the soluble FGF-R1 receptor blocks melanoma tumour cell proliferation and survival *in vitro* and blocks tumour progression *in vivo*. Thus, a compound which has an FGF receptor antagonist activity, such as the compounds of the present invention, may represent a therapy of choice in these pathological conditions (Rofstad EK, Halsor EF., *Cancer Res.*, (2000); Yayon A, Ma Y-S, Safran M, Klagsbrun M, Halaban R., *Oncogene*, (1997), Vol. 14, pp. 2999-3009).

Glyoma cells produce a-FGF and b-FGF *in vitro* and *in vivo*, and have various FGF receptors at their surface. This therefore suggests that these two factors play a pivotal role, by means of an autocrine and paracrine effect, in the progression of this type of tumour. Furthermore, like most solid tumours, the progression of gliomas and their ability to induce metastases is highly dependent on the angiogenic processes in the primary tumour. It has also been shown that FGF-R1 receptor antisenses block human astrocytoma proliferation. In addition, naphthalenesulphonate derivatives are described for inhibiting the cellular effects of a-FGF and b-FGF *in vitro* and the angiogenesis induced by these growth factors *in vivo*. An intracerebral injection of these compounds induces a very significant increase in apoptosis and a considerable decrease in angiogenesis, reflected by a considerable regression of gliomas in rats. Thus, a compound which has an a-FGF

antagonist and/or b-FGF antagonist and/or FGF receptor antagonist activity, such as the compounds of the present invention, may represent a therapy of choice in these pathological conditions (Yamada SM, Yamaguchi F, Brown R, Berger MS, Morrison RS, *Glia*, (1999), Vol. 76, pp. 28-66; Auguste P, Gürsel DB, Lemièrre S, Reimers D, Cuevas P, Carceller F *et al.*, *Cancer Res.*, (2001), Vol. 26, pp. 61-1717) (Loilome *et al.*, 2008).

Active angiogenesis is also described for hepatocarcinomas or hepatocellular carcinoma (HCC). *In vivo*, tumour progression in HCCs requires a considerable supply of oxygen and nutrients. Hepatocarcinomas are tumours which are typically angiogenic, because a drastic modification is observed with respect to arterial vascularization, and this results in the acquisition of an invasive and metastatic potential (Tanaka *et al.*, 2006). FGFs participate actively in the development of tumour angiogenesis within HCCs and are frequently associated with the inflammatory process. They are also overexpressed in the context of chronic hepatitis and liver sclerosis (Uematsu *et al.*, 2005) and the serum FGF level has been correlated with the clinicopathological progression of HCCs. Furthermore, the FGF-R4 receptor, and also FGF-R1, have been described as participating actively in HCC tumour genesis (Huang *et al.*, 2006) (Nicholes *et al.*, 2002). The antagonists of the present invention may therefore be a treatment of choice for hepatocellular carcinomas or hepatocarcinomas.

In lung cancers of NSCLC (Non-Small Cell Lung Cancer) type, recent studies show that b-FGF, FGF-9, FGF-R1 and FGF-R2 are regularly coexpressed in NSCLC cancer lines and especially in those resistant to anti-EGFR treatment such as gefitinib. These expressions are connected to the capacity for proliferation via autocrine cell signalling and anchorage-independent growth of tumours of NSCLC type and mainly the type insensitive to treatment with gefitinib (Marek *et al.*, 2008). Furthermore, b-FGF has been suggested as playing an important role in the survival of NSCLC cells during treatment by chemotherapy, by inducing the overexpression of the anti-apoptotic proteins BCL-2, BCL-X, XIAP or BIRC3 (Pardo *et al.*, 2002, 2003 and 2006). Thus, an FGF receptor antagonist, such as those of the present invention, may represent a therapy of choice for lung cancers of NSCLC type, alone or in combination with EGF receptor inhibitors or chemotherapies.

In approximately 10% of gastric cancers, this FGF-R2 gene amplification is observed. This amplification is associated with a poor vital prognosis for cancers of diffuse type. The proliferation of tumour cells may be ligand-independent or dependent on paracrine activation by FGF-7 (Turner *et al.*, 2010). The antagonists
5 of the present invention may therefore be a treatment of choice for gastric cancers.

More recently, the potential role of pro-angiogenic agents in leukaemias and lymphomas has been documented. Indeed, in general, it has been reported that cell clones in these pathological conditions can be destroyed naturally by the
10 immune system or switch into an angiogenic phenotype which promotes their survival and then their proliferation. This change in phenotype is induced by an overexpression of angiogenic factors, in particular by macrophages, and/or a mobilization of these factors from the extracellular matrix (Thomas DA, Giles FJ, Cortes J, Albitar M, Kantarjian HM., *Acta Haematol.*, (2001), Vol. 207, pp.106-190).
15 Among the angiogenic factors, b-FGF has been detected in many lymphoblastic and hematopoietic tumour cell lines. FGF receptors are also present on the majority of these lines, suggesting a possible autocrine cellular effect of a-FGF and b-FGF inducing proliferation of these cells. Moreover, it has been reported that bone marrow angiogenesis via paracrine effects is correlated with the progression
20 of some of these pathological conditions.

More particularly, it has been shown, in CLL (chronic lymphocytic leukaemia) cells, that b-FGF induces an increase in anti-apoptotic protein (Bcl2) expression, resulting in an increase in the survival of these cells, and that it therefore
25 participates considerably in their cancerization. In addition, the b-FGF levels measured in these cells are very well-correlated with the stage of clinical advancement of the disease and the resistance to the chemotherapy applied in this pathological condition (fludarabine). Thus, a compound which has an FGF receptor antagonist activity, such as the compounds of the present invention, may represent
30 a therapy of choice, either alone or in combination with fludarabine or other products that are active in this pathological condition (Thomas DA, Giles FJ, Cortes J, Albitar M, Kantarjian HM., *Acta Haematol.*, (2001), Vol. 207, pp. 106-190; Gabrilove JL, *Oncologist*, (2001), Vol. 6, pp. 4-7).

35 Furthermore, it has been shown in many recent studies that FGFs and FGF-Rs participate actively in the resistance of tumour and/or endothelial cells to treatments by chemotherapy, radiotherapy or else anti-VEGF treatments. These

resistances use various cell mechanisms, such as protection against apoptosis by positive regulation of the Bcl-xl protein by FGF-R4 in the case of breast cancer resistance to doxorubicin (Roidl *et al.*, 2009) or by FGF-2 production in the case of resistance of bladder tumours to cisplatin (Miyake *et al.*, 1998), by activation of the PI3K/AKT pathway by the FGF2/FGF-R1 couple in the case of resistance of acute myeloid leukaemia cells to cytarabin (Karajannis *et al.*, 2006), by stimulation of the RAS/MAP-K, PI3-K and mTOR pathway by FGF-1 for certain breast tumours resistant to anti-oestrogen treatments (Manuvakhova *et al.*, 2006). The FGFs/FGF-Rs couple is also involved in resistance to anti-VEGF treatments in the case of pancreatic carcinomas (Casanovas *et al.*, 2005) or of glioblastomas (Batchelor *et al.*, 2007) or else in radiotherapy resistance phenomena (Gu *et al.*, 2004; Moyal *et al.*, 2009). Thus, the compounds of the present invention could be combined with existing therapies in order to limit the appearance of resistance phenomena.

Furthermore, tumour invasion, which is one of the marks of malignancy, consists of the translocation of tumour cells from the initial neoplastic locus to the surrounding host tissues, allowing the tumour to penetrate into the vascular endothelium in order to circulate and to form metastatic loci remote from the primary tumour. An increasing number of recent articles suggest that changes in the tissue architecture at the periphery of the tumour appear to be responsible for the epithelial-mesenchymal transition (EMT) process. EMT is a cell process by which epithelial cells modulate their phenotype and acquire mesenchymal cell properties through the disruption of intercellular adhesion and an increase in cell motility, thus playing an essential role in tumour progression by conferring an invasive and metastatic phenotype on carcinomas. Growth factors such as FGFs participate in this cell process by virtue of their stimulatory activity on cell migration and invasion, but also, as regards FGF receptors, by virtue of their ability to interact with cadherins, thus facilitating tumour cell migration (Cowin *et al.*, 2005). The FGF-R antagonists described herein may be used for preventing these metastatic phases in a large number of cancers.

