

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
10 January 2002 (10.01.2002)

PCT

(10) International Publication Number
WO 02/02500 A1

(51) International Patent Classification⁷: **C07C 67/293**,
51/36, 59/66

(21) International Application Number: PCT/CH01/00397

(22) International Filing Date: 26 June 2001 (26.06.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
1317/00 3 July 2000 (03.07.2000) CH

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(81) Designated States (national): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ,
DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,
TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

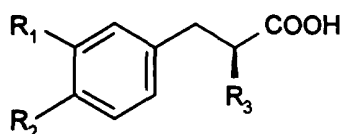
(84) Designated States (regional): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,
IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF,
CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

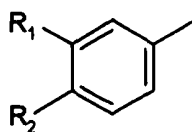
- with international search report
- before the expiration of the time limit for amending the
claims and to be republished in the event of receipt of
amendments

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: PREPARATION OF (R)-2-ALKYL-3-PHENYLPROPIONIC ACIDS



(I)



(a)

(57) Abstract: Compounds of
formula (I), wherein R₁ and R₂
are, independently of one another,
H, C₁-C₆alkyl, C₁-C₆halogenalkyl,
C₁-C₆alkoxy, C₁-C₆alkoxy-C₁-C₆alkyl,
or C₁-C₆alkoxy-C₁-C₆alkyloxy, and R₃
is C₁-C₆alkyl, are obtainable in high

yields by stereoselective addition of R₃-substituted propionic acid esters to R₁- and R₂-substituted benzaldehydes of formula R-CHO
to form corresponding 3-R-3-hydroxy-2-R₃-propionic acid esters, conversion of the OH group to a leaving group, subsequent
regioselective elimination to form 3-R-2-R₃-propenic acid esters, and their hydrolysis to form corresponding propenic carboxylic
acids and their enantioselective hydrogenation, wherein R is (a).

Preparation of (R)-2-alkyl-3-phenylpropionic acids

The invention relates to a stereoselective process for the preparation of (R)-2-alkyl-3-phenyl-propionic acids and intermediate products obtained in the process steps.

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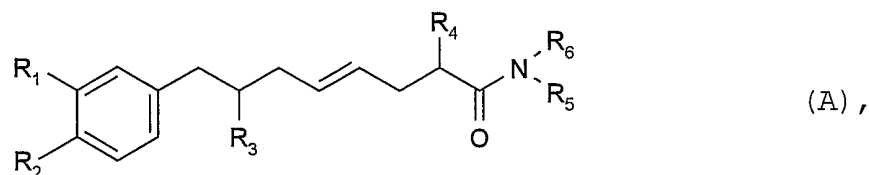
In EP-A-0 678 503, δ -amino- γ -hydroxy- ω -aryl-alkanecarboxamides are described which exhibit renin-inhibiting properties and could be used as antihypertensive agents in pharmaceutical preparations. The manufacturing processes described are unsatisfactory in terms of the number of process steps and yields and are not suitable for an industrial process. A disadvantage of these processes is also that the total yields of pure diastereomers that are obtainable are too small.

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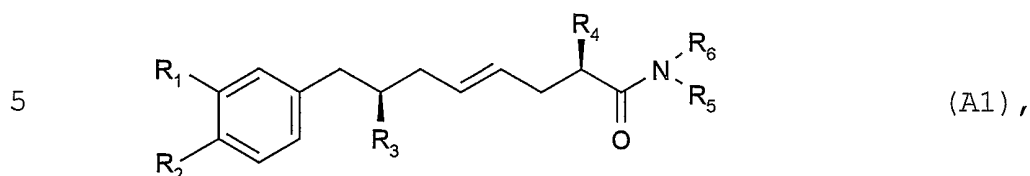
In a new process, one starts from 2,7-dialkyl-8-aryl-4-octenoyl amides, whose double bond is simultaneously halogenated in the 5-position and hydroxylated in the 4-position under lactonization, then the halogen is substituted by azide, the lactone amidated and the azide then transferred to the amine group. The desired alkanecarboxamides are obtained with the new process both in high total yields and in a high degree of purity, and selectively pure diastereomers can be prepared. The halolactonization of process step a), the azidation of process step b), and the azide reduction of process step d) are described by P. Herold in the Journal of Organic Chemistry, Vol. 54 (1989), pages 1178-1185.

30 The 2,7-dialkyl-8-aryl-4-octenoyl amides may correspond for example to formula A,

- 2 -



and especially to formula A1



wherein R_1 and R_2 are, independently of one another, H, C_1 - C_6 alkyl, C_1 - C_6 halogenalkyl, C_1 - C_6 alkoxy, C_1 - C_6 alkoxy- C_1 - C_6 alkyl, or C_1 - C_6 alkoxy- C_1 - C_6 alkyloxy, R_3 is C_1 - C_6 alkyl, R_4 is
 10 C_1 - C_6 alkyl, R_6 is C_1 - C_6 alkyl, R_5 is C_1 - C_6 alkyl or C_1 - C_6 alkoxy, or R_5 and R_6 together are tetramethylene, pentamethylene, 3-oxa-1,5-pentylene or $-CH_2CH_2O-C(O)-$ substituted if necessary with C_1 - C_4 alkyl, phenyl or benzyl.

15 The compounds of formulae A and A1 are obtainable by reacting a compound of formula B



20 as racemate or enantiomer, with a compound of formula C, as racemate or enantiomer,



- 3 -

wherein R_1 to R_4 , R_5 and R_6 are as defined above, Y is Cl, Br or I and Z is Cl, Br or I, in the presence of an alkali metal or alkaline earth metal. Y and Z are preferably Br and especially Cl.

5

The compounds of formula B are known from EP-A-0 678 503. The compounds of formula C may be prepared from amidation of the corresponding carbonic esters, amides, or halides. The formation of carboxamides from carbonic esters and amines in
10 the presence of trialkyl aluminium or dialkyl aluminium halide, for example using trimethyl aluminium or dimethyl aluminium chloride, is described by S. M. Weinreb in Org. Synthesis, VI, page 49 (1988). The carbonic esters are obtainable by the reaction of trans-1,3-dihalogenpropene
15 (for example, trans-1,3-dichlorepropene) with corresponding carbonic esters in the presence of strong bases, for example alkali metal amides.

A satisfactory solution for the stereoselective preparation
20 of compounds of formula B has not yet been found, especially with regard to an industrial process. Surprisingly it has now been found that 2-alkyl-3-phenylpropionic acids can be stereoselectively prepared with high yields in only three process steps. When suitably substituted benzaldehydes are
25 condensed with carbonic esters to form 2-alkyl-3-hydroxy-3-phenylpropionic acid esters, the desired diastereomers are obtainable in surprisingly high yields mostly as crystalline compounds which can be readily isolated. After conversion of the hydroxy group to a leaving group, 2-alkylcinnamic acid
30 esters are then formed by elimination with strong bases with surprisingly high regioselectivity. The carboxylic acids obtained after saponification can in turn be surprisingly hydrogenated in the presence of homogeneous, asymmetric hydrogenation catalysts to form practically enantiomer-pure
35 2-alkyl-3-phenylpropionic acids. These acids can then be

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reduced in a manner known per se to form enantiomer-pure alcohols, from which the compounds of formula B are obtainable by halogenation.

