

Process for the preparation of 5-arylnicotinaldehydes

The invention relates to a process for the preparation of 5-arylnicotinaldehydes by reduction of the corresponding 5-arylnicotinic acids by catalytic hydrogenation in the presence of carboxylic anhydrides in which the catalyst used is a palladium/ligand complex, characterized in that the molar ratio between palladium and ligand is from 1:5 to 1:15 in the case of monodentate ligands and from 1:2.5 to 1:7.5 in the case of bidentate ligands.

5-Arylnicotinaldehydes are important intermediates and end products in industrial organic chemistry. Appropriately substituted derivatives are, in particular, valuable intermediates in the synthesis of highly value-added end products or are themselves such end products, in particular for crop protection, such as, for example, fungicides, insecticides, herbicides or pesticides, or for the preparation of substances having high pharmaceutical activity. Production of the corresponding 5-arylnicotinaldehydes on an industrial scale makes highly economical and environmentally friendly preparation necessary.

The 5-arylnicotinaldehydes are distinguished by an unexpected sensitivity to oxidation and disproportionation. Consequently, the methods described in the literature for the preparation of these compounds by chemoselective reduction of aromatic acids or their esters or by oxidation of aromatic alcohols cannot be applied to a satisfactory extent to the corresponding 5-arylnicotinic acid or the corresponding 5-arylnicotinyl alcohol.

Methods for the reduction of nicotinic acid are known. Thus, for example, the reduction of nicotinic acid by palladium-catalysed hydrogenation is described in Chemistry Letters 1998, 1143-1144. The process disclosed therein is carried out in the presence of pivalic anhydride (trimethylacetic anhydride), giving, in the case of unsubstituted nicotinic acid, the corresponding aldehyde in good yields.

However, if substituted nicotinic acids and in particular aromatic-substituted nicotinic acids are employed in this process, considerable losses of yield occur. In addition, the use of pivalic acid is associated with considerable

costs, and consequently this process is not suitable for industrial application.

5 The object on which the invention is based was to discover a process for the preparation of 5-aryl nicotinaldehydes which makes inexpensive and environmentally friendly access even to this class of substances possible without the above-mentioned disadvantages.

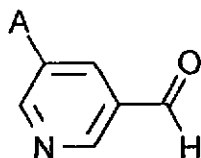
10 Surprisingly, it has now been found that 5-aryl nicotinaldehydes can be obtained by reduction of the corresponding 5-aryl nicotinic acids by catalytic hydrogenation in the presence of carboxylic anhydrides in which the catalyst used is a palladium/ligand complex if a molar ratio between palladium and ligand of from 1:5 to 1:15 is observed in the case of mono-

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The measure according to the invention also allows the use of other carboxylic anhydrides which are less expensive than pivalic anhydride and which, without an excess of ligand, give only a low yield of product.

20 The invention thus relates to a process for the preparation of 5-aryl nicotinaldehydes by reduction of the corresponding 5-aryl nicotinic acids by catalytic hydrogenation in the presence of carboxylic anhydrides in which the catalyst used is a palladium/ligand complex, characterized in that the molar ratio between palladium and ligand is from 1:5 to 1:15 in the case of
25 monodentate ligands and from 1:2.5 to 1:7.5 in the case of bidentate ligands.

30 The invention preferably relates, in particular, to a process for the preparation of 5-aryl nicotinaldehydes of the formula I



in which

A is a phenyl radical or naphthyl radical, each of which is unsubstituted or or monosubstituted or polysubstituted by R, F, Cl, Br, I, -CN, -NO₂, -CF₃, -OCF₃, -OCHF₂, -OCF₂CF₃, -OCHF₂CF₃ or -OH and in which, in addition, one or more CH groups may be replaced by N,

