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(54) Title  
**PALLADIUM CATALYST COMPOSITION, USE TO MAKE POLYKETONES**

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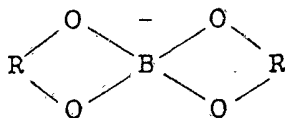
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(56) Prior Art Documents  
**AU 23319/88 C08G 67/02**  
**AU 596156 70492/87 C08G 67/02**  
**AU 589706 65069/86 C08G 02 C07F 9/00**

(57) Claim

1. A palladium catalyst prepared by reacting together:

- (a) a source of palladium,
- (b) a bidentate amine, phosphine, arsine or stibine having the formula  $(R^1)_2M-R^2-M(R^1)_2$  wherein the M atoms are independently selected from nitrogen, phosphorus, arsenic or antimony, the  $R^1$  groups are independently alkyl, cycloalkyl or aryl groups and  $R^2$  is an alkylene group, and
- (c) a source of an anion having the formula:

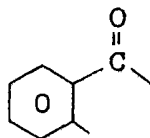


wherein the R groups are independently selected from the group consisting of  $C_1$  to  $C_6$  alkylene groups, orthophenylene groups or halogen, alkyl or sulphonic acid substituted

(11) AU-B-23320/88  
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derivatives thereof or groups having the  
formula:



or halogen, alkyl or sulphonic acid substituted  
derivatives thereof.

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COMPLETE SPECIFICATION

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This document contains the amendments made under Section 49 and is correct for printing.

TO BE COMPLETED BY APPLICANT

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Complete Specification for the invention entitled:

PROCESS FOR PREPARING POLYKETONES

The following statement is a full description of this invention, including the best method of performing it known to me/us:-

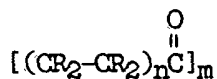
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PROCESS FOR PREPARING POLYKETONES

The present invention relates to a process for preparing  
interpolymers of olefins and carbon monoxide by polymerising a  
mixture of one or more olefins and carbon monoxide in the presence  
of a palladium catalyst. In particular, the present invention  
5 relates to new palladium catalysts for use in such processes.

The preparation of interpolymers of olefins and carbon monoxide  
having the formula:



where the R groups are independently hydrogen or hydrocarbyl groups,  
10 n is at least 1 and m is a large integer, is known from US 3694412.  
Such interpolymers, which hereafter will be called polyketones, are  
prepared according to US 3694412 by polymerising a mixture of one or  
more olefins and carbon monoxide in the presence of an aryl  
phosphine complex of a palladium halide and an inert solvent.  
15 However, the processes described in US 3694412 are slow even at  
elevated temperature and pressure.

An improved version of the process described in US 3694412 is  
described in European patent applications 181014 and 121965. It was  
subsequently found that the rate of the polymerisation process could  
20 be increased considerably by using a palladium catalyst with inter  
alia a bidentate phosphine and the anion of a carboxylic acid having  
a pKa of lower than 2 (as measured in aqueous solution). Examples  
of anions which can be used include trichloroacetate,  
dichloroacetate, tetrafluoroborate, hexafluorophosphate and  
25 p-toluene sulphonate, such anions being respectively the conjugate

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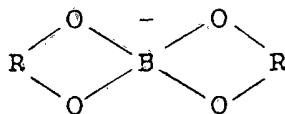
anions of trichloroacetic acid (pKa 0.70), dichloroacetic acid (pKa = 1.48), tetrafluoroboric acid, hexafluorophosphoric acid and p-toluenesulphonic acid.

More recently EP 222454 suggests that any acid having a pKa of  
5 less than 5 (determined in aqueous solution at 18°C) can be used.

It has now been found that if certain boron containing acids are employed then particularly high reaction rates can be achieved. Furthermore, if such acids are used then a lower catalyst  
10 deactivation on recycle is observed relative to previously described systems.

