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DESCRIPTION

MODIFIED PARTICLE, CARRIER, CATALYST COMPONENT FOR ADDITION
POLYMERIZATION, CATALYST FOR ADDITION POLYMERIZATION AND
5 PROCESS FOR PRODUCING ADDITION POLYMER

Technical Field

The present invention relates to a modified particle useful
for a carrier and a catalyst component for addition
10 polymerization, and also relates to a catalyst for addition
polymerization obtained using the same and a process for
producing an addition polymer. Further, the present invention
relates to a typical metal component used for producing the
modified particle mentioned above, a process for producing the
15 same, a catalyst component for addition polymerization
comprising the same, a catalyst for addition polymerization
obtained using the same, and a process for producing an addition
polymer.

20 Background Art

Addition polymers such as polypropylene and polyethylene
are superior in their mechanical properties and chemical
resistance and additionally superior in balance between their
properties and economies, and thereby they are extensively used
25 in various molding fields. As important quality items of a film,
which is one of their main applications, there are enumerated

appearance such as transparency and mechanical strength. Therefore, from a viewpoint of a quality of the film, it is desired to avoid, for example, occurrence of a fish eye due to a gel, according to which its appearance is remarkably deteriorated and moreover deterioration of strength is caused. On the other hand, at the time of mold-processing to obtain a film, an advanced processability such as low extrusion load and high melting tension is required.

These addition polymers have hitherto been produced by polymerizing an olefin and the like mainly using a conventional solid catalyst (multi-site catalyst) formed from a combination of the 4 Group metal compound such as titanium trichloride and titanium tetrachloride and the 13 Group metal compound typified by an organoaluminum compound.

In recent years, there has been proposed a process for producing an addition polymer comprising the step of polymerizing an olefin and the like using a so-called single site catalyst, which is a combination of a transition metal compound (for example, a metallocene complex and a non-metallocene complex) different from the solid catalyst component so far used, and an aluminoxane and the like. For example, a process comprising the step of using a combination of bis(cyclopentadienyl)zirconium dichloride and methylaluminoxane is reported in JP-A 58-19309. In addition, a combination of such a transition metal compound and a specific boron compound is also reported. For example, a process comprising the step of using bis(cyclopentadienyl)zirconium

dimethyl and tri(n-butyl) ammonium
tetrakis(pentafluorophenyl) borate is reported in JP-W 1-502036.
As known, an addition polymer obtained using such a single site
catalyst is generally narrower in its molecular weight
5 distribution than that obtained using the conventional solid
catalyst (multi-site catalyst), and in the case of a copolymer,
an addition polymer more homogeneous than that obtained using
the conventional solid catalyst can be obtained because the
comonomers are more homogeneously copolymerized.

10 While, these known single site catalysts are soluble in
a reaction system. Therefore, when these catalysts are applied
to polymerization accompanied by the formation of addition
polymer particles (for example, slurry polymerization, gas phase
polymerization and bulk polymerization), the obtained addition
15 polymer has an amorphous shape, and it is inevitable to result
in bulky particles of the addition polymer, mass of the addition
polymer or pulverized powders of the addition polymer, or to
decrease bulk density of the addition polymer, or it is inevitable
that the addition polymer adheres to a wall of a polymerization
20 reactor. Owing to any of these reasons, there has been a problem
of poor heat transfer or poor heat removal of the reactor, so
that stable operations have been made difficult and productivity
has been lowered. JP-A 11-193306 discloses a means to solve such
a problem. However, the means disclosed therein is not always
25 sufficient in respect to polymerization activity.

With respect to an α -olefin, particularly propylene,

there are known production examples of a high stereoregular isotactic propylene polymer using, for example, a metallocene catalyst cross-linked with two of a 1 to 3 substituted η^5 -cyclopentadienyl group (JP No. 2587251, JP No. 2627669 and JP No. 2668732). It is also reported to obtain a syndiotactic propylene polymer using a Cs symmetric metallocene complex, namely a combination of isopropylidene(cyclopentadienyl)(fluorenyl)zirconium dichloride and the like and an aluminum oxy compound (J. Am. Chem. Soc., 1988, 110, 6255). However, it has been difficult to say that a process for producing an α -olefin polymer according to the process using these known single site catalyst is sufficiently efficient.