A correlation exists between the bone marrow angiogenesis process and "extramedullar disease" in CML (chronic myelomonocytic leukaemia). Various

studies demonstrate that the inhibition of angiogenesis, in particular by means of a compound which has an FGF receptor antagonist activity, could represent a therapy of choice in this pathological condition.

- 5 The proliferation and migration of vascular smooth muscle cells contributes to intimal hypertrophy of the arteries and thus plays a predominant role in atherosclerosis and in restenosis after angioplasty and endarterectomy.

10 *In vivo* studies show, after lesion of the carotid "balloon injury", a local production of a-FGF and of b-FGF. In this same model, an anti-FGF2 neutralizing antibody inhibits vascular smooth muscle cell proliferation and thus decreases intimal hypertrophy.

15 A chimeric protein consisting of FGF2 linked to a molecule such as saporin inhibits vascular smooth muscle cell proliferation *in vitro* and intimal hypertrophy *in vivo* (Epstein CE, Siegall CB, Biro S, Fu YM, FitzGerald D., *Circulation*, (1991), Vol. 87, pp. 84-778; Waltenberger J., *Circulation*, (1997), pp. 96-4083).

20 Thus, FGF receptor antagonists, such as the compounds of the present invention, represent a therapy of choice, either alone or in combination with compounds that are antagonists of other growth factors involved in these pathological conditions, such as PDGF, in the treatment of pathological conditions related to vascular smooth muscle cell proliferation, such as atherosclerosis, post-angioplasty restenosis or restenosis following the implantation of endovascular prostheses
25 (stents) or during aortocoronary bypasses.

Cardiac hypertrophy occurs in response to a stress of the ventricular wall induced by an overload in terms of pressure or volume. This overload can be the consequence of numerous physiopathological states, such as hypertension, AC
30 (aortic coarctation), myocardial infarction, and various vascular disorders. The consequences of this pathological condition are morphological, molecular and functional changes such as cardiac myocyte hypertrophy, matrix protein accumulation and foetal gene reexpression. b-FGF is implicated in this pathological condition. Specifically, the addition of b-FGF to cultures of newborn rat
35 cardiomyocytes modifies the profile of the genes corresponding to the contractile proteins, resulting in a foetal-type gene profile. In a complementary manner, adult rat myocytes show a hypertrophic response under the effect of b-FGF, this

response being blocked by anti-b-FGF neutralizing antibodies. Experiments carried out *in vivo* in b-FGF-knock-out transgenic mice show that b-FGF is the major factor stimulating cardiac myocyte hypertrophy in this pathological condition (Schultz JeJ, Witt SA, Nieman ML, Reiser PJ, Engle SJ, Zhou M *et al.*, *J.Clin. Invest.*, (1999), Vol. 19, pp. 104-709). Thus, a compound, such as the compounds of the present invention, which has an FGF receptor antagonist activity represents a therapy of choice in the treatment of heart failure and any other pathological condition associated with cardiac tissue degeneration. This treatment could be carried out alone or in combination with the common treatments (beta-blockers, diuretics, angiotensic antagonists, antiarrhythmics, anti-calcium agents, antithrombotics, etc.).

Vascular disorders due to diabetes are characterized by an impairment of vascular reactivity and of blood flow, hyperpermeability, an exacerbated proliferative response and an increase in matrix protein deposits. More specifically, a-FGF and b-FGF are present in the preretinal membranes of patients having diabetic retinopathies, in the membranes of the underlying capillaries and in the vitreous humour of patients suffering from proliferative retinopathies. A soluble FGF receptor capable of binding both a-FGF and b-FGF is developed in diabetes-related vascular disorders (Tilton RG, Dixon RAF, Brock TA., *Exp. Opin. Invest. Drugs*, (1997), Vol. 84, pp. 6-1671). Thus, a compound, such as the compounds of formula I, which has an FGF receptor antagonist activity represents a therapy of choice, either alone or in combination with compounds that are antagonists of other growth factors involved in these pathological conditions, such as VEGF.

Fibrosis is the abnormal formation of scar tissues following a tissue lesion, and results in a chronic and progressive impairment of the affected organs that can result in serious dysfunction of the affected organ. It can occur in all tissues, but is mainly prevalent in organs exposed to chemical or biological attacks, such as the lungs, the skin, the kidneys, the digestive tract, the liver, etc. FGFs participate in this cell process by promoting the production and accumulation of extracellular matrices by fibroblasts, the proliferation of said fibroblasts and infiltration into many organs such as the kidneys or the lungs (Khalil *et al.*, 2005) (Strutz *et al.*, 2003). Antagonists of the activity of these FGFs, such as the molecules of the present invention, may be used alone or in combination in the treatment of fibrosis.

Rheumatoid arthritis (RA) is a chronic disease with an unknown etiology. Although it affects many organs, the most serious form of RA is a progressive synovial inflammation of the joints resulting in destruction. Angiogenesis appears to considerably affect the progression of this pathological condition. Thus, a-FGF and b-FGF have been detected in the synovial tissue and in the joint fluid of patients suffering from RA, indicating that this growth factor is involved in the initiation and/or the progression of this pathological condition. In models of AIA (adjuvant-induced model of arthritis) in rats, it has been shown that the overexpression of b-FGF increases the severity of the disease, whereas an anti-b-FGF neutralizing antibody blocks the progression of RA (Malemud, 2007) (Yamashita A, Yonemitsu Y, Okano S, Nakagawa K, Nakashima Y, Irida T *et al.*, *J.Immunol.*, (2002), Vol. 57, pp. 168-450 ; Manabe N, Oda H, Nakamura K, Kuga Y, Uchida S, Kawaguchi H, *Rheumatol*, (1999), Vol. 20, pp. 38-714). Thus, the compounds according to the invention represent a therapy of choice in this pathological condition.

Recent scientific articles document the involvement of b-FGF in neuropathic pain. Specifically, an increase in astroglial b-FGF production is observed in astrocytes following a spinal cord lesion (Madiat *et al.*, 2003). This b-FGF contributes to neuropathic pain due to contact or allodynia. Treatment using an anti-FGF2 neutralizing antibody reduces this mechanical allodynia (Madiat *et al.*, 2005). The antagonists of the present invention are treatments of choice for pain by inhibiting the effect of FGF-2 on these receptors.

It has also been described that the level of growth factors having a pro-angiogenic activity, such as FGF-1 and -2, are greatly increased in the synovial fluid of patients suffering from osteoarthritis. In this type of pathological condition, a considerable modification is recorded in the balance between the pro- and anti-angiogenic factors inducing the formation of new vessels, and consequently, the vascularization of nonvascularized structures, such as joint cartilages or intervertebral discs. Thus, angiogenesis represents a key factor in bone formation (osteophytes), thus contributing to the progression of the disease. Additionally, the innervation of the new vessels can also contribute to the chronic pain associated with this pathological condition (Walsh DA., *Curr Opin Rheumatol*. 2004 Sep;16(5):609-15). Thus, the compounds according to the invention represent a therapy of choice in this pathological condition.

IBD (inflammatory bowel disease) includes two forms of chronic inflammatory diseases of the intestine: UC (ulcerative colitis) and Crohn's disease (CD). IBD is characterized by an immune dysfunction reflected by an inappropriate production of inflammatory cytokines inducing the establishment of a local microvascular system. This angiogenesis of inflammatory origin results in an intestinal ischemia induced by vasoconstriction. High circulating and local levels of b-FGF have been measured in patients suffering from these pathological conditions (Kanazawa S, Tsunoda T, Onuma E, Majima T, Kagiya M, Kkuchi K., *American Journal of Gastroenterology*, (2001), Vol. 28, pp 96-822 ; Thorn M, Raab Y, Larsson A, Gerdin B, Hallgren R., *Scandinavian Journal of Gastroenterology*, (2000), Vol. 12, pp. 35-408). The compounds of the invention which exhibit a high anti-angiogenic activity in an inflammatory angiogenesis model represent a therapy of choice in these pathological conditions.