- 5 The object of the invention is a process for the preparation of compounds of formula I,



- 10 wherein R₁ and R₂ are, independently of one another, H, C₁-C₆alkyl, C₁-C₆halogenalkyl, C₁-C₆alkoxy, C₁-C₆alkoxy-C₁-C₆alkyl, or C₁-C₆alkoxy-C₁-C₆alkyloxy, and R₃ is C₁-C₆alkyl, comprising

- a) the reaction of a compound of formula II
- 15



- wherein R₁ and R₂ are as defined above, with a compound of formula III,
- 20



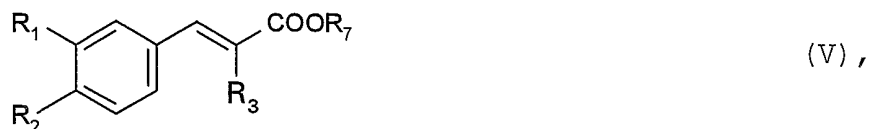
wherein R₃ is as defined above, to form a compound of IV,



- 25 wherein R₇ is C₁-C₁₂alkyl, C₃-C₈cycloalkyl, phenyl or benzyl,

b) the isolation of the crystalline compound of formula IV, the conversion of the OH group to a leaving group, and the reaction of a compound containing a leaving group in the presence of a strong base to form a compound of formula V,

5



c) the hydrolysis of carbonic esters of formula V to form the carboxylic acid of formula VI,

10



d) the hydrogenation of the carboxylic acid of formula VI in the presence of hydrogen and catalytic quantities of a metal complex as asymmetric hydrogenation catalyst, comprising metals from the group of ruthenium, rhodium and iridium, to which the chiral bidentate ligands are bonded, to form a compound of formula I.

20 R_1 and R_2 may be a linear or branched alkyl and preferably comprise 1 to 4 C atoms. Examples are methyl, ethyl, n- and i-propyl, n-, i- and t-butyl, pentyl and hexyl.

25 R_1 and R_2 may be a linear or branched halogenalkyl and preferably comprise 1 to 4 C atoms, 1 or 2 C atoms being especially preferred. Examples are fluoromethyl, difluoromethyl, trifluoromethyl, chloromethyl, dichloromethyl, trichloromethyl, 2-chloroethyl and 2,2,2-trifluoroethyl.

30

R₁ and R₂ may be a linear or branched alkoxy and preferably comprise 1 to 4 C atoms. Examples are methoxy, ethoxy, n- and i-propyloxy, n-, i- and t-butyloxy, pentyloxy and hexyloxy.

5

R₁ and R₂ may be a linear or branched alkoxyalkyl. The alkoxy group preferably comprises 1 to 4 and especially 1 or 2 C atoms, and the alkyl group preferably comprises 1 to 4 C atoms. Examples are methoxymethyl, 1-methoxyeth-2-yl, 10 1-methoxyprop-3-yl, 1-methoxybut-4-yl, methoxypentyl, methoxyhexyl, ethoxymethyl, 1-ethoxyeth-2-yl, 1-ethoxyprop-3-yl, 1-ethoxybut-4-yl, ethoxypentyl, ethoxyhexyl, propyloxymethyl, butyloxymethyl, 1-propyloxyeth-2-yl and 1-butyloxyeth-2-yl.

15

R₁ and R₂ may be linear or branched C₁-C₆alkoxy-C₁-C₆alkyloxy. The alkoxy group preferably comprises 1 to 4 and especially 1 or 2 C atoms, and the alkyloxy group preferably comprises 1 to 4 C atoms. Examples are methoxymethyloxy, 1-methoxyeth-20 2-yloxy, 1-methoxyprop-3-yloxy, 1-methoxybut-4-yloxy, methoxypentyloxy, methoxyhexyloxy, ethoxymethyloxy, 1-ethoxyeth-2-yloxy, 1-ethoxyprop-3-yloxy, 1-ethoxybut-4-yloxy, ethoxypentyloxy, ethoxyhexyloxy, propyloxymethyloxy, butyloxymethyloxy, 1-propyloxyeth-2-yloxy and 1-butyloxyeth-25 2-yloxy.

In a preferred embodiment, R₁ is methoxy-C₁-C₄alkyloxy or ethoxy-C₁-C₄alkyloxy, and R₂ is preferably methoxy or ethoxy. Quite especially preferred are compounds of formula I, 30 wherein R₁ is 1-methoxyprop-3-yloxy and R₂ is methoxy.

R₃ may be a linear or branched alkyl and preferably comprise 1 to 4 C atoms. Examples are methyl, ethyl, n- and i-propyl, n-, i- and t-butyl, pentyl and hexyl. In a preferred 35 embodiment, R₃ in compounds of formula I is isopropyl.

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Especially preferred are compounds of formula I wherein R₁ is methoxy-n-propoxy, R₂ is methoxy and R₃ is isopropyl.

R₇ is preferably C₁-C₆alkyl, C₁-C₄alkyl being especially preferred; some examples are methyl, ethyl, n-propyl and n-butyl.

The starting compounds of formulae II and III used in process step a) are known or can be prepared in a manner similar to known processes. Compounds of formula II are described in EP-A 0 678 503. The reaction is advantageously carried out at low temperatures, for example 0-40°C, in the presence of at least equivalent quantities of strong bases. The reaction is further expediently carried out in a solvent, ethers such as diethyl ether, tetrahydrofuran and dioxane being especially suitable. Suitable strong bases are in particular alkali metal alcoholates and secondary amides, such as lithium diisopropylamide.

The desired diastereomer of formula IV is surprisingly formed up to about 75%. The compounds of formula IV are surprisingly crystalline and can therefore be readily isolated without any substantial losses by means of extraction and crystallization.

The conversion of the OH group to a leaving group in reaction step b) is known per se. Reaction with carboxylic acids or sulfonic acids, or their anhydrides (acylation), is especially suitable. Some examples of carboxylic acids are formic acid, acetic acid, propionic acid, benzoic acid, benzenesulfonic acid, toluenesulfonic acid, methanesulfonic acid and trifluoromethanesulfonic acid. The use of acetic acid anhydride has proved especially successful. The elimination is expediently carried out in the presence of strong bases, alkali metal alcoholates such as potassium t-butylate being especially suitable. The presence of solvents

such as ethers is expedient. The reaction is advantageously carried out at low temperatures, for example 0-40°C. It is of advantage to conduct the elimination reaction directly in the reaction mixture for acylation. The elimination leads to the desired Z isomers with surprisingly high regioselectivity. These isomers are crystalline and can therefore be readily isolated without any substantial losses by means of extraction and crystallization. The yields are above 80%.