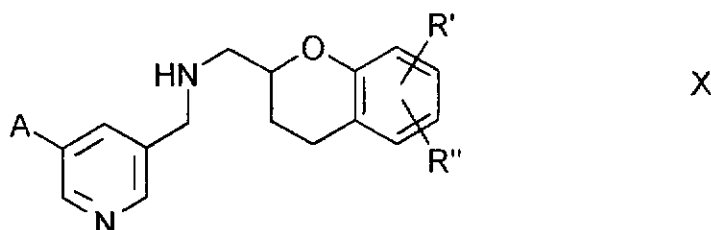
and

R is H, a straight-chain or branched alkyl radical having 1 to 15 carbon atoms which is monosubstituted by -CF₃ or at least monosubstituted by fluorine, where, in addition, one or more CH₂ groups in these radicals may each, independently of one another, be replaced by -S-, -O-, -O-CO-, -CO-O- or -CH=CH- in such a way that heteroatoms are not directly adjacent,

by reduction of the corresponding 5-arylnicotinic acids by catalytic hydrogenation in the presence of carboxylic anhydrides in which the catalyst used is a palladium/ligand complex, characterized in that the molar ratio between palladium and ligand is from 1:5 to 1:15 in the case of monodentate ligands and from 1:2.5 to 1:7.5 in the case of bidentate ligands.

The invention furthermore relates to the use of the 5-arylnicotinaldehydes for the preparation of N-alkylated amines by reaction of the 5-arylnicotinaldehydes with primary amines to give the corresponding imines, followed by reduction. The reaction of the aldehydes with the amines and also the reduction of the imines can be carried out by methods known per se, as described in the literature (for example in standard works, such as Houben-Weyl, Methoden der organischen Chemie [Methods of Organic Chemistry], Georg-Thieme-Verlag, Stuttgart, or Organikum [Organic Chemistry], 17th Edition, VEB Deutscher Verlag der Wissenschaften, Berlin, 1988), to be precise under reaction conditions which are known and are suitable for the said reactions.

The compounds of the formula I are, in particular, highly suitable for the preparation of N-alkylated amines of the formula X:



in which

R' and R'', independently of one another, are as defined for R or are F, Cl, Br, I, -CN, -NO₂, -CF₃, -OCF₃, -OCHF₂, -OCF₂CF₃, -OCHF₂CF₃ or -OH, and

- 5 A is as defined above, by reaction of the corresponding amines and reduction of the resultant imines by conventional reduction processes, in particular by catalytic hydrogenation or by using borohydrides, such as, for example, sodium borohydride.

- 10 Similar compounds are already known from EP 0707007.

Preference is given to the use of the aldehyde of the formula I in which in which A is 4-fluorophenyl for the preparation of (R)-2-[5-(4-fluorophenyl)-3-pyridylmethylaminomethyl]chroman by reaction with R-chromanamine and

15 reduction of the resultant imine.

In the preferred compounds of the formulae above and below, R is H, an alkyl or alkoxy group having 1 to 10 carbon atoms or an alkenyl group having 2 to 7 carbon atoms.

20

A is preferably a phenyl group, a phenyl group which is monosubstituted or disubstituted by F, a 4-fluorophenyl group, a pyrimidyl, a pyridyl, a pyrazyl or a pyridazyl group, in particular a 4-fluorophenyl group.

- 25 Preference is given to compounds which contain at least one of the preferred radicals.

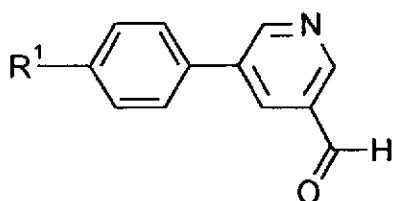
- If R is an alkyl group, in which, in addition, one CH₂ group (alkoxy or oxaalkyl) may be replaced by an O atom, it may be straight-chain or
- 30 branched. It preferably has 2, 3, 4, 5, 6, 7, 8, 9 or 12 carbon atoms and accordingly is preferably ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, nonyl, decyl, ethoxy, propoxy, butoxy, pentoxy, hexyloxy, heptyloxy, octyloxy, nonyloxy or decyloxy, furthermore also undecyl, dodecyl, undecyloxy,

dodecyloxy, 2-oxapropyl (= 2-methoxymethyl), 2-oxabutyl (= methoxyethyl) or 3-oxabutyl (= 2-ethoxymethyl), 2-, 3- or 4-oxapentyl, 2-, 3-, 4- or 5-oxahexyl, or 2-, 3-, 4-, 5- or 6-oxaheptyl. Particular preference is given to hexyl, pentyl, butyl, n-butyl, propyl, i-propyl, methyl and ethyl, in particular propyl and pentyl; particularly preferred alkoxy groups are hexyloxy, pentoxy, n-butoxy, propoxy, i-propoxy, methoxy and ethoxy, in particular ethoxy and n-butoxy. Compounds of the formulae above and below which have branched wing groups R may be of importance. Branched groups of this type generally contain not more than two chain branches. R is preferably a straight-chain group or a branched group having not more than one chain branch.

