

Methoxycarbonylation with formic acid as CO source

The invention relates to a process for methoxycarbonylation with formic acid as the CO source.

- 5 The methoxycarbonylation of alkenes is a process of increasing importance. In the classical methoxycarbonylation, an olefin is reacted with CO and MeOH in the presence of a catalyst comprising a ligand and a metal:



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Here, CO is introduced into the reaction vessel as a gas.

It is an object of the invention to provide a process that employs a CO source other than CO gas which is introduced into the reaction vessel. The process should achieve a high yield of methyl ester.

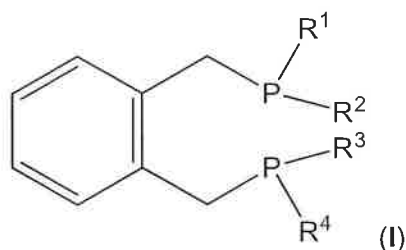
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The object is achieved by a process according to Claim 1.

Process comprising the process steps of:

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- a) addition of an olefin;
- b) addition of a compound comprising Pd, wherein the Pd is capable of forming a complex;
- c) addition of a compound of general formula (I):



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wherein R^1, R^2, R^3, R^4 are each independently selected from: -H, -(C₁-C₁₂)-alkyl, -O-(C₁-C₁₂)-alkyl, -(C₄-C₁₄)-aryl, -O-(C₄-C₁₄)-aryl, cycloalkyl, -(C₁-C₁₂)-heteroalkyl, -O-(C₁-C₁₂)-heteroalkyl, - (C₃-

C₁₄)-heteroaryl, -O-(C₃-C₁₄)-heteroaryl, -COO-alkyl, -COO-aryl, -C-O-alkyl, -C-O-aryl, NH₂, halogen and the residues are also capable of forming a larger condensed ring;

wherein the recited alkyl groups, aryl groups, cycloalkyl, heteroalkyl groups, heteroaryl groups may be substituted as follows:

- 5 -(C₁-C₁₂)-alkyl, -O-(C₁-C₁₂)-alkyl, halogen;

and at least one of the radicals R¹, R², R³, R⁴ does not represent phenyl;

d) addition of MeOH and HCOOH,

wherein the MeOH / HCOOH ratio based on the employed volume is in the range from 1.55 : 0.45 to 1.1 : 0.9;

- 10 e) heating the reaction mixture to convert the olefin into the methyl ester.

In one variant of the process, no CO gas is supplied to the reaction mixture.

In one variant of the process, HCOOH serves as the only CO source for the reaction.

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In one variant of the process, the compound in process step b) is selected from:

Pd(acac)₂, PdCl₂, Pd(dba)₃*CH₃Cl (dba = dibenzylideneacetone), Pd(OAc)₂, Pd(TFA)₂, Pd(CH₃CN)Cl₂.

- 20 In one variant of the process, the compound in process step b) is Pd(OAc)₂.

In one variant of the process, the process comprises the additional process step f):

f) addition of an acid.

- 25 In one variant of the process, the acid is selected from: H₂SO₄, CH₃SO₃H, CF₃SO₃H, PTSA (p-toluenesulfonic acid).

In one variant of the process, the acid is PTSA (p-toluenesulfonic acid).

- 30 In one variant of the process, the MeOH / HCOOH ratio based on the employed volume is in the range from 1.5 : 0.5 to 1.2 : 0.8.

In one variant of the process, R¹, R², R³, R⁴ are each independently selected from: -(C₁-C₁₂)-alkyl, -O-(C₁-C₁₂)-alkyl, -(C₄-C₁₄)-aryl, -O-(C₄-C₁₄)-aryl, cycloalkyl, -(C₁-C₁₂)-heteroalkyl, -O-(C₁-C₁₂)-

heteroalkyl, $-(C_3-C_{14})$ -heteroaryl, $-O-(C_3-C_{14})$ -heteroaryl, $-COO$ -alkyl, $-COO$ -aryl, $-C-O$ -alkyl, $-C-O$ -aryl, NH_2 , halogen and the residues are also capable of forming a larger condensed ring; wherein the recited alkyl groups, aryl groups, cycloalkyl, heteroalkyl groups, heteroaryl groups may be substituted as follows:

- 5 $-(C_1-C_{12})$ -alkyl, $-O-(C_1-C_{12})$ -alkyl, halogen;
and at least one of the radicals R^1 , R^2 , R^3 , R^4 does not represent phenyl.

