Novel process for the preparation of imidazolyl compounds

The present invention relates to a method for the preparation of an imidazolyl compound of the general formula (I), wherein: \( R_1 \) and \( R_2 \) each separately are \((C_1-C_6)\)alkyl, \((C_1-C_6)\)alkoxyalkyl, optionally substituted aryl or heteroaryl; or wherein \( R_3 \) and \( R_4 \) together form a further homocyclic or heterocyclic system comprising one or more rings; \( R_5 \) and \( R_6 \) each are hydrogen or together form a carbon-carbon double bond, said carbon-carbon double bond optionally being part of an aromatic system; \( R_7 \) is hydrogen, \((C_1-C_6)\)alkyl, \((C_1-C_6)\)alkoxy, \((C_1-C_6)\)alkoxyalkyl or halogen; \( R_8 \) is hydrogen or \((C_1-C_6)\)alkyl; \( R_9 \) is hydrogen or \((C_1-C_6)\)alkyl; \( m \) is 1 or 2; and \( R_{10} \) is hydrogen or \((C_1-C_6)\)alkyl; as well as its acid addition salt; characterized in that a compound of the general formula (II) is reacted with a compound of the general formula (III), wherein: \( R \) is a hydrogen, a \((C_1-C_6)\)alkyl group optionally substituted with a hydroxy group or an optionally substituted aryl group, \( R' \), \( R'' \), \( R''' \) and \( R'''' \) each individually are a hydrogen or a \((C_1-C_6)\)alkyl group; followed by a reaction with a compound of the formula (IV), wherein \( R_1 \), \( R_2 \) and \( R_3 \) have the meanings defined above; and optionally followed by a reaction with a suitable acid. De method according to the present invention is especially useful for the preparation of ondansetron and cilansetron.
For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

Published:
- with international search report
PROCESS FOR THE PREPARATION OF IMIDAZOLYL COMPOUNDS

The present invention relates to a process for the preparation of imidazolyl compounds.

1,2,3,9-Tetrahydro-9-methyl-3-[2-methyl-1H-imidazol-1-yl]methyl]-4H-carbazol-4-one (ondansetron) is known from EP191562 and US4,695,578. In these patent publications a general class of compounds including ondansetron and homologous compounds, their preparation and their use as potent selective antagonists at "neuronal" 5-hydroxytryptamine receptors and their use in the treatment of migraine and psychotic disorders is described.

(10R)-5,6,9,10-Tetrahydro-10-[(2-methyl-1H-imidazol-1-yl)methyl]-4H-pyrido[3,2,1-jk]-carbazol-11(8H)-one (cilansetron) (also known as (R)-(−)-4,5,6,8,9,10-hexahydro-10-[(2-methyl-1H-imidazol-1-yl)methyl]-11H-pyrido[3,2,1-jk]-carbazol-11-one) is known from EP-B-0297651, from EP-B-0601345 and from EP-B-768309. In the first patent publication a general class of compounds, including cilansetron and homologous compounds, their preparation and their use as 5-HT antagonists is described. The second patent publication describes the use of a selection of these type of compounds for the treatment of certain diseases and the third the preparation of enantiomerically pure compounds and their hydrochloride monohydrate.

It is a common feature of the above compounds that they contain a substituted imidazolyl group attached to the α-position with regard to the keto group of the carbazole system with a methylene bridge. Several possibilities for the synthesis of these compounds are described in the mentioned patent publications. It is a common feature in these syntheses that the substituted imidazolyl group is introduced by means of a Mannich reaction, followed by a deamination to yield an intermediate exomethylene compound which is reacted with the substituted imidazolyl group (see scheme 1 for an example).

A drawback of this route is that the yield in this sequence of reaction steps is rather low. In US 4,695,578 the first step which is normally giving the lowest yield is not described and the second step (Example 7 of US 4,695,578) gives a yield of 68%. In EP-B-0297651 the first step (Example 1c of EP-B-0297651) has a yield of 53% and the second step (Example 1d of EP-B-0297651) a yield of 87%. During scale-up it
appeared that this route gives rise to the formation of a considerable amount of tar-like side-products.

Scheme 1.

It is the objective of the present invention to provide an alternative process to prepare imidazolyl-compounds which process should be economically operative and meet one or more of the following requirements: a) a relatively high yield, b) short reaction times compared with prior art processes, c) less side reactions, d) higher quality of the final product and e) using non-diluted reaction conditions and an environmentally acceptable solvent.