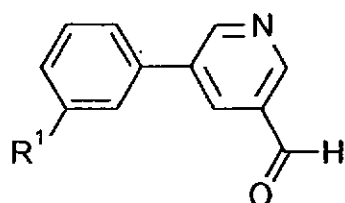
Preferred branched radicals are isopropyl, 2-butyl (= 1-methylpropyl), isobutyl (= 3-methylpropyl), tert-butyl, 2-methylbutyl, isopentyl (= 3-methylbutyl), 2-methylpentyl, 3-methylpentyl; 4-methylpentyl, 2-ethylhexyl, 5-methylhexyl, 2-propylpentyl, 6-methylheptyl, 7-methyloctyl, isopropoxy, 2-methylpropoxy, 2-methylbutoxy, 3-methylbutoxy, 2-methylpentoxy, 3-methylpentoxy, 2-ethylhexyloxy, 1-methylhexyloxy, 1-methylheptyloxy, 2-oxa-3-methylbutyl and 3-oxa-4-methylpentyl.

The radical R may also be an optically active organic radical containing one or more asymmetrical carbon atoms.

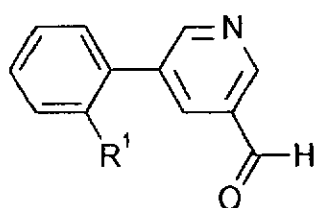
The process according to the invention is very particularly suitable for the preparation of the aldehydes of the formulae I1 to I10:



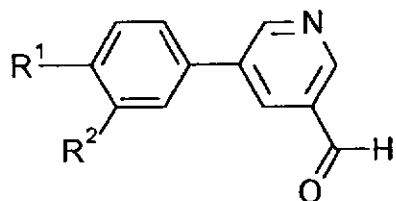
I1



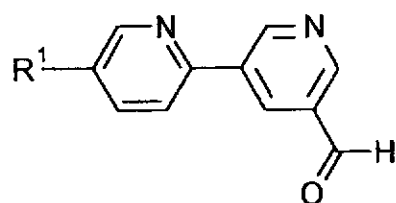
I2



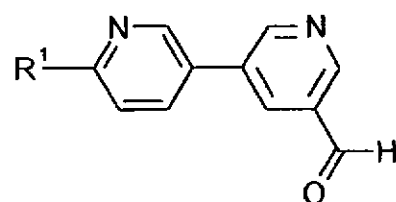
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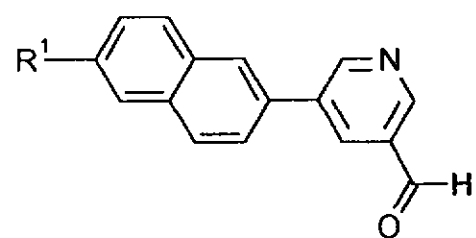
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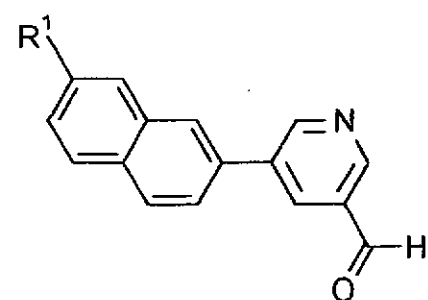
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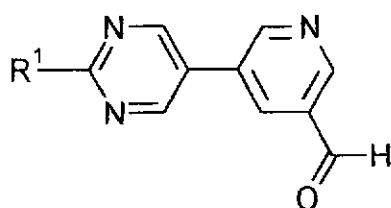
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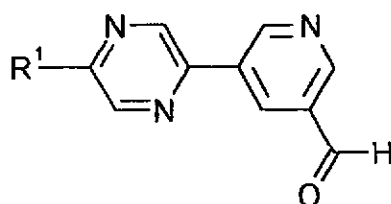
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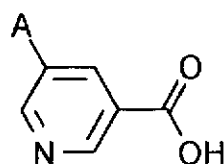
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In which R¹ and R², independently of one another, are as defined for R and are in particular F, -CF₃, -OCF₃, -OCHF₂ or H, and are very particularly preferably F.