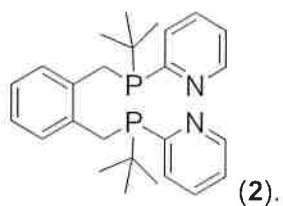
In one variant of the process, R^1 , R^2 , R^3 , R^4 are each independently selected from: $-(C_1-C_{12})$ -alkyl, $-(C_4-C_{14})$ -aryl, cycloalkyl, $-(C_1-C_{12})$ -heteroalkyl, $-(C_3-C_{14})$ -heteroaryl, halogen and the residues are also capable of forming a larger condensed ring; wherein the recited alkyl groups, aryl groups, cycloalkyl, heteroalkyl groups, heteroaryl groups may be substituted as follows:
10 $-(C_1-C_{12})$ -alkyl, $-O-(C_1-C_{12})$ -alkyl, halogen;
and at least one of the radicals R^1 , R^2 , R^3 , R^4 does not represent phenyl.

15 In one variant of the process, R^1 , R^2 , R^3 , R^4 are each independently selected from: $-(C_1-C_{12})$ -alkyl, cycloalkyl, $-(C_3-C_{14})$ -heteroaryl and the residues are also capable of forming a larger condensed ring;
wherein the recited alkyl groups, cycloalkyl, heteroaryl groups may be substituted as follows:
20 $-(C_1-C_{12})$ -alkyl, $-O-(C_1-C_{12})$ -alkyl, halogen,
and at least one of the radicals R^1 , R^2 , R^3 , R^4 does not represent phenyl.

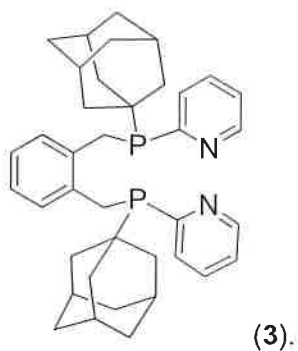
In one variant of the process, R^1 , R^4 are each independently selected from: $-(C_1-C_{12})$ -alkyl, cycloalkyl and the residues are also capable of forming a larger condensed ring;
25 wherein the recited alkyl groups, cycloalkyl may be substituted as follows:
 $-(C_1-C_{12})$ -alkyl, $-O-(C_1-C_{12})$ -alkyl, halogen.

In one variant of the process, R^2 , R^3 each independently represent $-(C_3-C_{14})$ -heteroaryl, wherein the recited heteroaryl groups may be substituted as follows:
30 $-(C_1-C_{12})$ -alkyl, $-O-(C_1-C_{12})$ -alkyl, halogen.

In one variant of the process, the compound of general formula (I) has the structure (2):



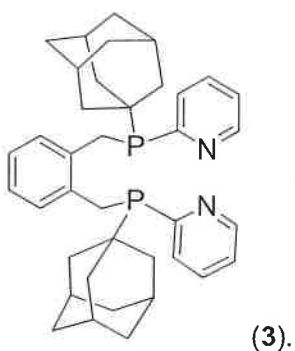
In one variant of the process, the compound of general formula (I) has the structure (3):



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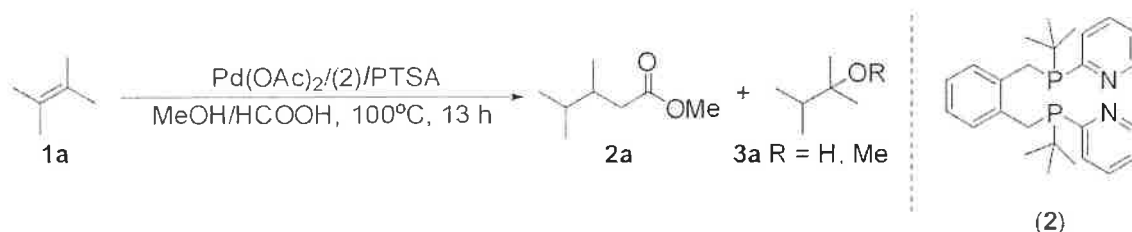
In addition to the process, a compound as such is also claimed.