According to the present invention there is provided a process for preparing polyketones by polymerising a mixture of carbon monoxide and one or more olefins in the presence of a palladium catalyst characterised in that the palladium catalyst is prepared by  
15 reacting together:

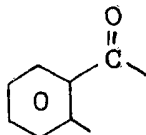
- (a) a source of palladium,
- (b) a bidentate amine, phosphine, arsine or stibine having the formula  $(R^1)_2M-R^2-M(R^1)_2$  wherein the M atoms are independently selected from nitrogen, phosphorus, arsenic or antimony, the  $R^1$   
20 are independently alkyl, cycloalkyl or aryl groups and  $R^2$  is an alkylene group, and
- (c) a source of an anion having the formula:



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wherein the R groups are independently selected from the group consisting of  $C_1$  to  $C_6$  alkylene groups, ortho-phenylene or biphenylene groups or halogen, alkyl or sulphonic acid substituted derivatives thereof or groups having the formula:

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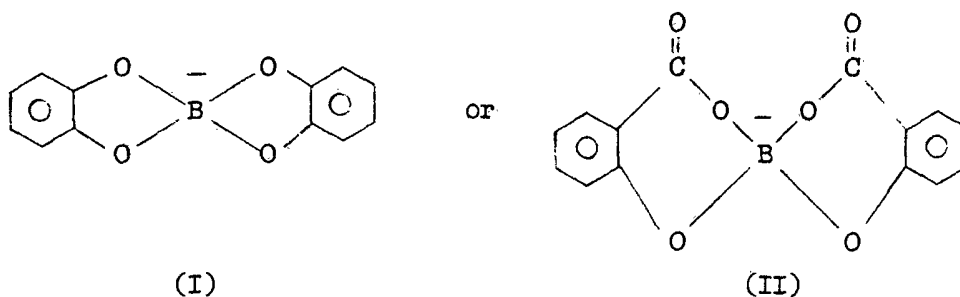
or halogen, alkyl or sulphonic acid substituted derivatives thereof.

35 The term polyketone is used herein to mean an interpolymer of



one or more olefins with carbon monoxide. The idealised structure of such a material would comprise a one, two or three dimensional network of strictly alternating olefin and carbon monoxide units. Although polyketones prepared according to the present invention correspond to this idealised structure, it is envisaged that materials corresponding to this structure in the main but containing small regimes (i.e. up to 10 wt %) of the corresponding polyolefin also fall within the definition.

Anions which are preferably employed in the process of the present invention include species such as:-



and analogues in which the benzene rings are partially or completely substituted with, e.g. halogen, alkyl or sulphonic acid moieties. Anions corresponding to (I) are hydrolysable and hence the corresponding acid has no measurable pka in aqueous solution. It is a feature of the present invention that palladium catalysts using anion (I) or (II) are more active than those using the anion of p-toluenesulphonic acid, a preferred anion according to EP 121965.

The palladium catalyst used in the process described above is itself prepared by reacting together a source of palladium, an appropriate amine, phosphine, arsine or stibine and anion having the formula defined above. As regards the source of palladium this can include simple inorganic and organic salts, e.g. halides, nitrates, carboxylates and the like as well as organometallic and coordination complexes. In some cases, by suitable choice of coordination complex, it may be possible to add the palladium and the amine, phosphine, arsine or stibine as a single entity.

Although any source of palladium can be used, it may be

necessary, when a palladium complex having strongly coordinating ligands is employed, to ensure that such ligands are removed. An example of such a complex is palladium acetate where the acetate anions bind strongly to the palladium. In such cases the acetate anions can be removed by adding component (c) above as its conjugate acid since such a conjugate acid will protonate the acetate anions and cause their removal.

Another approach which is useful when palladium halides are employed (halide anions also bind strongly to the palladium) is to use a thallium or silver salt of component (c). In such cases a metathesis reaction occurs and the insoluble silver or thallium halide precipitates and can be removed by filtration.