Hydrolysis of the ester of formula V to form the carboxylic acids of formula VI in process step c) is a generally known reaction. The hydrolysis may be carried out after isolation and purification of the compound of formula III. It is expedient to add water to the reaction mixture of process step b), to evaporate off the solvent and then to carry out alkaline or acidic hydrolysis. The carboxylic acids of formula VI are crystalline and can be readily isolated in yields of 80% or more.

The asymmetric hydrogenation in process step d) of α,β -unsaturated carboxylic acids with homogeneous, asymmetric hydrogenation catalysts is known per se and described for example by John M. Brown in E. Jacobsen, A. Pfaltz, H. Yamamoto (Eds.), Comprehensive Asymmetric Catalysis I to III, Springer Verlag, 1999, pages 121 to 182. Especially effective are ruthenium and rhodium catalysts. Chiral ditertiary diphosphines whose phosphine groups in the 1,2, 1,3 or 1,4 position are bonded to a C₂-C₄carbon chain are often used as ligands. The skeletal structures of the chiral ditertiary diphosphines may be acyclic, monocyclic or polycyclic. The phosphine groups may be substituted with the same or with different, preferably the same, substituents selected from the group of C₁-C₈alkyl, C₃-C₈cycloalkyl, C₆-C₁₂aryl, and C₆-C₁₂aryl- C₁-C₄alkyl. Cycloalkyl and aryl may be unsubstituted or substituted with C₁-C₄alkyl, C₁-C₄alkoxy,

C₁-C₄fluoroalkyl or C-C₁₂secondary amino. Suitable phosphine groups are also phosphanyl, preferably five-member phosphanyl, which if necessary is substituted in one or both α -positions with C₁-C₄alkyl or C₁-C₄alkoxy.

5

Some examples of chiral ditertiary diphosphines are (R"₂P is for example diphenylphosphino or dicyclohexylphosphino, substituted if necessary) 1,2-Di-R"₂P-propane, 2,3-Di-R"₂P-butane, 1,2-Di-R"₂P-norbornane or -norbornadiene, 1,2-Di-
10 R"₂P-cyclopentane, 1,2-Di-R"₂P-N-methylpyrrolidine, 2,2'-Di-R"₂P-biphenyl or -binaphthyl, 2,2'-Di-R"₂P-6-methyl or -6,6'-dimethylbiphenyl, 2,2'-Di-R"₂P-6-methoxy or -6,6'-dimethoxybiphenyl, and 1-(α -R"₂P-ethyl)-2-R"₂P-ferrocene.

15 Good optical yields are achieved using metal complexes of formula VII or VIIa,



20 wherein

Me is rhodium;

Y stands for two olefins or one diene;

Z is Cl, Br or I;

E⁻ is the anion of an oxygen acid or a complex acid; and

25 L is a chiral ligand from the group of ditertiary diphosphines, in which the phosphine groups are bonded to a C₂-C₄ chain of the diphosphine backbone chain, and the diphosphine forms a five to seven-member ring together with the rhodium atom.

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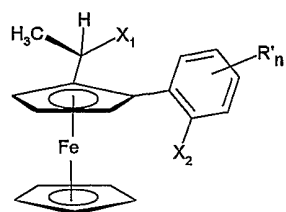
Where Y stands for two olefins, they may be C₂-C₁₂ olefins, C₂-C₆olefins being preferred and C₂-C₄olefins being especially preferred. Examples are propene, but-1-ene and especially ethylene. The diene may comprise 5 to 12 and preferably 5 to
35 8 C atoms and may be an acyclic, cyclic or polycyclic diene. The two olefin groups of the diene are preferably linked by

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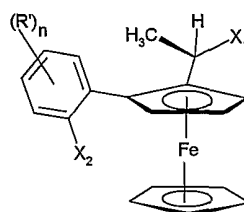
one or two CH_2 groups. Examples are 1,3-pentadiene, cyclopentadiene, 1,5-hexadiene, 1,4-cyclohexadiene, 1,4- or 1,5-heptadiene, 1,4- or 1,5-cycloheptadiene, 1,4- or 1,5-octadiene, 1,4- or 1,5-cyclooctadiene and norbornadiene.
 5 Y represents preferably two ethylene or 1,5- hexadiene, 1,5-cyclooctadiene or norbornadiene.

In formula VII, Z is preferably Cl or Br. Examples of E_1 are ClO_4^- , CF_3SO_3^- , CH_3SO_3^- , HSO_4^- , BF_4^- , $\text{B}(\text{phenyl})_4^-$, PF_6^- , SbCl_6^- ,
 10 AsF_6^- or SbF_6^- .

With known ligands for asymmetric catalysts, optical yields of up to about 80% ee can be achieved under optimized conditions. It was surprisingly found that new ligands with
 15 a ferrocenyl backbone are especially suitable for asymmetric hydrogenation of the compounds of formula VI. With these new ligands in the metal complexes of formulae VII and VIIa, optical yields of at least 95% ee can be achieved, which represents a substantial cost saving for manufacture on an
 20 industrial scale. In process step d), therefore, it is preferred to use metal complexes of formulae VII and VIIa which comprise ligands of formula VIII or VIIIa,



(VIII)



(VIIIa),

25

wherein

n is 0 or an integer from 1 to 4 and R' represents the same or different substituents selected from the C_1 - C_4 alkyl, $-\text{CF}_3$ and C_1 - C_4 alkoxy group; and

30 X_1 and X_2 are, independently of one another, secondary phosphino.

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As an alkyl, R' may preferably comprise 1 to 2 C atoms. Linear alkyl is preferred. Examples of R' as an alkyl are methyl, ethyl, n- and i-propyl, n-, i- and t-butyl. Methyl and ethyl are preferred, and methyl is especially preferred.

5

As an alkoxy, R' may preferably comprise 1 to 2 C atoms. Linear alkoxy is preferred. Examples of R' as an alkoxyl are methoxy, ethoxy, n- and i-propoxy, n-, i- and t-butoxy. Methoxy and ethoxy are preferred and methoxy is especially preferred.

10

The X₁ and X₂ groups may be different or preferably the same and correspond to formula PR₈R₉, wherein R₈ and R₉ are the same or different and represent branched C₃-C₈alkyl, C₃-C₈cycloalkyl, or unsubstituted or phenyl substituted with one to three C₁-C₄alkyl, C₁-C₄-alkoxy, or -CF₃.

15

Special preference is for ligands of formulae VIII and VIIIa, wherein n is 0, and X₁ and X₂ are a PR₈R₉ group, wherein R₈ and R₉ in each case are cyclohexyl, phenyl or phenyl substituted with 1 or 2 methyl, methoxy or CF₃.