It has surprisingly been found that these type of imidazolyl compounds can easily be prepared using a substituted oxazolidine compound for the introduction of the methylene bridge.

Therefore the present invention relates to a method for the preparation of an imidazolyl compound of the general formula
wherein:

- $R_a$ and $R_b$ each separately are (C$_1$-C$_6$)alkyl, (C$_1$-C$_6$) alkoxyalkyl, optionally substituted aryl or heteroaryl;
- or wherein $R_a$ and $R_b$ together form a further homocyclic or heterocyclic system comprising one or more rings;
- $R_x$ and $R_y$ each are hydrogen or together form a carbon-carbon double bond, said carbon-carbon double bond optionally being part of an aromatic system;
- $R_c$ is hydrogen, (C$_1$-C$_6$)alkyl, (C$_1$-C$_6$)alkoxy, (C$_1$-C$_6$)alkoxyalkyl or halogen;
- $R_d$ is hydrogen or (C$_1$-C$_6$)alkyl;
- $R_e$ is hydrogen or (C$_1$-C$_6$)alkyl;
- $m$ is 1 or 2; and
- $R_f$ is hydrogen or (C$_1$-C$_6$)alkyl;

as well as its acid addition salt;

characterized in that a compound of the general formula

is reacted with a compound of the formula

$$\text{(III)}$$
wherein:

R is a hydrogen, a (C₁-C₆)alkyl group optionally substituted with a
dihydroxy group or an optionally substituted aryl group,

R', R", R*' and R** each individually are a hydrogen or a (C₁-C₆)alkyl group;

followed by a reaction with a compound of the formula

![Chemical Structure](image)

wherein R₁, R₄ and R₅ have the meanings defined above;

and optionally followed by a reaction with a suitable acid.

Alkyl groups of the present invention include straight-chained, branched and cyclic alkyl radicals containing up to 6 carbon atoms. Suitable alkyl groups may be saturated or unsaturated. Further, an alkyl group may also be substituted one or more times with substituents selected from the group consisting of aryl, halo, hydroxy, cyano or one- or di-alkyl substituted amino.

Aryl groups of the present invention include aryl radicals which may contain up to 6 hetero atoms. An aryl group may also be optionally substituted one or more times with a substituent selected from the group consisting of aryl, (C₁-C₆) alkyl, halo, hydroxy, cyano or one- or di-alkyl substituted amino, and it may be also fused with an aryl group or cycloalkyl rings. Suitable aryl groups include, e.g. phenyl, naphthyl, tolyl, imidazolyl, pyridyl, pyrryl, thienyl, pyrimidyl, thiazolyl and furyl groups.

With a homocyclic system is meant a system containing at least one saturated or unsaturated cyclic group containing only carbon atoms and hydrogen atoms.

With a heterocyclic system is meant a system containing at least one saturated or unsaturated cyclic group containing also one or more heteroatoms such as N, O or S.

Both the homocyclic and heterocyclic system may optionally be substituted with a substituent selected from the group consisting of alkyl, aryl, cyano, halogen, hydroxy or one- or di-alkyl substituted amino.

In a preferred embodiment of the invention R₈ is hydrogen or (C₁-C₆)alkyl, R₉ is hydrogen or (C₁-C₆)alkyl; R₈ is hydrogen or (C₁-C₆)alkyl; R₉ is hydrogen or (C₁-C₆)alkyl; and R₁ is hydrogen, methyl or ethyl.
The reaction according to the invention is especially useful for the preparation of compounds of the general formula

\[ R_1 \text{N} \text{CH} \text{(CH}_m \text{)}_n \text{O} \]

wherein:

- \( m \) is 1 or 2;
- \( R_1 \) is hydrogen, methyl or ethyl; and
- \( R_5 \) is a \((C_1-C_4)\) alkyl;
- \( R_6 \) is a hydrogen or a \((C_1-C_4)\) alkyl, or
- \( R_5 \) and \( R_6 \) together with the intermediate atoms form a 5, 6, or 7 membered ring, optionally substituted with one or two substituents selected from the group consisting of halogen, hydroxy, \((C_1-C_4)\) alkyl, \((C_1-C_4)\) alkoxyalkyl and \((C_1-C_4)\) alkoxy.

In this case the starting compound is a compound of the general formula

\[ R_5 \text{N} \text{CH} \text{(CH}_m \text{)}_n \text{O} \]

This compound is further referred to as a carbazole compound.

