

SUPERINTENDENCIA DE INDUSTRIA Y COMERCIO



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Alix Carmenza Céspedes de Vergel

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SUPERINTENDENCIA DE INDUSTRIA Y COMERCIO

E. S. D.

Referencia: Expediente No. 06.095.699

Solicitud de Patente de Invención titulada: "Derivados de 2-
piridiniletilcarboxamida y procesos para su preparación"

Solicitante: Bayer CropScience AG

RESPUESTA AL AUTO NOTIFICADO POR FIJACIÓN EN LISTA NO.350**DEL 8 DE OCTUBRE DE 2010, OFICIO NO. 13166 (EXAMEN DE FONDO)**

ÁLVARO CORREA ORDÓÑEZ, mayor de edad y vecino de Bogotá D.C., identificado tal como aparece al pie de mi firma, en mi carácter de apoderado de la sociedad en referencia, doy respuesta al auto emanado por ese Despacho dentro del término legal

1. Objeciones del Examinador**1.1 De la solicitud de patente**

Solicita su Despacho que se haga modificación del título no aludiendo al uso.

Descripción

Se menciona por parte de su Despacho que en la página 49 de la descripción se cita el compuesto D1 al cual no se vuelve a hacer referencia en ninguna parte de la solicitud y por lo tanto se solicita corregir y aclarar.

Reivindicaciones

Considera su Despacho que el capítulo reivindicatorio es una generalización demasiado amplia de lo divulgado en la descripción y que por lo tanto no sería conciso.

Se sugiere por parte de su Despacho que el capítulo reivindicatorio se limite de acuerdo con los significados preferidos de las tablas A, B y C y para los cuales se evidenció una eventual mejora.

De otro lado se manifiesta que la reivindicación 15 hace referencia al uso por lo que no sería protegible por vía de patente.

1.2 Nivel Inventivo

Sobre el requisito del nivel inventivo, el Examinador establece que una persona del nivel técnico medio, versada en la materia, ensayaría las soluciones que se proponen en el documento WO 01/11965 (D1) y llegaría de manera obvia a la solución que se propone en la presente solicitud. Particularmente, el Examinador considera que las reivindicaciones 1 a 15 carecen de nivel inventivo debido a las enseñanzas contenidas en D1.

El Examinador establece:

"Al comparar los significados de los sustituyentes de esta solicitud con los significados de los sustituyentes de D1 se observan estructuras equivalentes. En la tabla comparativa No. 1 se presenta un ejemplo para evidenciar la equivalencia estructural:(...).

De modo tal que una persona medianamente versada en la materia encuentra sugerencia suficiente en D1 para plantear derivados como los acá presentados; y esperar igualmente una actividad fungicida de ellos.

En consecuencia y teniendo en cuenta además la amplitud del capítulo reivindicatorio se concluye que el material reclamado en las reivindicaciones 1-14 son alternativas obvias para un técnico medio y por lo tanto no tienen nivel inventivo."

2. Argumentos a favor de la patentabilidad de la solicitud

2.1 De la solicitud de patente

Título

En respuesta a su objeción sobre el título el solicitante ha decidido modificar el dicho título por el de: “Derivados de 2-piridiniletilcarboxamida y procesos para su preparación”

Descripción

En relación con la observación hecha al compuesto D1, por favor note que dicho compuesto fue mencionado por error y por lo tanto dicha referencia ha sido eliminada de la descripción. Allego nueva página con la respectiva corrección.

Reivindicaciones

En respuesta a las observaciones presentadas a las reivindicaciones de la presente solicitud el solicitante ha decidido presentar un nuevo capítulo reivindicatorio en el cual se han tenido en cuenta todas y cada una de las mencionadas observaciones.

El nuevo capítulo reivindicatorio consta de 9 reivindicaciones.

La reivindicación 1 se ha corregido al limitar X a la materia objeto de la reivindicación 3, limitar R^1 y R^2 a la materia objeto de la reivindicación 5 y al limitar R^3 y R^4 a la materia objeto de la reivindicación 7, además, el grupo alquilcarbonilamino C_1-C_6 se ha eliminado de la lista de residuos que representan R^3 y R^4 .

Las reivindicaciones originales 3, 5 y 7 se han eliminado.

Las reivindicaciones 6, 13 y la reivindicación de uso 15 se ha cancelado.

2.2 Nivel inventivo

El nivel inventivo de una solicitud se encuentra definido en el Artículo 18 de la Decisión 486 y se establece cuando *“si para una persona del oficio normalmente versada en la materia técnica correspondiente, esa invención no hubiese resultado obvia ni se hubiese derivado de manera evidente del estado de la técnica”*.

Contrario a lo que plantea el Examinador, el documento del arte previo citado no afecta el nivel inventivo de la presente solicitud por cuanto no es posible derivar la presente invención a partir de las enseñanzas presentadas en el documento D1.

Teniendo en cuenta las modificaciones al capítulo reivindicatorio el documento WO 01/11965 (D1) no tiene ninguna relevancia. En dicho documento el grupo 2-piridilo (A1) se debe sustituir con al menos un grupo haloalquilo. La presente invención no suministra tal sustituyente, en razón a que el radical X no puede ser haloalquilo. Además, los ejemplos en la tabla D del documento D1 le enseñan a la persona medianamente versada, que en caso de que L sea $-\text{CH}(\text{R}^3)\text{NHC}(\text{O})-$, el radical R^3 debe ser una clase de grupo carbonilo sustituido, lo cual de nuevo no es posible en la presente invención de acuerdo con los significados de R^3 y R^4 .

De acuerdo con lo anterior D1 no hace obvia la presente invención y por lo tanto queda claro que la presente solicitud si posee nivel inventivo.

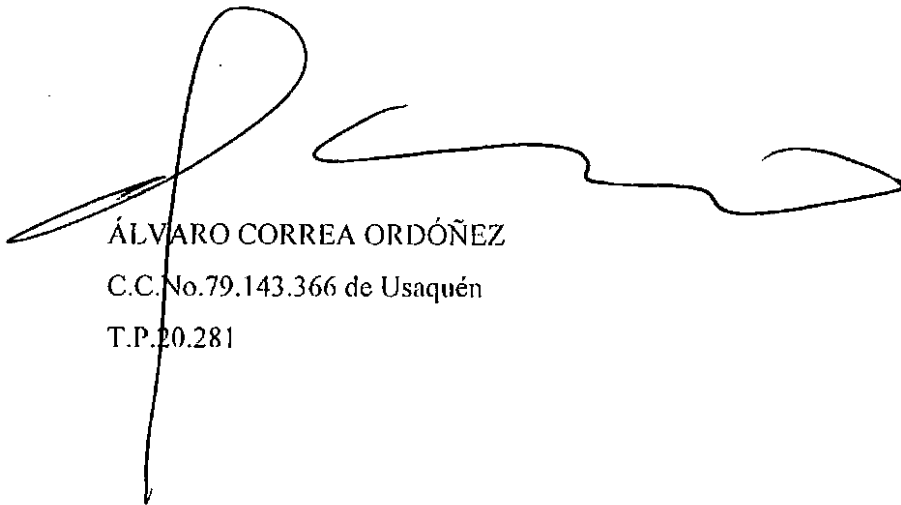
En este orden de ideas, tenemos que la invención propuesta por el solicitante si cumple con los requisitos de patentabilidad y de esta manera es susceptible de ostentar el título de patente de invención.

3. Solicitudes concedidas en otros países para esta misma invención.

A manera informativa, pongo en conocimiento del Examinador que la patente U.S. 7.723.363 equivalente a la presente solicitud colombiana ha sido otorgadas. Esta patente es más amplia que la que se esta tramitando en Colombia. Aún cuando entiendo que estas decisiones de ninguna manera obligan al Despacho para decidir de igual forma, considero que por tratarse de una pregunta fáctica la determinación de la existencia de nivel inventivo, las decisiones de otras oficinas de patentes constituyen un antecedente persuasivo en la materia.

En virtud de lo anterior, solicito que se acepten las razones expuestas en el presente documento y en consecuencia, se otorgue el privilegio de patente de invención para el invento que se tramita en este expediente.

Atentamente,



ÁLVARO CORREA ORDÓÑEZ

C.C. No. 79.143.366 de Usaquén

T.P. 20.281

Anexos: Hoja 49 modificada
Capítulo reivindicatorio modificado
Copia de las Patente U.S. 7.723.363.

RAC/BOGIDMS-#544177-V1-P2006_000144-0710.DOC

hoja (10 cm de alto), pulverizándolas con la suspensión acuosa descrita anteriormente. Las plantas, usadas como controles, se trataron con una disolución acuosa que no contenía el material activo.

Después de 24 horas, las plantas se contaminaron pulverizándolas con una suspensión acuosa de esporas de *Pyrenophora teres* (12.000 esporas por ml). Las esporas se obtuvieron de un cultivo de 12 días. Las plantas de cebada contaminadas se incubaron durante 24 horas a aproximadamente 20°C y humedad relativa de 100% y a continuación durante 12 días a humedad relativa de 80%.

La clasificación se llevó a cabo 12 días después de la contaminación, en comparación con las plantas de control.

En estas condiciones, se observó protección buena (al menos 50%) o total a una dosis de 330 ppm con los siguientes compuestos: A1, B1, C1, C2.

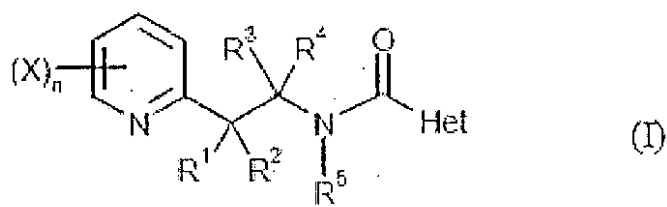
En estas condiciones, se observó protección buena (al menos 50%) o total a una dosis de 125 ppm con el siguiente compuesto: C4.

La N- {1-metilcarbamoil-2- [3-cloro-5-(trifluorometil) -2-piridinil] -etil} -4-fenil benzamida descrita en la Solicitud de Patente Internacional WO 01/11965 (véase el compuesto 316 en la Tabla D) mostró una eficacia deficiente sobre la *Alternaria brassicae* y una eficacia nula sobre la *Botrytis cinerea* a 330 ppm; la N-{1-etilcarbamoil-2-[3-cloro-5-(trifluorometil)-2-piridinil]-etil}-3-nitrobenzamida descrita también en la Solicitud de Patente Internacional WO 01/11965 (véase el compuesto 307 en la Tabla D) mostró una eficacia deficiente sobre la *Alternaria brassicae* y una eficacia nula sobre la *Botrytis cinerea* a 330 ppm; la N-{1-etilcarbamoil-2-[3-cloro-5-(trifluorometil)-2-piridinil]-etil}-benzamida y la N-{1-metilcarbamoil-2-[3-cloro-5-(trifluorometil)-2-piridinil]etil}-benzamida también descritas en la Solicitud de Patente Internacional WO 01/11965 (véanse los compuestos 304 y 314 en la Tabla D), mostraron una eficacia nula sobre la *Botrytis cinerea* a 330 ppm y la N-{1-etilcarbamoil-2-[3-cloro-5-(trifluorometil)-2-piridinil]etil}-4-clorobenzamida, la N-{1-etilcarbamoil-2-[3-cloro-5-(trifluorometil)-2-piridinil]etil}-2-bromobenzamida y la N-{1-metilcarbamoil-2-[3-cloro-5-(trifluorometil)-2-piridinil]etil}-4-metoxibenzamida, también descritas en la solicitud de patente internacional WO 01/11965 (véanse los compuestos 306, 310 y 315 en la Tabla D) mostraron una eficacia nula sobre la *Botrytis cinerea* a 330 ppm.

La N-{{3-cloro-5-(trifluorometil)-2-piridinil}metil}-5-tienilacetamida descrita en la solicitud de patente internacional WO 01/11965 (véase el compuesto 101 en la tabla B) mostró una eficacia deficiente frente a la *Alternaria brassicae* y ninguna eficacia frente a la *Botrytis cinerea* y la *Peronospora parasitica* a 330 ppm.

REIVINDICACIONES

1. Un compuesto de fórmula general (I):



En la cual:

- n es 1, 2, 3 o 4;
 - X es igual o diferente y es un átomo de halógeno, un alquilo C₁-C₈, un alcoxiimino C₁-C₆, un (alcoxiimino C₁-C₆)-alquilo C₁-C₆, o un alcoxi-C₁-C₆-alquilcarbonilo C₁-C₆;
 - R¹ y R² son iguales o diferentes y se seleccionan, independientemente uno del otro, de un átomo de hidrógeno, un átomo de halógeno, un grupo ciano, un grupo hidroxí, un alquilo C₁-C₆, un halogenoalquilo C₁-C₆ que tiene 1 a 5 átomos de halógeno, un alquenilo C₂-C₆, un alcoxi C₁-C₆, un alquilsulfanilo C₁-C₆, un alquilsulfenilo C₁-C₆, un alquilsulfinilo C₁-C₆, un alcoxicarbonilo C₁-C₆, un alquilcarbonilamino C₁-C₆, un alcoxicarboniloxi C₁-C₆, un alcoxicarbonilamino C₁-C₆ o un grupo fenilo;
 - o R¹ y R² pueden formar juntos un ciclopropilo, un ciclobutilo, un ciclopentilo o un ciclohexilo;
 - R³ y R⁴ son iguales o diferentes y se seleccionan, independientemente uno del otro, de un átomo de hidrógeno, un átomo de halógeno, un grupo ciano, un alquilo C₁-C₆, un halogenoalquilo C₁-C₆ que tiene 1 a 5 átomos de halógeno, un alquilcarbonilamino C₁-C₆ o un grupo fenilo;
 - Con la condición de que cuando tres de los cuatro sustituyentes R¹, R², R³ y R⁴ son un átomo de hidrógeno, entonces el cuarto sustituyente no es un átomo de hidrógeno;
 - R⁵ es un átomo de hidrógeno;
 - Het representa un heterociclo de 5, 6 o 7 miembros con uno, dos o tres heteroátomos que pueden ser iguales o diferentes; Het esta ligado por un átomo de carbono y esta, al menos, sustituido en la posición orto;
- Así como también sus sales, N-óxidos, complejos metálicos y metaloides.

2. Un compuesto de acuerdo con la reivindicación 1, caracterizado porque n es 1, 2 o

3. Un compuesto de acuerdo con la reivindicación 1 o 2, caracterizado porque el 2-piridilo se sustituye en la posición 3, 5 y/o 6.

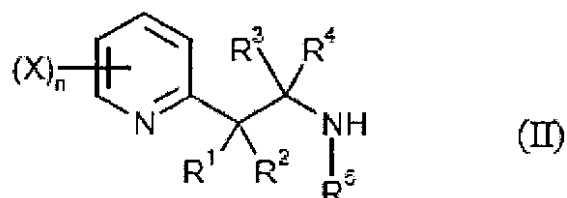
4. Un compuesto de acuerdo con cualquiera de las reivindicaciones 1 a 3, caracterizado porque R¹ y R² se seleccionan, independientemente uno del otro, por ser un átomo de halógeno, un alquilo C₁-C₆, un halogenoalquilo C₁-C₆, que tiene 1 a 5 átomos de halógeno o un alquilcarbonilamino C₁-C₆.

5. Un compuesto de acuerdo con cualquiera de las reivindicaciones 1 a 4, caracterizado porque R³ y R⁴ se seleccionan, independientemente uno del otro, por ser un átomo de halógeno, un alquilo C₁-C₆, un halogenoalquilo C₁-C₆ que tiene 1 a 5 átomos de halógeno o un grupo fenilo.

6. Un compuesto de acuerdo con cualquiera de las reivindicaciones 1 a 5, caracterizado porque Het es un anillo heterocíclico de cinco miembros.

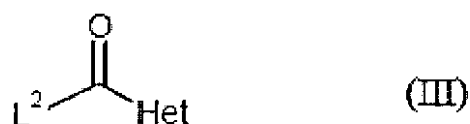
7. Un compuesto de acuerdo con cualquiera de las reivindicaciones 1 a 5, caracterizado porque Het es un anillo heterocíclico de seis miembros.

8. Un proceso para la preparación de un compuesto de fórmula general (I) tal como se definió en cualquiera de las reivindicaciones 1 a 7, que comprende hacer reaccionar un derivado de 2-piridina de fórmula general (II) o una de sus sales:



en la cual X, n, R¹, R², R³, R⁴ y R⁵ son como en cualquiera de las reivindicaciones precedentes;

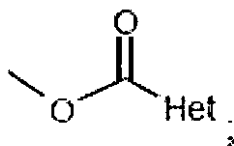
con un derivado de ácido carboxílico de la fórmula general (III)



en la cual:

Het es como se definió en cualquiera de las reivindicaciones precedentes; y

L² es un grupo saliente seleccionado por ser un átomo de halógeno, un grupo hidroxilo, -OR⁶, -OCOR⁶, R⁶ es un alquilo C₁-C₆, un haloalquilo C₁-C₆, un benzilo, 4-metoxibenzilo, pentafluorofenilo o un grupo de fórmula



En la presencia de un catalizador y, si L² es un grupo hidroxilo, en la presencia de un agente de condensación.

9. Una composición fungicida que comprende una cantidad efectiva de un compuesto de acuerdo con cualquiera de las reivindicaciones 1 a 7 y un soporte agrícolamente aceptable.



US007723363B2

(12) **United States Patent**
Mansfield et al.