Another disease which has a considerable inflammatory component and for which a strong implication of FGFs and FGF-Rs is described is benign prostatic hyperplasia (BPH). BPH is a disease related to ageing which is characterized by hyperplasia of the glandular tissues and of the stroma around the urethra until it becomes obstructed. At the cellular level, this pathological condition involves hyperplasia of the basal cells, an increase in the stromal mass, amplified matrix deposit or else a reduction in tissue elasticity (Untergasser *et al.*, 2005). FGFs participate in the development of this disease by stimulating the proliferation of the prostatic stroma and epithelial cells, and in particular FGF-7 or KGF, but also FGF-2 or FGF-17 (Wang 2008 , Boget 2001, Giri 2001). In addition, FGFs promote the transdifferentiation step by modifying epithelial cell/stromal cell interactions, in combination with TGF- β (Untergasser 2005). Finally, certain receptors, such as FGF-R1, are overexpressed in BPH, promoting induction of the pathological condition and potentiating the paracrine effects of FGF-2 (Boget 2001). An antagonist of the effect of these FGFs is therefore a treatment of choice for benign prostatic hyperplasia.

Psoriasis is a chronic skin disease caused by a hyperproliferation of the epidermal keratinocytes, while clear cell acanthoma (CCA) is a benign neoplasm of the epidermis which also involves an abnormal proliferation of keratinocytes. These two skin diseases have similar histological characteristics despite different underlying causes: a thickening of the epidermis, inflammatory infiltrations of lymphocytes and neutrophils, dilation and tortuosity of the papillary capillaries. In

both cases, KGF or FGF-7 plays a predominant role in the development of the pathological condition (Kovacs *et al.*, 2006) (Finch *et al.*, 1997). The use of the antagonists of the present invention may make it possible to slow down the development of such skin diseases.

5

FGF-R1, -R2 and -R3 receptors are involved in chronogenesis and osteogenesis processes. Mutations resulting in the expression of FGF-Rs that are always activated have been connected to a large number of human genetic diseases reflected by malformations of the skeleton, such as Pfeiffer syndrome, Crouzon syndrome, Apert syndrome, Jackson-Weiss syndrome and Bear-Stevenson cutis gyrate syndrome. Some of these mutations affect more particularly the FGF-R3 receptor, resulting in particular in achondroplasias (ACH), hyperchondroplasias (HCH) and TD (thanatophoric dysplasia); ACH being the most common form of dwarfism. From a biochemical point of view, the sustained activation of these receptors takes place via a dimerization of the receptor in the absence of ligand (Chen L., Adar R., Yang X. Monson E.O., Li C., Hauschka P.V, Yagon A. and Deng C.X., (1999), *The Journ. Of Clin. Invest.*, Vol. 104, n° 11, pp. 1517-1525). Thus, the compounds of the invention which exhibit an FGF antagonist or FGF receptor antagonist activity and which inhibit FGF-R-dependent intracellular signalling represent a therapy of choice in these pathological conditions.

It is also known that adipose tissue is one of the rare tissues that, in adults, can develop or regress. This tissue is highly vascularized and a very dense network of microvessels surrounds each adipocyte. These observations have resulted in the testing of the effect of anti-angiogenic agents on adipose tissue development in adults. Thus, it appears that, in pharmacological models in ob/ob mice, the inhibition of angiogenesis is reflected by significant weight loss in the mice (Rupnick MA *et al.*, (2002), *PNAS*, Vol. 99, n°16, pp. 10730-10735). Furthermore, FGFs appear to be key regulators of adipogenesis in humans (Hutley *et al.*, 2004). Thus, an FGF receptor antagonist compound which has a powerful anti-angiogenic activity may represent a therapy of choice in obesity-related pathological conditions.

By virtue of their low toxicity and their pharmacological and biological properties, the compounds of the present invention are of use in the treatment and prevention of any carcinoma which has a high degree of vascularization, such as lung, breast, prostate, oesophageal, pancreatic, liver, colon or kidney carcinomas, or which

induces metastases, such as colon, breast, liver or stomach carcinomas, or melanomas, or which is sensitive to a-FGF or to b-FGF in an autocrine manner or else in pathological conditions of glioma type, lymphomas and leukaemias or, finally, in any therapy-resistance phenomenon. These compounds represent a
5 therapy of choice, either alone or in combination with a chemotherapy, a radiotherapy or any other suitable treatment. The compounds according to the invention are also of use in the treatment and prevention of cardiovascular diseases, such as atherosclerosis, or restenosis post-angioplasty, in the treatment of diseases related to complications occurring following the implantation of
10 endovascular stents and/or aortocoronary bypasses or other vascular grafts, and cardiac hypertrophy or vascular complications of diabetes, such as diabetic retinopathies. The compounds according to the invention are also of use in the treatment and prevention of chronic inflammatory diseases such as rheumatoid arthritis, IBD or benign prostatic hyperplasia. Finally, the compounds according to
15 the invention can be used in the treatment and prevention of achondroplasias (ACH), hypochondroplasias (HCH) and TD (thanatophoric dysplasia), as also in the treatment of obesity.

The products according to the invention are also of use in the treatment and
20 prevention of macular degeneration, in particular age-related macular degeneration (or ARMD). A major characteristic of the loss of sight in adults is the neovascularization and the subsequent haemorrhages which cause considerable functional disorders in the eye and which are reflected by early blindness. Recently, studying the mechanisms involved in ocular neovascularization
25 phenomena has made it possible to demonstrate the involvement of pro-angiogenic factors in these pathological conditions. By using a laser-induced choroidal neoangiogenesis model, it has been possible to confirm that the products according to the invention also make it possible to modulate neovascularization of the choroid.

30 Moreover, the products of the invention can be used in the treatment or prevention of thrombopenias due in particular to anticancer chemotherapy. It has in fact been demonstrated that the products of the invention can improve circulating platelet levels during chemotherapy.

35 Finally, the products according to the invention are of use in the treatment and prevention of skin diseases, such as psoriasis or clear cell acanthoma, in

combating the progression of liver, kidney or lung fibrosis, and also in the treatment of neuropathic pain.

5 A subject of the invention is, according to another of its aspects, medicaments which comprise a compound of formula (I), or an addition salt thereof with a pharmaceutically acceptable acid or base, or else a hydrate or a solvate of the compound of formula (I).

10 According to another of its aspects, the present invention relates to pharmaceutical compositions comprising, as active ingredient, a compound of formula (I) according to the invention. These pharmaceutical compositions contain an effective dose of at least one compound according to the invention, or a pharmaceutically acceptable salt or a hydrate or solvate of said compound, and also at least one pharmaceutically acceptable excipient. Said excipients are selected, according to
15 the pharmaceutical form and the method of administration desired, from the usual excipients which are known to those skilled in the art.

In the pharmaceutical compositions of the present invention for oral, sublingual, subcutaneous, intramuscular, intravenous, topical, local, intratracheal, intranasal,
20 transdermal or rectal administration, the active ingredient of formula (I) above, or optional salt, solvate or hydrate thereof, can be administered in unit administration form, as a mixture with conventional pharmaceutical excipients, to animals and to human beings for the prophylaxis or the treatment of the disorders or the diseases mentioned above.

25 The suitable unit administration forms comprise forms for oral administration, such as tablets, soft or hard gel capsules, powders, granules and oral solutions or suspensions, sublingual, buccal, intratracheal, intraocular or intranasal administration forms, forms for administration by inhalation, topical, transdermal,
30 subcutaneous, intramuscular or intravenous administration forms, rectal administration forms, and implants. For topical application, the compounds according to the invention can be used in creams, gels, ointments or lotions.