Compound having the structure (3):



10 The invention is more particularly elucidated hereinbelow with reference to working examples.

A) Pd-catalyzed methoxycarbonylation of tetramethylethylene 1a with HCOOH: Effect of MeOH to HCOOH ratio



Added to a sealed 35 ml tube were $[\text{Pd}(\text{OAc})_2]$ (1.12 mg, 0.25 mol%), (2) (8.72 mg, 1.0 mol%), p-toluenesulfonic acid ($\text{PTSA} \cdot \text{H}_2\text{O}$) (15.2 mg, 4 mol%) and an oven-dried stirrer bar. The tube was placed together with the lid into a long Schlenk tube having a large opening. The Schlenk tube is evacuated three times and refilled with argon. Under an argon atmosphere **1a** (2 mmol), MeOH (X ml) and HCOOH (Y ml) ($X \text{ ml} + Y \text{ ml} = 2 \text{ ml}$) were injected into the 35 ml tube using a syringe. The 35 ml tube was then sealed with the lid. The reaction was carried out at 100°C over 13 h. After the reaction had ended, the tube was allowed to reach room temperature without additional cooling and carefully decompressed. Isooctane (100 μl) was then injected as internal standard. Conversion was measured by GC analysis.

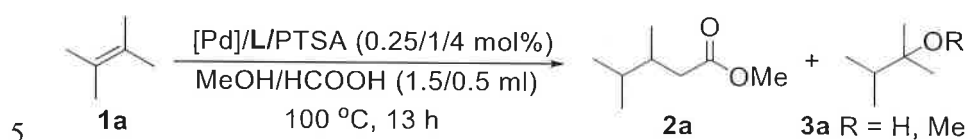
The results are summarized in table 1 which follows:

Table 1:

MeOH/HCOOH (altogether 2 ml)	Conversion %	Yield of 2a %	Yield of 3a %
1.8/0.2	63	36	24
1.6/0.4	70	49	19
1.5/0.5	76	62	13
1.4/0.6	76	61	13
1.2/0.8	78	66	10

1.0/1.0	74	48	10
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B) Pd-catalyzed methoxycarbonylation of tetramethylethylene **1a** with HCOOH: Effect of the ligand

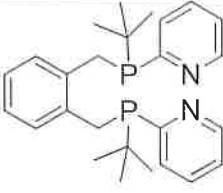
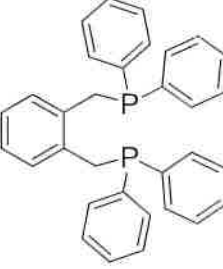
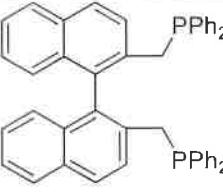


- Added to a sealed 35 ml tube under an argon atmosphere were $[\text{Pd}(\text{OAc})_2]$ (1.12 mg, 0.25 mol%), ligand (1 mol%), p-toluenesulfonic acid ($\text{PTSA} \cdot \text{H}_2\text{O}$) (15.2 mg, 4 mol%) and an oven-dried stirrer rod. The tube was placed together with the lid into a long Schlenk tube having a large opening.
- 10 The Schlenk tube was evacuated three times and refilled with argon. **1a** (2 mmol), HCOOH (0.5 ml) and MeOH (1.5 ml) were injected into the 35 ml tube using a syringe. The 35 ml tube was then sealed with the lid. The reaction was carried out at 100°C over 13 h. After the reaction had ended, the tube was allowed to reach room temperature without additional cooling (if very cold water used, tube can burst) and carefully decompressed. Isooctane (100 μl) was then injected as
- 15 internal standard. Conversion was measured by GC analysis.