(10) **Patent No.:** **US 7,723,363 B2**
(45) **Date of Patent:** ***May 25, 2010**

(54) **2-PYRIDINYLETHYLCARBOXAMIDE
DERIVATIVES AND THEIR USE AS
FUNGICIDES**

(75) **Inventors:** **Darren James Mansfield**, Lyons (FR);
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Pierre-Yves Coqueron, Lyons (FR);
Philippe Desbordes, Lyons (FR);
Marie-Claire Grosjean-Cournoyer,
Curis Au Mont d'Or (FR); **Pierre Genix**,
Lyons (FR); **Alain Villier**, Saint Cyr Au
Mont d'Or (FR)

(73) **Assignee:** **Bayer Cropscience SA (FR)**

(*) **Notice:** Subject to any disclaimer, the term of this
patent is extended or adjusted under 35
U.S.C. 154(b) by 739 days.

This patent is subject to a terminal dis-
claimer.

(21) **Appl. No.:** **10/588,985**

(22) **PCT Filed:** **Mar. 1, 2005**

(86) **PCT No.:** **PCT/EP2005/003282**

§ 371 (c)(1),
(2), (4) **Date:** **Oct. 12, 2006**

(87) **PCT Pub. No.:** **WO2005/085238**

PCT Pub. Date: **Sep. 15, 2005**

(65) **Prior Publication Data**

US 2007/0167491 A1 **Jul. 19, 2007**

(30) **Foreign Application Priority Data**

Mar. 3, 2004 (EP) **04356029**

(51) **Int. Cl.**

A01N 43/40 (2006.01)

C07D 401/12 (2006.01)

(52) **U.S. Cl.** **514/341; 546/275.4; 546/276.1**

(58) **Field of Classification Search** **546/275.4,**
546/276.1; 514/341

See application file for complete search history.

(56) **References Cited**

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WO **WO 01/05769** **1/2001**

WO **WO 01/11965** **2/2001**

WO **WO 2004/016088** **2/2004**

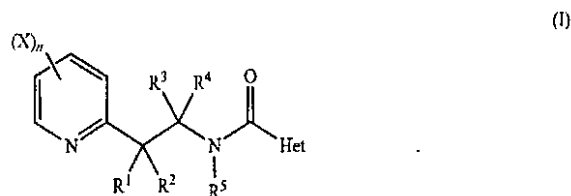
* cited by examiner

Primary Examiner—**Patricia L. Morris**

(74) *Attorney, Agent, or Firm*—**Ostrolenk Faber LLP**

(57) **ABSTRACT**

A compound of general formula (I). A process for preparing
this compound. A fungicidal composition comprising a com-
pound of general formula (I). A method for treating plants by
applying a compound of general formula (I) or a composition
comprising it.



11 Claims, No Drawings

1

2-PYRIDINYLETHYLCARBOXAMIDE
DERIVATIVES AND THEIR USE AS
FUNGICIDES

CROSS-REFERENCE TO RELATED
APPLICATION(S)

The present application is a 35 U.S.C. 371 national phase conversion of PCT/EP2005/003283 filed Mar. 1, 2005, which claims priority of European Application No. 04356029.1 filed Mar. 3, 2004.

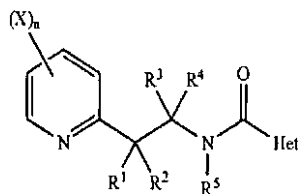
The present invention relates to novel N-[2-(2-pyridinyl)ethyl]carboxamides derivatives, their process of preparation, their use as fungicides, particularly in the form of fungicidal compositions, and methods for the control of phytopathogenic fungi of plants using these compounds or their compositions.

The international patent application WO 01/11965 discloses a broad family of fungicidal compounds in which the 2-pyridyl group is substituted by at least one halogenoalkyl group.

It is always of high-interest in the agriculture field to use novel pesticidal compounds in order to avoid or to fight the development of resistant strains to the active ingredients used by the farmer.

We have now found a new family of compounds which possess the above mentioned characteristics.

Accordingly, the present invention relates to N-[2-(2-pyridinyl)ethyl]carboxamide derivative of general formula (I):



in which:

n is 1, 2, 3 or 4;

X is the same or different and is a halogen atom, a nitro group, a cyano group, a hydroxy group, an amino group, a sulfanyl group, a pentafluoro- λ^6 -sulfanyl group, a formyl group, a formyloxy group, a formylamino group, a carboxy group, a carbamoyl group, a N-hydroxycarbamoyl group, a carbamate group, a (hydroxyimino)-C₁-C₆-alkyl group, a C₁-C₈-alkyl, a C₂-C₈-alkenyl, a C₂-C₈-alkynyl, a C₁-C₈-alkylamino, a di-C₁-C₈-alkylamino, a C₁-C₈-alkoxy, a C₁-C₈-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₈-alkylsulfanyl, a C₁-C₈-halogenoalkylsulfanyl having 1 to 5 halogen atoms, a C₂-C₈-alkenyloxy, a C₂-C₈-halogenoalkenyloxy having 1 to 5 halogen atoms, a C₃-C₈-alkynyloxy, a C₃-C₈-halogenoalkynyloxy having 1 to 5 halogen atoms, a C₃-C₈-cycloalkyl, a C₃-C₈-halogenocycloalkyl having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbonyl, a C₁-C₈-halogenoalkylcarbonyl having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbamoyl, a di-C₁-C₈-alkylcarbamoyl, a N-C₁-C₈-alkyloxycarbamoyl, a C₁-C₈-alkoxycarbamoyl, a N-C₁-C₈-alkyl-C₁-C₈-alkoxycarbamoyl, a C₁-C₈-alkoxycarbonyl, a C₁-C₈-halogenoalkoxycarbonyl having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbonyloxy, a C₁-C₈-halogenoalkylcarbonyloxy having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbonylamino, a C₁-C₈-halogenoalkylcarbonylamino

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having 1 to 5 halogen atoms, a C₁-C₈-alkylaminocarbonyloxy, a di-C₁-C₈-alkylaminocarbonyloxy, a C₁-C₈-alkyloxycarbonyloxy, a C₁-C₈-alkylsulphenyl, a C₁-C₈-halogenoalkylsulphenyl having 1 to 5 halogen atoms, a C₁-C₈-alkylsulphinyl, a C₁-C₈-halogenoalkylsulphinyl having 1 to 5 halogen atoms, a C₁-C₈-alkylsulphonyl, a C₁-C₈-halogenoalkylsulphonyl having 1 to 5 halogen atoms, a C₁-C₈-alkoxyimino, a (C₁-C₆-alkoxyimino)-C₁-C₆-alkyl, a (C₁-C₆-alkenyloxyimino)-C₁-C₆-alkyl, a (C₁-C₆-alkynyloxyimino)-C₁-C₆-alkyl, a (benzyloxyimino)-C₁-C₆-alkyl, a benzyloxy, a benzylsulfanyl, a benzylamino, a phenoxy, a phenylsulfanyl or a phenylamino;

R¹, R², R³ and R⁴ are the same or different and are a hydrogen atom, a halogen atom, a cyano group, a hydroxy group, an amino group, a sulfanyl group, a formyl group, a formyloxy group, a formylamino group, a carboxy group, a carbamoyl group, a N-hydroxycarbamoyl group, a carbamate group, a (hydroxyimino)-C₁-C₆-alkyl group, a C₁-C₈-alkyl, a C₁-C₈-halogenoalkyl having 1 to 5 halogen atoms, a C₂-C₈-alkenyl, a C₂-C₈-alkynyl, a C₁-C₈-alkylamino, a di-C₁-C₈-alkylamino, a C₁-C₈-alkoxy, a C₁-C₈-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₈-alkylsulfanyl, a C₁-C₈-halogenoalkylsulfanyl having 1 to 5 halogen atoms, a C₂-C₈-alkenyloxy, a C₂-C₈-halogenoalkenyloxy having 1 to 5 halogen atoms, a C₃-C₈-alkynyloxy, a C₃-C₈-halogenoalkynyloxy having 1 to 5 halogen atoms, a C₃-C₈-cycloalkyl, a C₃-C₈-halogenocycloalkyl having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbonyl, a C₁-C₈-halogenoalkylcarbonyl having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbamoyl, a di-C₁-C₈-alkylcarbamoyl, a N-C₁-C₈-alkyloxycarbamoyl, a C₁-C₈-alkoxycarbamoyl, a N-C₁-C₈-alkyl-C₁-C₈-alkoxycarbamoyl, a C₁-C₈-alkoxycarbonyl, a C₁-C₈-halogenoalkoxycarbonyl having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbonyloxy, a C₁-C₈-halogenoalkylcarbonyloxy having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbonylamino, a C₁-C₈-halogenoalkylcarbonylamino having 1 to 5 halogen atoms, a C₁-C₈-alkylaminocarbonyloxy, a di-C₁-C₈-alkylaminocarbonyloxy, a C₁-C₈-alkyloxycarbonyloxy, a C₁-C₈-alkylsulphenyl, a C₁-C₈-halogenoalkylsulphenyl having 1 to 5 halogen atoms, a C₁-C₈-alkylsulphinyl, a C₁-C₈-halogenoalkylsulphinyl having 1 to 5 halogen atoms, a C₁-C₈-alkylsulphonyl, a C₁-C₈-halogenoalkylsulphonyl having 1 to 5 halogen atoms, a benzyloxy, a benzylsulfanyl, a benzylamino, a phenoxy, a phenylsulfanyl or a phenylamino, a phenyl group, a phenyl sulphanyl group;

or R¹ and R² may form together a cyclopropyl, a cyclobutyl, a cyclopentyl or a cyclohexyl;

with the proviso that when three of the four substituents R¹, R², R³ and R⁴ are a hydrogen atom, then the fourth substituent is not a hydrogen atom;

R⁵ is a hydrogen atom, a cyano group, a formyl group, a hydroxy group, a C₁-C₆-alkyl, a C₁-C₆-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₆-alkoxy, a C₁-C₆-halogenoalkoxy having 1 to 5 halogen atoms, a C₃-C₆-cycloalkyl, a C₃-C₆-halogenocycloalkyl having 1 to 5 halogen atoms, a C₂-C₆-alkenyl, a C₂-C₆-alkynyl, a C₁-C₆-alkoxy-C₁-C₆-alkyl, a C₁-C₆-cyanoalkyl, a C₁-C₆-aminoalkyl, a C₁-C₆-alkylamino-C₁-C₆-alkyl, a di-C₁-C₆-alkylamino-C₁-C₆-alkyl, a C₁-C₆-alkylcarbonyl, a C₁-C₆-halogenoalkylcarbonyl having 1 to 5 halogen atoms, a C₁-C₆-alkyloxycarbonyl, a C₁-C₆-benzyloxycarbonyl, a C₁-C₆-alkoxy-C₁-C₆-alkylcarbonyl, a

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C_1 - C_6 -alkylsulfonyl or a C_1 - C_6 -halogenoalkylsulfonyl having 1 to 5 halogen atoms;

Het represents 5-, 6- or 7-membered heterocycle with one, two or three heteroatoms which may be the same or different; Het being linked by a carbon atom and being at least substituted in ortho position;

as well as its salts, N-oxides, metallic and metalloidic complexes.

In the context of the present invention:

halogen means fluorine, bromine, chlorine or iodine;

heteroatom means N, O or S;

carboxy means $-C(=O)OH$;

carbonyl means $-C(=O)-$;

carbamoyl means $-C(=O)NH_2$;

N-hydroxycarbamoyl means $-C(=O)NHOH$;

an alkyl group, an alkenyl group, and an alkynyl group as well as moieties containing these terms, can be linear or branched.

In the context of the present invention, it has also to be understood that in the case of di-substituted amino and of di-substituted carbamoyl radicals, the two substituents may form, together with the nitrogen atom bearing them, a saturated heterocyclic ring containing 3 to 7 atoms.

According to the present invention, the 2-pyridyl may be substituted in any position by $(X)_n$, in which X and n are as defined above. Preferably, the present invention relates to N-[2-(2-pyridinyl)ethyl]carboxamide derivative of general formula (I) in which the different characteristics may be chosen alone or in combination as being:

as regards n, n is 1, 2 or 3. More preferably n is 2 or 3.

as regards X, at least one of the X substituent is a halogen atom, a C_1 - C_8 -alkyl, a C_1 - C_6 -alkoxyimino, a $(C_1$ - C_6 -alkoxyimino)- C_1 - C_6 -alkyl, or a C_1 - C_6 -alkoxy- C_1 - C_6 -alkylcarbonyl.

as regards the positions in which the 2-pyridyl is substituted, the 2-pyridyl is substituted in 3-, 5- and/or in 6-position. More preferably, the 2-pyridyl is substituted in 3- and/or in 5-position

According to the present invention, the two carbon atoms and the nitrogen atom of the "ethylamide part" of the compound of formula (I) are respectively substituted by R^1 and R^2 , R^3 and R^4 , and R^5 , at least one the substituents R^1 , R^2 , R^3 and R^4 being different from hydrogen. Preferably, the present invention also relates to N-[2-(2-pyridinyl)ethyl]carboxamide derivative of general formula (I) in which the following characteristics may be chosen alone or in combination as being:

as regards R^1 and R^2 , R^1 and R^2 may be chosen, independently of each other, as being a hydrogen atom, a halogen atom, a cyano group, a hydroxy group, a C_1 - C_6 -alkyl, a C_1 - C_6 -halogenoalkyl having 1 to 5 halogen atoms, a C_2 - C_6 -alkenyl, a C_1 - C_6 -alkoxy, a C_1 - C_6 -alkylsulfonyl, a C_1 - C_6 -alkylsulfonyl, a C_1 - C_6 -alkoxycarbonyl, a C_1 - C_6 -alkylcarbonylamino, a C_1 - C_6 -alkoxycarbonyloxy, a C_1 - C_6 -alkoxycarbonylamino or a phenyl group. More preferably, R^1 and R^2 may be chosen, independently of each other, as being a halogen atom, a C_1 - C_6 -alkyl, a C_1 - C_6 -halogenoalkyl having 1 to 5 halogen atoms or a C_1 - C_6 -alkylcarbonylamino.

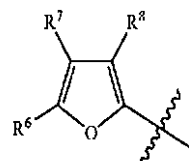
as regards R^3 and R^4 , R^3 and R^4 may be chosen, independently of each other, as being a hydrogen atom, a halogen atom, a cyano group, a C_1 - C_6 -alkyl, a C_1 - C_6 -halogenoalkyl having 1 to 5 halogen atoms, a C_1 - C_6 -alkylcarbonylamino or a phenyl group. More preferably, R^3 and R^4 may be chosen, independently of each other, as being a halogen atom, a C_1 - C_6 -alkyl, a C_1 - C_6 -halogenoalkyl having 1 to 5 halogen atoms or a phenyl group.

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as regards R^5 , R^5 may be chosen as being a hydrogen atom or a C_3 - C_7 -cycloalkyl.

According to the present invention, "Het" of the compound of general formula (I) may be a five membered ring heterocycle. Specific examples of compounds of the present invention where Het is a five membered heterocycle include:

Het may represent a heterocycle of the general formula (Het-1)



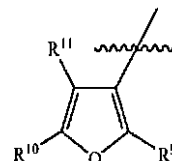
(Het-1)

in which:

R^6 and R^7 may be the same or different and may be a hydrogen atom, a halogen atom, an amino group, a nitro group, a C_1 - C_4 -alkyl or a C_1 - C_4 -halogenoalkyl having 1 to 5 halogen atoms; and

R^8 may be a halogen atom, a nitro group, a C_1 - C_4 -alkyl or a C_1 - C_4 -halogenoalkyl having 1 to 5 halogen atoms.

Het may represent a heterocycle of the general formula (Het-2)



(Het-2)

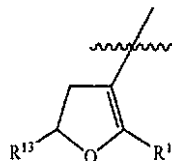
in which:

R^9 may be a hydrogen atom, a halogen atom, a C_1 - C_4 -alkyl or a C_1 - C_4 -halogenoalkyl having 1 to 5 halogen atoms; and

R^{10} and R^{11} may be the same or different and may be a hydrogen atom, a halogen atom, an amino group, a C_1 - C_4 -alkyl or a C_1 - C_4 -halogenoalkyl having 1 to 5 halogen atoms;

provided that the R^9 and R^{11} are not both a hydrogen atom.

Het may represent a heterocycle of the general formula (Het-3)



(Het-3)

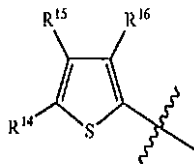
in which:

R^{12} may be a halogen atom, a C_1 - C_4 -alkyl or a C_1 - C_4 -halogenoalkyl having 1 to 5 halogen atoms; and

R^{13} may be a hydrogen atom, a C_1 - C_4 -alkyl or a C_1 - C_4 -halogenoalkyl having 1 to 5 halogen atoms.

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Het may represent a heterocycle of the general formula (Het-4)



(Het-4)

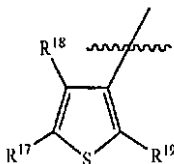
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in which:

R¹⁴ and R¹⁵ may be the same or different and may be a hydrogen atom, a halogen atom, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₄-alkylthio, a C₁-C₄-alkylsulphonyl, a phenyl optionally substituted by a halogen atom or a C₁-C₄-alkyl or a pyridyl optionally substituted by a halogen atom or a C₁-C₄-alkyl; and

R¹⁶ may be a halogen atom, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms or a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms.

