



Diphosphine

The present invention relates to a diphosphine of the formula

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where

10 R^1 is a divalent radical which together with the phosphorus atom to which it is linked forms an unsubstituted or substituted 2-phosphatricyclo[3.3.1.1{3,7}]decyl group or a derivative thereof in which one or more of the carbon atoms are replaced by heteroatoms,

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R^2 , R^3 are each, independently of one another, a monovalent radical having from 1 to 20 atoms or together form a divalent radical having from 2 to 20 atoms,

20 n is 4 or 5,

and mixtures of such diphosphines.

The invention further relates to a system which is suitable as
25 carbonylation catalyst and comprises palladium or a palladium compound and such a diphosphine, and also to a process for the carbonylation of a conjugated diene in the presence of such a system.

30 Alkenecarboxylic acids and their derivatives are important intermediates in organic synthesis, for example the synthesis of active compounds and polymers. Thus, for example, pentenoic acid or its derivatives, e.g. esters, can be converted by further functionalization into adipic acid derivatives or 6-aminocaproic
35 acid derivatives. Adipic acid and 6-aminocaproic acid are important starting compounds for the preparation of industrially important polymers, in particular polyamides.

Processes for preparing alkenecarboxylic acid derivatives by
40 carbonylation of conjugated dienes in the presence of a hydroxyl-containing compound in the liquid phase in the presence of a system which is suitable as carbonylation catalyst and comprises a palladium compound and a multidentate organic phosphorus ligand are known, for example from EP 273 489 A1 or EP
45 495 547 A2.

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A disadvantage of these processes is that palladium deposits as metal and is thus removed from the catalytically active system. In addition, the metallic palladium is technically difficult to recover from the reaction mixture.

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It is known from WO 98/42717 that in the carbonylation of olefins in the presence of catalyst systems comprising palladium or a palladium compound and a multidentate organic phosphorus ligand, the activity can be increased by using a diphosphine of the
10 formula $R^1>P-R^2-PR^3R^4$ in which R^1 is an unsubstituted or substituted 2-phosphatricyclo[3.3.1.1{3,7}]decyl group, R^2 is a bridging group, R^3 and R^4 are each, independently of one another, a monovalent radical having from 1 to 20 carbon atoms or together form a divalent radical having from 2 to 20 carbon atoms as
15 multidentate organic phosphorus ligand. Specific examples mentioned for R^2 are only the 1,2-ethane and 1,3-propane groups. According to the examples given in WO 98/42717, such diphosphines appear to lead to high activities or selectivities in the carbonylation of ethylene, propene, α -C₁₄-olefins and methyl
20 3-pentenoate. However, if such systems are employed in the case of conjugated dienes such as 1,3-butadiene, the selectivity is drastically worse than in the abovementioned EP 273 489 A1.

It is an object of the present invention to provide a system
25 which is suitable as carbonylation catalyst and comprises a palladium compound and a multidentate organic phosphorus ligand and makes it possible to carry out the carbonylation of a conjugated diene in the presence of a hydroxyl-containing compound in the liquid phase in good yields, with high
30 selectivities, with reduced deposition of palladium and in a technically simple and economical manner.

We have found that this object is achieved by the diphosphine defined at the outset, a system which is suitable as
35 carbonylation catalyst and comprises palladium or a palladium compound and such a diphosphine, and a process for the carbonylation of a conjugated diene in the presence of such a system.

40 According to the present invention, R^1 is a divalent radical which together with the phosphorus atom to which it is linked forms a 2-phosphatricyclo[3.3.1.1{3,7}]decyl group or a derivative thereof in which one or more of the carbon atoms are replaced by heteroatoms. To aid understanding of the present invention,
45 mention may be made of the fact that tricyclo[3.3.1.1{3,7}]decane

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is the systematic name for a compound which is generally known as "adamantane".

In an advantageous embodiment, R^1 together with the phosphorus atom to which it is linked can form an unsubstituted 2-phosphatricyclo[3.3.1.1{3,7}]decyl group.

In an advantageous embodiment, R^1 together with the phosphorus atom to which it is linked can form a substituted 2-phosphatricyclo[3.3.1.1{3,7}]decyl group, preferably a 2-phosphatricyclo[3.3.1.1{3,7}]decyl group substituted in one or more of the positions 1, 3, 5 and 7 by a monovalent radical R^4 . R^4 is advantageously a radical having from 1 to 20 atoms, for example methyl, trifluoromethyl, ethoxy, phenyl or 4-dodecylphenyl. In a particularly preferred embodiment, each of the positions 1, 3, 5 and 7 is substituted by radicals R^4 , preferably identical radicals R^4 .