Preferred compounds of the general formula (la) are the compound wherein \( m = 1 \) and \( R_5 \) and \( R_6 \) together with the intermediate atoms form a 6-membered ring and the compound wherein \( m = 1 \), \( R_5 \) is methyl and \( R_6 \) is hydrogen. For the first compound the yield for the process starting with 5,6,9,10-tetrahydro-4\(H\)-pyrido[3,2,1-\(jk\)]carbazol-11(8\(H\))-one and 3-oxazolidineethanol is 77% (see Example 2) compared with the overall yield of 46% in the process according to EP0297651 (Examples ic and id). Higher yields may be obtained at a production scale.
In the substituted oxazolidine preferably one of the R' and R" and one of the R" and R''' is hydrogen, as an oxazolidine disubstituted on the same carbon atom, such as 4,4-dimethyloxazolidine gives a lower yield in the reaction. Preferred oxazolidines are 3-oxazolidineethanol and 3-ethyl-oxazolidine. The most preferred oxazolidine is 3-oxazolidineethanol.

The reaction is carried out in acidic medium, and the grade of acidity depends on the activation of the system that has to react. In the case of carbazolone systems the medium should be highly acidic. Examples of suitable acids in the last case are methanesulfonic acid, trifluoromethanesulfonic acid, p-toluenesulfonic acid and HCl gas in alcoholic medium.

In order to get a high yield, the reaction solution should contain only a low amount of water. The amount of water should preferably be below 0.6% (V/V), more preferably below 0.3% V/V and most preferably below 0.1% V/V.

The optimal reaction temperature is dependent on the starting material and the solvent and differs for the two reaction steps. The first step of the reaction can be performed between 40°C and 110°C. For the carbazolone systems the preferred reaction temperature in the first step is between 50°C and 90°C and the most preferred temperature is approximately 70°C. The second step can generally be performed between 100°C and 140°C. For the carbazolone systems the preferred reaction temperature in the second step is between 110°C and 130°C and the most preferred temperature is approximately 120°C.

The reaction can be performed in different solvents such as dipolar aprotic solvents like DMF or in alcohols. Preferred solvents are C4-C7 alcohols and the choice may depend on the desired reaction temperature. Examples of suitable alcohols are 1-butanol, 1-hexanol and isoamyl alcohol. A preferred alcohol is 1-butanol. Also suitable are mixtures of aromatic hydrocarbons and alcohols, such as mixtures of toluene and an alcohol and monochlorobenzene and an alcohol. A preferred mixture is a mixture of monochlorobenzene and methanol. When solvent mixtures are used the lower boiling solvent can be distilled off before the second step in order to reach higher reflux temperatures of the solvent system in the second step.
The ratio of the solvent volume to the amount of reactants in the mixture can be varied over a relatively broad range and depends on the solubility of the reactants. In general the ratio of the amount of solvent to the amount of reactants can typically be about 1:1 to 15:1, where the ratio is expressed as the volume of solvent relative to the weight of the reactants in the solvent (in ml/g). Preferably the ratio is about 1:1 to about 10:1. In the case of the carbazolone systems the preferred ratio of the volume of solvent to the weight of reactants is about 4:1.

The products obtained can be crystallized from different solvents. Examples of solvents for the crystallisation of free bases are aromatic hydrocarbons such as toluene. The hydrochloric acid salts can e.g. be crystallized from alcoholic solvents, such as isopropanol or 1-butanol.

The following examples are only intended to further illustrate the invention, in more detail, and therefore these examples are not deemed to restrict the scope of the invention in any way.

Example 1: Materials and Methods

5,6,9,10-Tetrahydro-4H-pyrido[3,2,1-jk]carbazol-11(8H)-one was made according to EP0375045. 3,4-Dihydrop-1(2H)-napthalenone was obtained from a commercial source. 1,2,3,9-Tetrahydro-9-methyl-4H-carbazol-4-one was made according to US 3,892,766 from Warner-Lambert Company and Elz, S. and Heil, W., Bioorganic & Medicinal Chemistry Letters 1995, 5, 667-672. Methanesulfonic acid was obtained from a commercial source.

NMR spectra were measured on a Varian VXR 200 and MS spectra on a Finnigan TSQ 7000. HPLC analyses were performed on a HP1050 system with a Separations 757 detector (250 nm) and a Separations Marathon XT column oven at 35 °C. The column used was a Zorbax XDB C8 column 15x0.3 cm. The eluens was prepared as follows: mix 2 l water, 2 ml triethylamine and 5 ml 25% ammonia, buffer it at pH=4 with formic acid and add 0.5 l acetonitril. The flow was 1 ml/min.