Het may represent a heterocycle of the general formula (Het-5)



(Het-5)

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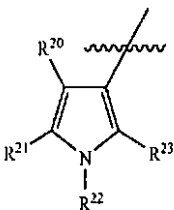
in which:

R¹⁷ and R¹⁸ may be the same or different and may be a hydrogen atom, a halogen atom, a C₁-C₄-alkyl, a C₁-C₄-alkyloxy or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms; and

R¹⁹ may be a hydrogen atom, a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms;

provided that the R¹⁸ and R¹⁹ are not both a hydrogen atom.

Het may represent a heterocycle of the general formula (Het-6)



(Het-6)

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in which:

R²⁰ may be a hydrogen atom, a halogen atom, a cyano group, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms;

R²¹ and R²³ may be the same or different and may be a hydrogen atom, a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms; and

R²² may be a hydrogen atom, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a

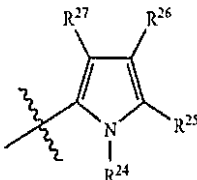
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C₁-C₄-alkoxy-C₁-C₄-alkyl, a hydroxy-C₁-C₄-alkyl, a C₁-C₄-alkylsulphonyl, a di(C₁-C₄-alkyl)aminosulphonyl, a C₁-C₆-alkyl-carbonyl, a phenylsulphonyl optionally substituted by a halogen atom or a C₁-C₄-alkyl, or a benzoyl optionally substituted by a halogen atom or a C₁-C₄-alkyl;

provided that the R²⁰ and R²³ are not both a hydrogen atom.

Het may represent a heterocycle of the general formula (Het-7) (Het-7)



(Het-7)

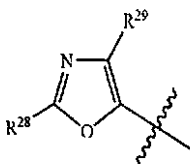
in which:

R²⁴ may be a hydrogen atom, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy-C₁-C₄-alkyl, a hydroxy-C₁-C₄-alkyl, a C₁-C₄-alkylsulphonyl, a di(C₁-C₄-alkyl)aminosulphonyl, a C₁-C₆-alkylcarbonyl, a phenylsulphonyl optionally substituted by a halogen atom or a C₁-C₄-alkyl, or a benzoyl optionally substituted by a halogen atom or a C₁-C₄-alkyl; and

R²⁵, R²⁶ and R²⁷ may be the same or different and may be a hydrogen atom, a halogen atom, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms or a C₁-C₄-alkylcarbonyl;

provided that R²⁴ and R²⁷ are not both a hydrogen atom.

Het may represent a heterocycle of the general formula (Het-8)



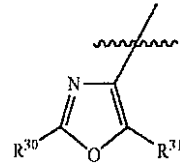
(Het-8)

in which:

R²⁸ may be a hydrogen atom or a C₁-C₄-alkyl; and

R²⁹ may be a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms.

Het may represent a heterocycle of the general formula (Het-9)



(Het-9)

in which:

R³⁰ may be a hydrogen atom or a C₁-C₄-alkyl; and

R³¹ may be a halogen atom, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms or a phenyl optionally substituted by a halogen atom or a C₁-C₄-alkyl.

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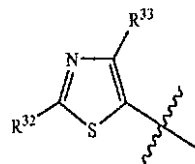
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Het may represent a heterocycle of the general formula (Het-10)



(Het-10) 5

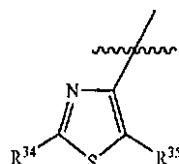
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in which:

R³² may be a hydrogen atom, a halogen atom, an amino group, a cyano group, a C₁-C₄-alkylamino, a di-(C₁-C₄-alkyl)amino, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms or a phenyl optionally substituted by a halogen atom or a C₁-C₄-alkyl; and

R³³ may be a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms.

Het may represent a heterocycle of the general formula (Het-11)



(Het-11) 25

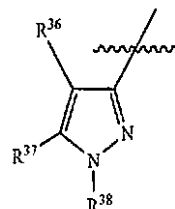
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in which:

R³⁴ may be a hydrogen atom, a halogen atom, an amino group, a cyano group, a C₁-C₄-alkylamino, a di-(C₁-C₄-alkyl)amino, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms; and

R³⁵ may be a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms.

Het may represent a heterocycle of the general formula (Het-12)



(Het-12) 45

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in which:

R³⁶ may be a halogen atom, a cyano group, a nitro group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₃-C₆-cycloalkyl, a C₁-C₄-alkoxy, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₄-alkylthio, a C₁-C₄-halogenoalkylthio having 1 to 5 halogen atoms, an aminocarbonyl group or an aminocarbonyl-C₁-C₄-alkyl;

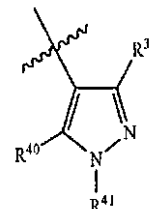
R³⁷ may be a hydrogen atom, a halogen atom, a cyano group, a nitro group, a C₁-C₄-alkyl, a C₁-C₄-alkoxy or a C₁-C₄-alkylthio; and

R³⁸ may be a hydrogen atom, a phenyl, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a hydroxy-C₁-C₄-alkyl, a C₂-C₆-alkenyl, a C₃-C₆-cy-

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cloalkyl, a C₁-C₄-alkylthio-C₁-C₄-alkyl, a C₁-C₄-halogenoalkylthio-C₁-C₄-alkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy-C₁-C₄-alkyl or a C₁-C₄-halogenoalkoxy-C₁-C₄-alkyl having 1 to 5 halogen atoms.

Het may represent a heterocycle of the general formula (Het-13)



(Het-13)

in which:

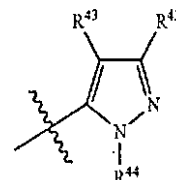
R³⁹ may be a hydrogen atom, a halogen atom, a cyano group, a nitro group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₃-C₆-cycloalkyl, a C₁-C₄-alkoxy, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₄-alkylthio, a C₁-C₄-halogenoalkylthio having 1 to 5 halogen atoms, an aminocarbonyl or an aminocarbonyl-C₁-C₄-alkyl;

R⁴⁰ may be a hydrogen atom, a halogen atom, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-alkoxy, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms or a C₁-C₄-alkylthio; and

R⁴¹ may be a hydrogen atom, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a hydroxy-C₁-C₄-alkyl, a C₂-C₆-alkenyl, a C₃-C₆-cycloalkyl, a C₁-C₄-alkylthio-C₁-C₄-alkyl, a C₁-C₄-halogenoalkylthio-C₁-C₄-alkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy-C₁-C₄-alkyl, a C₁-C₄-halogenoalkoxy-C₁-C₄-alkyl having 1 to 5 halogen atoms or a phenyl optionally substituted by a halogen atom, a C₁-C₄-alkyl, a C₁-C₄-alkoxyalkyl or a nitro group;

provided that the R³⁹ and R⁴⁰ are not both a hydrogen atom.

Het may represent a heterocycle of the general formula (Het-14)



(Het-14)

in which:

R⁴² may be a hydrogen atom, a halogen atom, a cyano group, a nitro group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₃-C₆-cycloalkyl, a C₁-C₄-alkoxy, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₄-alkylthio, a C₁-C₄-halogenoalkylthio having 1 to 5 halogen atoms, an aminocarbonyl, or an aminocarbonyl-C₁-C₄-alkyl;

R⁴³ may be a hydrogen atom, a halogen atom, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-alkoxy, a C₁-C₄-alkylthio or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms;

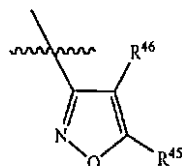
R⁴⁴ may be a hydrogen atom, a phenyl, a benzyl, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a hydroxy-C₁-C₄-alkyl, a C₂-C₆-alkenyl, a C₃-C₆-cycloalkyl, a C₁-C₄-alkylthio-C₁-C₄-alkyl, a

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C₁-C₄-halogenoalkylthio-C₁-C₄-alkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy-C₁-C₄-alkyl, a C₁-C₄-halogenoalkoxy-C₁-C₄-alkyl having 1 to 5 halogen atoms;

provided that R⁴³ and R⁴⁴ are not both a hydrogen atom.

Het may represent a heterocycle of the general formula (Het-15) (Het-15)



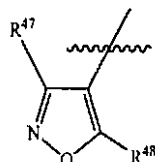
(Het-15)

in which:

R⁴⁵ may be a hydrogen atom, a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms; and

R⁴⁶ may be a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms.

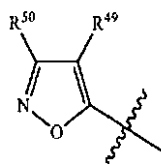
Het may represent a heterocycle of the general formula (Het-16) (Het-16)



in which R⁴⁷ and R⁴⁸ may be the same or different and may be a hydrogen atom, a halogen atom, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a phenyl optionally substituted by a halogen atom or a C₁-C₄-alkyl, or a heterocyclyl optionally substituted by a halogen atom or a C₁-C₄-alkyl;

provided that R⁴⁷ and R⁴⁸ are not both a hydrogen atom.

Het may represent a heterocycle of the general formula (Het-17) (Het-17)



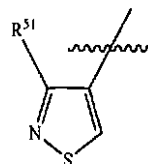
in which:

R⁴⁹ may be a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, and

R⁵⁰ may be a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms.

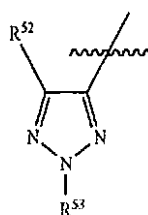
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Het may represent a heterocycle of the general formula (Het-18) (Het-18)



in which R⁵¹ may be a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms.

Het may represent a heterocycle of the general formula (Het-19) (Het-19)

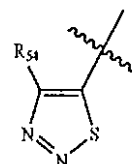


in which:

R⁵² may be a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms; and

R⁵³ may be a hydrogen atom, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, or a phenyl optionally substituted by a halogen atom or a C₁-C₄-alkyl.

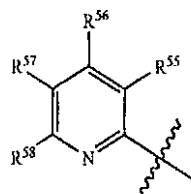
Het may represent a heterocycle of the general formula (Het-20) (Het-20)



in which R⁵⁴ may be a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms.

According to the present invention, "Het" of the compound of general formula (I) may be a six membered ring heterocycle. Specific examples of compounds of the present invention where Het is a six membered heterocycle include:

Het may represent a heterocycle of the general formula (Het-21) (Het-21)



in which:

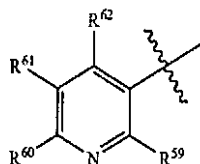
R⁵⁵ may be a halogen atom, a hydroxy group, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to

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5 halogen atoms, a C₁-C₄-alkoxy, a C₁-C₄-alkylthio, a C₁-C₄-halogenoalkylthio having 1 to 5 halogen atoms or a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms;

R⁵⁶, R¹⁷ and R⁵⁸, which may be the same or different, may be a hydrogen atom, a halogen atom, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy, a C₁-C₄-alkylthio, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₄-alkylsulphonyl or a C₁-C₄-alkylsulphonyl.

Het may represent a heterocycle of the general formula (Het-22)



(Het-22)

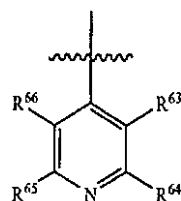
in which:

R⁵⁹ may be a hydrogen atom, a halogen atom, a hydroxy group, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy, a C₁-C₅-alkylthio, a C₂-C₅-alkenylthio, a C₁-C₄-halogenoalkylthio having 1 to 5 halogen atoms, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms, a phenyloxy optionally substituted by a halogen atom or a C₁-C₄-alkyl, or a phenylthio optionally substituted by a halogen atom or a C₁-C₄-alkyl;

R⁶⁰, R⁶¹ and R⁶², which may be the same or different, may be a hydrogen atom, a halogen atom, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy, a C₁-C₄-alkylthio, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₄-alkylsulphonyl, a C₁-C₄-alkylsulphonyl or a N-morpholine optionally substituted by a halogen atom or a C₁-C₄-alkyl, or a thienyl optionally substituted by a halogen atom or a C₁-C₄-alkyl;

provided that the R⁵⁹ and R⁶² are not both a hydrogen atom.

Het may represent a heterocycle of the general formula (Het-23)



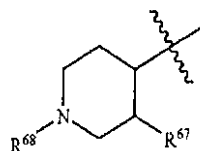
(Het-23)

in which R⁶³, R⁶⁴, R⁶⁵ and R⁶⁶, which may be the same or different, may be a hydrogen atom, a halogen atom, a hydroxy group, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy, a C₁-C₄-alkylthio, a C₁-C₄-halogenoalkylthio having 1 to 5 halogen atoms, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₄-alkylsulphonyl or a C₁-C₄-alkylsulphonyl;

provided that the R⁶³ and R⁶⁶ are not both a hydrogen atom.

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Het may represent a heterocycle of the general formula (Het-24)



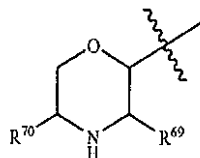
(Het-24)

in which:

R⁶⁷ may be a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms;

R⁶⁸ may be a hydrogen atom, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₆-alkoxycarbonyl, a benzyl optionally substituted by 1 to 3 halogen atoms, a benzyloxycarbonyl optionally substituted by 1 to 3 halogen atoms or a heterocyclyl.

Het may represent a heterocycle of the general formula (Het-25)



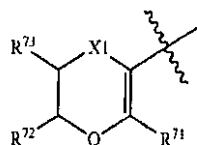
(Het-25)

in which:

R⁶⁹ may be a halogen atom, a hydroxy group, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy, a C₁-C₄-alkylthio, a C₁-C₄-halogenoalkylthio having 1 to 5 halogen atoms or a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms;

R⁷⁰ may be a hydrogen atom, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms or a benzyl.

Het may represent a heterocycle of the general formula (Het-26)



(Het-26)

in which:

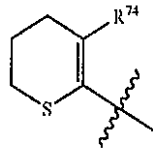
X¹ may be a sulphur atom, —SO—, —SO₂— or —CH₂—;

R⁷¹ may be a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms; and

R⁷² and R⁷³ may be the same or different and may be a hydrogen atom or a C₁-C₄-alkyl.

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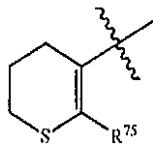
Het may represent a heterocycle of the general formula (Het-27)



in which:

R⁷⁴ may be a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms;

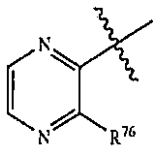
Het may represent a heterocycle of the general formula (Het-28)



in which:

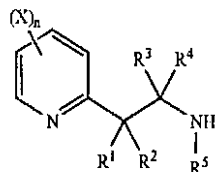
R⁷⁵ may be a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms.

Het may represent a heterocycle of the general formula (Het-29)



in which R⁷⁶ may be a halogen atom, a C₁-C₄-alkyl or a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms.

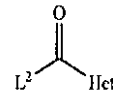
The present invention also relates to a process for the preparation of the compound of general formula (I). Thus, according to a further aspect of the present invention there is provided a process for the preparation of compound of general formula (I) as defined above, which comprises reacting a 2-pyridine derivative of general formula (II) or one of its salt:



in which X, n, R¹, R², R³, R⁴ and R⁵ are as defined above;

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with a carboxylic acid derivative of the general formula (III)



(Het-27) 5

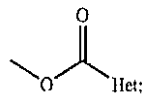
(III)

10 in which:

Het is as defined above; and

L² is a leaving group chosen as being a halogen atom, a hydroxyl group, —OR⁷⁷, —OCOR⁷⁷, R⁷⁷ being a C₁-C₆ alkyl, a C₁-C₆ haloalkyl, a benzyl, 4-methoxybenzyl, pentafluorophenyl or a group of formula

(Het-28) 20

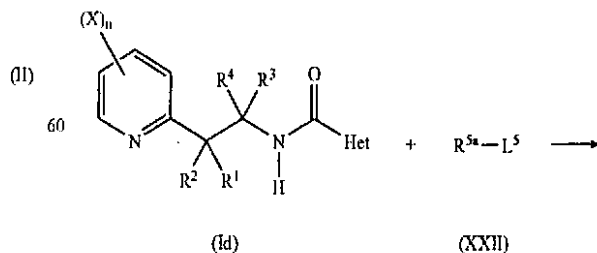


25 in the presence of a catalyst and, if L² is a hydroxyl group, in the presence of a condensing agent.

The process according to the present invention is conducted in the presence of a catalyst. Suitable catalyst may be chosen as being 4-dimethyl-aminopyridine, 1-hydroxy-benzotriazole or dimethylformamide.

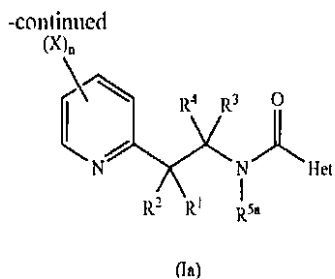
In case L² is a hydroxy group, the process according to the present invention is conducted in the presence of condensing agent. Suitable condensing agent may be chosen as being acid halide former, such as phosgene, phosphorous tribromide, phosphorous trichloride, phosphorous pentachloride, phosphorous trichloride oxide or thionyl chloride; anhydride former, such as ethyl chloroformate, methyl chloroformate, isopropyl chloroformate, isobutyl chloroformate or methane-sulfonyl-chloride; carbodiimides, such as N,N'-dicyclohexylcarbodiimide (DCC) or other customary condensing agents, such as phosphorous pentoxide, polyphosphoric acid, N,N'-carbonyl-diimidazole, 2-ethoxy-N-ethoxycarbonyl-1,2-dihydroquinoline (EEDQ), triphenylphosphine/tetrachloromethane, 4-(4,6-dimethoxy[1.3.5]triazin-2-yl)-4-methylmorpholinium chloride hydrate or bromotripyrrolidino-phosphonium-hexafluorophosphate.