In the skeleton of R^1 , one or more carbon atoms can advantageously be replaced by heteroatoms, in particular by oxygen or sulfur, preferably in the positions 6, 9 and 10.

In a preferred embodiment, R^1 together with the phosphorus atom to which it is linked is a 2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxodamantyl group.

According to the present invention, each of the groups R^2 and R^3 can be, independently of one another, a monovalent radical having from 1 to 20 atoms, e.g. alkyl such as methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, s-butyl, t-butyl, cyclohexyl, aryl such as phenyl, o,o-di(t-butoxy)phenyl, heteroaryl such as pyridyl, unsubstituted or substituted heterohydrocarbyl such as trimethylsilyl or alkoxy such as methoxy, ethoxy, n-propoxy, i-propoxy, n-butoxy, i-butoxy, s-butoxy, t-butoxy.

R^2 and R^3 may also together form a divalent radical having from 2 to 20 atoms, e.g. 1,5-pentylene, 1,6-hexylene, 1,3-cyclooctylene, 1,4-cyclooctylene. Preference is given to R^2 and R^3 together with the phosphorus atom to which they are linked together forming a 2-phosphatricyclo[3.3.1.1{3,7}]decyl group, with the embodiments described above for R^1 . In a particularly preferred embodiment, R^2 and R^3 together form a group identical to R^1 .

According to the present invention, n is 4 or 5.

In a preferred embodiment, n is 4.

In another preferred embodiment, n is 5.

Particularly preferred diphosphines are diphosphines selected from the group consisting of

- 5 1,4-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane,
 1,4-P,P'-diperfluoro(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane,
 1,4-P,P'-di(2-phospha-1,3,5,7-tetra(trifluoromethyl)-6,9,10-
 10 trioxatricyclo[3.3.1.1{3,7}]decyl)butane, preferably
 1,4-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane.

Further particularly preferred diphosphines are diphosphines

- 15 selected from the group consisting of
 1,5-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)pentane,
 1,5-P,P'-diperfluoro(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)pentane,
 20 1,5-P,P'-di(2-phospha-1,3,5,7-tetra(trifluoromethyl)-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)pentane, preferably
 1,5-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)pentane.

- 25 The preparation can be carried out in a manner similar to the process described in WO 98/42717, for example for
 1,2-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)ethane or
 1,3-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-

- 30 trioxatricyclo[3.3.1.1{3,7}]decyl)propane. Thus, for example,
 1,4-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane can be obtained by reacting 2,4-pentanedione with 1,4-diphosphinobutane or
 1,5-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-

- 35 trioxatricyclo[3.3.1.1{3,7}]decyl)pentane can be obtained by reacting 2,4-pentanedione with 1,5-diphosphinopentane.

Unsymmetrical diphosphines can be prepared, for example, by reaction of a 1,4-diphosphinobutane or 1,5-diphosphinopentane having one primary phosphino group and one tertiary phosphino

- 40 group. Alternatively, they can be prepared by reaction of a secondary 2-phosphatricyclo[3.3.1.1{3,7}]decane with another secondary monophosphine or by other known methods. Substituted diphosphines can be prepared, for example, by using substituted 2,4-pentanediones, e.g. perfluoro-2,4-pentanedione or
 45 1,1,1,5,5,5-hexafluoro-2,4-pentanedione, in the reaction with 1,4-diphosphinobutane or 1,5-diphosphinopentane.

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According to the present invention, the system which is suitable as carbonylation catalyst comprises

(I) palladium or a palladium compound and

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(II) one of the above-described diphosphines.

The component (II) can be a particular diphosphine of the present invention or a mixture of such diphosphines.

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It is likewise possible to use a particular diphosphine of the present invention together with another diphosphine or a mixture of such other diphosphines, or to use a mixture of diphosphines of the present invention together with another diphosphine or a

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mixture of such other diphosphines.

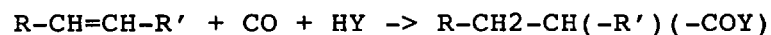
For the purposes of the present invention, the term "diphosphine" is used both for a particular diphosphine and for a mixture of diphosphines.