The results are summarized in table 2 which follows:

Table 2:

Ligand (L)	Conversion %	Yield of 2a %	Yield of 3a %
(1)	60	32	22

 (2)	76	62	13
 (3)	83	67	13
 (4)	40	0	35
 (5)	38	0	31

As is shown by the experiments described above, the object is achieved by a process according
5 to the invention.

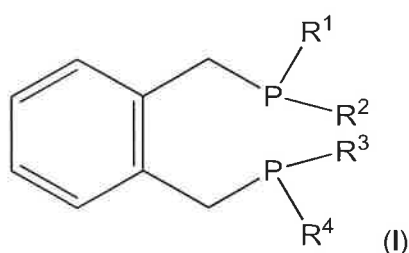
claims

1. Process comprising the process steps of:

a) addition of an olefin;

5 b) addition of a compound comprising Pd, wherein the Pd is capable of forming a complex;

c) addition of a compound of general formula (I):



wherein R^1 , R^2 , R^3 , R^4 are each independently selected from: -H, -(C₁-C₁₂)-alkyl, -O-(C₁-C₁₂)-alkyl, 10 -(C₄-C₁₄)-aryl, -O-(C₄-C₁₄)-aryl, cycloalkyl, -(C₁-C₁₂)-heteroalkyl, -O-(C₁-C₁₂)-heteroalkyl, -(C₃-C₁₄)-heteroaryl, -O-(C₃-C₁₄)-heteroaryl, -COO-alkyl, -COO-aryl, -C-O-alkyl, -C-O-aryl, NH₂, halogen and the residues are also capable of forming a larger condensed ring;

wherein the recited alkyl groups, aryl groups, cycloalkyl, heteroalkyl groups, heteroaryl groups may be substituted as follows:

15 -(C₁-C₁₂)-alkyl, -O-(C₁-C₁₂)-alkyl, halogen;

and at least one of the radicals R^1 , R^2 , R^3 , R^4 does not represent phenyl;

d) addition of MeOH and HCOOH,

wherein the MeOH / HCOOH ratio based on the total volume of MeOH and HCOOH employed is in the range from 1.55 : 0.45 to 1.1 : 0.9;

20 e) the reaction mixture is heated to convert the olefin into methyl ester.

2. Process according to Claim 1,

wherein no CO gas is supplied to the reaction mixture.

25 3. Process according to Claim 1 or 2,

wherein HCOOH serves as the only CO source for the reaction.

4. Process according to any of Claims 1 to 3,

wherein the compound in process step b) is selected from:

$\text{Pd}(\text{acac})_2$, PdCl_2 , $\text{Pd}(\text{dba})_3 \cdot \text{CH}_3\text{Cl}$ (dba = dibenzylideneacetone), $\text{Pd}(\text{OAc})_2$, $\text{Pd}(\text{TFA})_2$, $\text{Pd}(\text{CH}_3\text{CN})\text{Cl}_2$.

5. Process according to any of Claims 1 to 4,

5 wherein the process comprises additional process step f):

f) addition of an acid.

6. Process according to Claim 5,

wherein the acid is selected from: H_2SO_4 , $\text{CH}_3\text{SO}_3\text{H}$, $\text{CF}_3\text{SO}_3\text{H}$, PTSA.

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7. Process according to any of Claims 1 to 6,

wherein the MeOH / HCOOH ratio based on the total volume of MeOH and HCOOH employed is in the range from 1.5 : 0.5 to 1.2 : 0.8.

15 8. Process according to any of Claims 1 to 7,

wherein R^1 , R^2 , R^3 , R^4 are each independently selected from: $-(\text{C}_1\text{-C}_{12})\text{-alkyl}$, $-\text{O}-(\text{C}_1\text{-C}_{12})\text{-alkyl}$, $-(\text{C}_4\text{-C}_{14})\text{-aryl}$, $-\text{O}-(\text{C}_4\text{-C}_{14})\text{-aryl}$, cycloalkyl, $-(\text{C}_1\text{-C}_{12})\text{-heteroalkyl}$, $-\text{O}-(\text{C}_1\text{-C}_{12})\text{-heteroalkyl}$, $-(\text{C}_3\text{-C}_{14})\text{-heteroaryl}$, $-\text{O}-(\text{C}_3\text{-C}_{14})\text{-heteroaryl}$, $-\text{COO-alkyl}$, $-\text{COO-aryl}$, $-\text{C-O-alkyl}$, $-\text{C-O-aryl}$, NH_2 , halogen and the residues are also capable of forming a larger condensed ring;