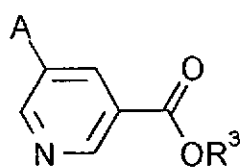
- 5 Very particularly preferred 5-arylnicotinaldehydes which can be prepared by the process according to the invention are the compounds of the formulae I1, I2, I9 and I10.

- 10 The 5-arylnicotinic acids used as starting materials for the process according to the invention are either known or are prepared by methods known per se, as described in the literature (for example in standard works, such as Houben-Weyl, Methoden der organischen Chemie [Methods of Organic Chemistry], Georg-Thieme-Verlag, Stuttgart), to be precise under reaction conditions which are known and suitable for said reactions.
- 15 However, use can also be made of variants which are known per se, but are not mentioned here in greater detail.

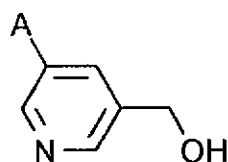
- The 5-arylnicotinic acids are preferably prepared by Suzuki coupling (N. Miyaura, A. Suzuki, Chem. Rev. 95, 2457 (1995)) by reacting 5-bromonictonic acid, an alkyl 5-bromonicotinate, 5-bromonicotinyl alcohol or the alkylcarboxylic acid ester thereof with the corresponding arylboronic acids under known reaction conditions to give a compound of the formula II, III, IV or V:
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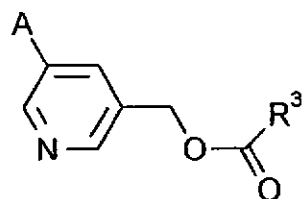
II



III



IV



V

in which A is as defined above, and R³ is methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl or tert-butyl,

- 5 subsequently hydrolysing the esters of the formulae III and V by conventional processes to give the free acid of the formula II or the alcohol of the formula IV, and oxidizing the alcohol of the formula IV by conventional methods to give the acid formula II.

- 10 The acids of the formula II obtained in this way can be used directly for the process according to the invention.

R³ is preferably methyl or ethyl, in particular methyl.

- 15 Compounds of the formulae I, II, III, IV and in which A is a 4-fluorophenyl radical are particularly preferred.

The invention likewise relates to the novel compounds of the formulae I, II, III and IV.

- 20 The reaction in the process according to the invention for the preparation of 5-aryl nicotinaldehydes is simple to carry out, with the 5-aryl nicotinic acids in question preferably being hydrogenated in organic solvents, at temperatures of from 10 to 180°C, preferably at from 20 to 150°C and very particularly preferably from at 40 to 100°C, and at a pressure of from 1 to
25 150 bar, preferably at from 1.5 to 120 bar and in particular at from 2 to

100 bar, with addition of a carboxylic anhydride in the presence of a palladium/ligand complex.

5 The solvents used for the process according to the invention are preferably ethers, such as, for example, diethyl ether or tetrahydrofuran, ketones, such as, for example, acetone, hydrocarbons, such as, for example, toluene, benzene, hexane or heptane, amides, such as, for example, dimethylformamide or N-methylpyrrolidone, or acid anhydrides, such as, for example, acetic anhydride.

10 Particular preference is given to ethers, in particular tetrahydrofuran.

Mixtures of the said solvents can likewise be used.

15 The amount of solvent is not crucial; from 10 g to 500 g of solvent can preferably be added per g of the compound of the formula II to be reacted.

The duration of the reaction depends on the selected reaction conditions. In general, the reaction duration is from 0.5 hour to 10 days, preferably from 1 to 24 hours.