15 Disclosure of Invention

According to the study of the present inventors, it has been found that one of problems of the addition polymer obtained by using such a single site catalyst is that many fish eyes are apt to appear on a film formed from the addition polymer.

20 An object of the present invention is to provide a particle, a carrier and a catalyst component for addition polymerization, which can be used for addition polymerization in combination with a transition metal compound capable of forming a single site catalyst, thereby producing an addition polymer, from which
25 a film having little fish eye and so good appearance can be obtained, a pre-polymerized catalyst component for addition

polymerization obtained using said particle, a catalyst for addition polymerization obtained using said particle, and a process for producing an addition polymer using said catalyst for addition polymerization.

5 Another object of the present invention is to provide a particle, a carrier and a catalyst component for addition polymerization, which are useful for the preparation of a high activity single site catalyst suitably applied for polymerization accompanied by the formation of addition polymer
10 particles (for example, slurry polymerization, gas phase polymerization and bulk polymerization), a high activity catalyst for addition polymerization and a process for producing an addition polymer efficiently.

An additional object of the present invention is to provide
15 a process for producing a stereoregular α -olefin polymer efficiently.

Other objects and advantages of the present invention will be apparent from the following descriptions.

That is, the present invention provides a modified particle
20 obtained by contacting the following (a), the following (b), the following (c) and a particle (d) with one another,

(a): a compound represented by the following general formula [1],



25 (b): a compound represented by the following general formula [2],

$R^1_{t-1}TH$ [2], and

(c): a compound represented by the following general formula [3],

$R^2_{t-2}TH_2$ [3].

5 (In the above general formulas [1] to [3], M^1 is a typical metal atom of the Group 1, 2, 12, 14 or 15 of the periodic table, m is a valence of M^1 , L^1 is a hydrogen atom, a halogen atom or a hydrocarbon group, when more than one L^1 exists, they may be the same or different from one another, R^1 is an
10 electron-withdrawing group or an electron-withdrawing group-containing group, when more than one R^1 exists, they may be the same or different from one another, R^2 is a hydrocarbon group or a halogenated hydrocarbon group, T in each compound is independently of each other a non-metal atom of the Group
15 15 or 16 of the periodic table, and t is a valence of T in each compound.)

The present invention also provides a carrier comprising said particle and a catalyst component for addition polymerization, which comprises said particle.

20 Further, the present invention provides a catalyst for addition polymerization, which is obtained by contacting the modified particle (A) and a compound (B) of a transition metal of the Groups 3 to 11 or a lanthanoide series with each other, and also provides a catalyst for addition polymerization, which
25 is obtained by contacting the modified particle (A), a compound (B) of a transition metal of the Groups 3 to 11 or a lanthanoide

series and an organoaluminum compound (C) with one another.

Still further, the present invention provides a pre-polymerized catalyst component for addition polymerization, which is obtained by pre-polymerizing an olefin in the presence of the catalyst obtained by contacting the modified particle (A) and a compound (B) of a transition metal of the Groups 3 to 11 or a lanthanoid series and optionally an organoaluminum compound (C) with one another, and also provides a catalyst for addition polymerization, which is obtained by contacting said pre-polymerized catalyst component for addition polymerization and the organoaluminum compound (C) with each other.

Additionally, the present invention provides a process for producing a stereoregular α -olefin polymer, which comprises the step of polymerizing an α -olefin in the presence of a catalyst obtained in advance by contacting the above-mentioned modified particle (A), a transition metal compound having a stereoregular polymerization ability of an α -olefin, which transition metal compound is selected from the above-mentioned transition metal compounds (B), and the organoaluminum compound (C) with one another.

The present invention is explained in more detail as follows.

Best Mode for Carrying out the Invention

Compound (a)

The compound (a) is a compound represented by the following

general formula [1].