20

The new ligands are prepared by means of reactions that are known per se or analogous to known reactions, such as those described in US-A-5,371,256, US-A-5,446,844 and US-A-5,583,241. Ligands with other phosphine groups may be prepared in a manner analogous to the method described in the example.

25

The metal complexes used as catalysts may be added as separately prepared isolated compounds, or also formed in situ before the reaction and then mixed with the substrate to be hydrogenated. It may be advantageous in the reaction using isolated metal complexes to add additional ligands, or in the in situ preparation to use surplus ligands. The surplus may for example be up to 10 moles and preferably

35

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0.001 to 5 moles, based on the metal complexes used for the preparation.

Process step d) may be carried out at low or elevated
5 temperatures, for example at temperatures from -20 to 150°C, preferably from -10 to 100°C, temperatures of 10 to 80°C being especially preferred. The optical yields are generally better at low temperatures than at high temperatures.

10 The process according to the invention may be carried out at normal pressure or preferably under positive pressure. The pressure may for example range from 10^5 to 2×10^7 Pa (Pascal).

Catalysts are preferably used in quantities from 0.0001 to
15 10 mol-% based on the compound to be hydrogenated, the range 0.001 to 10 mol-% being especially preferred and the range 0.01 to 5 mol-% being preferred in particular.

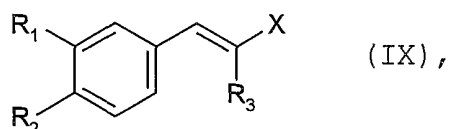
The preparation of catalysts as well as process step d) and
20 the other process steps may be carried out in the absence or the presence of an inert solvent, wherein one solvent or a mixture of solvents may be used. Suitable solvents are, for example, aliphatic, cycloaliphatic and aromatic hydrocarbons (pentane, hexane, petroleum ether, cyclohexane,
25 methylcyclohexane, benzene, toluene, xylene), aliphatic halogenated hydrocarbons (dichloromethane, chloroform, di- and tetrachloroethane), nitriles (acetonitrile, propionitrile, benzonitrile), ethers (diethyl ether, dibutyl ether, t-butyl methyl ether, ethylene glycol dimethyl ether,
30 ethylene glycol diethyl ether, diethylene glycol dimethyl ether, tetrahydrofuran, dioxane, diethylene glycol monomethyl or monoethyl ether), ketones (acetone, methyl isobutyl ketone), carbonic esters and lactones (ethyl or methyl acetate, valerolactone), N-substituted lactams
35 (N-methylpyrrolidone), carboxamides (dimethylamide, dimethylformamide), acyclic ureas (dimethylimidazoline), and

5 sulfoxides and sulfones (dimethyl sulfoxide, dimethyl sulfone, tetramethylene sulfoxide, tetramethylene sulfone) and alcohols (methanol, ethanol, propanol, butanol, ethylene glycol monomethyl ether, ethylene glycol monoethyl ether, diethylene glycol monomethyl ether) and water. The solvents may be used alone or in a combination of at least two solvents.

10 The reaction may be carried out in the presence of co-catalysts, for example quaternary ammonium halogenides (tetrabutylammonium iodide) and/or in the presence of protonic acids, for example mineral acids.

15 Using the regioselective and enantioselective process according to the invention, the intermediate products of formula (B) may be prepared via all process steps in yields of at least 50% by weight, based on the compounds of formula II. The high total yields make the process suitable for industrial use.

20 A further object of the invention relates to the compounds (intermediates) of formula IX,

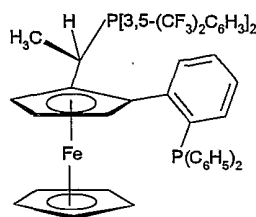


25 wherein R₁, R₂ and R₃ are as defined hereinbefore and X is the -COOH group.

The embodiments and preferences described hereinabove apply
30 for R₁, R₂, and R₃.

The following examples explain the invention in more detail.

A) Preparation of the ligands



Example A1: Preparation of

(L1)

5 a) Preparation of (R_C, S_P)-2-(2-bromophenyl)-1-[1-N,N-dimethylamino)ethyl]ferrocene, L2

At 0°C, 33 ml (43 mmol) of a 1.3 molar solution of s-butyl lithium in cyclohexane is added dropwise to a degassed solution of 10 g (38.9 mmol) (+)-(R)-1-N,N-dimethyl-
 10 aminoethyl ferrocene (L1) in 32 ml tetrahydrofuran (THF). After 30 minutes, still at 0°C, 44 ml of a 1 molar solution of ZnCl₂ in diethyl ether is added drop by drop. The reaction mixture is then stirred for one hour at room temperature. After the addition of 1.4 g (2 mmol) bis-diphenylphosphino-
 15 palladium(II) chloride and a solution of 22.64 g (80 mmol) 2-bromo-1-iodobenzene in 50 ml THF, the reaction mixture is heated for 3 days under reflux. The solvent is removed on a rotary evaporator, the residue taken up in CH₂Cl₂ and extracted with water. The aqueous phase is extracted 3 times
 20 with 30 ml CH₂Cl₂ and the combined organic phases are washed twice with 20 ml water. After drying over MgSO₄ and removal of the solvent in a vacuum, the residue is chromatographed on aluminium oxide 90. A mixture of petroleum ether, ether and triethylamine in a ratio of 60 : 1 : 3 is used as the
 25 mobile phase. The yield amounts to 4.65 g (11.3 mmol, 30%).

¹H-NMR: δ 1.61 (d, J=7.0 Hz, 3H), 1.75 (s, 6H), 3.54 (q, J=7.0 Hz, 1H), 4.13 (s, 5H, Cp), 4.23-4.25 (m, 1H, Cp), 4.32-4.34 (m, 1H, Cp), 4.59-4.61 (m, 1H, Cp), 7.07-7.11 (m, 1H, Ph),
 30 7.30-7.35 (m, 1H, Ph), 7.51-7.53 (m, 1H, Ph), 7.85-7.87 (m, 1H, Ph).

- 15 -

$[\alpha]^{20}$ (nm): +75.9 ° (589), + 61.4 ° (578), -45.4 ° (546) (c = 1, CHCl₃)

b) Preparation of (R_C,S_P)-1-[1-(N,N-dimethylamino)ethyl]-2-(2-diphenylphosphinophenyl)-ferrocene, L3

At -40°C, 4.5 ml of a 1.3 molar solution of s-butyl lithium in cyclohexane is slowly added drop by drop to a degassed solution of 2 g (4.87 mmol) of L2 in 25 ml THF. After 40 minutes, the reaction mixture is allowed to warm up to room temperature, and then 1.1 ml (6.6 mmol) diphenylchlorophosphine is added drop by drop. After 18 hours, 30 ml saturated NaHCO₃ solution is added. The organic phase is separated off and the aqueous phase is extracted twice with 20 ml CH₂Cl₂. The combined organic phases are washed twice with 20 ml water and dried over MgSO₄. After removal of the solvent in a vacuum and chromatography on silica gel 60 (petroleum ether / diethylamine = 95 : 5) a yield of 2.15 g (4.16 mmol, 85.4 %) of the product is obtained.