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The system of the present invention can be used as carbonylation catalyst as described, for example, in EP 273 489 A1, EP 495 547 A2 or WO 98/42717, particularly in the various reactions in which carbon monoxide is added onto an unsaturated compound, preferably

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onto an olefinic double bond. Such reactions can be schematically represented by



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Y is H, OH, OR'', NR''''R''''', and

R, R', R'', R''', R'''' are each an organic radical.

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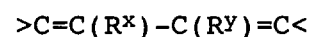
The system of the present invention can accordingly be used, for example, in hydroformylation reactions, hydrocarboxylation reactions, hydroesterification reactions, hydroamidation reactions. The system of the present invention has been found to

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be particularly advantageous in hydrocarboxylation reactions and hydroesterification reactions.

Advantageous unsaturated compounds are conjugated dienes, i.e. organic compounds comprising the structure

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where R^X , R^Y are each a monovalent radical,

in particular noncyclic compounds such as 1,3-butadiene,
2-methyl-1,3-butadiene, 1,3-pentadiene, 1,3-hexadiene,
5 2,4-hexadiene, 2,3-dimethyl-1,3-butadiene, preferably
1,3-butadiene.

In the unsaturated compounds, preferably conjugated dienes, one
or more hydrogen atoms may be replaced by other atoms, e.g.
10 halogen atoms, or groups of atoms, e.g. hydroxyl groups, cyano
groups, methoxy or ethoxy groups, amino groups such as
dimethylamino or diethylamino groups, carboxyl groups or ester
groups.

15 In a particularly preferred embodiment, such conjugated dienes
can be hydrocarboxylated to give alkenecarboxylic acids, for
example pentenoic acids such as cis-2-pentenoic acid,
trans-2-pentenoic acid, cis-3-pentenoic acid, trans-3-pentenoic
acid, and 4-pentenoic acid from 1,3-butadiene.

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In another particularly preferred embodiment, such conjugated
dienes can be hydroesterified to give alkenecarboxylic esters,
for example pentenoic esters such as cis-2-pentenoic esters,
trans-2-pentenoic esters, cis-3-pentenoic esters,

25 trans-3-pentenoic esters and 4-pentenoic esters from
1,3-butadiene.

In such carbonylation reactions, further components such as
coreactants or liquid diluents can be used. Likewise, anions can
30 be used as counterion to the palladium cation used in the case of
palladium compounds. Examples include anions which are the
conjugate base of an acid having a pK_a measured in water at 18°C
of less than 6, preferably less than 4. Such anions do not
coordinate or coordinate only weakly to palladium, i.e. so that
35 there is no interaction or only a weak interaction between the
anion and the palladium cation. Catalysts containing such an
anion generally display a good activity.

Suitable anions include anions derived from Brönstedt acids such
40 as phosphoric acid or sulfuric acid, preferably from carboxylic
acids, halogenated carboxylic acids or sulfonic acids such as
methanesulfonic acid, trifluoromethanesulfonic acid or
p-toluenesulfonic acid. Preference is given to anions derived
from carboxylic acids, e.g. 2,6-dichlorobenzoic acid,
45 2,6-dimethoxybenzoic acid and 2,4,6-trimethylbenzoic acid.

- It is likewise possible to use complex anions, for example anions obtained from the combination of a Lewis acid, e.g. BF_3 , $\text{B}(\text{C}_6\text{F}_5)_3$, AlCl_3 , SnF_2 , $\text{Sn}(\text{CF}_3\text{SO}_3)_2$, SnCl_2 or GeCl_2 , with a protic acid, preferably a protic acid having a pK_a of less than 5, e.g. a
- 5 sulfonic acid such as $\text{CF}_3\text{SO}_3\text{H}$, $\text{CH}_3\text{SO}_3\text{H}$ or a hydrohalic acid such as HF , HCl , or the combination of a Lewis acid with an alcohol. Examples of such complex anions are BF_4^- , SnCl_3^- , $[\text{SnCl}_2 \cdot \text{CF}_3\text{SO}_3]^-$ and PF_6^- .
- 10 The carbonylations can advantageously be carried out at from 30 to 200°C , preferably from 50 to 180°C . The pressure can vary within a wide range. The pressure can advantageously be in a range from 1 to 200 bar, in particular in the range from 5 to 90 bar. Particular preference is given to a pressure which results
- 15 in the carbonylation taking place in the liquid phase.

Carbon monoxide can advantageously be used in a molar excess over "HY".