The symbol M^1 is a typical metal atom of the Group 1, 2, 12, 14 or 15 of the periodic table of the elements (Revised
5 edition of IUPAC Inorganic Chemistry Nomenclature 1989). Specific examples thereof include a lithium atom, a sodium atom, a potassium atom, a rubidium atom, a cesium atom, a beryllium atom, a magnesium atom, a calcium atom, a strontium atom, a barium atom, a zinc atom, a cadmium atom, a mercury
10 atom, a germanium atom, a tin atom, a lead atom, an antimony atom and a bismuth atom. As M^1 , an atom of the Group 12 is particularly preferred, and a zinc atom is the most preferred.

The symbol m is a valence of M^1 , and for example, when M^1 is a zinc atom, m is 2.

15 The symbol L^1 is a hydrogen atom, a halogen atom or a hydrocarbon group, and when more than one L^1 exists, they may be the same or different from one another. Specific examples of the halogen atom as L^1 are a fluorine atom, a chlorine atom, a bromine atom, and an iodine atom. As a
20 hydrocarbon group as L^1 , an alkyl group, an aryl group or an aralkyl group is preferred.

As the alkyl group stated here, an alkyl group having 1 to 20 carbon atoms is preferred. Examples thereof are a methyl group, an ethyl group, a n-propyl group, an isopropyl
25 group, a n-butyl group, a sec-butyl group, a tert-butyl group, an isobutyl group, a n-pentyl group, a neopentyl group, a n-hexyl group, a n-octyl group, a n-decyl group, a n-dodecyl

group, a n-pentadecyl group and a n-eicosyl group. More preferred is a methyl group, an ethyl group, an isopropyl group, a tert-butyl group or an isobutyl group. These alkyl groups may be substituted with a halogen atom such as a fluorine atom, a chlorine atom, a bromine atom, or an iodine atom. Examples of the alkyl group having 1 to 20 carbon atoms substituted with the halogen atom are a fluoromethyl group, a difluoromethyl group, a trifluoromethyl group, a chloromethyl group, a dichloromethyl group, a trichloromethyl group, a bromomethyl group, a dibromomethyl group, a tribromomethyl group, an iodomethyl group, a diiodomethyl group, a triiodomethyl group, a fluoroethyl group, a difluoroethyl group, a trifluoroethyl group, a tetrafluoroethyl group, a pentafluoroethyl group, a chloroethyl group, a dichloroethyl group, a trichloroethyl group, a tetrachloroethyl group, a pentachloroethyl group, a bromoethyl group, a dibromoethyl group, a tribromoethyl group, a tetrabromoethyl group, a pentabromoethyl group, a perfluoropropyl group, a perfluorobutyl group, a perfluoropentyl group, a perfluorohexyl group, a perfluorooctyl group, a perfluorododecyl group, a perfluoropentadecyl group, a perfluoroeicosyl group, a perchloropropyl group, a perchlorobutyl group, a perchloropentyl group, a perchlorohexyl group, a perchlorooctyl group, a perchlorododecyl group, a perchloropentadecyl group, a perchloroeicosyl group, a perbromopropyl group, a perbromobutyl group, a

perbromopentyl group, a perbromohexyl group, a perbromooctyl group, a perbromododecyl group, a perbromopentadecyl group and a perbromoeicosyl group.

Further, these alkyl groups may be partially substituted
5 with an alkoxy group such as a methoxy group or an ethoxy group, an aryloxy group such as a phenoxy group, or an aralkyloxy group such as a benzyloxy group.

As the aryl group, an aryl group having 6 to 20 carbon atoms is preferred. Examples thereof are a phenyl group, a 2-tolyl
10 group, a 3-tolyl group, a 4-tolyl group, a 2,3-xylyl group, a 2,4-xylyl group, a 2,5-xylyl group, a 2,6-xylyl group, a 3,4-xylyl group, a 3,5-xylyl group, a 2,3,4-trimethylphenyl group, a 2,3,5-trimethylphenyl group, a 2,3,6-trimethylphenyl group, a 2,4,6-trimethylphenyl group, a 3,4,5-trimethylphenyl
15 group, a 2,3,4,5-tetramethylphenyl group, a 2,3,4,6-tetramethylphenyl group, a 2,3,5,6-tetramethylphenyl group, a pentamethylphenyl group, an ethylphenyl group, a n-propylphenyl group, an isopropylphenyl group, a n-butylphenyl group, a sec-butylphenyl group, a tert-butylphenyl group, an
20 isobutylphenyl group, a n-pentylphenyl group, a neopentylphenyl group, a n-hexylphenyl group, a n-octylphenyl group, a n-decylphenyl group, a n-dodecylphenyl group, a n-tetradecylphenyl group, a naphthyl group and an anthracenyl group, and more preferred is a phenyl group.