The other component of the catalyst is a bidentate amine, phosphine, arsine or stibine having the formula  $(R^1)_2M-R^2-M(R^1)_2$  where the M atoms are independently selected from nitrogen, phosphorus, arsenic or antimony. The  $R^1$  groups are independently selected from alkyl, cycloalkyl or aryl groups with  $C_1$  to  $C_4$  alkyl and phenyl groups being preferred. The  $R^2$  group is an alkylene group. In the context of this document alkylene groups are defined as being  $-(CH_2)_a(CHR^3)_b-$  groups in which the  $R^3$  groups are independently hydrogen, methyl, ethyl or propyl groups and a and b are either zero or integers such that  $a+b$  is at least 2, preferably between 2 and 10. Preferably the alkylene group is selected from  $-(CH_2)_2-$ ,  $-(CH_2)_3-$ ,  $-(CH_2)_4-$  and  $-(CH_2)_5-$ . Of these the most convenient species are the bidentate phosphines, 1,2-bis(diphenylphosphino)ethane (diphos), 1,3-bis(diphenylphosphino)propane and 1,4-bis(diphenylphosphino)butane.

The bidentate amines, phosphines, arsines or stibines defined above all either bond exclusively cis to the palladium or there is a reasonable concentration of this cis binding. Whilst not wishing to be held to any theory, it is believed that it is a cis type isomer which is catalytically active.

Considering next the feedstocks for the polymerisation feedstock, it is believed that any source of carbon monoxide can be

used. Thus the carbon monoxide may contain nitrogen, inert gases and up to 10% hydrogen.

Any olefin can in theory be used although the best reaction rates are obtained when either ethylene or a mixture of olefins which include ethylene, e.g. ethylene/propylene, ethylene/butylene and the like, is used. The lower rates obtained in the absence of ethylene should not be construed as indicating that the process can be used only with an ethylene feedstock since other olefins such as propylene, 4-methylpentene-1, styrene, acrylates, vinyl acetates and the like all undergo reaction to some extent.

The polymerisation process is suitably carried out in a solvent which is chemically inert under the conditions employed and one in which the palladium catalyst is soluble. Moreover, the solvent like the anion should be either weakly or non-coordinating. Examples of such solvents include alcohols, e.g. methanol, ethanol and propanol, ethers, glycols and glycol ethers. Preferred solvents are methanol or ethoxyethanol.

The polymerisation process is suitably carried out at a temperature in the range 20 to 150°C and at elevated pressure, (e.g. 1 to 100 bars). The overpressure of gas is suitably carbon monoxide or carbon monoxide and olefin, if the olefin is gaseous under the reaction conditions. It is possible to operate the polymerisation process either batchwise or continuously.

The following Examples illustrate the present invention.

Example 1

A stainless steel autoclave of 150 ml capacity was charged with palladium acetate (25 mg), 1,3-bis(diphenylphosphino)propane (65 mg),  $H[B(O_2C_6H_4)_2]_2DMF$  (800 mg), 1,2-dimethoxyethane (30 ml) and a magnetic stirrer bar. The autoclave was flushed with nitrogen, charged with ethylene (20 bar) followed by carbon monoxide (30 bar), and heated to 100°C. A pressure drop of 9.0 bar was recorded over 1 hour. The autoclave was allowed to cool over a period of 1½ hours, and was vented at 50°C, 4.96g polymer was subsequently recovered.

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Example 2

The method of Example 1 was repeated, except that the  $H[B(O_2C_6H_4)_2]_2DMF$  was replaced by  $H[B(OC_6H_4CO_2)_2]$  (600 mg) and the 1,2-dimethoxyethane by methanol (30 ml). After 40 minutes gas  
5 uptake had virtually ceased and pressure drop of 31.6 bar had been recorded. The autoclave was allowed to cool and was vented. 6.92g of polymer was then recovered by filtration under a nitrogen atmosphere. The filtrate was returned to the autoclave and the reaction repeated. A pressure drop of 19.3 bar occurred over  
10 1 hour. The autoclave was allowed to cool and was vented, and 4.62g of polymer recovered.