The pharmaceutical compositions according to the present invention are preferably
35 administered orally.

By way of example, a unit administration form of a compound according to the invention in tablet form may comprise the following components:

	Compound according to the invention	50.0 mg
	Mannitol	223.75 mg
5	Sodium croscarmellose	6.0 mg
	Maize starch	15.0 mg
	Hydroxypropylmethylcellulose	2.25 mg
	Magnesium stearate	3.0 mg

- 10 The present invention also relates to a pharmaceutical composition as defined above, as a medicament.

A subject of the present invention is also the use of a compound of formula (I), as defined above, for use thereof in the treatment and prevention of diseases
15 requiring a modulation of FGFs.

A subject of the present invention is also the use of a compound of formula (I), as defined above, for use thereof in the treatment and prevention of cancers, in particular carcinomas which have a high degree of vascularization, such as lung,
20 breast, prostate, pancreatic, colon, kidney and oesophageal carcinomas, cancers which induce metastases, such as colon cancer, liver cancer and stomach cancer, melanomas, gliomas, lymphomas and leukaemias.

A compound of formula (I) according to the present invention can be administered
25 alone or in combination with one or more compound(s) which has (have) an anti-angiogenic activity or with one or more cytotoxic compound(s) (chemotherapy), or else in combination with a radiation treatment. Thus, a subject of the present invention is also the use of a compound of formula (I), as defined above, in combination with one or more anticancer active ingredyent(s) and/or with
30 radiotherapy.

A subject of the present invention is also the use of a compound of formula (I), as defined above, in the treatment and prevention of cardiovascular diseases, such as atherosclerosis or post-angioplasty restenosis, diseases related to complications
35 occurring following the implantation of endovascular stents and/or aortocoronary bypasses or other vascular grafts, cardiac hypertrophy, or vascular complications of diabetes, such as diabetic retinopathies.

A subject of the present invention is also the use of a compound of formula (I), as defined above, in the treatment or prevention of chronic inflammatory diseases such as rheumatoid arthritis or IBD.

5

A subject of the present invention is also the use of a compound of formula (I), as defined above, in the treatment or prevention of osteoarthritis, achondroplasias (ACH), hypochondroplasias (HCH) and TD (thanatophoric dysplasia).

- 10 A subject of the present invention is also the use of a compound of formula (I), as defined above, in the treatment or prevention of obesity.

A subject of the present invention is also the use of a compound of formula (I), as defined above, in the treatment or prevention of macular degeneration, such as age-related macular degeneration (ARMD).

15

The compositions according to the invention, for oral administration, contain recommended doses of 0.01 to 700 mg. There may be particular cases where higher or lower dosages are appropriate; such dosages do not depart from the context of the invention. According to the usual practice, the dosage appropriate for each patient is determined by the physician according to the method of administration and the age, weight and response of the patient, and also according to the degree of progression of the disease.

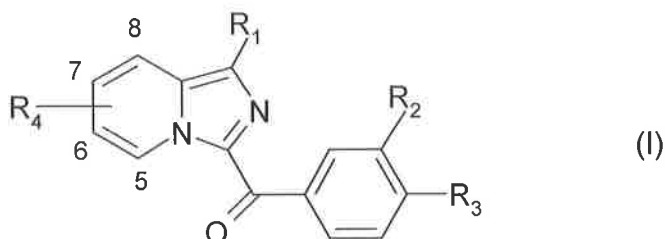
20

- 25 According to another of its aspects, the present invention also relates to a method for treating the pathological conditions indicated above, which comprises the administration, to a patient, of an effective dose of a compound according to the invention, or a pharmaceutically acceptable salt, hydrate or solvate thereof.

30

CLAIMS

1. A compound of formula (I):



in which:

- R_1 represents

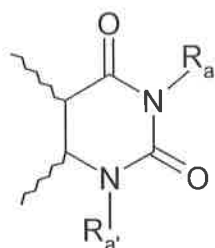
- . a hydrogen or halogen atom,
- . an alkyl group optionally substituted with $-\text{COOR}_5$,
- . an alkenyl group optionally substituted with $-\text{COOR}_5$,
- . a $-\text{COOR}_5$ or $-\text{CONR}_5\text{R}_6$ group,
- . an $-\text{NR}_5\text{COR}_6$ or $-\text{NR}_5\text{SO}_2\text{R}_6$ group,

or

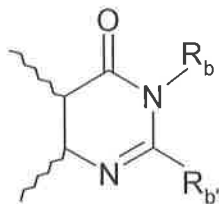
. an aryl group or a heteroaryl group, said aryl or heteroaryl group being optionally substituted with one or more groups selected from: halogen atoms, alkyl groups, cycloalkyl groups, $-\text{COOR}_5$, $-\text{CF}_3$, $-\text{OCF}_3$, $-\text{CN}$, $-\text{C}(\text{NH}_2)\text{NOH}$, $-\text{OR}_5$, $-\text{O-Alk-COOR}_5$, $-\text{O-Alk-NR}_5\text{R}_6$, $-\text{O-Alk-NR}_7\text{R}_8$, $-\text{Alk-OR}_5$, $-\text{Alk-COOR}_5$, $-\text{CONR}_5\text{R}_6$, $-\text{CO-NR}_5\text{OR}_6$, $-\text{CO-NR}_5\text{SO}_2\text{R}_7$, $-\text{CONR}_5\text{-Alk-NR}_5\text{R}_6$, $-\text{CONR}_5\text{-Alk-NR}_7\text{R}_8$, $-\text{Alk-NR}_5\text{R}_6$, $-\text{NR}_5\text{R}_6$, $-\text{NC}(\text{O})\text{N}(\text{CH}_3)_2$, $-\text{CO-Alk}$, $-\text{CO}(\text{OAlk})_n\text{OH}$, $\text{COO-Alk-NR}_5\text{R}_6$, $\text{COO-Alk-NR}_7\text{R}_8$ and 5-membered heteroaryl groups, said heteroaryl groups being optionally substituted with one or more groups selected from halogen atoms and alkyl, $-\text{CF}_3$, $-\text{CN}$, $-\text{COOR}_5$, $-\text{Alk-OR}_5$, $-\text{Alk-COOR}_5$, $-\text{CONR}_5\text{R}_6$, $-\text{CONR}_7\text{R}_8$, $-\text{CO-NR}_5\text{OR}_6$, $-\text{CO-NR}_5\text{SO}_2\text{R}_6$, $-\text{NR}_5\text{R}_6$ and $-\text{Alk-NR}_5\text{R}_6$ groups, or with a hydroxyl group or with an oxygen atom,

- n is an integer ranging from 1 to 3,

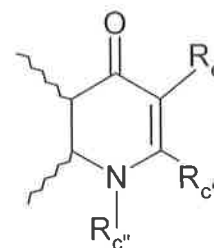
- R_2 and R_3 together form, with the carbon atoms of the phenyl nucleus to which they are attached, a 6-membered nitrogenous heterocycle corresponding to one of formula (A), (B) or (C) below:



(A)



(B)



(C)

in which the wavy lines represent the phenyl nucleus to which R_2 and R_3 are attached and:

. R_a represents a hydrogen atom or an alkyl, haloalkyl, $-\text{Alk}-\text{CF}_3$, $-\text{Alk}-\text{COOR}_5$, $-\text{Alk}'-\text{COOR}_5$, $-\text{Alk}-\text{CONR}_5\text{R}_6$, $-\text{Alk}'-\text{CONR}_5\text{R}_6$, $-\text{Alk}-\text{CONR}_7\text{R}_8$, $-\text{Alk}-\text{NR}_5\text{R}_6$, $-\text{AlkCONR}_5-\text{OR}_6$, $-\text{Alk}-\text{NR}_7\text{R}_8$, $-\text{Alk}-\text{cycloalkyl}$, $-\text{Alk}-\text{O}-\text{R}_5$, $-\text{Alk}-\text{S}-\text{R}_5$, $-\text{Alk}-\text{CN}$, $-\text{OR}_5$, $-\text{OAlkCOOR}_5$, $-\text{NR}_5\text{R}_6$, $-\text{NR}_5-\text{COOR}_6$, $-\text{Alk}-\text{aryl}$, $-\text{Alk}-\text{O}-\text{aryl}$, $-\text{Alk}-\text{O}-\text{heteroaryl}$, $-\text{Alk}-\text{heteroaryl}$ or heteroaryl group, where the aryl or heteroaryl group is optionally substituted with one or more halogen atoms and/or alkyl, cycloalkyl, $-\text{CF}_3$, $-\text{OCF}_3$, $-\text{O}-\text{R}_5$ or $-\text{S}-\text{R}_5$ groups,

. $R_{a'}$ represents a hydrogen atom or a linear, branched, cyclic or partially cyclic alkyl group, or an $-\text{Alk}-\text{OR}_5$, $-\text{Alk}-\text{NR}_5\text{R}_6$ or $-\text{Alk}-\text{NR}_7\text{R}_8$ group, $R_{a'}$ being optionally substituted with one or more halogen atoms,

. R_b represents a hydrogen atom or an alkyl or $-\text{Alk}-\text{COOR}_5$ group,

. $R_{b'}$ represents a hydrogen atom or an alkyl, haloalkyl, cycloalkyl, phenyl or $-\text{Alk}-\text{COOR}_5$ group,

. R_c represents a hydrogen atom or an alkyl, $-\text{CN}$, $-\text{COOR}_5$, $-\text{CO}-\text{NR}_5\text{R}_6$, $-\text{CONR}_7\text{R}_8$, $-\text{CO}-\text{NR}_5-\text{Alk}-\text{NR}_5\text{R}_6$, $-\text{CONR}_5-\text{Alk}-\text{OR}_5$, $-\text{CONR}_5\text{SO}_2\text{R}_5$, $-\text{Alk}-\text{aryl}$ or $-\text{Alk}-\text{heteroaryl}$ group, where the aryl or heteroaryl group is optionally substituted with one or more halogen atoms and/or alkyl, cycloalkyl, $-\text{CF}_3$, $-\text{OCF}_3$, $-\text{O}-\text{alkyl}$ or $-\text{S}-\text{alkyl}$ groups,

. $R_{c'}$ represents a hydrogen atom or an alkyl group,

. $R_{c''}$ represents a hydrogen atom or an alkyl, alkenyl, haloalkyl, cycloalkyl, $-\text{Alk}-\text{NR}_5\text{R}_6$, $-\text{Alk}-\text{NR}_7\text{R}_8$, $-\text{Alk}-\text{OR}_5$ or $-\text{Alk}-\text{SR}_5$ group,

- R_4 , located on position 6, 7 or 8 of the imidazopyridine nucleus, represents:

. a hydrogen atom,

. a $-\text{COOR}_5$ group,

- . a -CO-NR₅-Alk-NR₅R₆ group,
- . a -CO-NR₅-Alk-NR₇R₈ group, or
- . a -CO-NR₅-Alk-OR₆ group,

- R₅ and R₆, which may be identical or different, represent hydrogen atoms, haloalkyl groups or alkyl groups, cycloalkyl groups or an Ms group,

- R₇ and R₈, which may be identical or different, represent hydrogen atoms or alkyl or phenyl groups, or else R₇ and R₈ together form a 3- to 8-membered saturated ring which can optionally contain a heteroatom,

- Alk represents a linear or branched alkylene chain, and
- Alk' represents a linear, branched, cyclic or partially cyclic alkylene chain, optionally in the form of a pharmaceutically acceptable salt thereof.

2. The compound according to Claim 1, in which R₁ represents:

- . a hydrogen or halogen atom,
- . an alkyl group which is unsubstituted or substituted with -COOR₅,
- . an alkenyl group which is unsubstituted or substituted with -COOR₅,
- . a -COOR₅ group,
- . a -CONR₅R₆ group,
- . an -NR₅-SO₂R₆ group, or
- . a phenyl group optionally substituted with one or two groups selected from:
 - halogen atoms;
 - alkyl groups optionally substituted with -COOR₅;
 - -CN, -C(NH₂)NOH, -COOR₅, -CONR₅R₆, -CO-NR₅-OR₆, -CO-NR₅-SO₂R₆, -COAlk, -CO(OAlk)_nOH, -OR₅, -OCF₃, -O-Alk-COOR₅, -Alk-OR₅, -NR₅R₆ or -NC(O)N(CH₃)₂ groups,
 - 5-membered heteroaryls optionally substituted with an alkyl group and/or a hydroxyl group or an oxygen atom,

in which R₅ and R₆, which may be identical or different, represent hydrogen atoms, or alkyl groups optionally substituted with an -NR₇R₈ group,

R_7 represents a hydrogen atom, an alkyl group containing 1 or 2 carbon atoms or a phenyl group, n is an integer ranging from 1 to 3, or

. a heteroaryl group which is optionally condensed and/or optionally substituted with one or two groups selected from alkyl groups, OR_5 , $-COOR_5$, $-NR_5R_6$ and cycloalkyl groups, and an oxygen atom, in which R_5 and R_6 , which may be identical or different, represent hydrogen atoms or alkyl groups containing 1 or 2 carbon atoms,

optionally in the form of a pharmaceutically acceptable salt thereof.

3. The compound according to Claim 1, in which R_2 and R_3 together form, with the carbon atoms of the phenyl nucleus to which they are attached, a 6-membered nitrogenous heterocycle corresponding to either of formulae (A) and (B) as defined in Claim 1, optionally in the form of a pharmaceutically acceptable salt thereof.

4. The compound according to Claim 1, in which R_2 and R_3 together form, with the carbon atoms of the phenyl nucleus to which they are attached, a 6-membered nitrogenous heterocycle corresponding to formula (A) as defined in Claim 1, optionally in the form of a pharmaceutically acceptable salt thereof.

5. The compound according to Claim 1, in which formula (A) or (B) is such that:

. R_a represents a hydrogen atom or an alkyl group, optionally substituted with one or more halogens; $-AlkCONR_5R_6$; haloalkyl; $-CH_2-COOR_5$; $-Alk$ -heteroaryl, $-Alk$ -O-phenyl or $-Alk$ -phenyl, where the phenyl group is optionally substituted with one or two alkyl groups and/or OR_5 and/or halogen atoms; $-Alk$ -cycloalkyl,

. R_a' represents a hydrogen atom or a linear, branched, cyclic or partially cyclic alkyl group, or a $-CH_2-OR_5$ or $-Alk-NR_5R_6$ group,

. R_b represents a hydrogen atom or an alkyl group,

. R_b' represents a hydrogen atom or an alkyl, phenyl or $-CH_2-COOR_5$ group,

in which the alkyl groups contain 1 to 6 carbon atoms, R_5 being as described in Claim 1, optionally in the form of a pharmaceutically acceptable salt thereof.

6. The compound according to Claim 1, in which R_4 represents a hydrogen atom or a $-COOH$, $-CO-NH-Alk-NR_7R_8$ or $-CO-NH-Alk-OH$ group, or else an alkyl group which is unsubstituted,

in which Alk, R₇ and R₈ are as described in Claim 1.