¹H-NMR: δ 1.64 (d, J=7.0 Hz, 3H), 1.86 (s, 6H), 3.72 (q, J=7.0 Hz, 1H), 4.04-4.06 (m, 1H, Cp), 4.08 (s, 5H, Cp), 4.22 (m, 1H, Cp), 4.25 (m, 1H, Cp), 6.93-6.98 (m, 2H, Ph), 6.99-7.02 (m, 1H, Ph), 7.15-7.20 (m, 4H, Ph), 7.31-7.40 (m, 6H, Ph), 7.94-7.98 (m, 1H, Ph).

³¹P-NMR: δ -14.09.

$[\alpha]^{20}$ (nm): -23.7 ° (589), -47.5 ° (578), -203.2 ° (546) (c = 1, CHCl₃).

c) Preparation of (R_C,S_P)-1-[1-(N,N-dimethylamino)ethyl]-2-(2-diphenylphosphinylphenyl)-ferrocene, L4

To a solution of 1 g (1.93 mmol) L3 in 15 ml acetone, 0.8 ml 30% H₂O₂ is added dropwise. The solution is stirred for 45 minutes at room temperature, and then 20 ml saturated Na₂S₂O₅ solution is added. After extraction 3 x 25 ml CH₂Cl₂ the combined organic phases are washed with 2 x 20 ml water and dried over MgSO₄. The solvent is removed in a vacuum and the product purified by chromatography on aluminium oxide 90.

Non-polar impurities are removed by elution with a mixture of petroleum ether and ethyl acetate in a ratio of 80 : 20, and the product is then eluted with methanol. A yield of 990 mg (1.86 mmol, 96%) product is obtained.

5

$^1\text{H-NMR}$: δ 1.67 (d, $J=7.0$ Hz, 3H), 2.03 (s, 6H), 4.04 (s, 5H, Cp), 4.04 (q, $J=7.0$ Hz, 1H), 4.09-4.11 (m, 1H, Cp), 4.21-4.23 (m, 1H, Cp), 4.26 (m, 1H, Cp), 7.05-7.11 (m, 1H, Ph), 7.18-7.23 (m, 1H, Ph), 7.28-7.33 (m, 2H, Ph), 7.34-7.43 (m, 3H, Ph), 7.48-7.60 (m, 4H, Ph), 7.65-7.71 (m, 2H, Ph), 8.10-8.13 (m, 1H, Ph).

10

$^{31}\text{P-NMR}$: δ 31.67.

$[\alpha]^{20}$ (nm): -160° (589), -200.6° (578), -449.4° (546) ($c = 0.5$, CHCl_3).

15

d) Preparation of (R_C, S_P) -1-{1-[bis-(bis-3,5-trifluoromethylphenyl)phosphino]ethyl}-2-(2-diphenylphosphinylphenyl)ferrocene, L6

To a degassed solution of 1.25 g (2.35 mmol) L4 in 15 ml freshly distilled acetic acid, 1.6 g (3.5 mmol) bis-(3,5-trifluoromethylphenyl)phosphine is added. The reaction mixture is then agitated for 3 days at 100°C . The solvent is removed in a vacuum, the residue dissolved in CH_2Cl_2 and chromatographed on aluminium oxide 90. Non-polar impurities are removed by elution with hexane, and subsequent elution with a mixture of CH_2Cl_2 and methanol in a ratio of 99 : 1 yields 2.09 g (2.21 mmol, 88.9%) of product. Two diastereomers are formed in a ratio of 6 : 1 (determined by $^{31}\text{P-NMR}$), but these are not separated. The $^1\text{H-NMR}$ data are those of the principal isomer.

25

30

$^1\text{H-NMR}$: δ 1.32 (dd, $J_1=6.1$ Hz, $J_2=6.8$ Hz, 3H), 3.47 (m, 1H, Cp), 3.79 (dq, $J_1=2.8$ Hz, $J_2=7.1$ Hz, 1H), 3.96 (t, $J=2.8$ Hz, Cp), 4.08 (s, 5H, Cp), 5.03 (m, 1H, Cp), 7.1-7.15 (m, 2H, Ph), 7.20-7.30 (m, 3H, Ph), 7.42-7.53 (m, 3H, Ph), 7.56-7.75 (m, 8H, Ph), 7.86 (s, 2H, Ph), 8.24-8.28 (m, 1H, Ph).

35

³¹P-NMR: main component: δ 4.63, 30.29; secondary component: δ 4.77, 29.67.

5 e) Preparation of title compound (R_c,S_p)-1-{1-[bis-(bis-3,5-trifluoromethylphenyl)phosphino]-ethyl}-2-(2-diphenylphosphinophenyl)ferrocene, L1

To a degassed solution of 1.97 g (2.08 mmol) L6 in 20 ml THF, 9.2 ml polymethyl hydrosiloxane and 5.04 ml Ti(Oi-Propyl)₄ are added. The reaction mixture is heated under
10 reflux for 18 hours, during which the solution turns a dark violet colour. Then 15 ml hexane is added and heated for a further 2 hours under reflux. The reaction mixture is applied to an aluminium oxide column without any further preparation, and the product is eluted with a mixture of
15 petroleum ether, ethyl acetate and methanol in a ratio of 90 : 10 : 1. The yield amounts to 1.78 g (1.91 mmol, 91.8%). The two diastereomers are separated by chromatography on silica gel. A mixture of petroleum ether and CH₂Cl₂ in a ratio of 80:20 is used as the mobile phase.

20 ¹H-NMR: δ 1.32 (dd, $J_1=6.1$ Hz, $J_2=6.8$ Hz, 3H), 3.29 (s, 1H, Cp), 3.66 (dq, $J_1=J_2=7.1$ Hz, 1H), 3.86 (m, 1H, Cp), 4.02 (t, $J=2.5$ Hz, 1H, Cp), 4.16 (s, 5H, Cp), 7.06-7.11 (m, 2H, Ph), 7.13-7.19 (m, 3H, Ph), 7.25-7.29 (m, 2H, Ph), 7.38-7.50 (m, 6H, Ph), 7.59 (d, $J=4.0$ Hz, 2H, Ph), 7.80 (s, 1H, Ph), 7.90
25 (s, 1H, Ph), 7.97 (d, $J=6.1$ Hz, 2H, Ph), 8.12-8.15 (m, 1H, Ph).

³¹P-NMR: main component: δ -14.04 (d, $J=23.5$ Hz), 3.55 (d, $J=23.5$ Hz); secondary component: δ -15.19 (d, $J=28.5$ Hz), -5.16 (d, $J=28.5$ Hz).