25 These aryl groups may be partially substituted with a halogen atom such as a fluorine atom, a chlorine atom, a bromine atom or an iodine atom, an alkoxy group such as a methoxy group

or an ethoxy group, an aryloxy group such as a phenoxy group, or an aralkyloxy group such as a benzyloxy group.

As the aralkyl group, an aralkyl group having 7 to 20 carbon atoms is preferred. Examples thereof are a benzyl group, a
5 (2-methylphenyl)methyl group, a (3-methylphenyl)methyl group, a (4-methylphenyl)methyl group, a (2,3-dimethylphenyl)methyl group, a (2,4-dimethylphenyl)methyl group, a (2,5-dimethylphenyl)methyl group, a (2,6-dimethylphenyl)methyl group, a
10 (3,4-dimethylphenyl)methyl group, a (3,5-dimethylphenyl)methyl group, a (2,3,4-trimethylphenyl)methyl group, a (2,3,5-trimethylphenyl)methyl group, a (2,3,6-trimethylphenyl)methyl group, a
15 (3,4,5-trimethylphenyl)methyl group, a (2,4,6-trimethylphenyl)methyl group, a (2,3,4,5-tetramethylphenyl)methyl group, a (2,3,4,6-tetramethylphenyl)methyl group, a (2,3,5,6-tetramethylphenyl)methyl group, a
20 (pentamethylphenyl)methyl group, an (ethylphenyl)methyl group, a (n-propylphenyl)methyl group, an (isopropylphenyl)methyl group, a (n-butylphenyl)methyl group, a (sec-butylphenyl)methyl group, a (tert-butylphenyl)methyl group, an (isobutylphenyl)methyl group, a
25 (n-pentylphenyl)methyl group, a (neopentylphenyl)methyl group, a (n-hexylphenyl)methyl group, a (n-octylphenyl)methyl group, a (n-decylphenyl)methyl group, a naphthylmethyl group and an

anthracenylmethyl group, and more preferred is a benzyl.

These aralkyl groups may be partially substituted with a halogen atom such as a fluorine atom, a chlorine atom, a bromine atom or an iodine atom, an alkoxy group such as a methoxy group
5 or an ethoxy group, an aryloxy group such as a phenoxy group, or an aralkyloxy group such as a benzyloxy group.

As L^1 , preferred is a hydrogen atom, an alkyl group or an aryl group, more preferred is a hydrogen atom or an alkyl group, and particularly preferable is an alkyl group.

10 Compounds (b) and (c)

The compound (b) is a compound represented by the general formula [2],



the compound (c) is a compound represented by the following
15 general formula [3],



Each T in the general formula [2] or [3] is independently of each other a non-metal atom of Group 15 or Group 16 of the periodic table of elements (Revised edition of IUPAC Inorganic
20 Chemistry Nomenclature 1989). Each T in the general formula [2] or in the general formula [3] may be the same or different from each other. Specific examples of the non-metal atom of the Group 15 are a nitrogen atom and a phosphorous atom, and specific
25 examples of the non-metal atom of the Group 16 are an oxygen atom and a sulfur atom. As T, preferred is a nitrogen atom or an oxygen atom, and more preferred is an oxygen atom.

Each t in the above-mentioned general formula [2] or [3]

is a valence of T. For example, when T is an atom of the Group 15, t is 3, and when T is an atom of the Group 16, t is 2.

The symbol R^1 in the above-mentioned general formula [2] is an electron-withdrawing group or an electron-withdrawing group-containing group, and when more than one R^1 exists, they may be the same or different from one another. As an index of the electron-withdrawing property, there is known a substituent constant σ of the Hammett's rule. As the electron-withdrawing group, there is a functional group whose substituent constant σ of the Hammett's rule is positive.