7. The compound according to Claim 1 selected from the following compounds:

6-(imidazo[1,5-a]pyridin-3-ylcarbonyl)-3-propylquinazoline-2,4(1*H*,3*H*)-dione,

3-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid,

3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridine-6-carboxylic acid,

3-(3-{[3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoic acid,

3-{3[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzamide,

6-({1-[3-(5-methyl-1,3,4-oxadiazol-2-yl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)-3-propylquinazoline-2,4(1*H*, 3*H*)-dione,

6-({1-[3-(3-methyl-1,2,4-oxadiazol-5-yl)phenyl]imidazo[1,5-a]pyridin-3-yl}carbonyl)-3-propylquinazoline-2,4(1*H*, 3*H*)-dione,

N-{3-[(2,4-dioxo-3-propyl-1,2,3,4-tetrahydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}methanesulphonamide,

2-morpholin-4-yl-ethyl 3-(3-{[3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoate,

N-[2-(dimethylamino)ethyl]-3-(3-{[3-(4-fluorobenzyl)-1-methyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzamide,

3-(3-{[3-(4-fluorobenzyl)-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl]carbonyl}imidazo[1,5-a]pyridin-1-yl)benzoic acid,

3-(4-fluorobenzyl)-1-methyl-6-[(1-pyridin-3-ylimidazo[1,5-a]pyridin-3-yl)carbonyl]quinazoline-2,4(1*H*, 3*H*)-dione,

3-{3-[(2-methyl-4-oxo-3-propyl-3,4-dihydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzoic acid,

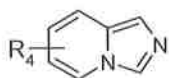
3-{3-[(2-methyl-4-oxo-3-propyl-3,4-dihydroquinazolin-6-yl)carbonyl]imidazo[1,5-a]pyridin-1-yl}benzamide,

6-(imidazo[1,5-a]pyridin-3-ylcarbonyl)quinazolin-4(3*H*)-one,

N,N,1,2-tetramethyl-4-oxo-6-{[1-(pyridin-3-yl)imidazo[1,5-*a*]pyridin-3-yl]carbonyl}-1,4-dihydroquinoline-3-carboxamide,
 3-[3-({3-[2-(4-fluorophenoxy)ethyl]-1-propyl-2,4-dioxo-1,2,3,4-tetrahydroquinazolin-6-yl}carbonyl)imidazo[1,5-*a*]pyridin-1-yl]benzoic acid,
 optionally in the form of a pharmaceutically acceptable salt thereof.

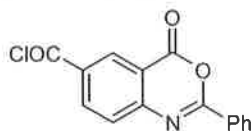
8. A process for preparing the compound according to Claim 1 in which R_2 and R_3 together form a nitrogenous heterocycle of formula (A) as defined in Claim 1, in which R_1 and R_a represent hydrogen atoms, comprising:

- condensing the compound of formula (IV)



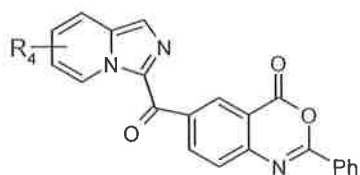
(IV)

in which R_4 is as defined in Claim 1, with the compound of formula (V)



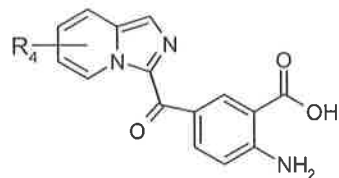
(V)

in order to obtain the compound of formula (VI)



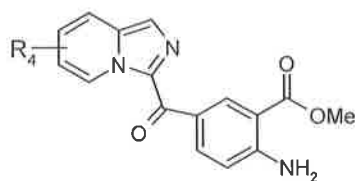
(VI)

- subjecting the compound of formula (VI) to a basic hydrolysis reaction in order to obtain the compound of formula (VII):



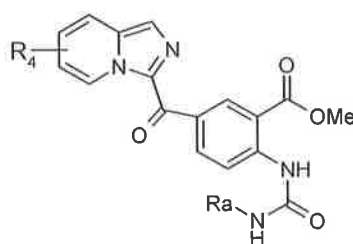
(VII)

- subjecting the compound of formula (VII) to a reaction of esterification to obtain the compound of formula (VIII)



(VIII)

- subjecting the compound of formula (VIII) to the action of triphosgene so as to form the isocyanate corresponding to the compound (VIII), and then condensing said isocyanate with an amine of formula R_aNH_2 , R_a being as defined in Claim 1, in order to obtain the urea of formula (IX),

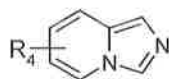


(IX)

-subjecting the urea of formula (IX) to a cyclization reaction in a basic medium.

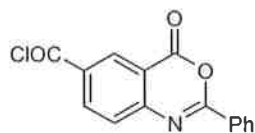
9. A process for preparing a derivative of formula (I) according to Claim 1 in which R_2 and R_3 together form a nitrogenous heterocycle of formula (A) as defined in Claim 1, R_1 is as defined in Claim 1, with the proviso that R_1 does not represent a hydrogen atom, and R_4 is as defined in Claim 1, comprising:

- condensing the compound of formula (IV)



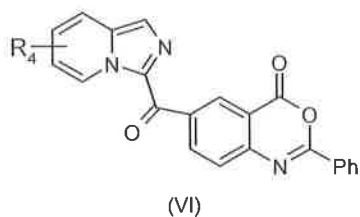
(IV)

in which R_4 is as defined in Claim 1, with the compound of formula (V)

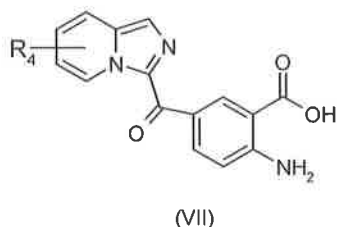


(V)

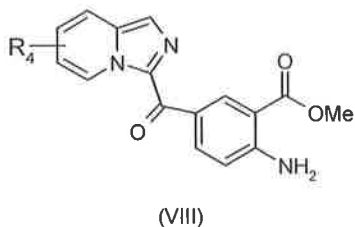
in order to obtain the compound of formula (VI)



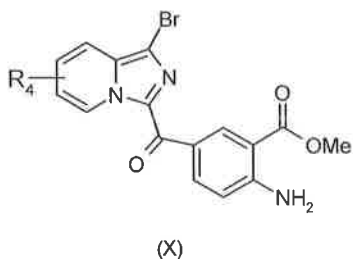
- subjecting the compound of formula (VI) to a basic hydrolysis reaction in order to obtain the compound of formula (VII):



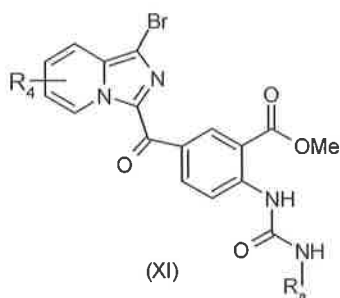
-subjecting the compound of formula (VII) to a reaction of esterification so as to obtain the compound of formula (VIII)



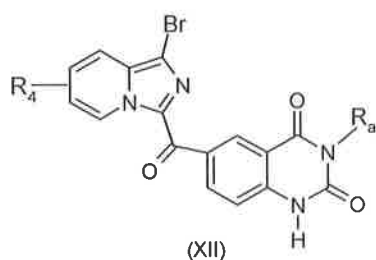
- subjecting the compound of formula (VIII) to a bromination reaction in order to obtain the compound of formula (X):



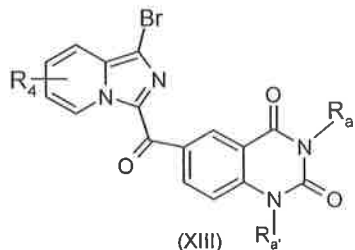
- subjecting the derivative of formula (X) to the action of triphosgene and forming the isocyanate corresponding to the compound of formula (X), which is condensed with an amine of formula R_aNH_2 , R_a being as defined in Claim 1, in order to obtain the urea of formula (XI):



- subjecting the compound of formula XI to a cyclization reaction in a basic medium in order to obtain the compound of formula (XII),



- subjecting the compound of formula (XII) to an alkylation reaction in the presence of a base and of a halogenated derivative $R_a'X$, R_a' being as defined in Claim 1, in order to obtain the compound of formula (XIII):



- subjecting the compound of formula (XIII), in the presence of a palladium catalyst, of a ligand and of a base:

. to a reaction with phenylboronic or heteroarylboronic or phenylboronate ester or heteroarylboronate ester derivatives according to a Suzuki coupling,

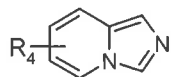
. or else to an imination reaction with benzophenoneimine, followed by an acid hydrolysis and by an alkylation reaction with a sulphonyl chloride of formula R_6SO_2Cl ,

. or else to a cyanation reaction with zinc cyanide, followed by an acid hydrolysis and by an esterification or a peptide coupling with an amine $R_5R_6NH_2$, R_5 and R_6 being defined in Claim 1.