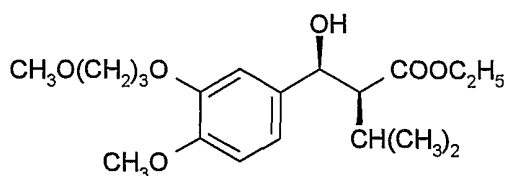
30 $[\alpha]_{20}$ (nm): -0.88 (589), -7.72 (578), -52.8 (546) (c=0.57, CHCl₃).

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B) Preparation of (R)-3-[4'-(CH₃O-3'-(CH₃O(CH₂)₃O)-phen-1-yl]-2-isopropylpropionic acid

Example B1: Preparation of

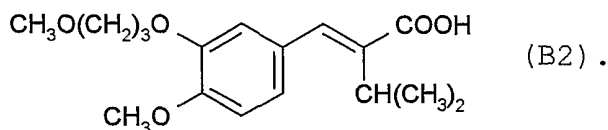
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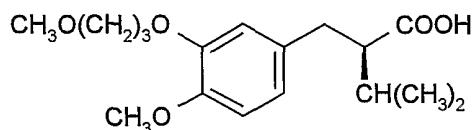
(B1)

A solution of 436 ml diisopropylamine and 2.6 l tetrahydrofuran is cooled to -20°C, and 1.234 l n-hexyl lithium (2.5 M in hexane) is added dropwise over a period of 15 minutes. A solution of 368 g ethyl isovalerate in 1.7 l tetrahydrofuran is added dropwise over a period of 15 minutes at -20°C. After a further 10 minutes, a solution of 584 g 4-methoxy-3-(3-methoxy-propoxy)benzaldehyde (EP 0 678 503) in 1.7 l tetrahydrofuran is added drop by drop and stirred for 40 minutes at -20°C. Then 2.15 l saturated aqueous ammonium chloride solution is added drop by drop and extracted with ethyl acetate (2 x 8 l). The organic phases are washed consecutively with 0.5 N hydrochloric acid (1x 4.3 l), water (1x 4.4 l) and brine (1x 4.4 l). The combined organic phases are dried over sodium sulfate (1.6 kg), filtered and boiled down in a rotary evaporator. By means of crystallization from ethyl acetate (1 l) and hexane (11 l), title compound B1 is obtained from the residue as a white solid (656 g, 72 %): ¹H-NMR (400 MHz, DMSO-d₆, δ): 0.90 - 1.04 (m, 9H), 1.97 (m, 2H), 2.32 (m, 1H), 2.58 (m, 1H), 3.28 (s, 3H), 3.50 (m, 2H), 3.74 (s, 3H), 3.82 (q, 2H), 3.98 (m, 2H), 4.57 (m, 1H), 5.30 (d, 1H), 6.75 - 6.90 (m, 3H) ppm.

30

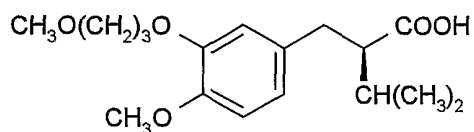
Example B2: Preparation of

5 A solution of 649 g (content: 98.3 %) B1 and 11.0 g 4-dimethylaminopyridine in 3.2 l tetrahydrofuran is cooled to 0°C, 187.2 ml acetic acid anhydride is added dropwise and the reaction mixture then stirred for 1 hour. A solution of 606 g potassium t-butyrate in 4.4 l tetrahydrofuran is added
10 drop by drop over a period of 30 minutes at -2°C to 0°C and then stirred for 2 hours at 0°C. After the addition of 2 l water and distilling off 7.6 l tetrahydrofuran at 35 °C, 6.5 l ethanol and 0.9 l 2N KOH are added to the aqueous residue. The mixture obtained is stirred for 20 hours under
15 reflux. The reaction solution is cooled off and concentrated by evaporation. At 0°C, 7.2 l t-butyl methyl ether and 3 l 2N HCl are added to the residue. The organic phase is separated off and the aqueous phase extracted again with 7.2 l t-butyl methyl ether. The organic phases are then
20 washed consecutively with 7.2 l water and 7.2 l brine. The combined organic phases are dried over magnesium sulfate (2 kg), filtered and concentrated in a rotary evaporator. By means of crystallization from diisopropyl ether (2.4 l) and hexane (2.4 l), 470 g of crude title compound is obtained
25 from the residue. After recrystallization from diisopropyl ether (2 l) and Hexan (2 l), pure title compound B2 (454.3 g, 81.8 %) is obtained: ¹H-NMR (400 MHz, DMSO-d₆, δ): 1.22 (d, 6H), 1.97 (m, 2H), 3.14 (m, 1H), 3.28 (s, 3H), 3.50 (m, 2H), 3.82 (s, 3H), 4.02 (m, 2H), 6.90 - 7.05 (m, 3H), 7.42
30 (s, 1H), COOH (exchanged) ppm.

Example B3: Preparation of

(B3).

5 In a flask with a magnetic stirrer, 5.83 mg (0.0156 mmol) [Rh(NBD)₂]BF₄ and 15.3 mg (0.0164 mmol) L1 are placed under an atmosphere of argon through repeated evacuation and purging with argon. Then 20 ml degassed methanol is added and stirred for 15 minutes, before 24 g (0.078 mol) B2 and
10 140 ml degassed methanol are introduced into a 250 ml flask fitted with a side stopcock and flushed with argon. With gentle heating, agitation is continued until a homogeneous solution is formed. The solution is forced under pressure via a steel capillary tube into a 300 ml steel autoclave
15 under cover of argon. In 3 purge cycles (argon 20 bar / hydrogen 20 bar) the hydrogen pressure is eventually increased to 50 bar. Hydrogenation is started by switching on the stirrer and carried out at room temperature. The reaction takes place via hydrogen consumption (fall of
20 pressure in the reservoir of hydrogen). After a reaction time of 8 hours, a full conversion is measured by HPLC (method 1). The reaction mixture is concentrated by evaporation and crude title compound B3 obtained as a slightly yellowish oil (24 g , quantitative): HPLC (method
25 2) optical yield > 95% R-Isomer; ¹H-NMR (400 MHz, CDCl₃, δ): 1.03 (m, 6H), 1.95 (m, 1H), 2.07 (m, 2H), 2.48 (m, 1H), 2.81 (m, 2H), 3.40 (s, 3H), 3.60 (m, 2H), 3.85 (s, 3H), 4.10 (m, 2H), 6.70 - 6.80 (m, 3H) ppm.

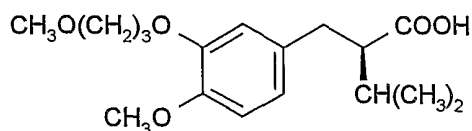
Example B4: Preparation of

(B3).