When R⁵ is a hydrogen atom, the above mentioned process for the preparation of compound of general formula (I) may optionally be completed by a further step according to the following reaction scheme:



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in which:

R¹, R², R³, R⁴, X, n and Het are as defined above;

R^{5a} is a cyano group, a formyl group, a hydroxy group, a C₁-C₆-alkyl, a C₁-C₆-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₆-alkoxy, a C₁-C₆-halogenoalkoxy having 1 to 5 halogen atoms, a C₃-C₆-cycloalkyl, a C₃-C₆-halogenocycloalkyl having 1 to 5 halogen atoms, a C₂-C₆-alkenyl, a C₂-C₆-alkynyl, a C₁-C₆-alkoxy-C₁-C₆-alkyl, a C₁-C₆-cyanoalkyl, a C₁-C₆-aminoalkyl, a C₁-C₆-alkylamino-C₁-C₆-alkyl, a di-C₁-C₆-alkylamino-C₁-C₆-alkyl, a C₁-C₆-alkylcarbonyl, a C₁-C₆-halogenalkylcarbonyl having 1 to 5 halogen atoms, a C₁-C₆-alkyloxycarbonyl, a C₁-C₆-benzyloxycarbonyl, a C₁-C₆-alkoxy-C₁-C₆-alkylcarbonyl, a C₁-C₆-alkylsulfonyl or a C₁-C₆-halogenoalkylsulfonyl having 1 to 5 halogen atoms; and

L⁵ is a leaving group chosen as being a halogen atom, a 4-methyl phenylsulfonyloxy or a methylsulfonyloxy;

comprising the reaction of a compound of general formula (Id) with a compound of general formula (XXII) to provide a compound of general formula (Ia).

Depending on the definition of R¹, R², R³, R⁴ or R⁵, amine derivatives of general formula (II) may be prepared by different processes. One example (A) of such a process may be when:

X, n are as defined above;

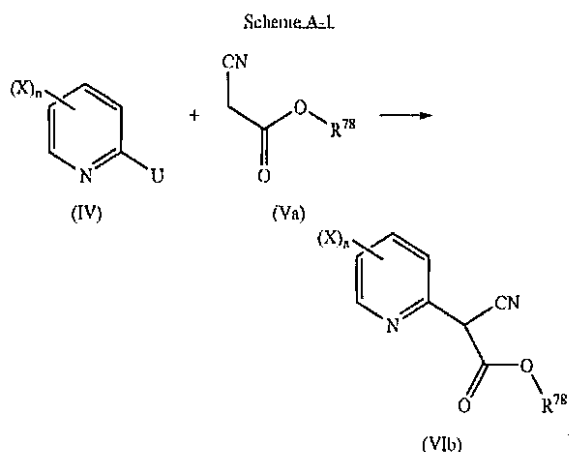
R¹ is a C₁-C₆ alkyl;

R² is a hydrogen atom or a C₁-C₆ alkyl; and

R³, R⁴, R⁵ are hydrogen atoms;

then, the amine derivative of general formula (II) may be prepared according to a process which comprises

a first step according to reaction scheme A-1:



in which:

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X and n are as defined above;

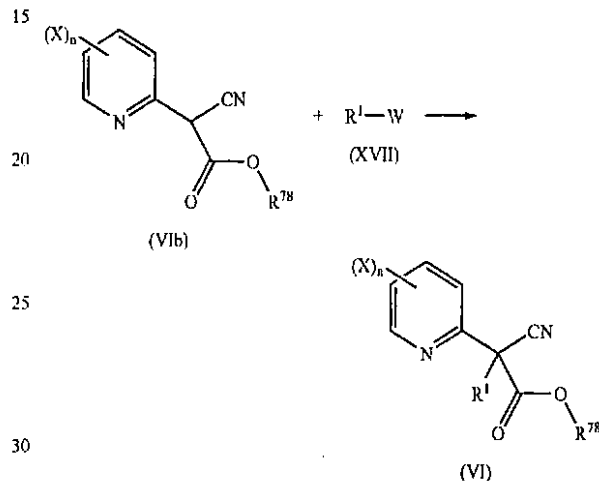
R⁷⁸ is a C₁-C₆ alkyl, a C₁-C₆ haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

U is a leaving group chosen as being a halogen, a C₁-C₆ alkylsulfonate or a C₁-C₆ haloalkylsulfonate;

comprising the arylation of a cyanoacetate derivative of general formula (Va) by a pyridine derivative of general formula (IV), to provide a 2-(pyridyl)cyanoacetate derivative of general formula (VI b), in the presence of a base, at a temperature of from 0° C. to 200° C.;

a second step according to reaction scheme A-2:

Scheme A-2



in which:

X, n are as defined above;

R¹ is a C₁-C₆ alkyl;

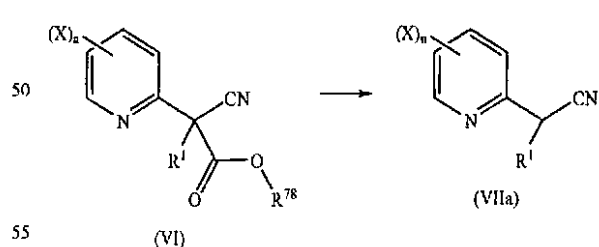
R⁷⁸ is a C₁-C₆ alkyl, a C₁-C₆ haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

W is a halogen atom, a C₁-C₆ alkylsulfonate, a C₁-C₆ haloalkylsulfonate or a 4-methyl-phenylsulfonate;

comprising the alkylation of a compound of general formula (VI b) by a reagent of general formula (XVII) to provide a compound of general formula (VI);

a third step according to reaction scheme A-3:

Scheme A-3



in which:

X, n are as defined above;

R¹ is a C₁-C₆ alkyl;

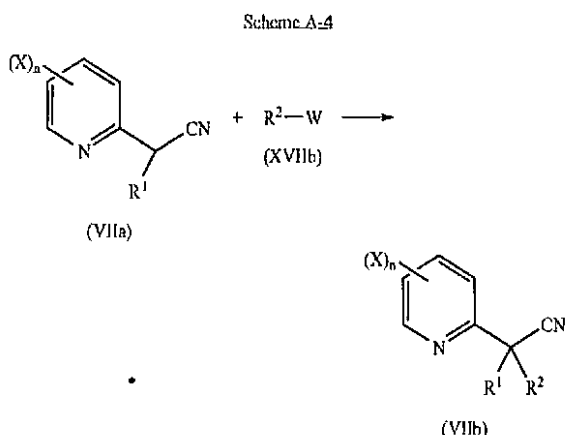
R⁷⁸ is a C₁-C₆ alkyl, a C₁-C₆ haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

comprising a basic hydrolysis, an acidic hydrolysis or a displacement by an halide of a compound of general formula (VI) in the same or a different pot to provide, upon heating at a temperature of from 40° C. to reflux, a 2-pyridylacetone derivative of general formula (VIIa);

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a fourth step according to reaction scheme A-4:



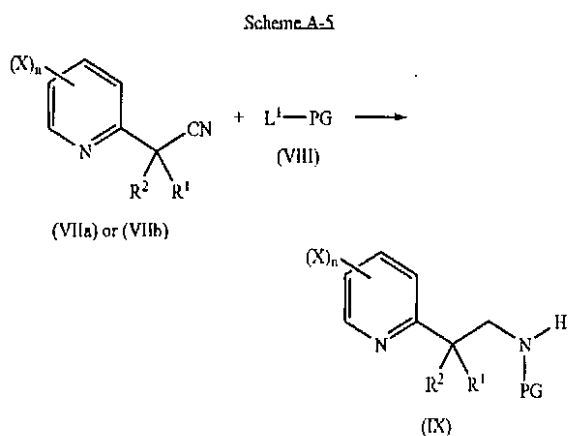
in which:

X, n are as defined above;

 R^1 is a C_1 - C_6 alkyl; R^2 is a C_1 - C_6 alkyl;W is a halogen atom, a C_1 - C_6 alkylsulfonate, a C_1 - C_6 haloalkylsulfonate or a 4-methyl-phenylsulfonate;

comprising the alkylation of a compound of general formula (VII a) by a reagent of general formula (XVII b) to provide a compound of general formula (VII b);

a fifth step according to reaction scheme A-5:



in which:

X, n are as defined above;

 R^1 is a C_1 - C_6 alkyl; R^2 is a hydrogen atom or a C_1 - C_6 alkyl;

L^1 is a leaving group chosen as being a $-OR^{77}$ group or a $-OCOR^{77}$ group, R^{77} being a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

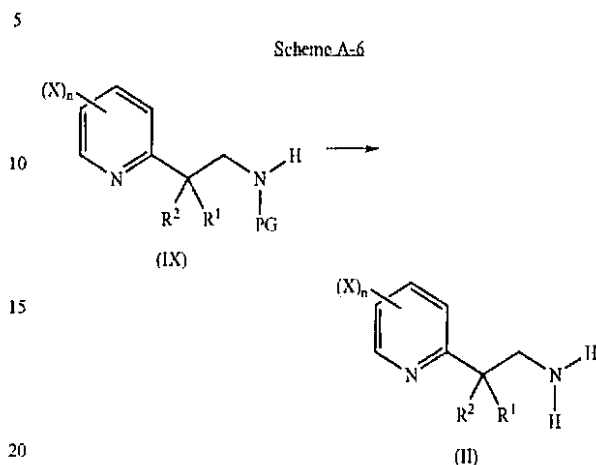
PG represents a protecting group which may be a $-COOR^{77}$ group or $-COR^{77}$ group, R^{77} being a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

comprising the reduction, by hydrogenation or by an hydride donor, of a compound of general formula (VIIa) or (VIIb), in the presence of a catalyst and in the presence of a compound of general formula (VIII) to produce a compound of general

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formula (IX), at a temperature of from 0° C. to 150° C. and under a pressure of from 1 bar and 100 bar;

a sixth step according to reaction scheme A-6:



in which:

X, n are as defined above;

 R^1 is a C_1 - C_6 alkyl; R^2 is a hydrogen atom or a C_1 - C_6 alkyl

PG represents a protecting group which may be a $-COOR^{77}$ group or $-COR^{77}$ group, R^{77} being a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

comprising a deprotection reaction, in an acidic or in a basic medium, of a compound of general formula (IX) to provide an amine derivative of general formula (II) or one of its salt.

The first step (step A-1) is conducted in the presence of a base. Preferably, the base will be chosen as being an inorganic or an organic base. Suitable examples of such bases may for example be alkaline earth metal or alkali metal hydrides, hydroxides, amides, alcoholates, carbonates or hydrogen carbonates, acetates or tertiary amines.

The first step (step A-1) according to the present invention is conducted at a temperature of from 0° C. to 200° C. Preferably, first step (step A-1) is conducted at a temperature of from 0° C. to 120° C., more preferably at a temperature of from 0° C. to 80° C.

The first step (step A-1) according to the present invention may be conducted in the presence of a solvent. Preferably, the solvent is chosen as being water, an organic solvent or a mixture of both. Suitable organic solvents may for example be aliphatic, alicyclic or aromatic solvent.

The first step (step A-1) according to the present invention may also be conducted in the presence of a catalyst. Preferably, the catalyst is chosen as being palladium salts or complexes. More preferably, the catalyst is chosen as being a palladium complex. Suitable palladium complex catalyst may for example be generated directly in the reaction mixture by separately adding to the reaction mixture a palladium salt and a complex ligand. Suitable ligands may for example be bulky phosphines or arsines ligands, such as (R)-(-)-1-[(S)-2-(dicyclohexylphosphino)ferrocenyl]ethyldicyclohexylphosphine and its corresponding enantiomer, or a mixture of both; (R)-(-)-1-[(S)-2-(dicyclohexylphosphino)ferrocenyl]ethyldiphenylphosphine and its corresponding enantiomer, or a mixture of both; (R)-(-)-1-[(S)-2-(diphenylphosphino)ferrocenyl]ethyldi-*i*-butylphosphine and its corresponding enantiomer, or a mixture of both; or (R)-(-)-1-[(S)-2-(diphe-

nylphosphino)ferrocenyl]ethyldicyclohexylphosphine and its corresponding enantiomer, or a mixture of both.

The fifth step (step A-5) according to the present invention is conducted in the presence of a hydride donor. Preferably, the hydride donor is chosen as being metal or metallorid hydrides such as LiAlH_4 , NaBH_4 , KBH_4 , B_2H_6 .

The fifth step (step A-5) according to the present invention is conducted in the presence of a catalyst. Preferably, the catalyst is chosen as being Co(II)-Chloride , Ni(II)-chloride , ammonia or one of its salt, Palladium on charcoal, Raney Nickel, Raney Cobalt or Platinum.

The fifth step (step A-5) according to the present invention is conducted at a temperature of from 0°C . to 150°C . Preferably the temperature is of from 10°C . to 120°C . More preferably, the temperature is of from 10°C . to 80°C .

The fifth step (step A-5) according to the present invention is conducted under a pressure of from 1 bar to 100 bar. Preferably the pressure is of from 1 bar to 50 bar.

The fifth step (step A-5) according to the present invention may be conducted in the presence of an organic solvent, of water or of a mixture thereof. Preferably, the solvent is chosen as being ether, alcohol, carboxylic acid, or a mixture thereof with water or pure water.

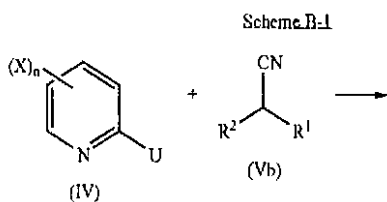
A second example (B) of such a process may be when:

R^1 , R^2 , X, n are as defined above; and

R^3 , R^4 , R^5 are hydrogen atoms;

then, the amine derivative of general formula (II) may be prepared according to a process which comprises:

a first step according to reaction scheme B-1:



in which:

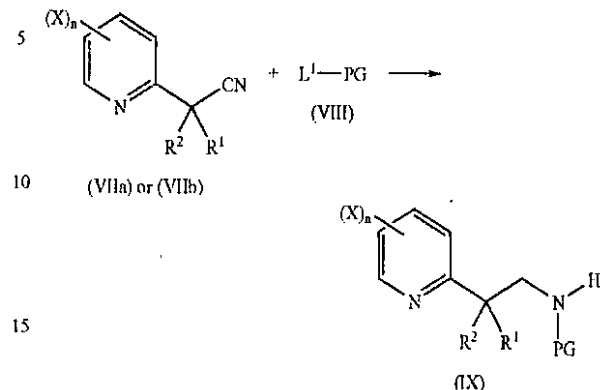
R^1 , R^2 , X and n are as defined above;

U is a leaving group chosen as being a halogen atom, a $\text{C}_1\text{-C}_6$ alkylsulfonate or a $\text{C}_1\text{-C}_6$ haloalkylsulfonate;

comprising the arylation of a compound of general formula (Vb) by a pyridine derivative of general formula (IV) to provide a 2-pyridylacetonitrile derivative of general formula (VIIb), in the presence of a base and at a temperature of from -100°C . to 200°C ;

a second step according to reaction scheme B-2:

Scheme B-2



in which:

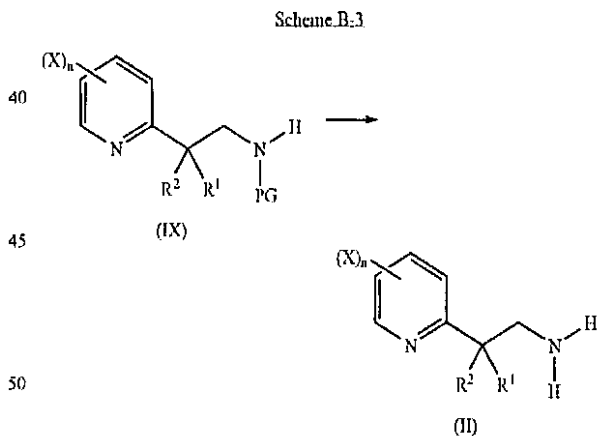
R^1 , R^2 , X and n are as defined above;

L^1 is a leaving group chosen as being a $-\text{OR}^{79}$ group or a $-\text{OCOR}^{79}$ group, R^{79} being a $\text{C}_1\text{-C}_6$ alkyl, a $\text{C}_1\text{-C}_6$ haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

PG represents a protecting group which may be a $-\text{COOR}^{79}$ group or $-\text{COR}^{79}$ group, R^{79} being a $\text{C}_1\text{-C}_6$ alkyl, a $\text{C}_1\text{-C}_6$ haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

comprising the reduction, by hydrogenation or by an hydride donor, of a compound of general formula (VIIa) or (VIIb), in the presence of a compound of general formula (VIII) to produce a compound of general formula (IX),

a third step according to reaction scheme B-3:



in which:

R^1 , R^2 , X and n are as defined above;

PG represents a protecting group which may be a $-\text{COOR}^{79}$ group or $-\text{COR}^{79}$ group, R^{79} being a $\text{C}_1\text{-C}_6$ alkyl, a $\text{C}_1\text{-C}_6$ haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

comprising a deprotection reaction, in an acidic or in a basic medium, of a compound of general formula (IX) to provide an amine derivative of general formula (II) or one of its salt.

The first step (step B-1) is conducted at a temperature of from -100°C . to 200°C . Preferably, first step (step A-1) is conducted at a temperature of from -80°C . to 120°C ., more preferably at a temperature of from -80°C . to 80°C .

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The first step (step B-1) according to the present invention is conducted in the presence of a base. Preferably, the base will be chosen as being an inorganic or an organic base. Suitable examples of such bases may for example be alkaline earth metal or alkali metal hydrides, hydroxides, amides, alcoholates, carbonates or hydrogen carbonates, acetates or tertiary amines.