Example 1a: Preparation of oxazolidines.

3-Oxazolidineethanol was made as follows:

Equimolar amounts of diethanolamine and paraformaldehyde in 1-butanol were heated to 70°C. After 1 hour reaction time the water formed was removed by azeotropic distillation with 1-butanol.
3-Ethyl-oxazolidine was made according to Heany, H. et al., Tetrahedron 1997, 53, 14381-96.

4,4-Dimethyl-oxazolidine is commercially available and was purchased as a 75% w/w solution in water. The 4,4-dimethyl-oxazolidine was extracted from the water layer by washing with dichloromethane/ saturated NaCl-solution. The dichloromethane layer was dried on anhydrous sodium sulfate and subsequently evaporated.

**Example 2. Reaction of 5,6,9,10-tetrahydro-4H-pyrido[3,2,1-jk]carbazol-11(8H)-one with 3-oxazolidineethanol.**

5,6,9,10-tetrahydro-4H-pyrido[3,2,1-jk]carbazol-11(8H)-one (25.00 g = 111.0 mmole) and methanesulfoinic acid (17.06 g = 177.5 mmole) in 1-butanol (100 ml) were heated to 70°C. In 3 minutes a solution was added of 3-oxazolidineethanol (19.49 g = 166.4 mmole) in 1-butanol (39 ml).

After 50 minutes at 80°C 2-methylimidazole (45.55 g = 554.8 mmole) and 1-butanol (10 ml) were added. After 1.5 hours at 120°C the reaction mixture was partly evaporated till 30 ml of 1-butanol was left over.

At 70°C, 75 ml of toluene and 50 ml of water were added to the residue. The layers were separated. The water layer was extracted with 75 ml of toluene. The combined toluene layers were washed three times with 100 ml of water.

The organic layer was evaporated to dryness and subsequently 125 ml of 1-butanol was added. To the resulting solution 12.5 ml of 36% m/m hydrochloric acid was added. After stirring for 2 hours at room temperature the formed solid was filtered off and washed with 1-butanol and MTBE. Yield after drying: 30.40 g (77.0%) 5,6,9,10-tetrahydro-10-[(2-methyl-1H-imidazol-1-yl)methyl]-4H-pyrido[3,2,1-jk]carbazol-11(8H)-one hydrochloride (77.0%). HPLC: ≥ 95% . 1H NMR [200 MHz, DMSO-d_{6}:CDCl_{3} 4:1] δ 1.97(1H,m), 2.18 (3H,m), 2.68(3H,s), 2.95(2H,t), 3.00(1H,dd), 3.12(2H,m), 4.13(2H,m), 4.29(1H,dd), 4.66(1H,dd), 6.97(1H,d), 7.09(1H,t), 7.55(1H,d), 7.68(1H,d) and 7.71(1H,d). MS [ESI] MH^+ = 320.
Example 3. Reaction of 5,6,9,10-tetrahydro-4H-pyrido[3,2,1-jk]carbazol-11(8H)-one with 4,4-dimethyl-oxazolidine.

5,6,9,10-tetrahydro-4H-pyrido[3,2,1-jk]carbazol-11(8H)-one (20.00 g = 88.8 mmole) and methanesulfonic acid (13.65 g = 142.0 mmole) in 1-butanol (60 ml) were heated to 70°C. In 2 minutes 4,4-dimethyl-oxazolidine (13.47 g = 133.2 mmole) in 1-butanol (10 ml) was added.

After 50 minutes at 80°C 2-methylimidazole (36.45 g = 444.0 mmole) and butanol (10 ml) were added. After 2 hours at 120°C the reaction mixture was partly evaporated till 20 ml of 1-butanol was left over.

At 70°C, 60 ml of toluene and 40 ml of water were added to the residue. The layers were separated. The water layer was extracted with 60 ml of toluene. The combined toluene layers were washed three times with 80 ml of water.

The organic layer was evaporated to dryness and subsequently 100 ml of 1-butanol was added. To the resulting solution 10.0 ml of 36% m/m hydrochloric acid was added. After stirring for 2 hours at room temperature the formed solid was filtered off and washed with 1-butanol and MTBE. Yield after drying: 12.38 g 5,6,9,10-tetrahydro-10-[(2-methyl-1H-imidazol-1-yl)methyl]-4H-pyrido[3,2,1-jk]carbazol-11(8H)-one hydrochloride (39.2%). HPLC: ≥ 95%. 1H NMR and MS: see Example 2.