10. A process for preparing the compounds of formula (I) according to Claim 1 in which R_2 and

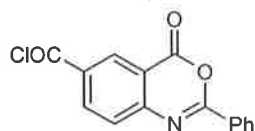
R_3 together form a nitrogenous heterocycle of formula (A) as defined in Claim 1, and in which R_1 and R_4 represent groups as defined in Claim 1, with the proviso that R_1 is not a hydrogen atom, comprising:

- condensing the compound of formula (IV)



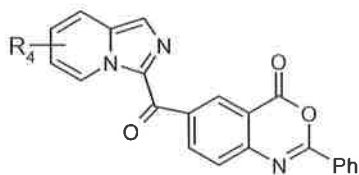
(IV)

in which R_4 is as defined in Claim 1, with the compound of formula (V)



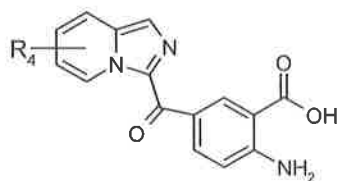
(V)

in order to obtain the compound of formula (VI)



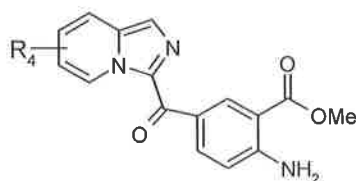
(VI)

- subjecting the compound of formula (VI) to a basic hydrolysis reaction in order to obtain the compound of formula (VII):



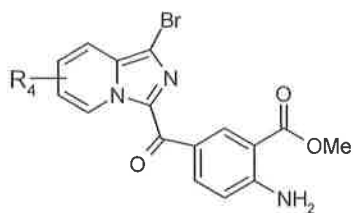
(VII)

- subjecting the compound of formula (VII) to a reaction of esterification so as to obtain the compound of formula (VIII):



(VIII)

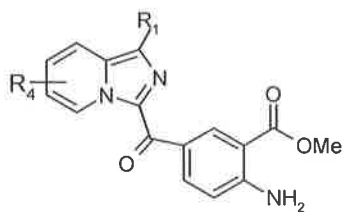
-subjecting the compound of formula (VIII) to a bromination reaction in order to obtain the compound of formula (X):



(X)

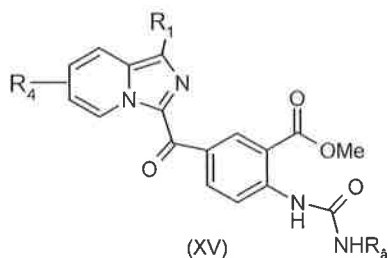
- subjecting the compound of formula (X), in the presence of a palladium catalyst, of a ligand and of a base:

- . to a reaction with phenylboronic or heteroarylboronic or phenylboronate ester or heteroarylboronate ester derivatives according to a Suzuki coupling,
- . or else to an imination reaction with benzophenoneimine, followed by an acid hydrolysis and by an alkylation reaction with a sulphonyl chloride of formula R_6SO_2Cl ,
- . or else to a cyanation reaction with zinc chloride, followed by an acid hydrolysis and by an esterification or a peptide coupling with an amine $R_5R_6NH_2$, R_5 and R_6 being defined in Claim 1, in order to obtain the compound of formula (XIV):



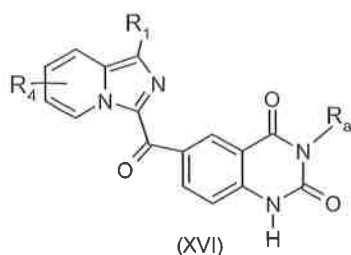
(XIV)

-subjecting the derivative of formula (XIV) to the action of triphosgene so as to form the corresponding isocyanate, condensing the isocyanate obtained with an amine of formula R_aNH_2 in order to obtain the urea of formula (XV), R_a being as defined in Claim 1:



(XV)

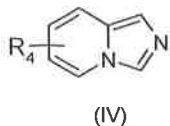
-subjecting the derivative of formula (XV) to a cyclization reaction in a basic medium in order to obtain the compound of formula (XVI):



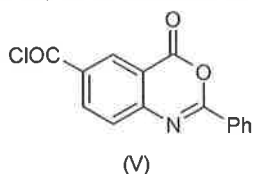
- subjecting the compound of formula (XVI) to an alkylation reaction in the presence of a base and of a halogenated derivative $R_a \cdot X$, R_a being as defined in Claim 1 and X being a halogen.

11. A process for preparing the compounds of formula (I) according to Claim 1 in which R_2 and R_3 together form a nitrogenous heterocycle of formula (A) and in which R_1 represents a group as defined in Claim 1, with the proviso that R_1 does not represent a hydrogen atom, R_4 being as defined in claim 1, comprising:

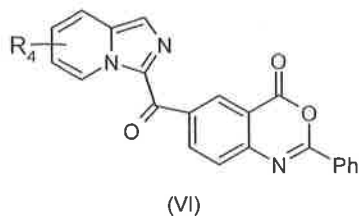
- condensing the compound of formula (IV)



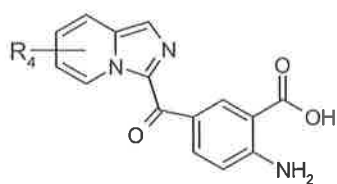
in which R_4 is as defined in Claim 1, with the compound of formula (V)



in order to obtain the compound of formula (VI)

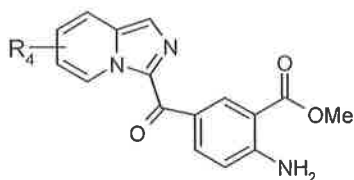


- subjecting the compound of formula (VI) to a basic hydrolysis reaction in order to obtain the compound of formula (VII):



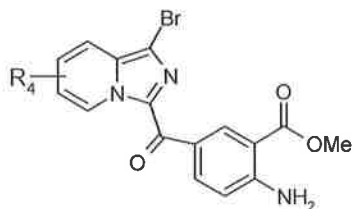
(VII)

- subjecting the compound of formula (VII) to a reaction of esterification so as to obtain the compound of formula (VIII)



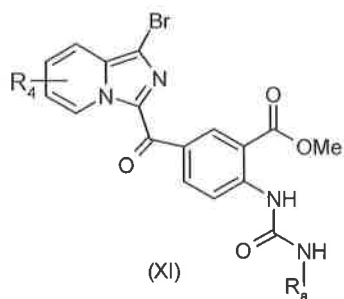
(VIII)

- subjecting the compound of formula (VIII) to a bromination reaction in order to obtain the compound of formula (X):



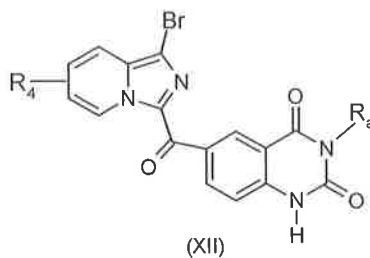
(X)

-subjecting the derivative of formula (X) to the action of triphosgene so as to form the corresponding isocyanate condensing said isocyanate with an amine of formula R_aNH_2 , R_a being as defined in Claim 1, in order to obtain the urea of formula (XI):

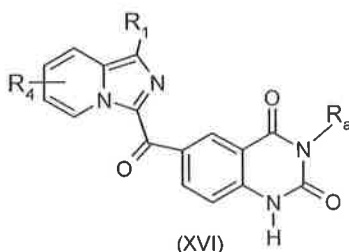


(XI)