The first step (step B-1) according to the present invention may be conducted in the presence of a solvent. Preferably, the solvent is chosen as being water, an organic solvent or a mixture of both. Suitable organic solvents may for example be aliphatic, alicyclic or aromatic solvent.

The first step (step B-1) according to the present invention may also be conducted in the presence of a catalyst. Preferably, the catalyst is chosen as being palladium salts or complexes. More preferably, the catalyst is chosen as being a palladium complex. Suitable palladium complex catalyst may for example be generated directly in the reaction mixture by separately adding to the reaction mixture a palladium salt and a complex ligand. Suitable ligands may for example be bulky phosphines or arsines ligands, such as (R)-(-)-1-[(S)-2-(dicyclohexylphosphino)ferrocenyl]ethyl-dicyclohexylphosphine and its corresponding enantiomer, or a mixture of both; (R)-(-)-1-[(S)-2-(dicyclohexylphosphino)ferrocenyl]ethyl-diphenylphosphine and its corresponding enantiomer, or a mixture of both; (R)-(-)-1-[(S)-2-(diphenylphosphino)ferrocenyl]ethyl-di-t-butylphosphine and its corresponding enantiomer, or a mixture of both; or (R)-(-)-1-[(S)-2-(diphenylphosphino)ferrocenyl]ethyl-dicyclohexylphosphine and its corresponding enantiomer, or a mixture of both.

The preferred conditions under which step B-2 is conducted are the same than the preferred conditions under which step A-4 of the above mentioned process A is conducted.

The preferred conditions under which step B-3 is conducted are the same than the preferred conditions under which step A-5 of the above mentioned process A is conducted.

A third example (C) of such a process may be when:

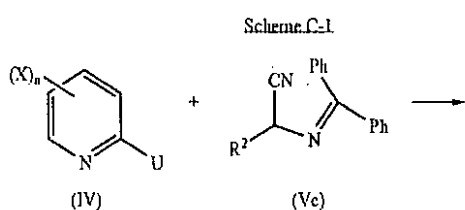
R^2 , X, n are as defined above;

R^1 is a formylamino, a C_1 - C_8 -alkylcarbonylamino, a C_1 - C_8 -halogenoalkylcarbonylamino having 1 to 5 halogen atoms, a phenylcarbonylamino or a 2,6-dichlorophenylcarbonylamino; and

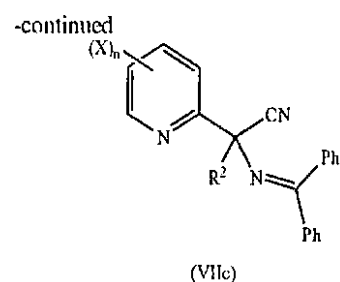
R^3 , R^4 , R^5 are hydrogen atoms;

then, the amine derivative of general formula (II) may be prepared according to a process comprising:

a first step according to reaction scheme C-1:



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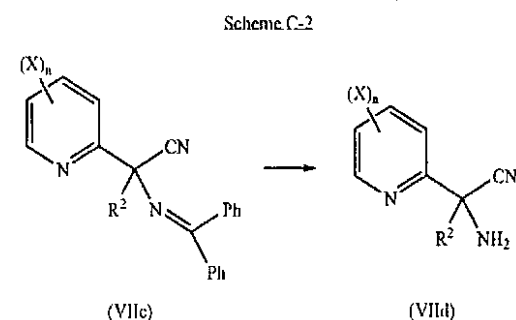
in which:

R^2 , X, n are as defined above;

U is a leaving group chosen as being a halogen atom, a C_1 - C_6 alkylsulfonate or a C_1 - C_6 haloalkylsulfonate;

comprising the arylation of a compound of general formula (Vc) by a pyridine derivative of general formula (IV) to provide a 2-pyridylacetone nitrile derivative of general formula (VIIc), in the presence of a base and at a temperature of from -100° C. to 200° C.;

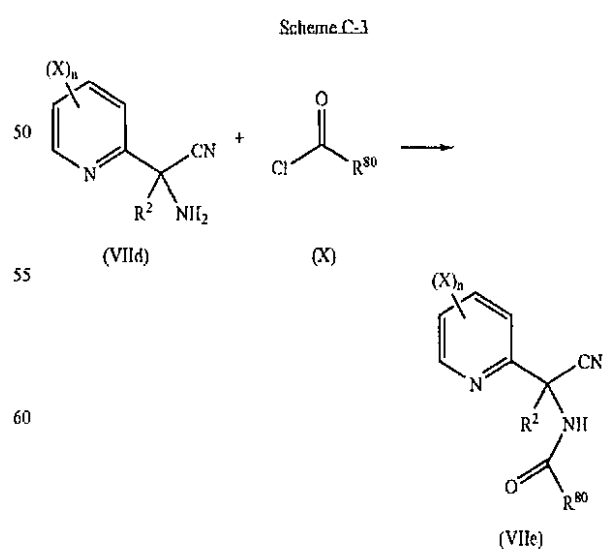
a second step according to reaction scheme C-2:



in which R^2 , X, n are as defined above;

comprising the de-protection, by acidic hydrolysis, of a compound of general formula (VIIc), to produce a compound of general formula (VIIId) or one of its salt;

a third step according to reaction scheme C-3:



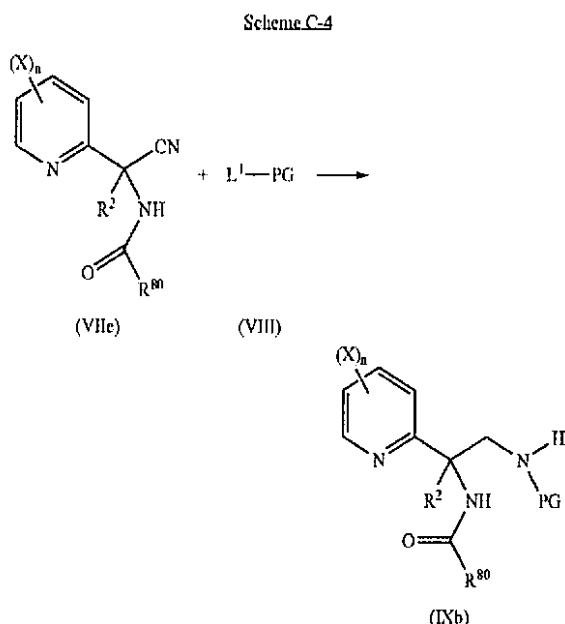
in which:

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R^2 , X, n are as defined above;
 R^{80} is a hydrogen atom, a C_1 - C_6 alkyl group, a C_1 - C_8 -halogenoalkyl having 1 to 5 halogen atoms, a phenyl or 2,6 dichlorophenyl;

comprising the coupling between a compound of general formula (VIIc) and an acyl chloride of general formula (X) to produce a compound of general formula (VIIe).

a fourth step according to reaction scheme C-4:



in which:

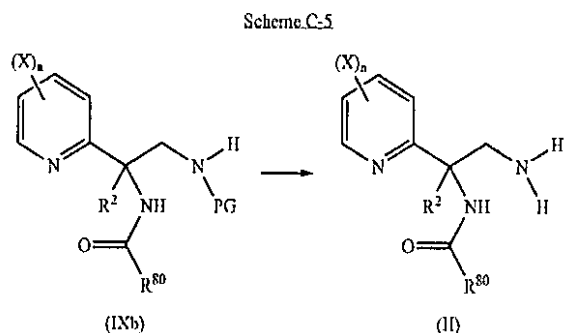
R^2 , X, n are as defined above;
 R^{80} is a hydrogen atom, a C_1 - C_6 alkyl group, a C_1 - C_8 -halogenoalkyl having 1 to 5 halogen atoms, a phenyl or 2,6 dichlorophenyl;

L^1 is a leaving group chosen as being a $-OR^{79}$ group or a $OCOR^{79}$ group, R^{79} being a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

PG represents a protecting group which may be a $-COOR^{79}$ group or $-COR^{79}$ group, R^{79} being a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

comprising the reduction, by hydrogenation or by an hydride donor, of a compound of general formula (VIIe), in the presence of a compound of general formula (VIII) to produce a compound of general formula (IXb);

a fifth step according to scheme C-5



in which:

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R^2 , X, n are as defined above;

R^{80} is a hydrogen atom, a C_1 - C_6 alkyl group, a C_1 - C_8 -halogenoalkyl having 1 to 5 halogen atoms, a phenyl or 2,6 dichlorophenyl;

L^1 is a leaving group chosen as being a $-OR^{79}$ group or a $OCOR^{79}$ group, R^{79} being a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

PG represents a protecting group which may be a $-COOR^{79}$ group or $-COR^{79}$ group, R^{79} being a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

comprising a deprotection reaction, in an acidic or in a basic medium, of a compound of general formula (IXb) to provide an amine derivative of general formula (II) or one of its salt.

A fourth example (D) of such a process may be when:

R^1 , X, Y, n and p are as defined above;

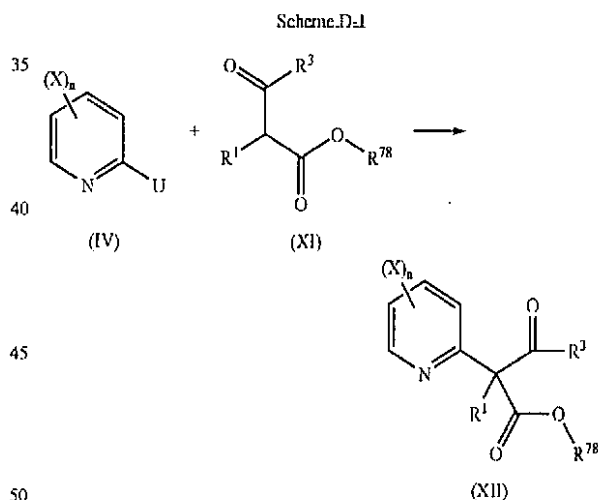
R^2 is a hydrogen atom;

R^3 is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl or a phenyl; and

R^5 is a hydrogen atom, a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a C_1 - C_6 alkoxy or a C_3 - C_7 cycloalkyl;

then, the amine derivative of general formula (II) may be prepared according to a process comprising:

a first step according to reaction scheme D-1:



in which:

R^1 , X and n are as defined above;

R^3 is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl or a phenyl;

R^{78} is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

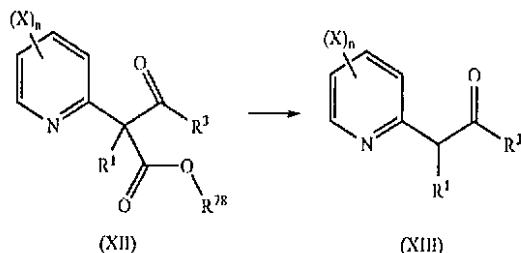
U is a leaving group chosen as being a halogen, a C_1 - C_6 alkylsulfonate or a C_1 - C_6 haloalkylsulfonate;

comprising the arylation of a ketoacetate derivative of general formula (XI) by a pyridine derivative of general formula (IV), to provide a 2-(pyridyl)ketoacetate derivative of general formula (XII), in the presence of a base, at a temperature of from 0°C . to 200°C ;

a second step according to reaction scheme D-2:

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Scheme D-2



in which:

R^1 , X , n are as defined above;

R^2 is a hydrogen atom;

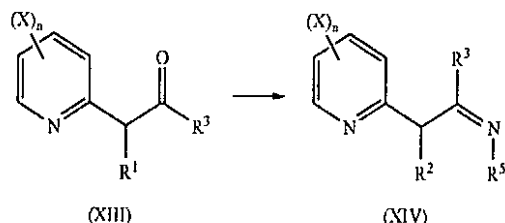
R^3 is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl or a phenyl;

R^{78} is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

comprising a basic hydrolysis, an acidic hydrolysis or a displacement by an halide of a compound of general formula (XII) in the same or a different pot to provide, upon heating at a temperature of from 40° C. to reflux, a 2-pyridylketone derivative of general formula (XIII);

a third step according to reaction scheme D-3:

Scheme D-3



in which:

R^1 , X and n are as defined above;

R^2 is a hydrogen atom;

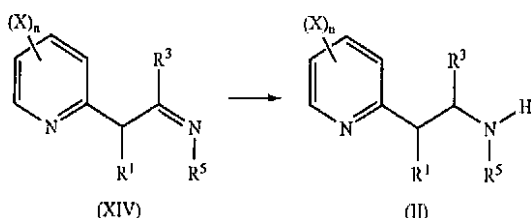
R^3 is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl or a phenyl;

R^5 is a hydrogen atom, a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a C_1 - C_6 alkoxy or a C_3 - C_7 cycloalkyl;

comprising the reaction of a compound of general formula (XIII) with an amine of formula R^5 -NH₂ to provide an imine derivative of general formula (XIV);

a fourth step according to scheme D-4:

Scheme D-4



in which:

R^1 , X and n are as defined above;

R^2 is a hydrogen atom;

R^3 is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, a phenyl;

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R^5 is a hydrogen atom, a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a C_1 - C_6 alkoxy or a C_3 - C_7 cycloalkyl;

comprising the reduction of an imine derivative of general formula (XIV) by hydrogenation or by an hydride donor, in the same or a different pot to provide an amine derivative of general formula (II) or one of its salt.

A fifth example (E) of such a process may be when:

R^1 , R^2 , X , Y , n and p are as defined above;

R^3 is a hydrogen atom

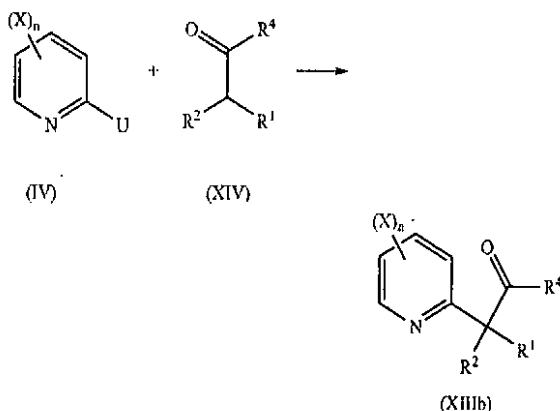
R^4 is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, a phenyl; and

R^5 is a hydrogen atom, a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a C_1 - C_6 alkoxy or a C_3 - C_7 cycloalkyl;

then, the amine derivative of general formula (II) may be prepared according to a process comprising:

a first step according to reaction scheme E-1:

Scheme E-1



in which:

R^1 , R^2 , X and n are as defined above;

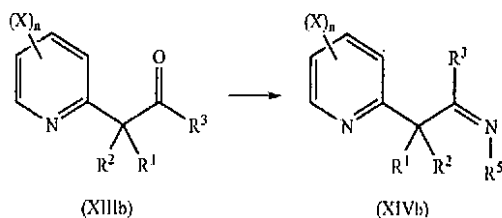
R^4 is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl or a phenyl;

U is a leaving group chosen as being a halogen, a C_1 - C_6 alkylsulfonate or a C_1 - C_6 haloalkylsulfonate;

comprising the arylation of a ketone derivative of general formula (XIV) by a pyridine derivative of general formula (IV), to provide a 2-(pyridyl)ketone derivative of general formula (XIIIb), in the presence of a base, at a temperature of from 0° C. to 200° C.;

a second step according to reaction scheme E-2:

Scheme E-2



in which:

R^1 , R^2 , X and n are as defined above;

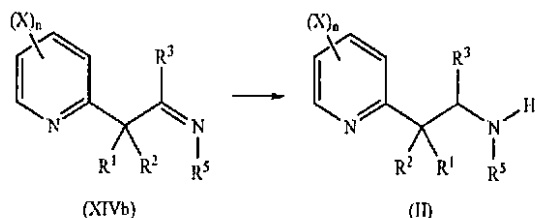
R^4 is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl or a phenyl;

R^5 is a hydrogen atom, a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a C_1 - C_6 alkoxy or a C_3 - C_7 cycloalkyl;

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comprising the reaction of a compound of general formula (XIVb) with an amine of formula R^5-NH_2 to provide an imine derivative of general formula (XIVb);
a third step according to scheme E-3;

Scheme E-3



in which:

R^1 , R^2 , X and n are as defined above;
 R^3 is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl or a phenyl;
 R^4 is a hydrogen atom;
 R^5 is a hydrogen atom, a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a C_1 - C_6 alkoxy or a C_3 - C_7 cycloalkyl;

comprising the reduction of an imine derivative of general formula (XIVb) by hydrogenation or by an hydride donor, in the same or a different pot to provide an amine derivative of general formula (II) or one of its salt.