- 20 The compound "HY" can advantageously be used in a molar ratio to the unsaturated compound of from 10:1 to 1:10, preferably from 5:1 to 1:5, in particular from 2:1 to 1:2.

- The amount of catalyst is not critical per se and can vary within
- 25 a wide range. In general, an amount in the range from 10^{-8} to 10^{-1} mol, preferably from 10^{-7} to 10^{-2} mol, of palladium per mol of unsaturated compound has been found to be advantageous.

- In the preparation of the system of the present invention, the
- 30 molar amount of component (II) can generally be somewhat higher than the molar amount of palladium or palladium cation.

- A molar ratio of ligand to palladium in the range from 0.5 to 10 has been found to be particularly advantageous.
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In the case of particularly active systems, use of diphosphine and palladium in equimolar amounts is possible.

- The molar ratio of ligand to palladium is advantageously in the
- 40 range from 1 to 3, preferably from 1 to 2.

If oxygen is present, a slightly larger amount can be advantageous.

- 45 The amount of anion source can advantageously be in the range from 0.5 to 200 mol, preferably from 1 to 80 mol, per mol of palladium.

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In the process of the present invention, starting material and product of the carbonylation reaction can act as liquid diluent. In such a case, the use of a further liquid diluent may be able to be dispensed with. It may be advantageous to use a further liquid diluent such as a hydrocarbon, for example an alkane such as a branched or unbranched alkane, or an ether such as 2,5,8-trioxanonane (diglyme), diethyl ether, diphenyl ether or anisole, a sulfone such as sulfolane, or an aromatic hydrocarbon such as toluene.

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Suitable coreactants in the carbonylation include compounds having a nucleophilic center and a mobile hydrogen atom.

Preferred nucleophilic compounds are molecular hydrogen, water, alcohols such as monoalcohols, for example, methanol, ethanol, i-propanol, n-propanol, 1-n-butanol, s-butanol, t-butanol, i-butanol, polyalcohols such as ethylene glycol, 1,2-propanediol, 1,3-propanediol, 1,4-butanediol, glycerol, thiols, primary and secondary amines, polyamines, amides and polyamides, for example diethylamine, N,N-dimethylethylenediamine, aromatic alcohols, carboxylic acids such as acetic acid, pivalic acid and propionic acid, in particular molecular hydrogen, monoalcohols having from 1 to 6 carbon atoms, dialcohols having from 2 to 6 carbon atoms, particularly preferably monoalcohols having from 1 to 6 carbon atoms, e.g. methanol, ethanol, i-propanol, n-propanol, 1-n-butanol, s-butanol, t-butanol, i-butanol.

The use of such monoalcohols makes it possible to prepare valuable intermediates such as methyl cis-2-pentenoate, methyl trans-2-pentenoate, methyl cis-3-pentenoate, methyl trans-3-pentenoate, methyl 4-pentenoate. These intermediates can be used, for example, as mentioned above in the preparation of polymers, in particular polyamide 6 and polyamide 66.

Another group of preferred coreactants encompasses alkylphenols, in which one or more of the alkyl groups linked to the phenol part preferably contain(s) up to 30 carbon atoms, preferably from 6 to 22 carbon atoms. The carbonylation described leads in this case to alkylphenyl esters which can be employed as synthetic lubricants, for example in industrial applications or in the automobile sector.

The invention is illustrated by the nonlimiting examples below.

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Examples

Example 1

- 5 107 g of diphenyl ether, 20 ml of methanol, 20 ml of butadiene, 0.5 mmol of palladium acetate, 1 mmol of 1,4-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane and 30 mmol of 2,4,6-trimethylbenzoic acid were placed under a nitrogen
- 10 atmosphere in a 270 ml autoclave fitted with a magnetic stirrer. The autoclave was closed, pressurized with 40 bar of carbon monoxide and heated to 150°C. After this temperature had been reached, the pressure was kept constant for 10 hours by
- 15 introduction of further carbon monoxide. After the autoclave had been cooled and vented, the reaction solution was taken out. This was a clear yellow solution without a palladium precipitate. The conversion and selectivity to the various products was measured by GC analysis of the solution.
- 20 The conversion of butadiene was 82%, and the selectivity to methyl pentenoate was 80.5%.

Comparative Example A

- 25 Example 1 was repeated using 1 mmol of 1,3-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)propane instead of 1 mmol of 1,4-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane. The reaction solution
- 30 was clear yellow without a palladium precipitate.