Specific examples of the electron-withdrawing group are a fluorine atom, a chlorine atom, a bromine atom, an iodine atom, a cyano group, a nitro group, a carbonyl group, a sulfone group and a phenyl group. Examples of the electron-withdrawing group-containing group are a halogenated alkyl group, a halogenated aryl group, a (halogenated alkyl)aryl group, a cyanated aryl group, a nitrated aryl group, an ester group and an acyl group.

Specific examples of the halogenated alkyl group are a fluoromethyl group, a chloromethyl group, a bromomethyl group, an iodomethyl group, a difluoromethyl group, a dichloromethyl group, a dibromomethyl group, a diiodomethyl group, a trifluoromethyl group, a trichloromethyl group, a tribromomethyl group, a triiodomethyl group, a 2,2,2-trifluoroethyl group, a 2,2,2-trichloroethyl group, a 2,2,2-tribromoethyl group, a 2,2,2-triiodoethyl group, a 2,2,3,3,3-pentafluoropropyl group, a

2,2,3,3,3-pentachloropropyl group, a
 2,2,3,3,3-pentabromopropyl group, a 2,2,3,3,3-pentaiodopropyl
 group, a 2,2,2-trifluoro-1-trifluoromethylethyl group, a
 2,2,2-trichloro-1-trichloromethylethyl group, a
 5 2,2,2-tribromo-1-tribromomethylethyl group, a
 2,2,2-triiodo-1-triiodomethylethyl group, a
 1,1-bis(trifluoromethyl)-2,2,2-trifluoroethyl group, a
 1,1-bis(trichloromethyl)-2,2,2-trichloroethyl group, a
 1,1-bis(tribromomethyl)-2,2,2-tribromoethyl group and a
 10 1,1-bis(triiodomethyl)-2,2,2-triiodoethyl group.

Specific examples of the halogenated aryl group are a
 2-fluorophenyl group, a 3-fluorophenyl group, a 4-fluorophenyl
 group, a 2,4-difluorophenyl group, a 2,6-difluorophenyl group,
 a 3,4-difluorophenyl group, a 3,5-difluorophenyl group, a
 15 2,4,6-trifluorophenyl group, a 2,3,5,6-tetrafluorophenyl group,
 a pentafluorophenyl group, a
 2,3,5,6-tetrafluoro-4-trifluoromethylphenyl group, a
 2,3,5,6-tetrafluoro-4-pentafluorophenylphenyl group, a
 perfluoro-1-naphthyl group, a perfluoro-2-naphthyl group, a
 20 2-chlorophenyl group, a 3-chlorophenyl group, a 4-chlorophenyl
 group, a 2,4-dichlorophenyl group, a 2,6-dichlorophenyl group,
 a 3,4-dichlorophenyl group, a 3,5-dichlorophenyl group, a
 2,4,6-trichlorophenyl group, a 2,3,5,6-tetrachlorophenyl group,
 a pentachlorophenyl group, a
 25 2,3,5,6-tetrachloro-4-trichloromethylphenyl group, a
 2,3,5,6-tetrachloro-4-pentachlorophenylphenyl group, a
 perchloro-1-naphthyl group, a perchloro-2-naphthyl group, a

2-bromophenyl group, a 3-bromophenyl group, a 4-bromophenyl group, a 2,4-dibromophenyl group, a 2,6-dibromophenyl group, a 3,4-dibromophenyl group, a 3,5-dibromophenyl group, a 2,4,6-tribromophenyl group, a 2,3,5,6-tetrabromophenyl group, a pentabromophenyl group, a 2,3,5,6-tetrabromo-4-tribromomethylphenyl group, a 2,3,5,6-tetrabromo-4-pentabromophenylphenyl group, a perbromo-1-naphthyl group, a perbromo-2-naphthyl group, a 2-iodophenyl group, a 3-iodophenyl group, a 4-iodophenyl group, a 2,4-diiodophenyl group, a 2,6-diiodophenyl group, a 3,4-diiodophenyl group, a 3,5-diiodophenyl group, a 2,4,6-triiodophenyl group, a 2,3,5,6-tetraiodophenyl group, a pentaiodophenyl group, a 2,3,5,6-tetraiodo-4-triiodomethylphenyl group, a 2,3,5,6-tetraiodo-4-pentaiodophenylphenyl group, a periodo-1-naphthyl group and a periodo-2-naphthyl group.