20 wherein the recited alkyl groups, aryl groups, cycloalkyl, heteroalkyl groups, heteroaryl groups may be substituted as follows:

$-(\text{C}_1\text{-C}_{12})\text{-alkyl}$, $-\text{O}-(\text{C}_1\text{-C}_{12})\text{-alkyl}$, halogen;

and at least one of the radicals R^1 , R^2 , R^3 , R^4 does not represent phenyl.

25 9. Process according to any of Claims 1 to 8,

wherein R^1 , R^2 , R^3 , R^4 are each independently selected from: $-(\text{C}_1\text{-C}_{12})\text{-alkyl}$, $-(\text{C}_4\text{-C}_{14})\text{-aryl}$, cycloalkyl, $-(\text{C}_1\text{-C}_{12})\text{-heteroalkyl}$, $-(\text{C}_3\text{-C}_{14})\text{-heteroaryl}$, halogen and the residues are also capable of forming a larger condensed ring;

wherein the recited alkyl groups, aryl groups, cycloalkyl, heteroalkyl groups, heteroaryl groups may be substituted as follows:

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$-(\text{C}_1\text{-C}_{12})\text{-alkyl}$, $-\text{O}-(\text{C}_1\text{-C}_{12})\text{-alkyl}$, halogen;

and at least one of the radicals R^1 , R^2 , R^3 , R^4 does not represent phenyl.

10. Process according to any of Claims 1 to 9,

wherein R^1 , R^2 , R^3 , R^4 are each independently selected from: $-(C_1-C_{12})$ -alkyl, cycloalkyl, $-(C_3-C_{14})$ -heteroaryl and the residues are also capable of forming a larger condensed ring;

wherein the recited alkyl groups, cycloalkyl, heteroaryl groups may be substituted as follows:

$-(C_1-C_{12})$ -alkyl, $-O-(C_1-C_{12})$ -alkyl, halogen,

5 and at least one of the radicals R^1 , R^2 , R^3 , R^4 does not represent phenyl.

11. Process according to any of Claims 1 to 10,

wherein R^1 , R^4 are each independently selected from: $-(C_1-C_{12})$ -alkyl, cycloalkyl and the residues are also capable of forming a larger condensed ring;

10 wherein the recited alkyl groups, cycloalkyl may be substituted as follows:

$-(C_1-C_{12})$ -alkyl, $-O-(C_1-C_{12})$ -alkyl, halogen.

12. Process according to any of Claims 1 to 11,

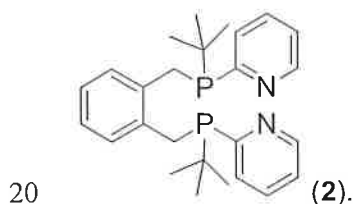
wherein R^2 , R^3 each independently represent $-(C_3-C_{14})$ -heteroaryl,

15 wherein the recited heteroaryl groups may be substituted as follows:

$-(C_1-C_{12})$ -alkyl, $-O-(C_1-C_{12})$ -alkyl, halogen.

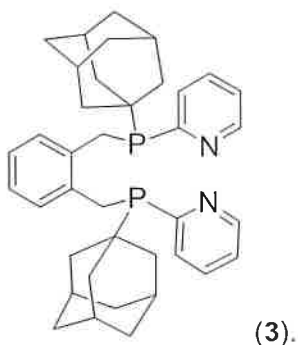
13. Process according to any of Claims 1 to 12,

wherein the compound of general formula (I) has the structure (2):



14. Process according to any of Claims 1 to 12,

wherein the compound of general formula (I) has the structure (3):



15. Compound having the structure **(3)**:

