EUROPEAN PATENT SPECIFICATION

PROCESS FOR THE PREPARATION OF 5-(2-ETHYL-DIHYDRO-1H-INDEN-2-YL)-1H-IMIDAZOLE AND SALTS THEREOF

VERFAHREN ZUR HERSTELLUNG VON 5-(2-ETHYL-DIHYDRO-1H-INDEN-2-YL)-1H-IMIDAZOL UND SALZEN DAVON

PROCÉDÉ DE PRÉPARATION DU 5-(2-ÉTHYL-DIHYDRO-1H-INDÈN-2-YL)-1H-IMIDAZOLE ET DE SES SELS

Designated Contracting States:
AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HR HU IE IS IT LV MC MT NL NO PL PT RO SE SI SK TR

Designated Extension States:
RS

Priority: 05.12.2007 EP 07122400

Date of publication of application: 08.09.2010 Bulletin 2010/36

Proprietor: Grindeks, a joint stock company Riga 1057 (LV)

Inventors:
• LUSIS, Viesturs
  LV-1048 Riga (LV)
• MUCENIECE, Dzintra
  LV-1048 Riga (LV)
• REINE, Inese
  LV-1055 Riga (LV)
• ZANDERSONS, Armands
  LV-1010 Riga (LV)

References cited:
EP-A-0 247 764
WO-A-2004/063168
WO-A-2006/134219

• DATABASE BEILSTEIN [Online] BEILSTEIN INSTITUTE FOR ORGANIC CHEMISTRY, FRANKFURT-MAIN, DE; XP002470549 Database accession no. 79113, 79114 (Reaction IDs) & CHEM. BER., vol. 47, 1914, page 1439,
• DATABASE BEILSTEIN [Online] BEILSTEIN INSTITUTE FOR ORGANIC CHEMISTRY, FRANKFURT-MAIN, DE; XP002470550 Database accession no. 755379 (BRN) & J. MED. CHEM., vol. 14, 1971, pages 883-885,
• DATABASE BEILSTEIN [Online] BEILSTEIN INSTITUTE FOR ORGANIC CHEMISTRY, FRANKFURT-MAIN, DE; XP002518331 Database accession no. 7780814 (BRN) & J. MED. CHEM., vol. 40, no. 19, 1997, pages 3014-3024,
The present invention provides processes and intermediates useful in the preparation of 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole (international non-proprietary name - "Atipamezole") and salts, in particular pharmaceutically acceptable salts, thereof, a potent and selective $\alpha_2$-receptor antagonist.


First synthetic route as starting material was used 2-acetyl-1-indanone, which was alkylated with ethylbromide in acetone in the presence of sodium carbonate to 2-acetyl-2-ethyl-1-indanone. The acetyl group was brominated with bromine in methanol and to imidazole by heating in formamide. Then the intermediate was hydrogenated in 2N hydrochloric acid in the presence of 10% palladium on carbon.

Second synthetic route disclosed in the same patent is following, as starting material was used 2,3-dihydro-1H-indene-2-carboxylic acid methyl ester, which was prepared by methylation of 2,3-dihydro-1H-indene-2-carboxylic acid in the presence of sulphuric acid. The 2,3-dihydro-1H-indene-2-carboxylic acid methyl ester was reacted with N-isopropylcyclohexylamide and ethylbromide yielding 2,3-dihydro-2-ethyl-1H-indene-2-carboxylic acid chloride and reaction mixture was treated with sulphuric acid, and 1-(2,3-dihydro-2-ethyl-1H-inden-2-yl)ethanone was obtained, then the intermediate was stirred in methylene chloride and bromine was added by giving a new intermediate 2-bromo-1-(2,3-dihydro-2-ethyl-1H-inden-2-yl)ethanone, to which was thereafter added formamide and hydrochloric acid yielding crude product of 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole. The last step involved hydrogenation of the crude product of 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole with 10% palladium on carbon.

EP 0247764 B (ORION-YHTYMÄ OY) 1987.02.12. disclosed the following process for preparation of 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole hydrochloride. The process starts by reaction of alpha,alpha-dibromo-o-xylene with 4-penten-2-one to obtain 1-(2,3-dihydro-2-vinyl-1H-inden-2-yl)ethanone. The obtained intermediate was brominated, e.g. with bromine, methylene chloride was used as solvent and 2-bromo-1-(2,3-dihydro-2-vinyl-1H-inden-2-yl)ethanone was obtained, which is thereafter reacted with formamide in excess formamide to give a 4(5)-(2,3-dihydro-2-vinyl-1H-inden-2-yl)imidazole hydrochloride. As the last step the vinyl group was catalytically hydrogenated to an ethyl group so as to form a product 4(5)-(2,3-dihydro-2-ethyl-1H-inden-2-yl)imidazole.

Another synthetic route for obtaining 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole is disclosed in WAI, Wonf, et al. A Concise Synthesis of Atipamezole. Synthesis. 1995, no.2, p.139-140. The cyclization of alpha,alpha'-dibromo-o-xylene with acetylacetone by means of NaOH and tetrabutylammonium bromide in toluene/water at 80°C under phase-transfer conditions gives the unstable diacetyl derivative, which presumably undergoes cleavage to afford 2-acetylindane. The alkylation of 2-acetylindane with ethyl iodide and potassium tert-butoxide yields 2-acetyl-2-ethylindan, which is brominated with Br$_2$ to give 2-bromoacetyl-2-ethylindan. Finally, this compound is cyclised with formamide at 160°C (some 2-ethyl-2-(4-oxazolyl)indane is also formed but easily eliminated); the cyclization can also be carried out with formamidine in liquid ammonia. Although the substitution of formamide by formamidine acetate eliminates the oxazole formation, it does not increase the yield of Atipamezole (<30%) WAI, Wonf, et al. A Concise Synthesis of Atipamezole. Synthesis. 1995, no.2, p.139-140 in the final step.

The bromination of ketone by using bromine in dichloromethane was sluggish and led to many byproducts; Converting bromo ketone to Atipamezole using formamide at 160°C, Atipamezole was formed as the minor product. The major product in this reaction was the oxazole. The EP 0310745 B (FARMOS OY) 1989.04.12. which describes this reaction did not mention the formation of any oxazole, and the yield of the reaction was not disclosed. The instant invention provides processes useful in the preparation of Atipamezole. The invention further provides intermediates useful in the preparation of Atipamezole, and processes useful in the production of such intermediates. Moreover it will be evident for the person skilled in the art that this novel method provides convenient route to new analogues of Atipamezole not attainable by the known methods, as well as to new derivatives of indane-1,3-dione.

The above objective is achieved according to the present invention by sequential combination of four process steps, starting from 1-trityl-1H-imidazole-4-carboxaldehyde:
(a) condensation of phtalide to 1-trityl-1H-imidazole-4-carboxaldehyde (I), and, thus preparation of 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione (II);
(b) alkylating of 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione with ethyl iodide to produce 2-ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione (III);
(c) removing the trityl group of 2-ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione by acid hydrolysis to yield the deprotected 2-ethyl-2-(1H-imidazol-2-yl)indan-1,3-dione (IV);
(d) reducing the product of step (c) by catalytic hydrogenation to form the desired 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole hydrochloride (V).

In another aspect of the present invention, there is provided new valuable intermediates for preparing Atipamezole and analogues thereof, as well as new derivatives of indane-1,3-dione, namely 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione (II) and 2-ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione (III).

Reaction Scheme 1
Method includes the following advantages:

1. We have found an essential process for obtaining 5-(2-ethyl-2,3,1H-inden-2-yl)-1H-imidazole, without bromination in any step of process, thus preventing the possibility of brominated by-products;
2. This process has given superior yields, compared to patents cited above;
3. This process is amenable to large scale production which does not require specialized equipment.

The condensing of commercially available 1-trityl-1H-imidazole-4-carboxaldehyde (I) with phtalide to form 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione (II) is performed under the conditions that are similar to those used for synthesis of 4-(indane-1,3-dionyl) pyridine J. Org. Chem. 1971, vol. 36, p.1563. surprisingly, the bulky 1-trityl-1H-imidazole-4-carboxaldehyde (I) reacted as expected and produced 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione (II) in over 67% yield. Both ethyl acetate and dioxane can be used as reaction media.

The alkylation of (II) by ethyl iodide is performed in boiling acetone with potassium carbonate as basic agent. 2-Ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione (III) is formed in over 67% yield and easily isolated from the acetone solution by concentrating it and diluting with water. A high purity (III) is obtained after crystallization from methanol or ethanol.
Removing the trityl group of 2-ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione by acid hydrolysis to yield the deprotected 2-ethyl-2-(1H-imidazol-2-yl)indan-1,3-dione.

The reduction of (IV) to 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole hydrochloride (V) is performed in hydrogenation apparatus with Pd/C catalyst under hydrogen pressure in HCl solution. The reaction proceeds under variable pressure and temperature conditions, but a pressure of about 3 bar and the temperature of about 80-85°C is preferable. After removing the catalyst the product crystallizes on chilling in over 77% yield. It can be purified by additional crystallization.

The present invention will be described in more detail by referring to the following non-limiting examples.

### Example 1
#### Preparation of 1-trityl-1H-imidazole-4-carboxaldehyde

1-Trityl-4-iodoimidazole (87.3g, 0.200mol) was added to stirred methylene chloride (525mL) in a 4-neck round bottom flask fitted with a mechanical stirrer, a thermometer, a dropping funnel and a tube for argon introduction into the reaction mixture.

The reaction mixture was cooled to 10°C, at which point isopropylmagnesium chloride solution in tetrahydrofuran (112mL, 0.213 mol) was added dropwise under an argon atmosphere. After addition of isopropylmagnesium chloride the reaction mixture was warmed to 20°C.

N,N-Dimethylformamide (47mL, 0,608mol) was added to methylene chloride (300mL) in a 4-neck round bottom flask fitted with a mechanical stirrer, a thermometer, a dropping funnel and a tube for argon introduction into the reaction mixture. The reaction mixture was stirred and cooled to (-5) °C and solution of the imidazole Grignard derivative, which was prepared above, was added to the reaction mixture. The reaction mixture was stirred at (-5) °C for half an hour and then at 20°C for 10 hours, at which point 10% aqueous ammonium chloride solution (300mL) was added to the reaction mixture.

The aqueous layer was extracted with methylene chloride (550mL). The organic layer was separated and washed with saturated sodium chloride solution, and then the organic layer was stirred with anhydrous magnesium sulphate for 2 hours. The precipitate of magnesium sulphate was separated by filtration.

The solvent was removed by distillation at a reduced pressure. Ethanol (200mL) was added to the distillation residue and the reaction mixture was cooled to (-5)°C for 2 hours.

The precipitates were separated by filtration. The obtained intermediate 1-trityl-1H-imidazole-4-carboxaldehyde was dried at under reduced pressure. The yield was 56.6g (73.2%).

### Example 2
#### Preparation of 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione

1-Trityl-1H-imidazole-4-carboxaldehyde (120g, 0.319mol) and phthalide (42.8g, 0.319mol) were added to stirred ethyl acetate (1000mL) in a 4-neck round bottom flask fitted with a mechanical stirrer, a thermometer, a dropping funnel and a reflux condenser.

Meanwhile sodium methoxide (51.7g, 0.957mol) was added to cooled methanol (500mL) in a separate vessel.

Thereafter the methanolic solution of sodium methoxide was added to the reaction mixture at 60°C and was heated at this temperature for 3 hours and then cooled to 30°C, at which point the solvent was removed by distillation at a reduced pressure.

The distillation residue was poured into water and aqueous hydrochloric acid solution was added, until pH=4-5. The solid orange-brownish precipitates were filtered, the crude product of 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione was washed on filter with water.

After the recrystallization of crude 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione from ethanol the yield of the 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione intermediate product was 97.4g (67.2%), having a melting temperature of 213 to 215°C.

### Example 3
#### Preparation of 2-ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione

1-Trityl-1H-imidazole-4-carboxaldehyde (120g, 0.319mol) and phthalide (42.8g, 0.319mol) were added to stirred ethyl acetate (1000mL) in a 4-neck round bottom flask fitted with a mechanical stirrer, a thermometer, a dropping funnel and a reflux condenser.

Meanwhile sodium methoxide (51.7g, 0.957mol) was added to cooled methanol (500mL) in a separate vessel.

Thereafter the methanolic solution of sodium methoxide was added to the reaction mixture at 60°C and was heated at this temperature for 3 hours and then cooled to 30°C, at which point the solvent was removed by distillation at a reduced pressure.

The distillation residue was poured into water and aqueous hydrochloric acid solution was added, until pH=4-5. The solid orange-brownish precipitates were filtered, the crude product of 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione was washed on filter with water.

After the recrystallization of crude 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione from ethanol the yield of the 2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione intermediate product was 97.4g (67.2%), having a melting temperature of 213 to 215°C.
then cooled to 20 °C and filtered. The inorganic residue on the filter was washed with acetone. The filtrate was concentrated in vacuo and poured into water.

The mixture was stirred at 20°C, the solid yellow-reddish precipitate was filtered, and the crude 2-ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione was washed on the filter with water.

After the recrystallization of crude of 2-ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione from ethanol, the yield of 2-ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione was 43.6g (50.9%), having a melting temperature of 196-197°C.

Example 4

Preparation of 2-ethyl-2-(1H-imidazol-1-yl)indan-1,3-dione

2-Ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione (93.4g, 0.193mol) was added to stirred 2N hydrochloric acid (650mL) in a 3-neck round bottom flask fitted with a mechanical stirrer, a thermometer and a reflux condenser. The reaction mixture was heated at 100°C for 3 hours.

Thereafter the reaction mixture was cooled to 25°C, at which point the reaction mixture was filtered to remove triphenylmethanol. The filtrate was cooled and 20% sodium hydroxide (276mL) was added.

The precipitate was separated by filtration and washed with water (700 mL). The obtained intermediate 2-ethyl-2-(1H-imidazol-4-yl)indan-1,3-dione was dried under reduced pressure. The yield was 35.7g (76.2%) of white crystalline 2-ethyl-2-(1H-imidazol-4-yl)indan-1,3-dione.

Example 5

Preparation of crude 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole hydrochloride

2-Ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione (26.8g, 0.056mol) was added to stirred 5N hydrochloric acid (200mL) in a 3-neck round bottom flask fitted with a mechanical stirrer, a thermometer and a reflux condenser. The reaction mixture was heated at 100°C for 3 hours.

Thereafter the reaction mixture was cooled to 25°C, at which point the reaction mixture was filtered to remove triphenylmethanol. The triphenylmethanol cake on the filter was washed with 5N hydrochloric acid (100mL). The filtrate was mixed with activated charcoal and stirred for 10 minutes; the charcoal was separated by filtration.

A hydrochloric acid solution of 2-ethyl-2-(1H-imidazol-4-yl)indan-1,3-dione hydrochloride was obtained.

Palladium catalyst (1.4g of 10% Pd/C) was suspended in the hydrochloric acid solution of 2-ethyl-2-(1H-imidazol-4-yl)indan-1,3-dione hydrochloride. The obtained suspension was poured into a hydrogenation autoclave. Hydrogen was supplied to autoclave to 3.0 bar. The reaction mixture was stirred and heated to 80-85°C. The typical hydrogenation time was 6 hours. After hydrogenation the reaction mixture was filtered to remove the catalyst. The filtrate was cooled to (-5)°C, the precipitates were separated by filtration. The obtained crude 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole hydrochloride was dried at 50°C under reduced pressure. The yield was 3.7g of a white powder.

Alternatively, the filtrate of reaction mixture was concentrated in vacuo almost to dryness. Acetone (50mL) was added to the stirred reaction mixture and thereafter the solvent was removed by distillation at a reduced pressure. Another portion of acetone (40mL) was added to the reaction mixture. The reaction mixture was cooled to 0°C for 4 hours.

The suspension was filtered; the crude product was washed with acetone on the filter.

Example 6

Preparation of 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole hydrochloride.

The crude 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole hydrochloride (12.5g) was added to stirred acetonitrile (162mL) and water (14mL) in a 3-neck round bottom flask fitted with a mechanical stirrer, a thermometer and a reflux condenser.

The reaction mixture was stirred at 75-80°C the reaction mixture was filtered, and the filtrate was cooled to 0-5°C.

The precipitates were separated by filtration and the product cake on the filter was washed with acetone (50mL). The 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H-imidazole hydrochloride was dried at 80-90°C under reduced pressure. The yield was 9.8g (77.3%) of colourless crystalline powder.

Claims

1. A process for preparing 5-(2-ethyl-2,3-dihydro-1H-inden-2-yl)-1H imidazole hydrochloride of formula (V)
comprising the steps of:

a) condensing the 1-trityl-1H-imidazole-4-carboxaldehyde with phtalide to form 2-(1-trityl-1H-imidazole-4-yl) indan-1,3-dione,

b) alkylating the product of step (a) with ethyl iodide to form 2-ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione,

c) removing the trityl group from the product of step (b) by acid hydrolysis and form 2-ethyl-2-(1H-imidazol-4- yl)indan-1,3-dione,

d) reducing the product of step (c) by catalytic hydrogenation to form the desired 5-(2-ethyl-2,3-dihydro-1H- inden-2-yl)-1H imidazole hydrochloride.

2. A process according to claim 1 wherein step a) is carried out in an organic solvent selected from the group of ethyl acetate in the presence methanolic solution of sodium methoxide.

3. A process according to claim 1 or 2 wherein step b) is carried out by alkylation of a compound of formula (II) by ethyl iodide and alkali metal carbonate.

4. A process according to claim 3 wherein alkali metal is potassium carbonate.

5. A process according to claim 1 to 4 wherein step c) is carried out by using hydrochloric acid.

6. A process according to claim 1 to 5 wherein step d) is carried out by catalytic hydrogenation using hydrochloric acid solution in the presence of Pd/C catalyst.

7. 2-(1-Trityl-1H-imidazole-4-yl)indan-1,3-dione.

8. 2-Ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione.

Patentansprüche

1. Ein Verfahren zur Herstellung von 5-(2-Ethyl-2,3-Dihydro-1H-Inden-2-yl)-1H-Imidazol-Hydrochlorid mit der Formel (V),

2. 2-(1-Trityl-1H-imidazole-4-yl)indan-1,3-dione.

3. 2-Ethyl-2-(1-trityl-1H-imidazole-4-yl)indan-1,3-dione.
2-(1H-Imidazol-4-yl)inden-1,3-dion,

d) Reduzierung des Produkts aus Schritt (c) durch katalytische Hydrierung, um das gewünschte 5-(2-Ethyl-2,3-Dihydro-1H-Inden-2-yl)-1H-Imidazol-Hydrochlorid zu bilden.

2. Ein Verfahren gemäß Anspruch 1, wobei Schritt a) in einem organischen Lösungsmittel, das aus der Gruppe der Ethylacetate ausgesucht wurde, in Gegenwart einer methanolischen Lösung von Natriummethoxid durchgeführt wird.


5. Ein Verfahren gemäß Ansprüche 1 bis 4, wobei Schritt c) durch die Verwendung von Salzsäure durchgeführt wird.


7. 2-(1-Trityl-1H-Imidazol-4-yl)inden-1,3-dion.

8. 2-Ethyl-2-(1-Trityl-1H-Imidazol-4-yl)inden-1,3-dion.

Revendications

1. Le procédé de préparation de 5-(2-éthyl-2,3-dihydro-1H-inden-2-yl)-1H hydrochloride imidazole avec la formule (V)

\[
\text{\( \text{V} \)}
\]

comprend les étapes suivantes :

a) la condensation de 1-trityl-1H-imidazole-4-carboxaldehyde avec la phtalide afin de former 2-(1-trityl-1H-imidazole-4-yl)inden-1,3-dione,

b) l'alcoylation du produit de l'étape (a) avec l'iode d'éthyle afin de former la 2-éthyl-2-(1-trityl-1H-imidazole-4-yl)inden-1,3-dione,

c) l'évacuation du groupe trityl du produit de l'étape (b) par l'hydrolyse acide afin de former la 2-éthyl-2-(1H-imidazol-4-yl)inden-1,3-dione,

d) la réduction du produit de l'étape (c) par l'hydrogénation catalytique afin de former la 5-(2-éthyl-2,3-dihydro-1H-inden-2-yl)-1H hydrochloride d'imidazole désirée.

2. Un procédé conformément à la revendication 1 où l'étape a) est réalisée dans un dissolvant organique choisi du groupe de l'acétate d'éthyle dans la présence de la solution méthanolique du méthoxide de sodium.

3. Un procédé conformément aux revendications 1 ou 2 où l'étape b) est réalisée par l'alcoylation du composé de la formule (II) par l'iode d'éthyle et le carbonate du métal alcalin.

4. Un procédé conformément à la revendication 3 où le métal alcalin est le carbonate de potassium.

5. Un procédé conformément aux revendications 1 à 4 où l'étape c) est réalisée en utilisant l'acide hydrochlorique.

6. Un procédé conformément aux revendications 1 à 5 où l'étape d) est réalisée par l'hydrogénation catalytique en
utilisant la dissolution de l'acide hydrochlorique dans la présence du catalyseur Pd/C.

7. La 2-(1-Trityl-1H-imidazole-4-yl)indan-1,3-dione.

8. La 2-éthyl-2-(trityl-1H-imidazole-4-yl)indan-1,3-dione.
REFERENCES CITED IN THE DESCRIPTION

This list of references cited by the applicant is for the reader’s convenience only. It does not form part of the European patent document. Even though great care has been taken in compiling the references, errors or omissions cannot be excluded and the EPO disclaims all liability in this regard.

Patent documents cited in the description

- EP 0310745 B [0002] [0008]
- EP 0247764 B [0005]

Non-patent literature cited in the description