The mother liquor contained 3.45 g (10.9%) of product.

Example 4. Reaction of 5,6,9,10-tetrahydro-4H-pyrido[3,2,1-jk]carbazol-11(8H)-one with 3-ethyl-oxazolidine.

5,6,9,10-tetrahydro-4H-pyrido[3,2,1-jk]carbazol-11(8H)-one (20.00 g = 88.8 mmole) and methanesulfonic acid (13.65 g = 142.0 mmole) in 1-butanol (60 ml) were heated to 70°C. In 2 minutes 3-ethyl-oxazolidine (13.46 g = 133.2 mmole) in 1-butanol (10 ml) was added.

After 50 minutes at 80°C 2-methylimidazole (36.45 g = 444.0 mmole) and 1-butanol (10 ml) were added. After 2 hours at 120°C the reaction mixture was partly evaporated till 20 ml of 1-butanol was left over.

At 70°C, 60 ml of toluene and 40 ml of water were added to the residue. The layers were separated. The water layer was extracted with 60 ml of toluene. The combined toluene layers were washed three times with 80 ml of water.
The organic layer was evaporated to dryness and subsequently 100 ml of 1-butanol was added. To the resulting solution 10.0 ml of 36% m/m hydrochloric acid was added. After stirring for 2 hours at room temperature the formed solid was filtered off and washed with 1-butanol and MTBE. Yield after drying: 22.10 g (70.0 %) 5,6,9,10-tetrahydro-10-[(2-methyl-1H-imidazol-1-yl)methyl]-4H-pyrido[3,2,1-jk]carbazol-11(8H)-one hydrochloride (70.0%). HPLC: ≥ 95%. 1H NMR and MS: see Example 2.

Example 5. Reaction of 3,4-dihydro-1(2H)-naphthalenone with 3-oxazolidineethanol.

3,4-dihydro-1(2H)-naphthalenone (12.98 g = 88.8 mmole) and methanesulfonic acid (13.65 g = 142.0 mmole) in 1-butanol (60 ml) were heated to 50°C. In 2 minutes a solution was added of 3-oxazolidineethanol (15.59 g = 133.1 mmole) in 1-butanol (14 ml).

After 50 minutes at 80°C 2-methylimidazole (36.45 g = 444.0 mmole) and 1-butanol (10 ml) were added. After 2 hours at 120°C the reaction mixture was partly evaporated till 20 ml of 1-butanol was left over.

At 70°C, 60 ml of toluene and 40 ml of water were added to the residue. The layers were separated. The water layer was extracted with 60 ml of toluene. The combined toluene layers were washed three times with 80 ml of water.

The organic layer was evaporated to dryness and subsequently 100 ml of 1-butanol was added. To the resulting solution 10.0 ml of 36% m/m hydrochloric acid was added. The resulting solution was evaporated till an end volume of 60 ml. After stirring for 2 hours at room temperature the formed solid was filtered off and washed with 1-butanol and MTBE. Yield after drying: 15.28 g (62.2%) 3,4-dihydro-2-[(2-methyl-1H-imidazol-1-yl)methyl]-1(2H)-naphthalenone hydrochloride. HPLC: ≥ 95%. 1H NMR[200 MHz, DMSO-d6;CDCl3 4:1] δ 2.00 (2H,m), 2.73 (3H,s), 3.20 (3H,m), 4.27 (1H,dd), 4.68 (1H,dd), 7.35 (2H,t), 7.55 (2H,m), 7.70 (1H,d) and 7.90 (1H,d). MS [ESI] MH* = 241. The mother liquor contained 3.28 g (13.3%) of product.

Example 6. Reaction of 3,4-dihydro-1(2H)-naphthalenone with 4,4-dimethyl-oxazolidine.

3,4-dihydro-1(2H)-naphthalenone (12.98 g = 88.8 mmole) and methanesulfonic acid (13.65 g = 142.0 mmole) in 1-butanol (60 ml) were heated to 70°C. In 2 minutes 4,4-dimethyl-oxazolidine (13.46 g = 133.1 mmole) in 1-butanol (10 ml) was added. After
50 minutes at 80°C 2-methylimidazole (36.45 g = 444.0 mmole) and 1-butanol (10 ml) were added. After 2 hours at 120°C the reaction mixture was partly evaporated till 20 ml of 1-butanol was left over. At 70°C, 60 ml of toluene and 40 ml of water were added to the residue. The layers were separated. The water layer was extracted with 60 ml of toluene. The combined toluene layers were washed three times with 80 ml of water.