A sixth example (F) of such a process may be when:

R^2 , X, n are as defined above;

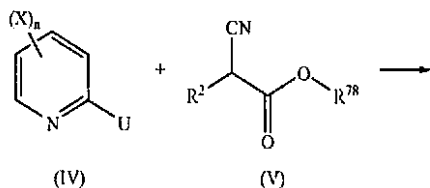
R^1 is a cyano group, a hydroxy group, a C_2 - C_6 -alkynyl, a C_1 - C_6 -alkylamino, a di- C_1 - C_6 -alkylamino, a C_1 - C_6 -alkoxy, a C_1 - C_6 -halogenoalkoxy having 1 to 5 halogen atoms, a C_1 - C_6 -alkylsulfanyl, a C_1 - C_6 -halogenoalkylsulfanyl having 1 to 5 halogen atoms, a C_2 - C_6 -alkenyloxy, a C_2 - C_6 -halogenoalkenyloxy having 1 to 5 halogen atoms, a C_3 - C_6 -alkynyloxy, a C_3 - C_6 -halogenoalkynyloxy having 1 to 5 halogen atoms, a benzyloxy, a benzylsulfanyl, a benzylamino, a phenoxy, a phenylsulfanyl, a phenylamino, or a phenyl sulphanyl group, a C_1 - C_6 -alkylcarbonyloxy, a C_1 - C_6 -halogenoalkylcarbonyloxy having 1 to 5 halogen atoms; and

R^3 and R^4 are hydrogen atoms;

then, the amine derivative of general formula (II) may be prepared according to a process comprising:

a first step according to reaction scheme F-1;

Scheme F-1



in which:

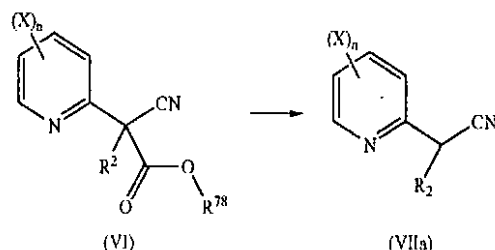
28

R^2 , X and n are as defined above;
 R^{78} is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;
U is a leaving group chosen as being a halogen, a C_1 - C_6 alkylsulfonate or a C_1 - C_6 haloalkylsulfonate;

comprising the arylation of a cyanoacetate derivative of general formula (V) by a pyridine derivative of general formula (IV), to provide a 2-(pyridyl)cyanoacetate derivative of general formula (VI), in the presence of a base, at a temperature of from 0° C. to 200° C.;

a second step according to reaction scheme F-2;

Scheme F-2



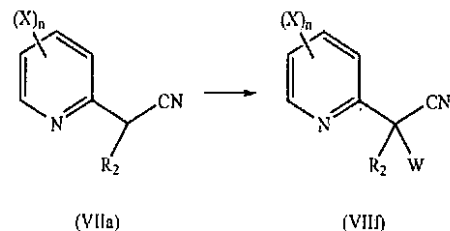
in which:

R^2 , X, n are as defined above;
 R^1 is a hydrogen atom;
 R^{78} is a C_1 - C_6 alkyl, a C_1 - C_6 haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

comprising a basic hydrolysis, an acidic hydrolysis or a displacement by an halide of a compound of general formula (VI) in the same or a different pot to provide, upon heating at a temperature of from 40° C. to reflux, a 2-pyridylacetonitrile derivative of general formula (VIIa);

a third step according to reaction scheme F-3;

Scheme F-3

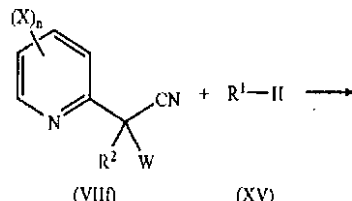


in which:

R^2 , X, n are as defined above;
W is a halogen atom;

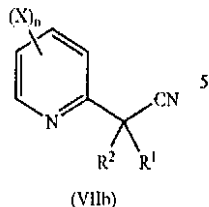
comprising the halogenation of a compound of general formula (VIIa) to provide a compound of general formula (VIIb);
a fourth step according to reaction scheme F-4;

Scheme F-4



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-continued



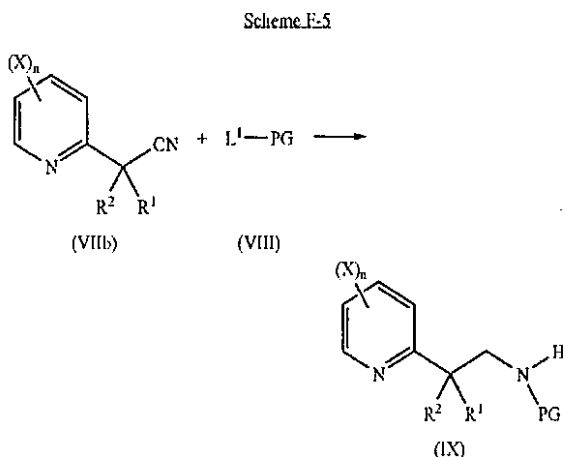
in which:

R², X, n are as defined above;

W is a halogen atom;

R¹ is a cyano group, a hydroxy group, a C₂-C₆-alkynyl, a C₁-C₆-alkylamino, a di-C₁-C₆-alkylamino, a C₁-C₆-alkoxy, a C₁-C₆-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₆-alkylsulfanyl, a C₁-C₆-halogenoalkylsulfanyl having 1 to 5 halogen atoms, a C₂-C₆-alkenyl, a C₂-C₆-halogenoalkenyl having 1 to 5 halogen atoms, a C₃-C₆-alkynyl, a C₃-C₆-halogenoalkynyl having 1 to 5 halogen atoms, a benzyloxy, a benzylsulfanyl, a benzylamino, a phenoxy, a phenylsulfanyl, a phenylamino, or a phenyl sulphanyl group, a C₁-C₆-alkylcarbonyloxy or a C₁-C₆-halogenoalkylcarbonyloxy having 1 to 5 halogen atoms; comprising the nucleophilic substitution, in the presence of a base, of a compound of general formula (VIIb) in the presence of a compound of general formula (VIII) to produce a compound of general formula (IX), at a temperature of from -78° C. to 150° C.;

a fifth step according to reaction scheme F-5:



in which:

R², X, n are as defined above;

R¹ is a cyano group, a hydroxy group, a C₂-C₆-alkynyl, a C₁-C₆-alkylamino, a di-C₁-C₆-alkylamino, a C₁-C₆-alkoxy, a C₁-C₆-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₆-alkylsulfanyl, a C₁-C₆-halogenoalkylsulfanyl having 1 to 5 halogen atoms, a C₂-C₆-alkenyl, a C₂-C₆-halogenoalkenyl having 1 to 5 halogen atoms, a C₃-C₆-alkynyl, a C₃-C₆-halogenoalkynyl having 1 to 5 halogen atoms, a benzyloxy, a benzylsulfanyl, a benzylamino, a phenoxy, a phenylsulfanyl, a phenylamino, or a phenyl sulphanyl group, a C₁-C₆-alkylcarbonyloxy or a C₁-C₆-halogenoalkylcarbonyloxy having 1 to 5 halogen atoms;

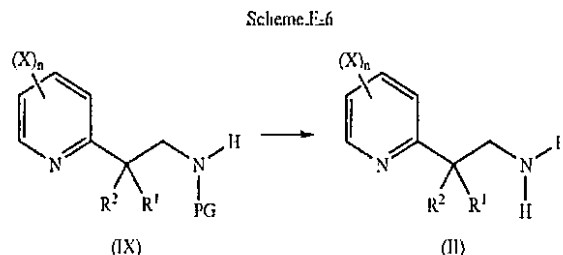
30

L¹ is a leaving group chosen as being a —OR⁷⁹ group or a —OCOR⁷⁹ group, R⁷⁹ being a C₁-C₆ alkyl, a C₁-C₆ haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

PG represents a protecting group which may be a —COOR⁷⁹ group or —COR⁷⁹ group, R⁷⁹ being a C₁-C₆ alkyl, a C₁-C₆ haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

comprising the reduction, by hydrogenation or by an hydride donor, of a compound of general formula (VIIb), in the presence of a catalyst and in the presence of a compound of general formula (VIII) to produce a compound of general formula (IX), at a temperature of from 0° C. to 150° C. and under a pressure of from 1 bar and 100 bar;

a sixth step according to reaction scheme F-6:



in which:

R², X, n are as defined above;

R¹ is a cyano group, a hydroxy group, a C₂-C₆-alkynyl, a C₁-C₆-alkylamino, a di-C₁-C₆-alkylamino, a C₁-C₆-alkoxy, a C₁-C₆-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₆-alkylsulfanyl, a C₁-C₆-halogenoalkylsulfanyl having 1 to 5 halogen atoms, a C₂-C₆-alkenyl, a C₂-C₆-halogenoalkenyl having 1 to 5 halogen atoms, a C₃-C₆-alkynyl, a C₃-C₆-halogenoalkynyl having 1 to 5 halogen atoms, a benzyloxy, a benzylsulfanyl, a benzylamino, a phenoxy, a phenylsulfanyl, a phenylamino, or a phenyl sulphanyl group, a C₁-C₆-alkylcarbonyloxy or a C₁-C₆-halogenoalkylcarbonyloxy having 1 to 5 halogen atoms;

PG represents a protecting group which may be a —COOR⁷⁹ group or —COR⁷⁹ group, R⁷⁹ being a C₁-C₆ alkyl, a C₁-C₆ haloalkyl, a benzyl, 4-methoxybenzyl or pentafluorophenyl;

comprising a deprotection reaction, in an acidic or in a basic medium, of a compound of general formula (IX) to provide an amine derivative of general formula (II) or one of its salt.

A seventh example (G) of such a process may be when:

X, n are as defined above;

R¹ is a hydroxy group; and

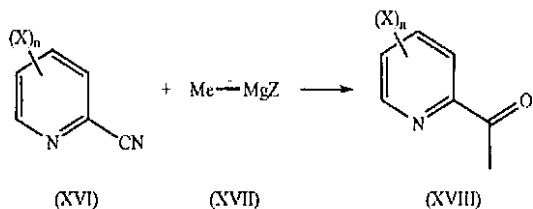
R³, R², and R⁴ are hydrogen atoms;

then, the amine derivative of general formula (II) may be prepared according to a process comprising:

a first step according to reaction scheme G-1:

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Scheme G-1



in which:

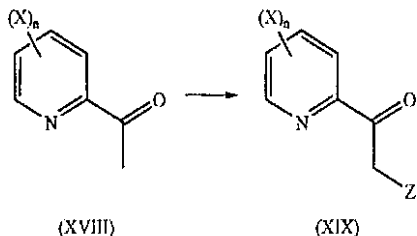
X and n are as defined above;

Z is a halogen atom;

comprising the addition of a methyl magnesium halogenide of general formula (XVII) on a pyridine derivative of general formula (XVI) to provide a compound of general formula (XVIII);

a second step according to reaction scheme G-2:

Scheme G-2



in which:

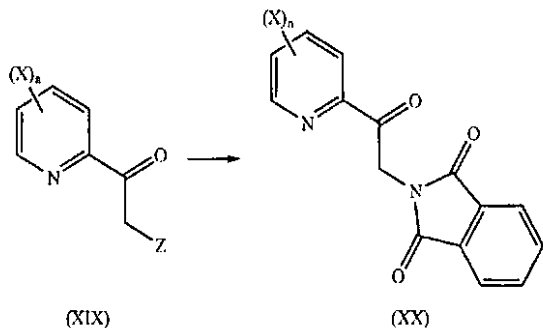
X and n are as defined above;

Z is a halogen atom;

comprising the halogenation of a compound of general formula (XVIII) into a compound of general formula (XIX) by use of a halogenating agent such as chlorine, bromine, hypochlorite ions, hypobromite ions, trichloride ions, tribromide ions, N-chloro imides, N-chloro amides, N-chloro amines, N-bromo imides, N-bromo amides or N-bromo amines;

a third step according to reaction scheme G-3:

Scheme G-3



in which:

X and n are as defined above;

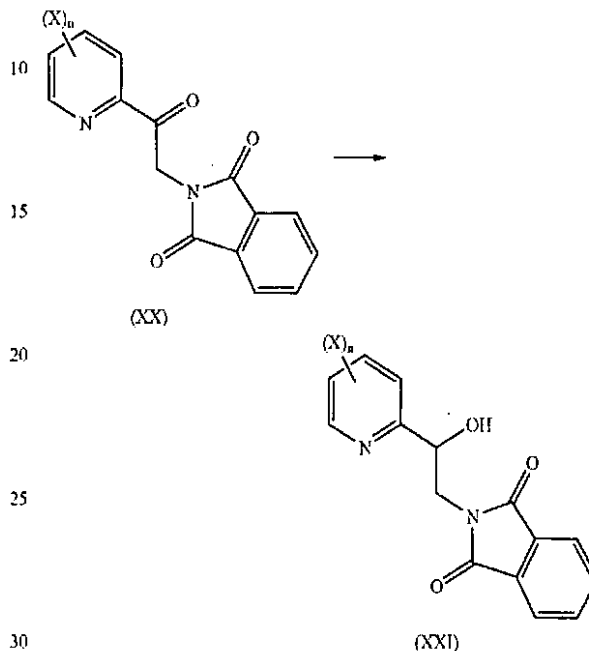
Z is a halogen atom;

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comprising the nucleophilic substitution of a compound of general formula (XIX) by a phthalimide salt to produce a compound of general formula (XX);

a fourth step according to reaction scheme G-4:

Scheme G-4

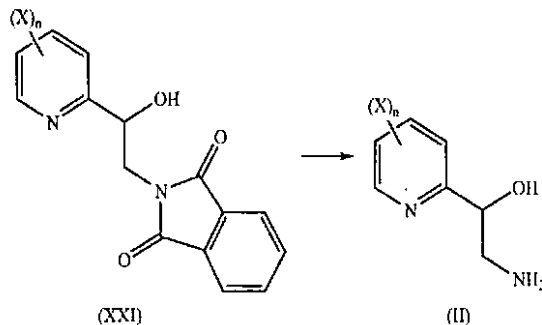


in which X and n are as defined above;

comprising the reduction of a compound of general formula (XX) by a hydride donor to produce a compound of general formula (XXI);

a fifth step according to reaction scheme G-5:

Scheme G-5



in which X and n are as defined above;

comprising the de-protection of a compound of general formula (XXI) by reacting it with hydrazine hydrate or a hydrazine salt to provide an amine derivative of general formula (II) or one of its salt;

Compounds according to the present invention can be prepared according to the above described processes. It will nevertheless be understood that, on the basis of his general knowledge and of available publications, the skilled worker will be able to adapt these processes according to the specifics of each of the compounds which it is desired to synthesise.

The present invention also relates to a fungicidal composition comprising an effective amount of an active material of

general formula (I). Thus, according to the present invention, there is provided a fungicidal composition comprising, as an active ingredient, an effective amount of a compound of general formula (I) as defined above and an agriculturally acceptable support, carrier or filler.

In the present specification, the term "support" denotes a natural or synthetic, organic or inorganic material with which the active material is combined to make it easier to apply, notably to the parts of the plant. This support is thus generally inert and should be agriculturally acceptable. The support may be a solid or a liquid. Examples of suitable supports include clays, natural or synthetic silicates, silica, resins, waxes, solid fertilisers, water, alcohols, in particular butanol, organic solvents, mineral and plant oils and derivatives thereof. Mixtures of such supports may also be used.

The composition may also comprise additional components. In particular, the composition may further comprise a surfactant. The surfactant can be an emulsifier, a dispersing agent or a wetting agent of ionic or non-ionic type or a mixture of such surfactants. Mention may be made, for example, of polyacrylic acid salts, lignosulphonic acid salts, phenolsulphonic or naphthalenesulphonic acid salts, polycondensates of ethylene oxide with fatty alcohols or with fatty acids or with fatty amines, substituted phenols (in particular alkylphenols or arylphenols), salts of sulphosuccinic acid esters, taurine derivatives (in particular alkyl taurates), phosphoric esters of polyoxyethylated alcohols or phenols, fatty acid esters of polyols, and derivatives of the above compounds containing sulphate, sulphonate and phosphate functions. The presence of at least one surfactant is generally essential when the active material and/or the inert support are water-insoluble and when the vector agent for the application is water. Preferably, surfactant content may be comprised between 5% and 40% by weight of the composition.

Optionally, additional components may also be included, e.g. protective colloids, adhesives, thickeners, thixotropic agents, penetration agents, stabilisers, sequestering agents. More generally, the active materials can be combined with any solid or liquid additive, which complies with the usual formulation techniques.

In general, the composition according to the invention may contain from 0.05 to 99% (by weight) of active material, preferably 10 to 70% by weight.

Compositions according to the present invention can be used in various forms such as aerosol dispenser, capsule suspension, cold fogging concentrate, dustable powder, emulsifiable concentrate, emulsion oil in water, emulsion water in oil, encapsulated granule, fine granule, flowable concentrate for seed treatment, gas (under pressure), gas generating product, granule, hot fogging concentrate, macrogranule, microgranule, oil dispersible powder, oil miscible flowable concentrate, oil miscible liquid, paste, plant rodlet, powder for dry seed treatment, seed coated with a pesticide, soluble concentrate, soluble powder, solution for seed treatment, suspension concentrate (flowable concentrate), ultra low volume (ulv) liquid, ultra low volume (ulv) suspension, water dispersible granules or tablets, water dispersible powder for slurry treatment, water soluble granules or tablets, water soluble powder for seed treatment and wettable powder.

These compositions include not only compositions which are ready to be applied to the plant or seed to be treated by means of a suitable device, such as a spraying or dusting device, but also concentrated commercial compositions which must be diluted before application to the crop.

The compounds of the invention can also be mixed with one or more insecticides, fungicides, bactericides, attractant acaricides or pheromones or other compounds with biological

activity. The mixtures thus obtained have a broadened spectrum of activity. The mixtures with other fungicides are particularly advantageous.

The fungicidal compositions of the present invention can be used to curatively or preventively control the phytopathogenic fungi of crops. Thus, according to a further aspect of the present invention, there is provided a method for curatively or preventively controlling the phytopathogenic fungi of crops characterised in that a fungicidal composition as hereinbefore defined is applied to the seed, the plant and/or to the fruit of the plant or to the soil in which the plant is growing or in which it is desired to grow.

The composition as used against phytopathogenic fungi of crops comprises an effective and non-phytotoxic amount of an active material of general formula (I).