Comparative Test A

The method of Example 2 was repeated, except that the  $H[B(OC_6H_4CO_2)_2]$  was replaced by para-toluenesulphonic acid (380mg).  
15 In the first instance, a pressure drop of 25.6 bar and polymer yield of 5.07g were recorded. On repeating the reaction with the recovered filtrate, a pressure drop of 7.4 bar and a polymer yield of 2.11g were recorded.

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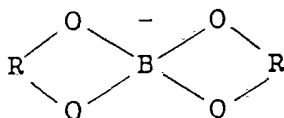
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THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

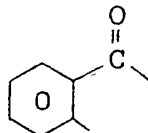
1. A palladium catalyst prepared by reacting together:

- (a) a source of palladium,
- 5 (b) a bidentate amine, phosphine, arsine or stibine having the formula  $(R^1)_2M-R^2-M(R^1)_2$  wherein the M atoms are independently selected from nitrogen, phosphorus, arsenic or antimony, the  $R^1$  groups are independently alkyl, cycloalkyl or aryl groups and  $R^2$  is an alkylene group, and
- 10 (c) a source of an anion having the formula:



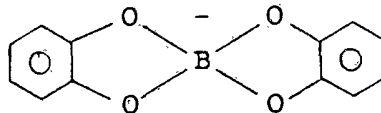
wherein the R groups are independently selected from the group consisting of  $C_1$  to  $C_6$  alkylene groups, orthophenylene groups or halogen, alkyl or sulphonic acid substituted derivatives thereof or groups having the formula:

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or halogen, alkyl or sulphonic acid substituted derivatives thereof.

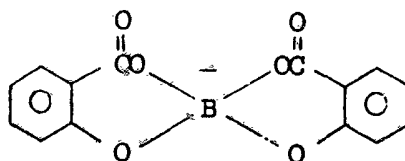
2. A catalyst as claimed in claim 1 wherein the anion is:



or halogen, alkyl or sulphonic acid substituted derivatives thereof.

3. A catalyst as claimed in claim 1 wherein the anion is:

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or halogen, alkyl or sulphonic acid substituted derivatives thereof.

4. A catalyst as claimed in any one of the preceding claims wherein the source of the anion is the conjugate acid of the anion.

5. A catalyst as claimed in any one of claims 1-3 wherein the source of palladium is a palladium halide and the source of the anion is the silver or thallium salt.

6. A catalyst as claimed in any one of the preceding claims wherein the reaction is carried out in either methanol or ethoxyethanol.

7. A palladium catalyst substantially as herein described with reference to Example 1 or Example 2.

8. A process for preparing polyketones by polymerising a mixture of carbon monoxide and one or more olefins in the presence of a palladium catalyst as claimed in any one of the preceding claims.

9. A process for preparing polyketones by polymerising a mixture of carbon monoxide and one or more olefins in the presence of a palladium catalyst as claimed in any one of claims 1-7 which comprises the steps of

- (a) contacting the palladium catalyst with the mixture of carbon monoxide and one or more olefins,
- (b) allowing the palladium catalyst to polymerise or polymerise partially the mixture of carbon monoxide and one or more olefins,
- (c) separating the palladium catalyst from the products of step (b),
- (d) contacting the palladium catalyst with further carbon monoxide and one or more olefins.

10. A process as claimed in claim 8 or claim 9 wherein the one or more olefins are selected from the group consisting of ethylene, ethylene/propylene and ethylene/butylene.



11. A process as claimed in claim 9 or claim 10  
carried out continuously.

12. A process for preparing polyketones  
5 substantially as herein described with reference to Example  
1 or Example 2.

DATED this 2nd day of November 1990

10 THE BRITISH PETROLEUM COMPANY p.l.c.  
By their Patent Attorneys  
GRIFFITH HACK & CO.

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