The organic layer was evaporated to dryness and subsequently 100 ml of 1-butanol was added. To the resulting solution 10.0 ml of 36% m/m hydrochloric acid was added. The resulting solution was evaporated till an end volume of 50 ml. After stirring for 2 hours at 0°C the formed solid was filtered off and washed with 1-butanol and MTBE. Yield after drying: 14.13 g (57.5%) 3,4-dihydro-2-[(2-methyl-1H-imidazol-1-yl)methyl]-1(2H)-naphthalenone hydrochloride. HPLC: ≥ 95%. ¹H NMR and MS: see Example 5. The mother liquor contained 2.33 g (9.5%) of product.

Example 7. Reaction of 3,4-dihydro-1(2H)-naphthalenone with 3-ethyl-oxazolidine.

3,4-dihydro-1(2H)-naphthalenone (12.98 g = 88.8 mmole) and methanesulfonic acid (13.65 g = 142.0 mmole) in 1-butanol (60 ml) were heated to 50°C. In 2 minutes 3-ethyl-oxazolidine (13.46 g = 133.1 mmole) in 1-butanol (10 ml) was added. After 50 minutes at 80°C 2-methylimidazole (36.45 g = 444.0 mmole) and 1-butanol (10 ml) were added. After 2 hours at 120°C the reaction mixture was partly evaporated till 20 ml of 1-butanol was left over.

At 70°C, 60 ml of toluene and 40 ml of water were added to the residue. The layers were separated. The water layer was extracted with 60 ml of toluene. The combined toluene layers were washed three times with 80 ml of water.

The organic layer was evaporated to dryness and subsequently 100 ml of 1-butanol was added. To the resulting solution 10.0 ml of 36% m/m hydrochloric acid was added. The resulting solution was evaporated till an end volume of 50 ml. After stirring for 2 hours at 0°C the formed solid was filtered off and washed with 1-butanol and MTBE. Yield after drying: 17.30 g 3,4-dihydro-2-[(2-methyl-1H-imidazol-1-yl)methyl]-1(2H)-naphthalenone hydrochloride (70.4%). HPLC: ≥ 95%. ¹H NMR and MS: see Example 5.
Example 8. Reaction of 1,2,3,9-tetrahydro-9-methyl-4H-carbazol-4-one with 3-oxazolidineethanol.

1,2,3,9-tetrahydro-9-methyl-4H-carbazol-4-one (13.26 g = 66.5 mmole) and methanesulfonic acid (10.23 g = 106.4 mmole) in 1-butanol (45 ml) were heated to 90°C. In 2 minutes 11.68 g (99.8 mmole) of 3-oxazolidineethanol in 1-butanol (11 ml) was added.

After 50 minutes at 80°C 2-methylimidazole (27.32 g = 332.5 mmole) and 1-butanol (8 ml) were added. After 2 hours at 120°C 180 ml of toluene and 120 ml of water were added at 80°C. The layers were separated. The water layer was extracted with 180 ml of toluene and 60 ml of 1-butanol. The combined organic layers were washed twice with 240 ml of water. The organic layer was evaporated to dryness. 150 ml of 1-butanol and 10 ml of 36% m/m hydrochloric acid were added to the residue. At 0°C crystallization soon occurred. After 1 hour at 0°C the formed crystals were filtered off, washed with 1-butanol and MTBE and subsequently dried: 15.39 g (70.1%) of 1,2,3,9-tetrahydro-9-methyl-3-[(2-methyl-1H-imidazol-1-yl)methyl]-4H-carbazol-4-one hydrochloride was isolated. HPLC: ≥ 95%. 1H NMR [200 MHz, DMSO-d6;CDCl3 4:1] δ 2.00 (1H, m), 2.20 (1H, m), 3.69 (3H, s), 3.09 (3H, m), 3.75 (3H, s), 4.30 (1H, dd), 4.67 (1H, dd), 7.23 (2H, m), 7.53 (2H, m), 7.69 (1H, d), 8.01 (1H, d). MS [ESI] MH+ = 294.