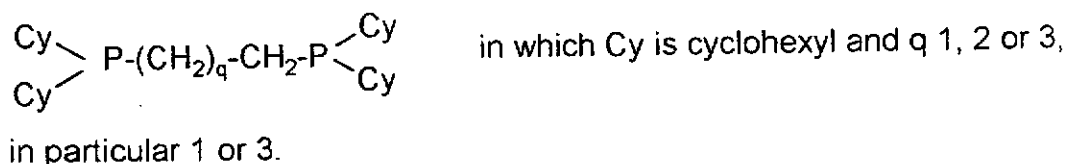
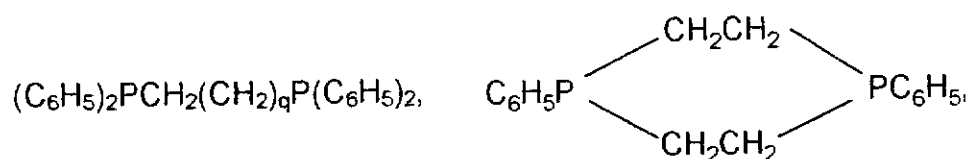
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In a preferred embodiment of the invention, the end of the reduction to the aldehyde is determined by suitable analytical methods, for example HPLC, and the reduction is interrupted.

25 The hydrogenation catalyst used is a palladium/ligand complex.

The ligands used are phosphines, preferably having three organic radicals, in particular alkyl and aryl radicals. In general, the ligands employed can be monodentate or bidentate phosphines. Suitable monodentate phosphines
30 are, for example, $(\text{CH}_3)_3\text{P}$, $(\text{C}_2\text{H}_5)_3\text{P}$, $(\text{C}_3\text{H}_7)_3\text{P}$, $(\text{C}_4\text{H}_9)_3\text{P}$, $(\text{C}_5\text{H}_{11})_3\text{P}$, $(\text{C}_6\text{H}_{13})_3\text{P}$, $(\text{C}_7\text{H}_{15})_3\text{P}$, $(\text{C}_8\text{H}_{17})_3\text{P}$, $(\text{C}_6\text{H}_{11})_3\text{P}$, $(\text{C}_6\text{H}_5)_3\text{P}$, $(\text{CH}_3\text{C}_6\text{H}_4)_3\text{P}$, $(\text{C}_6\text{H}_5\text{CH}_2)_3\text{P}$, $(\text{C}_4\text{H}_9)_2(\text{C}_6\text{H}_5)\text{P}$, $(\text{C}_4\text{H}_9)(\text{C}_6\text{H}_5)_2\text{P}$. The above empirical formulae include the branched and cyclic hydrocarbon radicals. Preference is given to n-propyl, n-butyl, sec-butyl, tert-butyl and cyclohexyl as hydro-
35 carbon radicals. Phosphines having identical radicals are preferred.

Examples of suitable bidentate ligands are the following:



The ligands employed are preferably the following: triphenylphosphine, diphenylphosphinoferrocene, tri-*o*-tolylphosphine, 1,2-bis(diphenyl-
5 phosphino)ethane, 2,4-bis(diphenylphosphino)pentane, bis(diphenylphosphino)methane and tris(tert-butyl)phosphine. It is likewise possible to use mixtures of the ligands.

10 Palladium compounds can be employed in oxidation state II, i.e. as a salt, or in oxidation state 0. It is preferably employed in oxidation state 0, preferably complexed by phosphines.

15 Palladium is generally employed as a palladium compound, from which the corresponding catalyst is prepared by addition of ligands. It is likewise possible to employ palladium as a complex having the correct stoichiometric composition to which, in accordance with the invention, an amount of ligand is added so that the ligand excess defined in accordance with the invention is observed.

20 Suitable palladium compounds employed in the presence of the excess of ligand are preferably the following:

Tetrakis(triphenylphosphine)palladium(0), dibenzylidenepalladium(0) complexes, palladium on carbon (preferably 5%), PdCl₂dppf, palladium acetate/(tri-*o*-tolylphosphine) complex, Pd(0)*dppe, Pd(0)*dppp,

25 Pd(0)*dppm, Pd(COD)Cl₂, PdCl₂, Pd(OAc)₂ and PdBr₂.

Particular preference is given to tetrakis(triphenylphosphine)palladium(0), dibenzylidenepalladium(0) complexes, palladium on carbon (preferably 5%), PdCl₂dppf, palladium acetate/(tri-*o*-tolylphosphine) complex,

30 Pd(0)*dppe, Pd(0)*dppp and Pd(0)*dppm.

The molar ratio between palladium and the acid to be hydrogenated is preferably from 1:500 to 1:200, in particular from 1:50 to 1:100.