The conversion of butadiene was 81%, and the selectivity to methyl pentenoate was 22.8%.

35 Comparative Example B

- Example 1 was repeated using 1 mmol of 1,4-bis(diphenylphosphino)butane in place of 1 mmol of 1,4-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane. After the reaction, the
- 40 reaction mixture contained a black palladium precipitate. The conversion of butadiene was 97%, and the selectivity to methyl pentenoate was 68.8%.

45 Comparative Example C

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Example 1 was repeated using 1 mmol of 1,3-bis(dicyclohexylphosphino)ethane in place of 1 mmol of 1,4-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane and using 5 mmol of 2,4,6-trimethylbenzoic acid instead of 30 mmol. The reaction solution after the reaction was clear yellow without a palladium precipitate.

The conversion of butadiene was 55%, and the selectivity to methyl pentenoate was 8.9 %.

Comparative Example D

Example 1 was repeated using 1 mmol of 1,3-bis(1,5-cyclooctylenephosphino)propane in place of 1 mmol of 1,4-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane. After the reaction, the reaction mixture contained a black palladium precipitate.

The conversion of butadiene was 73%, and the selectivity to methyl pentenoate was 64.5%.

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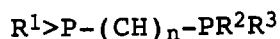
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We claim:

1. A diphosphine of the formula

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where

- 10 R^1 is a divalent radical which together with the phosphorus atom to which it is linked forms an unsubstituted or substituted 2-phosphatricyclo[3.3.1.1{3,7}]decyl group or a derivative thereof in which one or more of the carbon atoms are replaced by heteroatoms,

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R^2 , R^3 are each, independently of one another, a monovalent radical having from 1 to 20 atoms or together form a divalent radical having from 2 to 20 atoms,

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n is 4 or 5,

or a mixture of such diphosphines.

2. A diphosphine as claimed in claim 1, wherein R^1 is
25 substituted in one or more of the positions 1, 3, 5 and 7 by a monovalent radical R^4 having from 1 to 20 atoms.

3. A diphosphine as claimed in claim 2, wherein R^4 is a radical
30 selected from the group consisting of methyl, trifluoromethyl, ethoxy, phenyl and 4-dodecylphenyl.

4. A diphosphine as claimed in any of claims 1 to 3, wherein R^1
is substituted in each of the positions 1, 3, 5, and 7 by preferably identical radicals R^4 .

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5. A diphosphine as claimed in any of claims 1 to 4, wherein R^1 contains oxygen or sulfur in the skeleton, preferably in the positions 6, 9 and 10.

- 40 6. A diphosphine as claimed in any of claims 1 to 5, wherein R^1 is a 2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxoadamantyl group.

7. A diphosphine as claimed in any of claims 1 to 6, wherein R^2
45 and R^3 together form a group R^1 as set forth in any of claims 1 to 6, preferably a group identical to R^1 .

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8. A diphosphine as claimed in any of claims 1 to 7 in which n is 4.
9. A diphosphine as claimed in any of claims 1 to 7 in which n is 5.
10. A diphosphine as claimed in claim 1 selected from the group consisting of
1,4-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane,
1,4-P,P'-diperfluoro(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane,
1,4-P,P'-di(2-phospha-1,3,5,7-tetra(trifluoromethyl)-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane,
preferably
1,4-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)butane.
11. A diphosphine as claimed in claim 1 selected from the group consisting of
1,5-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)pentane,
1,5-P,P'-diperfluoro(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)pentane,
1,5-P,P'-di(2-phospha-1,3,5,7-tetra(trifluoromethyl)-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)pentane,
preferably 1,5-P,P'-di(2-phospha-1,3,5,7-tetramethyl-6,9,10-trioxatricyclo[3.3.1.1{3,7}]decyl)pentane.
12. A system which is suitable as carbonylation catalyst and comprises
(I) palladium or a palladium compound and
(II) a diphosphine as claimed in any of claims 1 to 11.
13. A process for preparing an alkenecarboxylic acid or a derivative thereof by carbonylation of a conjugated diene in the presence of a hydroxyl-containing compound in the liquid phase, wherein the carbonylation is carried out in the presence of a system which is suitable as carbonylation catalyst and comprises
(I) palladium or a palladium compound and
(II) a diphosphine as claimed in any of claims 1 to 11.

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14. A process as claimed in claim 13, wherein 1,3-butadiene is used as conjugated diene to give pentenoic acid or a derivative thereof, preferably a pentenoic ester.

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