The expression "effective and non-phytotoxic amount" means an amount of composition according to the invention which is sufficient to control or destroy the fungi present or liable to appear on the crops, and which does not entail any appreciable symptom of phytotoxicity for the said crops. Such an amount can vary within a wide range depending on the fungus to be controlled, the type of crop, the climatic conditions and the compounds included in the fungicidal composition according to the invention.

This amount can be determined by systematic field trials, which are within the capabilities of a person skilled in the art.

The method of treatment according to the present invention is useful to treat propagation material such as tubers or rhizomes, but also seeds, seedlings or seedlings pricking out and plants or plants pricking out. This method of treatment can also be useful to treat roots. The method of treatment according to the present invention can also be useful to treat the overground parts of the plant such as trunks, stems or stalks, leaves, flowers and fruits of the concerned plant.

Among the plants that can be protected by the method according to the invention, mention may be made of cotton; flax; vine; fruit crops such as Rosaceae sp. (for instance pip fruits such as apples and pears, but also stone fruits such as apricots, almonds and peaches), Ribesioideae sp., Juglandaceae sp., Betulaceae sp., Anacardiaceae sp., Fagaceae sp., Moraceae sp., Oleaceae sp., Actinidiaceae sp., Lauraceae sp., Musaceae sp. (for instance banana trees and plantains), Rubiaceae sp., Theaceae sp., Sterculiaceae sp., Rutaceae sp. (for instance lemons, oranges and grapefruits); leguminous crops such as Solanaceae sp. (for instance tomatoes), Liliaceae sp., Asteraceae sp. (for instance lettuces), Umbelliferae sp., Cruciferae sp., Chenopodiaceae sp., Cucurbitaceae sp., Papilionaceae sp. (for instance peas), Rosaceae sp. (for instance strawberries); big crops such as Graminae sp. (for instance maize, cereals such as wheat, rice, barley and triticale), Asteraceae sp. (for instance sunflower), Cruciferae sp. (for instance colza), Papilionaceae sp. (for instance soja), Solanaceae sp. (for instance potatoes), Chenopodiaceae sp. (for instance beetroots); horticultural and forest crops; as well as genetically modified homologues of these crops.

Among the plants and the possible diseases of these plants protected by the method according to the present invention, mention may be made of:

wheat, as regards controlling the following seed diseases: fusaria (*Microdochium nivale* and *Fusarium roseum*), stinking smut (*Tilletia caries*, *Tilletia controversa* or *Tilletia indica*), septoria disease (*Septoria nodorum*) and loose smut;

wheat, as regards controlling the following diseases of the aerial parts of the plant: cereal eyespot (*Tapesia yellunda*, *Tapesia acuiformis*), take-all (*Gaeumannomyces graminis*), foot blight (*F. culmorum*, *F. graminearum*),

black speck (*Rhizoctonia cerealis*), powdery mildew (*Erysiphe graminis forma specie tritici*), rusts (*Puccinia striiformis* and *Puccinia recondita*) and septoria diseases (*Septoria tritici* and *Septoria nodorum*);

wheat and barley, as regards controlling bacterial and viral diseases, for example barley yellow mosaic;

barley, as regards controlling the following seed diseases: net blotch (*Pyrenophora graminea*, *Pyrenophora teres* and *Cochliobolus sativus*), loose smut (*Ustilago nuda*) and fusaria (*Microdochium nivale* and *Fusarium roseum*);

barley, as regards controlling the following diseases of the aerial parts of the plant: cercal eyespot (*Tapesia yellundae*), net blotch (*Pyrenophora teres* and *Cochliobolus sativus*), powdery mildew (*Erysiphe graminis forma specie hordei*), dwarf leaf rust (*Puccinia hordei*) and leaf blotch (*Rhynchosporium secalis*);

potato, as regards controlling tuber diseases (in particular *Helminthosporium solani*, *Phoma tuberosa*, *Rhizoctonia solani*, *Fusarium solani*), mildew (*Phytophthora infestans*) and certain viruses (virus Y);

potato, as regards controlling the following foliage diseases: early blight (*Alternaria solani*), mildew (*Phytophthora infestans*);

cotton, as regards controlling the following diseases of young plants grown from seeds: damping-off and collar rot (*Rhizoctonia solani*, *Fusarium oxysporum*) and black root rot (*Thielaviopsis basicola*);

protein yielding crops, for example peas, as regards controlling the following seed diseases: anthracnose (*Ascochyta pisi*, *Mycosphaerella pinodes*), fusaria (*Fusarium oxysporum*), grey mould (*Botrytis cinerea*) and mildew (*Peronospora pisi*);

oil-bearing crops, for example rape, as regards controlling the following seed diseases: *Phoma lingam*, *Alternaria brassicae* and *Sclerotinia sclerotiorum*;

corn, as regards controlling seed diseases: (*Rhizopus* sp., *Penicillium* sp., *Trichoderma* sp., *Aspergillus* sp., and *Gibberella fujikuroi*);

flax, as regards controlling the seed disease: *Alternaria linicola*;

forest trees, as regards controlling damping-off (*Fusarium oxysporum*, *Rhizoctonia solani*);

rice, as regards controlling the following diseases of the aerial parts: blast disease (*Magnaporthe grisea*), bordered sheath spot (*Rhizoctonia solani*);

leguminous crops, as regards controlling the following diseases of seeds or of young plants grown from seeds: damping-off and collar rot (*Fusarium oxysporum*, *Fusarium roseum*, *Rhizoctonia solani*, *Pythium* sp.);

leguminous crops, as regards controlling the following diseases of the aerial parts: grey mould (*Botrytis* sp.), powdery mildews (in particular *Erysiphe cichoracearum*, *Sphaerotheca fuliginea* and *Leveillula taurica*), fusaria (*Fusarium oxysporum*, *Fusarium roseum*), leaf spot (*Cladosporium* sp.), *alternaria* leaf spot (*Alternaria* sp.), anthracnose (*Colletotrichum* sp.), *septoria* leaf spot (*Septoria* sp.), black speck (*Rhizoctonia solani*), mildews (for example *Bremia lactucae*, *Peronospora* sp., *Pseudoperonospora* sp., *Phytophthora* sp.);

fruit trees, as regards diseases of the aerial parts: *monilia* disease (*Monilia fructigenae*, *M. laxa*), scab (*Venturia inaequalis*), powdery mildew (*Podosphaera leucotricha*);

vine, as regards diseases of the foliage: in particular grey mould (*Botrytis cinerea*), powdery mildew (*Uncinula necator*), black rot (*Gignardia biwelli*) and mildew (*Plasmopara viticola*);

beetroot, as regards the following diseases of the aerial parts: *cercospora* blight (*Cercospora beticola*), powdery mildew (*Erysiphe beticola*), leaf spot (*Ramularia beticola*).

The fungicide composition according to the present invention may also be used against fungal diseases liable to grow on or inside timber. The term "timber" means all types of species of wood, and all types of working of this wood intended for construction, for example solid wood, high-density wood, laminated wood, and plywood. The method for treating timber according to the invention mainly consists in contacting one or more compounds of the present invention, or a composition according to the invention; this includes for example direct application, spraying, dipping, injection or any other suitable means.

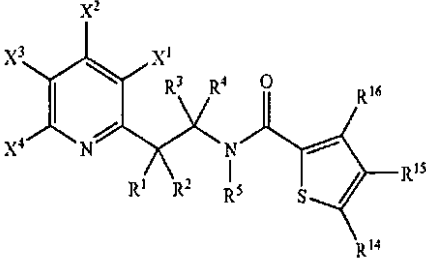
The dose of active material usually applied in the treatment according to the present invention is generally and advantageously between 10 and 800 g/ha, preferably between 50 and 300 g/ha for applications in foliar treatment. The dose of active substance applied is generally and advantageously between 2 and 200 g per 100 kg of seed, preferably between 3 and 150 g per 100 kg of seed in the case of seed treatment. It is clearly understood that the doses indicated above are given as illustrative examples of the invention. A person skilled in the art will know how to adapt the application doses according to the nature of the crop to be treated.

The fungicidal composition according to the present invention may also be used in the treatment of genetically modified organisms with the compounds according to the invention or the agrochemical compositions according to the invention. Genetically modified plants are plants into whose genome a heterologous gene encoding a protein of interest has been stably integrated. The expression "heterologous gene encoding a protein of interest" essentially means genes which give the transformed plant new agronomic properties, or genes for improving the agronomic quality of the transformed plant.

The compositions according to the present invention may also be used for the preparation of composition useful to curatively or preventively treat human and animal fungal diseases such as, for example, mycoses, dermatoses, *trichophyton* diseases and candidiasis or diseases caused by *Aspergillus* spp., for example *Aspergillus fumigatus*.

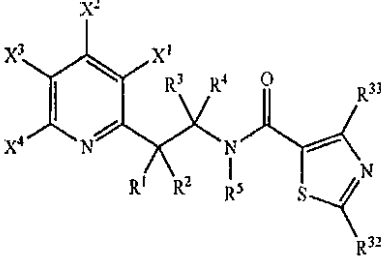
The aspects of the present invention will now be illustrated with reference to the following tables of compounds and examples. The following Tables A to C illustrate in a non-limiting manner examples of fungicidal compounds according to the present invention. In the following Examples, M+1 (or M-1) means the molecular ion peak, plus or minus 1 a.m.u. (atomic mass units) respectively, as observed in mass spectroscopy and m.p. means melting point.

TABLE A



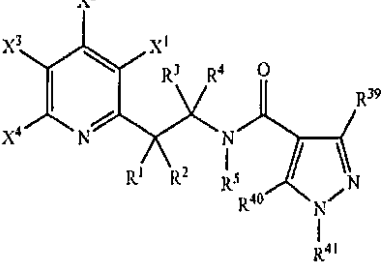
Compound	R ¹	R ²	R ³	R ⁴	R ⁵	X ¹	X ²	X ³	X ⁴	R ¹⁴	R ¹⁵	R ¹⁶	M + 1	m.p. (° C.)
A-1	H	Me	H	H	H	Cl	H	Cl	H	H	H	I	441 at 1 ³⁵ Cl	—
A-2	H	Me	H	H	H	H	H	Cl	H	H	H	I	407 at 1 ³⁵ Cl	—
A-3	H	CO ₂ H	H	H	H	F	H	Cl	F	H	H	I	—	192

TABLE B



Compound	R ¹	R ²	R ³	R ⁴	R ⁵	X ¹	X ²	X ³	X ⁴	R ³²	R ³³	M + 1
B-1	H	Me	H	H	H	Cl	H	Cl	H	Me	CF ₃	398 at 1 ³⁵ Cl
B-2	H	Me	H	H	H	H	H	Cl	H	Me	CF ₃	364 at 1 ³⁵ Cl

TABLE C



Compound	R ¹	R ²	R ³	R ⁴	R ⁵	X ¹	X ²	X ³	X ⁴	R ³⁹	R ⁴⁰	R ⁴¹	M + 1
C-1	H	Me	H	H	H	Cl	H	Cl	H	CF ₃	H	Me	381 at 1 ³⁵ Cl
C-2	H	Me	H	H	H	H	H	Cl	H	CF ₃	H	Me	347 at 1 ³⁵ Cl
C-3	Cyclopentyl		H	H	H	H	H	Cl	H	CHF ₂	H	Me	381 at 1 ³⁵ Cl
C-4	H	Et	H	H	H	H	H	Cl	H	CHF ₂	H	Me	347 at 1 ³⁵ Cl

Examples of Process for the Preparation of the Compound of General Formula (I)

Preparation of N-[2-(5-chloro-2-pyridinyl)propyl]-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxamide

Preparation of ter-butyl 2-cyano-2-(5-chloro-2-pyridinyl)propanoate

To 150 ml of dimethoxyethane was slowly added portion-wise at 0° C., 11.6 g (0.29 mol) of sodium hydride (60% dispersion in mineral oil). To this suspension, was further added dropwise at 5° C., 21.1 g (0.149 mol) of ter-butyl cyanoacetate in 50 ml of dimethoxyethane. The suspension was stirred for 45 min at room temperature. To the suspension were successively added 20 g (0.136 mol) of 2,5-dichloropyridine, 0.73 g (1.36 mmol) of (S)-(+)-1-[(R)-2-(diphenylphosphino)ferrocenyl]ethyl-ter-butylphosphine, and 1.56 g (2.7 mmol) of bis(dibenzylideneacetone)palladium(0). The black mixture was heated at reflux for 4 hours. After cooling at room temperature, 12.7 ml (0.2 mol) of methyl iodide were added dropwise and the mixture was further stirred at room temperature for 17 hours. The reaction mixture was poured into 100 ml of 1N chlorhydric acid and the aqueous phase was filtered on supersel and was extracted with ethyl acetate (3x200 ml).

The organic phase was washed with brine and dried over magnesium sulphate. The solvent was evaporated under reduced pressure to give 44.56 g of the crude product as a brown oil. The crude product was purified by flash chromatography on silica gel (eluent: heptane/ethyl acetate: 8/2) to give ter-butyl 2-cyano-2-(5-chloro-2-pyridinyl)propanoate: 23.44 g (64%) as a yellow oil.

Mass spectrum: [M+1]=267.

Preparation of 2-(5-chloro-2-pyridinyl)propanenitrile

To a solution of 23 g (0.086 mol) of ter-butyl 2-cyano-2-(5-chloro-2-pyridinyl)propanoate in 150 ml of toluene, was added 7.4 g (0.043 mol) of p-toluenesulfonic acid monohydrate. The mixture was stirred for 1.5 hours at 110° C. After cooling, the reaction mixture was poured into ice water and the pH was brought to 8 by addition of aqueous sodium hydrogenocarbonate.

The aqueous phase was extracted with ethyl acetate (3x250 ml) and the organic phase was washed with brine and dried over magnesium sulphate. The solvent was evaporated under reduced pressure to give 15.4 g of the crude product as an orange oil. The crude product was purified by flash chromatography on silica gel (eluent: heptane/ethyl acetate: 8/2) to give 2-(5-chloro-2-pyridinyl)propanenitrile: 7.8 g (54%) as a yellow oil.

Mass spectrum: [M+1]=167.

Preparation of ter-butyl 2-(5-chloro-2-pyridinyl)propylcarbamate

To a solution of 5.2 g (0.031 mol) of 2-(5-chloro-2-pyridinyl)propanenitrile in 60 ml of methanol were rapidly added 8 g (0.034 mol) of cobalt(II) chloride hexahydrate and 13.5 g (0.062 mol) of di-ter-butyl dicarbonate. The dark solution was cooled to -5° C. and 8.2 g (0.217 mol) of sodium borohydride was added portion-wise at 0° C. The reaction mixture was stirred at room temperature for 1 hour. The reaction mixture was neutralized by 100 ml of 1 N hydrochloric acid and methanol was removed under reduced pressure. The aqueous

ous phase was reextracted by ethyl acetate (2x150 ml) and the organic phase was washed with brine and dried over magnesium sulphate.

The solvent was evaporated under reduced pressure to give 11.7 g of the crude product as a black oil. The crude product was purified by flash chromatography on silica gel (eluent: heptane/ethyl acetate: 9/1) to give ter-butyl 2-(5-chloro-2-pyridinyl)propyl carbamate: 4.8 g (57%) as a yellow oil.

Mass spectrum: [M+1]=271.

Preparation of 2-(5-chloro-2-pyridinyl)propylamine hydrochloride

To a solution of 4.8 g (0.018 mol) of ter-butyl 2-(5-chloro-2-pyridinyl)propyl carbamate in 30 ml of diethyl ether were added 44 ml (0.089 mol) of a 2M solution of hydrogen chloride in diethyl ether. The mixture was stirred 3 hours at room temperature.

The solid was filtered and dried overnight under vacuum to give 2-(5-chloro-2-pyridinyl)propylamine hydrochloride: 3.5 g (94%) as a yellow solid.

Mass spectrum: [M+1]=169.

Preparation of N-[2-(5-chloro-2-pyridinyl)propyl]-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxamide

To a suspension of 100 mg (0.48 mmol) of 2-(5-chloro-2-pyridinyl)propylamine hydrochloride in 3 ml of dichloromethane was added successively 240 mg (1.68 mmol) of triethylamine and 113 mg (0.58 mmol) of 1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carbonyl chloride. The mixture was stirred 4 hours at room temperature. The reaction mixture was poured into water and the pH brought to 4.

The solvent was evaporated and the residue was purified by flash chromatography on silica gel (eluent: heptane/ethyl acetate: 8/2) to give N-[2-(5-chloro-2-pyridinyl)propyl]-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxamide as a yellow oil: 40 mg (24%) as a yellow oil.

Mass spectrum: [M+1]=347.

Examples of Biological Activity of the Compound of General Formula (I)

Example A

In Vivo Test on *Alternaria brassicae* (Leaf Spot of Crucifers)

The active ingredient tested is prepared by potter homogenisation in a concentrated suspension type formulation at 100 g/l. This suspension is then diluted with water to obtain the desired active material concentration.

Radish plants (Pernot variety) in starter cups, sown on a 50/50 peat soil-pozzolana substrate and grown at 18-20° C., are treated at the cotyledon stage by spraying with the aqueous suspension described above.

Plants, used as controls, are treated with an aqueous solution not containing the active material.

After 24 hours, the plants are contaminated by spraying them with an aqueous suspension of *Alternaria brassicae* spores (40,000 spores per cm³). The spores are collected from a 12-13-day-old culture.

The contaminated radish plants are incubated for 6-7 days at about 18° C., under a humid atmosphere.

Grading is carried out 6 to 7 days after the contamination, in comparison with the control plants.