- subjecting the compound of formula (XI) to a cyclization reaction in a basic medium in order to obtain the compound of formula (XII)



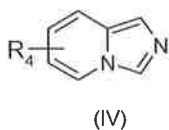
- subjecting the compound of formula (XII), in the presence of a palladium catalyst, of a ligand and of a base,
- to a reaction with phenylboronic or heteroarylboronic or phenylboronate ester or heteroarylboronate ester derivatives according to a Suzuki coupling,
- or else to an imination reaction with benzophenoneimine, followed by an acid hydrolysis and by a sulphonylation reaction with a sulphonyl chloride of formula R_6SO_2Cl ,
- or else to a cyanation reaction with zinc cyanide, followed by an acid hydrolysis and by an esterification or a peptide coupling with an amine $R_5R_6NH_2$, R_5 and R_6 being as defined in Claim 1, in order to obtain the compound of formula (XVI) in which R_1 is as defined in Claim 1, with the proviso that R_1 does not represent a hydrogen atom,



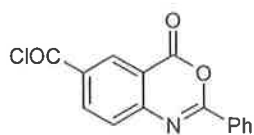
- subjecting the compound of formula (XVI) to an alkylation reaction in the presence of a base and of a halogenated derivative $R_a'X$, R_a' being as defined in Claim 1 and X being a halogen.

12. A process for preparing the compound according to Claim 1, in which R_2 and R_3 together form a nitrogenous heterocycle of formula (B) as defined in Claim 1, R_4 being as defined in Claim 1, and R_1 represents a hydrogen atom, comprising:

- condensing the compound of formula (IV)

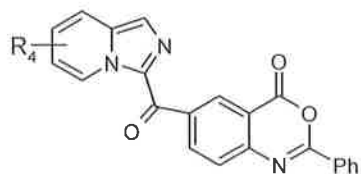


in which R_4 is as defined in Claim 1, with the compound of formula (V)



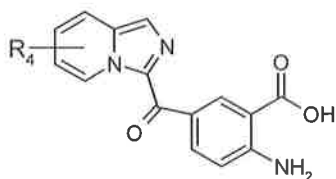
(V)

in order to obtain the compound of formula (VI)



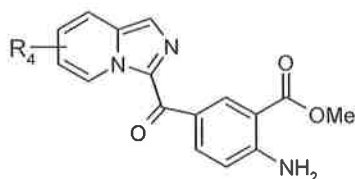
(VI)

-subjecting the compound of formula (VI) to a basic hydrolysis reaction in order to obtain the compound of formula (VII):



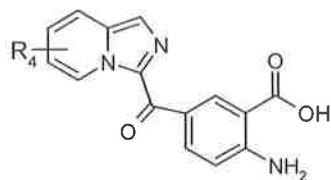
(VII)

- subjecting the compound of formula (VII) to a reaction of esterification so as to obtain the compound of formula (VIII)



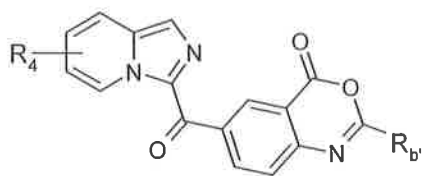
(VIII)

- subjecting the compound (VIII) to a saponification reaction in order to obtain the compound (XXIV):



(XXIV)

-subjecting the compound (XXIV) to a condensation reaction with an alkyl or aryl anhydride $(R_bCO)_2O$ in order to obtain the compound of formula (XVII),

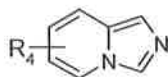


(XVII)

- subjecting the compound of formula (XVII) to a condensation reaction with an amine R_bNH_2 , R_b and $R_{b'}$ being as defined in Claim 1.

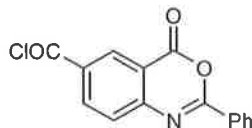
13. A process for preparing the compound according to Claim 1 in which R_2 and R_3 together form a nitrogenous heterocycle of formula (B) as defined in Claim 1, R_4 being as defined in Claim 1 and R_1 being as defined in Claim 1, with the proviso that R_1 does not represent a hydrogen atom, comprising:

-condensing the compound of formula (IV)



(IV)

in which R_4 is as defined in Claim 1, with the compound of formula (V):



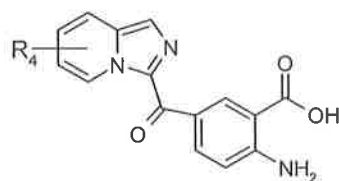
(V)

in order to obtain the compound of formula (VI):



(VI)

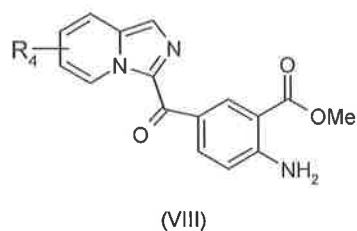
- subjecting the compound of formula (VI) to a basic hydrolysis reaction in order to obtain the compound of formula (VII):



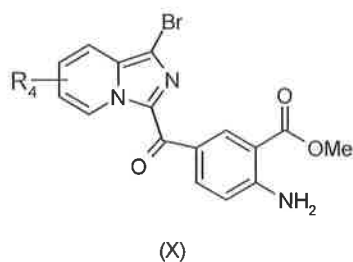
(VII)

-subjecting the compound of formula (VII) to a reaction of esterification so as to

obtain the compound of formula (VIII):



-subjecting the compound of formula (VIII) to a bromination reaction in order to obtain the compound of formula (X):



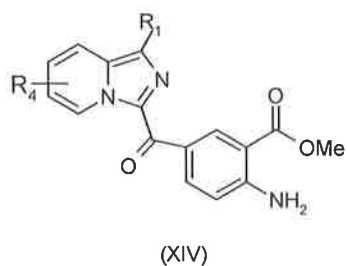
-subjecting the compound of formula (X), in the presence of a palladium catalyst, of a ligand and of a base,

- to a reaction with phenylboronic or heteroarylboronic or phenylboronate ester or heteroarylboronate ester derivatives according to a Suzuki coupling,

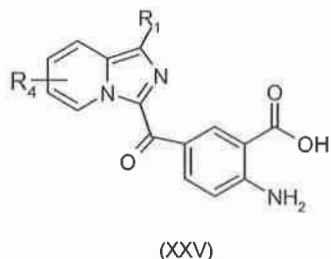
- or else to an imination reaction with benzophenoneimine, followed by an acid hydrolysis and by an alkylation reaction with a sulphonyl chloride of formula R_6SO_2Cl ,

- or else to a cyanation reaction with zinc chloride, followed by an acid hydrolysis and by an esterification or a peptide coupling with an amine $R_5R_6NH_2$, R_5 and R_6 being defined in Claim 1,

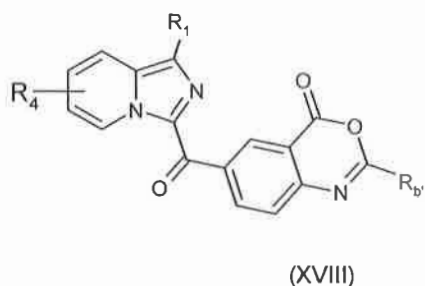
- in order to obtain the compound of formula XIV:



- subjecting the compound (XIV) to a saponification reaction in order to obtain the compound (XXV):



- subjecting the compound (XXV) to a condensation reaction with an alkyl or aryl anhydride $(R_bCO)_2O$, R_b being as defined in Claim 1, in order to obtain the compound of formula (XVIII):



- subjecting the compound of formula (XVIII) a condensation reaction with an amine R_bNH_2 , R_b being as defined in Claim 1.

14. A pharmaceutical composition containing, as an active ingredient, a derivative of formula (I) according to Claim 1, optionally in combination with one or more suitable inert excipients.

15. The compound according to claim 1, wherein said aryl group is phenyl.

16. The compound according to Claim 6, wherein said alkyl group contains 1 to 3 carbon atoms.