5 In a flask with a magnetic stirrer, 1.50 mg (0.0024 mmol) [Rh(NBD)(OOCCH₃)₂] and 4.75 mg (0.0051 mmol) L1 are placed under an atmosphere of argon through repeated evacuation and purging with argon. Then 5 ml degassed methanol is added and stirred for 15 minutes, before 3.0 g (9.73 mmol) B2 and
10 15 ml degassed methanol are introduced into a 50 ml flask fitted with a side stopcock and flushed with argon. With gentle heating, agitation is continued until a homogeneous solution is formed. The solution is forced under pressure via a steel capillary tube into a 50 ml steel autoclave
15 under cover of argon. In 3 purge cycles (argon 20 bar / hydrogen 20 bar) the hydrogen pressure is eventually increased to 20 bar. Hydrogenation is started by switching on the stirrer and carried out at room temperature. The reaction takes place via hydrogen consumption (fall of
20 pressure in the reservoir of hydrogen). After a reaction time of 20 hours, a full conversion is measured. The optical yield amounts to >95% (R)-compound.

Example B5: Preparation of

25



(B3).

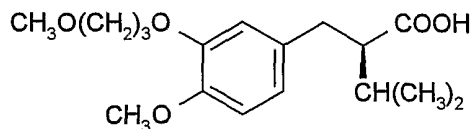
In a flask with a magnetic stirrer, 1.50 mg (0.0024 mmol) [Rh(NBD)(OOCCH₃)₂] and 4.75 mg (0.0051 mmol) L1 are placed
30 under an atmosphere of argon through repeated evacuation and purging with argon. Then 5 ml degassed toluene is added and

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stirred for 15 minutes, before 150 mg (0.486 mmol) B2 and 15 ml degassed toluene are introduced into a 50 ml flask fitted with a side stopcock and flushed with argon. With gentle heating, agitation is continued until a homogeneous solution is formed. The solution is forced under pressure via a steel capillary tube into a 50 ml steel autoclave under cover of argon. In 3 purge cycles (argon 20 bar / hydrogen 20 bar) the hydrogen pressure is eventually increased to 100 bar. Hydrogenation is started by switching on the stirrer and carried out at room temperature. The reaction takes place via hydrogen consumption (fall of pressure in the reservoir of hydrogen). After a reaction time of 72 hours, a full conversion is measured. The optical yield amounts to 95% (R)-compound.

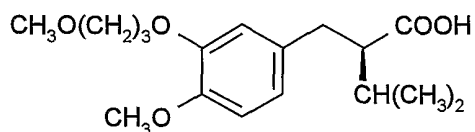
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Example B6: Preparation of



(B3) .

The procedure is analogous to that described under Example B5. For preparation of the catalyst, 1.5 mg (0.004 mmol) $[\text{Rh}(\text{NBD})\text{Cl}]_2$ and 3.86 mg (0.004 mmol) (2S,4S)-N-(t-butyl-oxycarbonyl)-4-(dicyclohexylphosphino)-2-(diphenylphosphino-methyl)pyrrolidine are used. 0.205 g (0.65 mmol) of educt B2 is hydrogenated in 10 ml toluene. After 20 hours at 60 bar hydrogen and 50°C , the reaction is stopped and the conversion and enantiomeric purity are determined. The conversion amounts to 98.9% and the optical yield is 80% ee.

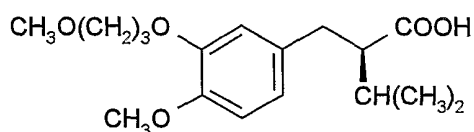
Example B7: Preparation of

(B3).

5 The procedure is analogous to that described under Example B6, and 2.2 mg (0.01 mmol) [Rh(NBD)Cl]₂ and 9.6 mg (0.01 mmol) (R)-1,1'-(di-3,4,5-methoxyphenylphosphino)-6,6'-di-methoxybiphenyl are used for the preparation of the catalyst. 0.304 g (0.99 mmol) of educt B2 is hydrogenated in
10 10 ml toluene. After a reaction time of 18 hours at 60 bar hydrogen and 50°C, the conversion amounts to 93.5% and the optical yield is 73.5% ee.

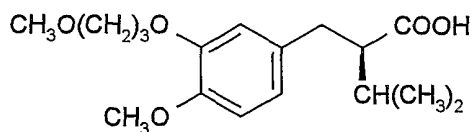
Example B8: Preparation of

15



(B3).

The procedure is analogous to that described under Example B6, and 3.0 mg (0.01 mmol) [Rh(NBD)Cl]₂ and 6.19 mg (0.01
20 mmol) (R)-(S)-1-[1-[bis-(bis-t-butylphenyl)phosphino]ethyl]-2-(diphenylphosphino)ferrocene are used for the preparation of the catalyst. 0.227 g (0.74 mmol) of educt B2 is hydrogenated in 10 ml toluene. After a reaction time of 90 hours at 60 bar hydrogen and 30°C, the conversion amounts to
25 98.6 % and the optical yield is 49% ee.

Example B9: Preparation of

(B3)

5 In a 50 l steel autoclave are placed 25 l of methanol under an atmosphere of argon through repeated evacuation and purging with argon. Then 12.00 kg B2 are added under argon. The suspension is heated under argon (2 bar) to 50°C , cooled to 35°C and degassed. 2.634 g (6.486 mmol)

10 $[\text{Rh}(\text{COD})_2]\text{BF}_4$ and 6.337 g (6.810 mmol) L1 are placed in a flask with magnetic stirrer under an atmosphere of argon through repeated evacuation and purging with argon. Then 700 ml degassed methanol are added and stirred for 45 minutes. This catalyst solution is forced under argon

15 atmosphere through a steel capillary in the autoclave. In 3 purge cycles (argon 20 bar / hydrogen 20 bar) the hydrogen pressure is increased to 50 bar. The hydrogenation is started by switching on the stirrer and carried out at 35°C . The reaction is monitored via hydrogen consumption

20 (decrease of pressure in the reservoir of hydrogen). After a reaction time of 21 hours (15 hours take-up of hydrogen) a full conversion is measured according to HPLC (methode 1). The reaction mixture is concentrated by evaporation and crude title compound B3 obtained as slightly yellowish oil

25 (12,08 kg, quantitative): HPLC (Methode 2) optical yield > 95% R-Isomer.

Determination of conversion and optical yield:

For the HPLC analysis, B1 and B2 are derivatized

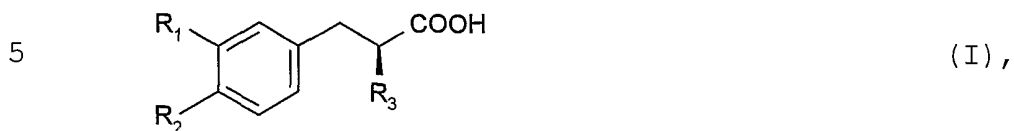
30 (preparation of the respective methyl esters): a sample of the residue in diethyl ether is mixed with excess diazo methane in diethyl ether. The solvent is then evaporated off, and the residue obtained is the corresponding methyl ester.