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Under these conditions, good (at least 50%) or total protection is observed at a dose of 330 ppm with the following compounds: A1.

Under these conditions, good (at least 50%) or total protection is observed at a dose of 125 ppm with the following compound: C4.

Example B

In Vivo Test on *Pyrenophora teres* (Barley Net Blotch)

The active ingredient tested is prepared by potter homogenisation in a concentrated suspension type formulation at 100 g/l. This suspension is then diluted with water to obtain the desired active material concentration.

Barley plants (Express variety) in starter cups, sown on a 50/50 peat soil-pozzolana substrate and grown at 12° C., are treated at the 1-leaf stage (10 cm tall) by spraying with the aqueous suspension described above. Plants, used as controls, are treated with an aqueous solution not containing the active material.

After 24 hours, the plants are contaminated by spraying them with an aqueous suspension of *Pyrenophora teres* spores (12,000 spores per ml). The spores are collected from a 12-day-old culture. The contaminated barley plants are incubated for 24 hours at about 20° C. and at 100% relative humidity, and then for 12 days at 80% relative humidity.

Grading is carried out 12 days after the contamination, in comparison with the control plants.

Under these conditions, good (at least 50%) or total protection is observed at a dose of 330 ppm with the following compounds: A1, B1, C1, C2 and D1.

Under these conditions, good (at least 50%) or total protection is observed at a dose of 125 ppm with the following compound: C4.

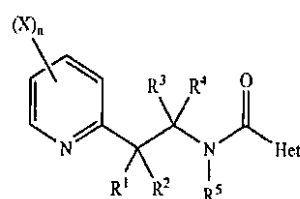
The N-{1-methylcarbamoyl-2-[3-chloro-5-(trifluoromethyl)-2-pyridinyl]-ethyl}-4-phenylbenzamide disclosed by Patent Application WO 01/11965 (see compound 316 in Table D) showed poor effectiveness on *Alternaria brassicae*, and zero effectiveness on *Botrytis cinerea* at 330 ppm; the N-{1-ethylcarbamoyl-2-[3-chloro-5-(trifluoromethyl)-2-pyridinyl]ethyl}-3-nitrobenzamide also disclosed by Patent Application WO 01/11965 (see compound 307 in Table D) showed poor effectiveness on *Alternaria brassicae* and zero effectiveness on *Botrytis cinerea* at 330 ppm; the N-{1-ethylcarbamoyl-2-[3-chloro-5-(trifluoromethyl)-2-pyridinyl]ethyl}-benzamide and the N-{1-methylcarbamoyl-2-[3-chloro-5-(trifluoromethyl)-2-pyridinyl]ethyl}-benzamide also disclosed by Patent Application WO 01/11965 (see compounds 304 and 314 in Table D) showed zero effectiveness on *Botrytis cinerea* at 330 ppm; and the N-{1-ethylcarbamoyl-2-[3-chloro-5-(trifluoromethyl)-2-pyridinyl]ethyl}-4-chlorobenzamide, the N-{1-ethylcarbamoyl-2-[3-chloro-5-(trifluoromethyl)-2-pyridinyl]ethyl}-2-bromobenzamide and the N-{1-methylcarbamoyl-2-[3-chloro-5-(trifluoromethyl)-2-pyridinyl]ethyl}-4-methoxybenzamide also disclosed by Patent Application WO 01/11965 (see compounds 306, 310 and 315 in Table D) showed zero effectiveness on *Botrytis cinerea* at 330 ppm.

The N-{[3-chloro-5-(trifluoromethyl)-2-pyridinyl]methyl}-5-thienylacetamide disclosed by Patent Application WO 01/11965 (see compound 101 in table B) showed poor efficacy against *Alternaria brassicae* and no efficacy against *Botrytis cinerea* and *Peronospora parasitica* at 330 ppm.

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The invention claimed is:

1. A compound of formula (I):



in which:

n is 1, 2, 3 or 4;

each X is independently selected from the group consisting of a halogen atom, a nitro group, a cyano group, a hydroxy group, an amino group, a sulfonyl group, a pentafluoro- λ^6 -sulfonyl group, a formyl group, a formyloxy group, a formylamino group, a carboxy group, a carbamoyl group, a N-hydroxycarbamoyl group, a carbamate group, a (hydroxyimino)-C₁-C₆-alkyl group, a C₁-C₈-alkyl, a C₂-C₈-alkenyl, a C₁-C₈-alkynyl, a C₁-C₈-alkylamino, a di-C₁-C₈-alkylamino, a C₁-C₈-alkoxy, a C₁-C₈-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₈-alkylsulfonyl, a C₁-C₈-halogenoalkylsulfonyl having 1 to 5 halogen atoms, a C₂-C₈-alkenyloxy, a C₂-C₈-halogenoalkenyloxy having 1 to 5 halogen atoms, a C₃-C₈-alkynyloxy, a C₃-C₈-halogenoalkynyloxy having 1 to 5 halogen atoms, a C₃-C₈-cycloalkyl, a C₃-C₈-halogenocycloalkyl having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbonyl, a C₁-C₈-halogenoalkylcarbonyl having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbamoyl, a di-C₁-C₈-alkylcarbamoyl, a N-C₁-C₈-alkyloxycarbamoyl, a C₁-C₈-alkoxycarbamoyl, a N-C₁-C₈-alkyl-C₁-C₈-alkoxycarbamoyl, a C₁-C₈-alkoxycarbonyl, a C₁-C₈-halogenoalkoxycarbonyl having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbonyloxy, a C₁-C₈-halogenoalkylcarbonyloxy having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbonylamino, a C₁-C₈-halogenoalkylcarbonylamino having 1 to 5 halogen atoms, a C₁-C₈-alkylaminocarbonyloxy, a di-C₁-C₈-alkylaminocarbonyloxy, a C₁-C₈-alkyloxycarbonyloxy, a C₁-C₈-alkylsulphenyl, a C₁-C₈-halogenoalkylsulphenyl having 1 to 5 halogen atoms, a C₁-C₈-alkylsulphinyl, a C₁-C₈-halogenoalkylsulphinyl having 1 to 5 halogen atoms, a C₁-C₈-alkylsulphonyl, a C₁-C₈-halogenoalkylsulphonyl having 1 to 5 halogen atoms, a C₁-C₆-alkoxyimino, a (C₁-C₆-alkoxyimino)-C₁-C₆-alkyl, a (C₁-C₆-alkynyloxyimino)-C₁-C₆-alkyl, a (benzyloxyimino)-C₁-C₆-alkyl, a benzyloxy, a benzylsulfonyl, a benzylamino, a phenoxy, a phenylsulfonyl and a phenylamino;

R¹, R², R³ and R⁴ are independently selected from the group consisting of a hydrogen atom, a halogen atom, a cyano group, a hydroxy group, an amino group, a sulfonyl group, a formyl group, a formyloxy group, a formylamino group, a carboxy group, a carbamoyl group, a N-hydroxycarbamoyl group, a carbamate group, a (hydroxyimino)-C₁-C₆-alkyl group, a C₁-C₈-alkyl, a C₁-C₈-halogenoalkyl having 1 to 5 halogen atoms, a C₂-C₈-alkenyl, a C₂-C₈-alkynyl, a C₁-C₈-alkylamino, a di-C₁-C₈-alkylamino, a C₁-C₈-alkoxy, a C₁-C₈-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₈-alkylsulfonyl, a C₁-C₈-halogenoalkylsulfonyl having 1 to 5 halogen atoms, a C₂-C₈-alkenyloxy, a C₂-C₈-halo-

43

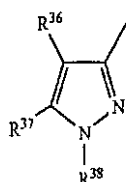
genoalkenyloxy having 1 to 5 halogen atoms, a C₃-C₈-alkynyloxy, a C₃-C₈-halogenoalkynyloxy having 1 to 5 halogen atoms, a C₃-C₈-cycloalkyl, a C₃-C₈-halogeno-cycloalkyl having 1 to 5 halogen atoms, a C₁-C₈-alkyl-carbonyl, a C₁-C₈-halogenoalkylcarbonyl having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbamoyl, a di-C₁-C₈-alkylcarbamoyl, a N—C₁-C₈-alkyloxycarbamoyl, a C₁-C₈-alkoxycarbamoyl, a N—C₁-C₈-alkyl-C₁-C₈-alkoxycarbamoyl, a C₁-C₈-alkoxycarbonyl, a C₁-C₈-halogenoalkoxycarbonyl having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbonyloxy, a C₁-C₈-halogenoalkylcarbonyloxy having 1 to 5 halogen atoms, a C₁-C₈-alkylcarbonylamino, a C₁-C₈-halogenoalkylcarbonylamino having 1 to 5 halogen atoms, a C₁-C₈-alkylaminocarbonyloxy, a di-C₁-C₈-alkylaminocarbonyloxy, a C₁-C₈-alkyloxycarbonyloxy, a C₁-C₈-alkylsulphenyl, a C₁-C₈-halogenoalkylsulphenyl having 1 to 5 halogen atoms, a C₁-C₈-alkylsulphinyl, a C₁-C₈-halogenoalkylsulphinyl having 1 to 5 halogen atoms, a C₁-C₈-alkylsulphonyl, a C₁-C₈-halogenoalkylsulphonyl having 1 to 5 halogen atoms, a benzyloxy, a benzylsulfonyl, a benzylamino, a phenoxy, a phenylsulfonyl or a phenylamino, a phenyl group, and a phenyl sulphonyl group;

or R¹ and R² may form together a cyclopropyl, a cyclobutyl, a cyclopentyl or a cyclohexyl;

with the proviso that when three of the four substituents R¹, R², R³ and R⁴ are a hydrogen atom, then the fourth substituent is not a hydrogen atom;

R⁵ is selected from the group consisting of a hydrogen atom, a cyano group, a formyl group, a hydroxy group, a C₁-C₆-alkyl, a C₁-C₆-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₆-alkoxy, a C₁-C₆-halogenoalkoxy having 1 to 5 halogen atoms, a C₃-C₆-cycloalkyl, a C₃-C₆-halogenocycloalkyl having 1 to 5 halogen atoms, a C₂-C₆-alkenyl, a C₂-C₆-alkynyl, a C₁-C₆-alkoxy-C₁-C₆-alkyl, a C₁-C₆-cyanoalkyl, a C₁-C₆-aminoalkyl, a C₁-C₆-alkylamino-C₁-C₆-alkyl, a di-C₁-C₆-alkylamino-C₁-C₆-alkyl, a C₁-C₆-alkylcarbonyl, a C₁-C₆-halogenalkylcarbonyl having 1 to 5 halogen atoms, a C₁-C₆-alkyloxycarbonyl, a C₁-C₆-benzyloxycarbonyl, a C₁-C₆-alkoxy-C₁-C₆-alkylcarbonyl, a C₁-C₆-alkylsulfonyl and a C₁-C₆-halogenoalkylsulfonyl having 1 to 5 halogen atoms;

Het represents a substituted pyrazole ring selected from the group consisting of:



(A) 50

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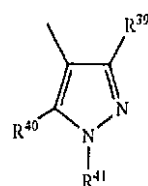
wherein:

R³⁶ is selected from the group consisting of a halogen atom, a cyano group, a nitro group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₃-C₆-cycloalkyl, a C₁-C₄-alkoxy, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₄-alkylthio, a C₁-C₄-halogenoalkylthio having 1 to 5 halogen atoms, an aminocarbonyl group, and an aminocarbonyl-C₁-C₄-alkyl;

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R³⁷ is selected from the group consisting of a hydrogen atom, a halogen atom, a cyano group, a nitro group, a C₁-C₄-alkyl, a C₁-C₄-alkoxy, and a C₁-C₄-alkylthio; and

R³⁸ is selected from the group consisting of a hydrogen atom, a phenyl, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a hydroxy-C₁-C₄-alkyl, a C₂-C₆-alkenyl, a C₃-C₆-cycloalkyl, a C₁-C₄-alkylthio-C₁-C₄-alkyl, a C₁-C₄-halogenoalkylthio-C₁-C₄-alkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy-C₁-C₄-alkyl, and a C₁-C₄-halogenoalkoxy-C₁-C₄-alkyl having 1 to 5 halogen atoms;



(B)

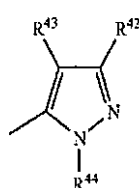
wherein:

R³⁹ is selected from the group consisting of a hydrogen atom, a halogen atom, a cyano group, a nitro group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₃-C₆-cycloalkyl, a C₁-C₄-alkoxy, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₄-alkylthio, a C₁-C₄-halogenoalkylthio having 1 to 5 halogen atoms, an aminocarbonyl, and an aminocarbonyl-C₁-C₄-alkyl;

R⁴⁰ is selected from the group consisting of a hydrogen atom, a halogen atom, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-alkoxy, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms, and a C₁-C₄-alkylthio; and

R⁴¹ is selected from the group consisting of a hydrogen atom, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a hydroxy-C₁-C₄-alkyl, a C₂-C₆-alkenyl, a C₃-C₆-cycloalkyl, a C₁-C₄-alkylthio-C₁-C₄-alkyl, a C₁-C₄-halogenoalkylthio-C₁-C₄-alkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy-C₁-C₄-alkyl, a C₁-C₄-halogenoalkoxy-C₁-C₄-alkyl having 1 to 5 halogen atoms, and a phenyl optionally substituted by a halogen atom, a C₁-C₄-alkyl, a C₁-C₄-alkoxyalkyl or a nitro group;

provided that the R³⁹ and R⁴⁰ are not both a hydrogen atom; and



(C)

wherein:

R⁴² is selected from the group consisting of a hydrogen atom, a halogen atom, a cyano group, a nitro group, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a C₃-C₆-cycloalkyl, a C₁-C₄-alkoxy, a C₁-C₄-halogenoalkoxy having 1 to 5 halogen atoms, a C₁-C₄-alkylthio, a C₁-C₄-halogenoalkylthio having 1 to 5 halogen atoms, an aminocarbonyl, and an aminocarbonyl-C₁-C₄-alkyl;

R⁴³ is selected from the group consisting of a hydrogen atom, a halogen atom, a cyano group, a C₁-C₄-alkyl, a C₁-C₄-alkoxy, a C₁-C₄-alkylthio, and a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms;

R⁴⁴ is selected from the group consisting of a hydrogen atom, a phenyl, a benzyl, a C₁-C₄-alkyl, a C₁-C₄-halogenoalkyl having 1 to 5 halogen atoms, a hydroxy-C₁-C₄-alkyl, a C₂-C₆-alkenyl, a C₃-C₆-cycloalkyl, a C₁-C₄-alkylthio-C₁-C₄-alkyl, a C₁-C₄-halogenoalkylthio-C₁-C₄-alkyl having 1 to 5 halogen atoms, a C₁-C₄-alkoxy-C₁-C₄-alkyl, and a C₁-C₄-halogenoalkoxy-C₁-C₄-alkyl having 1 to 5 halogen atoms;

provided that R⁴³ and R⁴⁴ are not both a hydrogen atom; as well as its salt and N-oxides.

2. The compound of claim 1 wherein n is 1, 2 or 3.

3. The compound of claim 1 wherein at least one of the X substituents is selected from the group consisting of a halogen atom, a C₁-C₆-alkyl, a C₁-C₆-alkoxyimino, a (C₁-C₆-alkoxyimino)-C₁-C₆-alkyl, and a C₁-C₆-alkoxy-C₁-C₆-alkylcarbo-

4. The compound of claim 1 wherein the 2-pyridyl is substituted in 3-, 5- and/or in 6-position.

5. The compound of claim 1 wherein R¹ and R² are independently selected from the group consisting of a hydrogen atom, a halogen atom, a cyano group, a hydroxy group, a C₁-C₆-alkyl, a C₁-C₆-halogenoalkyl having 1 to 5 halogen atoms, a C₂-C₆-alkenyl, a C₁-C₆-alkoxy, a C₁-C₆-alkylsulfa-

nyl, a C₁-C₆-alkylsulfenyl, a C₁-C₆-alkylsulfinyl, a C₁-C₆-

alkoxycarbonyl, a C₁-C₆-alkylcarbonylamino, a C₁-C₆-alkoxycarbonyloxy, a C₁-C₆-alkoxycarbonylamino and a phenyl group.

6. The compound of claim 5 wherein R¹ and R² are independently selected from the group consisting of a halogen atom, a C₁-C₆-alkyl, a C₁-C₆-halogenoalkyl having 1 to 5 halogen atoms and a C₁-C₆-alkylcarbonylamino.

7. The compound of claim 1 wherein R³ and R⁴ are independently selected from the group consisting of a hydrogen atom, a halogen atom, a cyano group, a C₁-C₆-alkyl, a C₁-C₆-halogenoalkyl having 1 to 5 halogen atoms, a C₁-C₆-alkylcarbonylamino and a phenyl group.

8. The compound of claim 7 wherein R³ and R⁴ are independently selected from the group consisting of a halogen atom, a C₁-C₆-alkyl, a C₁-C₆-halogenoalkyl having 1 to 5 halogen atoms and a phenyl group.

9. The compound of claim 1 wherein R⁵ is selected from the group consisting of a hydrogen atom and a C₃-C₇-cycloalkyl.

10. A fungicidal composition comprising an effective amount of a compound according to claim 1 and an agriculturally acceptable support.

11. A method for combating the phytopathogenic fungi of crops comprising applying an effective and non-phytotoxic amount of the composition of claim 10 to the plant seeds or to the plant leaves and/or to the fruits of the plants or to the soil in which the plants are growing or in which it is desired to grow them.

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