The mother liquor contained 3.19 g (14.5%) of product.

Example 9. Reaction of 1,2,3,9-tetrahydro-9-methyl-4H-carbazol-4-one with 4,4-dimethyl-oxazolidine.

1,2,3,9-tetrahydro-9-methyl-4H-carbazol-4-one (13.26 g = 66.5 mmole) and methanesulfonic acid (10.23 g = 106.4 mmole) in 1-butanol (45 ml) were heated to 90°C. In 2 minutes 4,4-dimethyl-oxazolidine (10.09 g = 99.9 mmole) in 1-butanol (8 ml) was added.

After 50 minutes at 80°C 2-methylimidazole (27.32 g = 332.5 mmole) and 1-butanol (8 ml) were added. After 2 hours at 120°C 180 ml of toluene and 120 ml of water were added at 80°C. The layers were separated. The water layer was extracted with 180 ml of toluene and 60 ml of 1-butanol. The combined organic layers were washed twice with 240 ml of water. The organic layer was evaporated to dryness. 150 ml of 1-butanol and 10 ml of 36% m/m hydrochloric acid were added to the residue. At 0°C crystallization soon occurred. After 1 hour at 0°C the formed crystals were filtered off, washed with 1-butanol and MTBE and subsequently dried: 10.02 g (45.7%) of
1,2,3,9-tetrahydro-9-methyl-3-[(2-methyl-1H-imidazol-1-yl)methyl]-4H-carbazol-4-one hydrochloride. The mother liquor contained 2.70 g (12.3%) of product.
HPLC: ≥ 95%. NMR and MS: see Example 8.

Example 10. Reaction of 1,2,3,9-tetrahydro-9-methyl-4H-carbazol-4-one with 3-ethyl-oxazolidine.

1,2,3,9-tetrahydro-9-methyl-4H-carbazol-4-one (13.26 g = 66.5 mmole) and methane-sulfonic acid (10.23 g = 106.4 mmole) in 1-butanol (45 ml) were heated to 90°C. In 2 minutes 3-ethyl-oxazolidine (10.09 g = 99.9 mmole) in 1-butanol (8 ml) was added. After 50 minutes at 80°C 2-methylimidazole (27.32 g = 332.5 mmole) and 1-butanol (8 ml) were added. After 2 hours at 120°C 180 ml of toluene and 120 ml of water were added at 80°C. The layers were separated. The water layer was extracted with 180 ml of toluene and 60 ml of 1-butanol. The combined organic layers were washed twice with 240 ml of water. The organic layer was evaporated to dryness. 150 ml of 1-butanol and 10 ml of 36% m/m hydrochloric acid were added to the residue. At 0°C crystallization soon occurred. After 1 hour at 0°C the formed crystals were filtered off, washed with 1-butanol and MTBE and subsequently dried: 15.67 g (71.4%) of 1,2,3,9-tetrahydro-9-methyl-3-[(2-methyl-1H-imidazol-1-yl)methyl]-4H-carbazol-4-one hydrochloride was isolated. HPLC: ≥ 95%. NMR and MS: see Example 8. The mother liquor contained 2.06 g (9.4%) of product.
Claims

1. Method for the preparation of an imidazolyl compound of the general formula

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\begin{align*}
\text{II}(l) & \\
\text{wherein:} & \\
R_a & \text{and } R_b \text{ each separately are } (C_{1-6})\text{alkyl, } (C_{1-6})\text{alkoxyalkyl, optionally} \\
& \text{substituted aryl or heteroaryl; } \\
& \text{or wherein } R_a \text{ and } R_b \text{ together form a further homocyclic or heterocyclic system} \\
& \text{comprising one or more rings; } \\
R_a & \text{and } R_b \text{ each are hydrogen or together form a carbon-carbon double bond,} \\
& \text{said carbon-carbon double bond optionally being part of an aromatic system; } \\
R_c & \text{is hydrogen, } (C_{1-6})\text{alkyl, } (C_{1-6})\text{alkoxy, } (C_{1-6})\text{alkoxyalkyl or halogen; } \\
R_d & \text{is hydrogen or } (C_{1-6})\text{alkyl; } \\
R_l & \text{is hydrogen or } (C_{1-6})\text{alkyl; } \\
m & \text{is } 1 \text{ or } 2; \text{ and } \\
& \text{as well as its acid addition salt; } \\
& \text{characterized in that a compound of the general formula} \\
\end{align*}
\]

\[
\begin{align*}
\text{(II)} & \\
\text{wherein } R_a, R_d, R_b \text{ and } R_b \text{ have the meanings defined above; }
\end{align*}
\]
is reacted with a compound of the formula