Method 1 (determination of conversion): column HP Hypersil BDS-C 18 125 x 4mm ; acetonitrile and water 5% to 100%; 40 minutes flow: 0.8 ml.

- 5 Method 2 (determination of optical yield): column: Daicel OJ-R 0.45 x 15 cm; solvent 30% acetonitrile and 70% water.

What is claimed is:

1. A process for the preparation of compounds of formula I,



wherein R_1 and R_2 are, independently of one another, H, C_1 - C_6 alkyl, C_1 - C_6 halogenalkyl, C_1 - C_6 alkoxy, C_1 - C_6 alkoxy- C_1 - C_6 alkyl, or C_1 - C_6 alkoxy- C_1 - C_6 alkyloxy, and R_3 is C_1 - C_6 alkyl
 10 comprising

a) the reaction of a compound of formula II



15 wherein R_1 and R_2 are as defined hereinbefore, with a compound of formula III,



wherein R_3 is as defined hereinbefore, to form a compound of formula IV,

20



wherein R_7 is C_1 - C_{12} alkyl, C_3 - C_8 cycloalkyl, phenyl or benzyl,
 b) the isolation of the crystalline compound of formula IV,
 25 the conversion of the OH group to a leaving group, and the

reaction of a compound containing a leaving group in the presence of a strong base to form a compound of formula V,



5

c) the hydrolysis of the carbonic esters of formula V to form the carboxylic acid of formula VI,



10

d) the hydrogenation of the carboxylic acid of formula VI in the presence of hydrogen and catalytic quantities of a metal complex as asymmetric hydrogenation catalyst, comprising metals from the group of ruthenium, rhodium and iridium, to which the chiral bidentate ligands are bonded, to form a compound of formula I.

2. A process according to claim 1, comprising R₁ as methoxy-C₁-C₄alkyloxy or ethoxy-C₁-C₄alkyloxy and R₂ as methoxy or ethoxy.

3. A process according to claim 2, comprising R₁ as 1-methoxyprop-3-yloxy and R₂ as methoxy.

4. A process according to claim 1, comprising R₃ as a linear or branched C₁-C₄alkyl.

5. A process according to claim 4, comprising R₃ as isopropyl.

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6. A process according to claim 1, comprising R_1 as 1-methoxy-n-propyloxy, R_2 as methoxy, and R_3 as isopropyl.
7. A process according to claim 1, comprising the processing
5 of step a) at low temperatures in the presence of a secondary lithium amide.
8. A process according to claim 1, comprising in step b) first acylation of the hydroxyl group and then elimination
10 at low temperatures in the presence of an alkali metal alcoholate in the reaction mixture of the acylation process.
9. A process according to claim 1, comprising step c) being carried out in the reaction mixture of step b).
- 15 10. A process according to claim 1, comprising step d) being carried out in the presence of metal complexes of formula VII or VIIa as hydrogenation catalysts,



wherein

Me is rhodium;

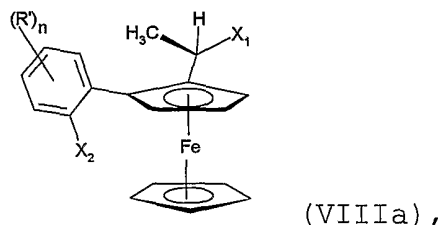
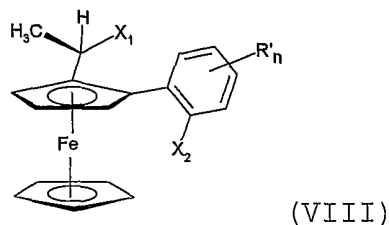
Y stands for two olefins or one diene;

25 Z is Cl, Br or I;

E^- is the anion of an oxygen acid or a complex acid; and

L is a chiral ligand from the ditertiary diphosphine group, in which the phosphine groups are bonded to a C_2-C_4 chain of the diphosphine backbone chain, and the diphosphine forms
30 a five to seven-member ring together with the rhodium atom.

11. A process according to claim 10, comprising L as formula VIII or VIIIa,



5

wherein

n is 0 or an integer from 1 to 4 and R' represents the same or different substituents from the C₁-C₄alkyl, -CF₃ and C₁-C₄alkoxy group; and

10 X₁ and X₂ are, independently of one another, secondary phosphino.

12. A process according to claim 11, comprising the X₁ and X₂ groups being the same or different and corresponding to
 15 formula -PR₈R₉, wherein R₈ and R₉ are the same or different and are branched C₃-C₈alkyl, C₃-C₈cycloalkyl, or unsubstituted phenyl or phenyl substituted with one to three C₁-C₄alkyl, C₁-C₄alkoxy, or -CF₃.

20 13. A process according to claim 11, comprising in formulae VIII and VIIIa n as 0, and X₁ and X₂ as a PR₈R₉ group, wherein R₈ and R₉ are in each case cyclohexyl, phenyl or phenyl substituted with 1 or 2 methyl, methoxy or CF₃.

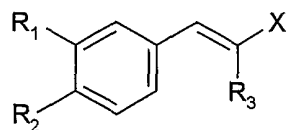
25 14. A process according to claim 1, comprising step d) being carried out at temperatures of -20 to 150 °C.

15. A process according to claim 1, comprising step d) being carried out under positive pressure.

30

16. A process according to claim 1, comprising pressure conditions at 10⁵ to 2x10⁷ Pa (Pascal).

17. Compounds of formula IX,



(IX),

wherein

R₁ and R₂, independently of one another, are H, C₁-C₆alkyl,
5 C₁-C₆halogenalkyl, C₁-C₆alkoxy, C₁-C₆alkoxy-C₁-C₆alkyl, or
C₁-C₆alkoxy-C₁-C₆alkyloxy, R₃ is C₁-C₆alkyl, and X is the -COOH
group.

18. Compounds according to claim 17, comprising R₁ as
10 methoxy-C₁-C₄alkyloxy or ethoxy-C₁-C₄alkyloxy and R₂ as
methoxy or ethoxy, R₃ as C₁-C₄alkyl, and X as the -COOH group.

19. Compounds according to claim 18, comprising R₁ as
1-methoxy-n-propyloxy and R₂ as methoxy, R₃ as isopropyl, and
15 X as the -COOH group.

INTERNATIONAL SEARCH REPORT

Int. Application No

PCT/CH 01/00397

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07C67/293 C07C51/36 C07C59/66

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 C07C

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

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	MIYANO, SOTARO ET AL: "Axially dissymmetric bis(aminophosphine)s derived from 2,2'-diamino-1,1'-binaphthyl. Synthesis and application to rhodium (I)-catalyzed asymmetric hydrogenations" BULL. CHEM. SOC. JPN. (1984), 57(8), 2171-6 , XP002183784	17
Y	abstract table 2 ----- -/--	1,9,10, 14-16

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *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

30 November 2001

Date of mailing of the international search report

14/12/2001

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International Application No

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