5 The molar ratio between palladium and ligand is from 1:5 to 1:15, preferably from 1:5 to 1:10, in particular from 1:5 to 1:6, in the case of monodentate ligands and from 1:2.5 to 1:7.5, preferably from 1:2.5 to 1:5, in particular from 1:2.5 to 1:3, in the case of bidentate ligands.

10 Non-solvent anhydrides for the process are preferably pivalic anhydride, acetic anhydride, isobutyric anhydride, 5-norbornene-2,3-dicarboxylic anhydride, 1,2,3,6-tetrahydrophthalic anhydride and naphthalene-1,8-dicarboxylic anhydride.

15 The molar ratio between the carboxylic anhydride and the 5-aryl nicotinic acid is preferably from 10:1 to 8:1, preferably from 7:1 to 6:1 and in particular from 5:1 to 2:1.

20 The compounds of the formula I can be obtained, after removal of the solvent, by conventional work-up steps, such as, for example, addition of water to the reaction mixture and extraction. It may be advantageous, for further purification of the product, subsequently to carry out a distillation or crystallization.

25 The compounds of the formula I which can be prepared by the process according to the invention are important intermediates or end products in industrial organic chemistry. Appropriately substituted derivatives are, in particular, valuable intermediates in the synthesis of highly value-added end products or are such end products themselves, for crop protection, such as, for example, fungicides, insecticides, herbicides or pesticides, or
30 for the preparation of substances having high pharmaceutical activity.

Even without further embodiments, it is assumed that a person skilled in the art can utilize the above description in the broadest scope. The preferred embodiments should therefore merely be regarded as a
35 descriptive disclosure which is not limiting in any way.

The following example is intended to illustrate the invention without representing a limitation. Unless otherwise stated, percentages are per cent by weight. All temperatures are given in degrees Celsius.

5 Example 1:

13.2 g of 4-fluorobenzeneboronic acid and 10.8 g of sodium hydrogen-carbonate are initially introduced, and a mixture of 16.1 g of 5-bromo-3-hydroxymethylpyridine, 30 ml of water and 60 ml of toluene is added with stirring and under a protective gas. A further 14 ml of water, 28 ml of
10 toluene and 0.5 g of tetrakis(triphenylphosphine)palladium(0) are subsequently added. The reaction mixture is heated to reflux and held at this temperature for 2 hours. After cooling, the mixture is subjected to conventional work-up, giving 5-(4-fluorophenyl)nicotinyl alcohol. The alcohol is subsequently oxidized by conventional methods to give 5-(4-fluorophenyl)-
15 nicotinic acid.

Example 2:

1.38 kg of 5-bromonicotinic acid, 1.10 kg of 4-fluorobenzeneboronic acid and 0.40 kg of Pd/activated carbon (5%) are introduced into 8.5 litres of
20 water. The black suspension is rendered alkaline using 1.39 kg of 32% sodium hydroxide solution and warmed to 100°C. After 4 hours, the mixture is subjected to conventional work-up, giving 5-(4-fluorophenyl)nicotinic acid.

Melting range: 198.5°C to 203°C.

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Example 3:

15.3 g of methyl 5-(4-fluorophenyl)nicotinate (obtainable by Suzuki coupling of methyl 5-bromopyridine-3-carboxylate with 4-fluorobenzeneboronic acid analogously to Example 1 or 2) are saponified by conventional
30 methods to give 5-(4-fluorophenyl)nicotinic acid.

Example 4:

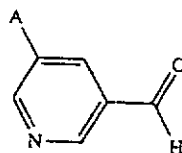
5-Bromo-3-acetoxymethylpyridine is reacted with 4-fluorobenzeneboronic acid as described in Example 1. The 5-(4-fluorophenyl)pyridin-3-ylmethyl acetate obtained is saponified by conventional methods and subsequently
35 oxidized to give 5-(4-fluorophenyl)nicotinic acid.

Example 5:

- 5-(4-Fluorophenyl)nicotinic acid is dissolved in tetrahydrofuran in an autoclave, three equivalents of the respective acid anhydride are added, and the mixture is rendered inert by repeatedly injecting a protective gas. A solution of $\text{Pd}(\text{OAc})_2$ and PPh_3 in tetrahydrofuran is added in a counter-current of protective gas, and the reaction solution is subsequently hydrogenated at 80°C under a hydrogen pressure of 8 bar with stirring. Conventional work-up gives the aldehyde in a yield of 98%.

Claims:

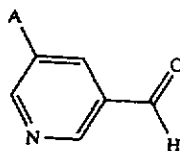
1. A process for the preparation of a 5-arylnicotinaldehyde of formula I



in which

A is a phenyl radical or naphthyl radical, each of which is unsubstituted, monosubstituted or polysubstituted by R, F, Cl, Br, I, -CN, -NO₂, -CF₃, -OCF₃, -OCHF₂, -OCF₂CF₃ or -OH and in which or more CH groups is optionally replaced by N, and R is H, a straight-chain or branched alkyl radical having 1 to 15 carbon atoms which is monosubstituted by -CF₃ or at least monosubstituted by fluorine, where, in addition, one more CH₂ groups is optionally independently replaced by -S-, -O-, -O-CO-, -CO-O- or -CH=CH- in such a way that heteroatoms are not directly adjacent, said process comprising reduction of a corresponding 5-arylnicotinic acid by catalytic hydrogenation in the presence of carboxylic anhydrides and a palladium/ligand complex, wherein molar ratio between palladium and ligand is from 1:5 to 1:15 in the case of monodentate ligands and from 1:2.5 to 1:7.5 in the case of bidentate ligands.

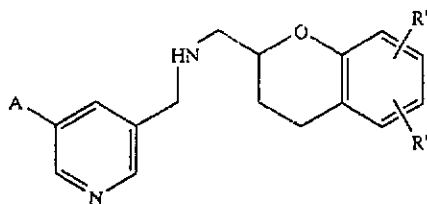
2. A compound of formula I



in which A is a 4-fluorophenyl group.

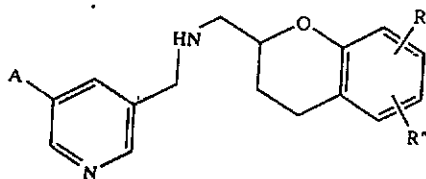
3. A process according to claim 1, further comprising reacting said 5-arylnicotinaldehyde with a primary amine to give a corresponding imine, and reducing to produce an N-alkylated amine.
4. A process according to claim 1, further comprising reacting a compound of formula I with a primary amine to give a corresponding imine, and reducing to produce an N-alkylated amine.

5. A process according to claim 3, wherein the N-alkylated amine is one of formula X



in which R' and R'', independently or one another, are H, straight-chain or branched alkyl radical having 1 to 15 carbon atoms which is monosubstituted by $-\text{CF}_3$ or at least monosubstituted by fluorine, where, in addition, one more CH_2 groups is optionally independently replaced by $-\text{S}-$, $-\text{O}-$, $-\text{O}-\text{CO}-$, $-\text{CO}-\text{O}-$ or $-\text{CH}=\text{CH}-$ in such a way that heteroatoms are not directly adjacent, or are F, Cl, Br, I, $-\text{CN}$, $-\text{NO}_2$, $-\text{CF}_3$, $-\text{OCF}_3$, $-\text{OCHF}_2$, $-\text{OCF}_2\text{CF}_3$, $-\text{OCHFCF}_3$ or $-\text{OH}$, and A is as defined above.

6. A process according to claim 4, wherein the N-alkylated amine is one of formula X



in which R' and R'', independently or one another, are as defined for R or are F, Cl, Br, I, $-\text{CN}$, $-\text{NO}_2$, $-\text{CF}_3$, $-\text{OCF}_3$, $-\text{OCHF}_2$, $-\text{OCF}_2\text{CF}_3$, $-\text{OCHFCF}_3$ or $-\text{OH}$, and A is as defined above.

7. A process according to claim 5, wherein the N-alkylated amine is (R)-2-[5-(4-fluorophenyl)-3-pyridylmethylaminomethyl]chroman.
8. A process according to claim 6, wherein the N-alkylated amine is (R)-2-[5-(4-fluorophenyl)-3-pyridylmethylaminomethyl]chroman.