![Chemical structure image]

(III)

wherein:

R is a hydrogen, a (C<sub>1</sub>-C<sub>4</sub>) alkyl group optionally substituted with a hydroxy group or an optionally substituted aryl group,

R', R", R" and R"" each individually are a hydrogen or a (C<sub>1</sub>-C<sub>4</sub>) alkyl group; followed by a reaction with a compound of the formula

![Chemical structure image]

(IV)

wherein R<sub>i</sub>, R<sub>d</sub> and R<sub>e</sub> have the meanings defined above;

and optionally followed by a reaction with a suitable acid

2. Method according to claim 1, wherein

R<sub>a</sub>, R<sub>b</sub>, R<sub>c</sub>, R<sub>d</sub>, R<sub>e</sub>, R<sub>f</sub>, R<sub>g</sub>, R<sub>h</sub>, R<sub>i</sub>, R" and R"" have the same meanings as in claim 1;

R<sub>a</sub> is hydrogen or (C<sub>1</sub>-C<sub>4</sub>) alkyl;

R<sub>i</sub> is hydrogen or (C<sub>1</sub>-C<sub>4</sub>) alkyl;

m is 1 or 2; and

R<sub>i</sub> is hydrogen, methyl or ethyl.

3. Method according to claim 1-2 for the preparation of an imidazolyl compound of the general formula
wherein:

- $m$ is 1 or 2;
- $R_1$ is hydrogen, methyl or ethyl; and
- $R_3$ is a $(C_1-C_4)$ alkyl
- $R_6$ is a hydrogen or a $(C_1-C_4)$ alkyl, or
- $R_5$ and $R_6$ together with the intermediate atoms form a 5, 6, or 7 membered ring, optionally substituted with one or two substituents selected from the group consisting of halogen, hydroxyl, $(C_1-C_4)$ alkyl, $(C_1-C_4)$ alkoxyalkyl and $(C_1-C_4)$ alkoxy

as well as its pharmaceutically acceptable acid addition salt;

characterized in that a compound of the general formula

wherein $R_5$, $R_6$ and $m$ have the meanings defined above;

is reacted with a compound of the formula

wherein $R$, $R'$, $R''$, $R'''$ and $R''''$ have the same meanings as in claim 1;
followed by a reaction with a compound of the formula

![Chemical Structure](image)

$\text{R}_1$

$\text{HN}$

$\text{N}$

(IVa)

wherein $\text{R}_1$ has the same meaning as in claim 1.

4. Method according to claims 1-2, characterized in that $\text{R}_1$, $\text{R}_1^\text{'}$, $\text{R}_1^\text{''}$, $\text{R}_1^\text{'''}$ and $\text{R}_1^\text{''''}$ in formula (III) are 2-hydroxyethyl, hydrogen, hydrogen, hydrogen and hydrogen respectively.

5. Method as claimed in all preceding claims, characterized in that $m=1$ and that $\text{R}_6$ and $\text{R}_6$ together with the intermediate atoms form a 6-membered ring.

6. Method as claimed in claims 1-4, characterized in that $m=1$, that $\text{R}_6$ is methyl and that $\text{R}_6$ is hydrogen.

7. Method as claimed in any of the preceding claims, characterized in that the reaction is performed in an alcoholic solvent.

8. Method as claimed in claim 7, characterized in that the alcoholic solvent is 1-butanol.

9. Method as claimed in claims 1-6, characterized in that the reaction is performed in a mixture of an alcoholic solvent and an aromatic hydrocarbon

10. Method as claimed in claim 9, characterized in that said mixture is a mixture of methanol and chlorobenzene.
INTERNATIONAL SEARCH REPORT

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D233/60 C07D403/06 C07D471/06 A61K31/4174 A61K31/4178
A61P25/18

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data, EPO-Internal, PAJ, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

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<td>EP 0 297 651 B (DUPHAR INT RES) 4 January 1989 (1989-01-04) cited in the application page 2, line 3-5,26 * Example 1c), 1d) * claim 4</td>
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  *O* document referring to an oral disclosure, use, exhibition or other means
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Date of the actual completion of the international search

29 January 2004

Date of mailing of the international search report

06/02/2004

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Hoepfner, W
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