Specific examples of the (halogenated alkyl)aryl group are a 2-(trifluoromethyl)phenyl group, a 3-(trifluoromethyl)phenyl group, a 4-(trifluoromethyl)phenyl group, a 2,6-bis(trifluoromethyl)phenyl group, a 3,5-bis(trifluoromethyl)phenyl group, a 2,4,6-tris(trifluoromethyl)phenyl group and a 3,4,5-tris(trifluoromethyl)phenyl group.

Specific examples of the cyanated aryl group are a 2-cyanophenyl group, a 3-cyanophenyl group and a 4-cyanophenyl group.

Specific examples of the nitrated aryl group are a

2-nitrophenyl group, a 3-nitrophenyl group and a 4-nitrophenyl group.

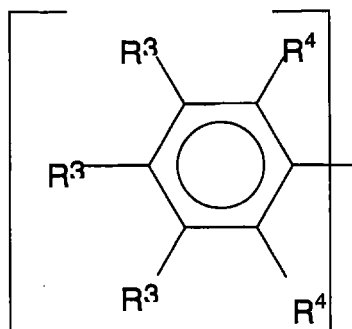
Specific examples of the ester group are a methoxycarbonyl group, an ethoxycarbonyl group, a n-propoxycarbonyl group, an
5 isopropoxycarbonyl group, a phenoxycarbonyl group, a trifluoromethoxycarbonyl group and a pentafluorophenoxycarbonyl group.

Specific examples of the acyl group are a formyl group, an ethanoyl group, a propanoyl group, a butanoyl group, a
10 trifluoroethanoyl group, a benzoyl group and a pentafluorobenzoyl group.

As R¹, preferred is a halogenated hydrocarbon group, and more preferred is a halogenated alkyl group or a halogenated aryl group. Much more preferred is a fluoromethyl group, a
15 difluoromethyl group, a trifluoromethyl group, a 2,2,2-trifluoroethyl group, a 2,2,3,3,3-pentafluoropropyl group, a 2,2,2-trifluoro-1-trifluoromethylethyl group, a 1,1-bis(trifluoromethyl)-2,2,2-trifluoroethyl group, a 2-fluorophenyl group, a 3-fluorophenyl group, a 4-fluorophenyl
20 group, a 2,4-difluorophenyl group, a 2,6-difluorophenyl group, a 3,4-difluorophenyl group, a 3,5-difluorophenyl group, a 2,4,6-trifluorophenyl group, a 2,3,5,6-tetrafluorophenyl group, a pentafluorophenyl group, a 2,3,5,6-tetrafluoro-4-trifluoromethylphenyl group, a
25 2,3,5,6-tetrafluoro-4-pentafluorophenylphenyl group, a perfluoro-1-naphtyl group, a perfluoro-2-naphtyl group, a chloromethyl group, a dichloromethyl group, a trichloromethyl

group, a 2,2,2-trichloroethyl group, a
 2,2,3,3,3-pentachloropropyl group, a
 2,2,2-trichloro-1-trichloromethylethyl group, a
 1,1-bis(trichloromethyl)-2,2,2-trichloroethyl group, a
 5 4-chlorophenyl group, a 2,6-dichlorophenyl group, a
 3,5-dichlorophenyl group, a 2,4,6-trichlorophenyl group or a
 pentachlorophenyl group. Particularly preferred is a
 fluoroalkyl group or a fluoroaryl group, and the most preferred
 is a trifluoromethyl group, a
 10 2,2,2-trifluoro-1-trifluoromethylethyl group, a
 1,1-bis(trifluoromethyl)-2,2,2-trifluoroethyl group, a
 3,5-difluorophenyl group, or a pentafluorophenyl group.

Further, when R^1 is an aryl group having an electron-drawing
 group, as such an aryl group, a group represented by the following
 15 general formula [4] is preferred from a viewpoint of increase
 of activity rather than the viewpoint of the fish eye.



[4]

20 The symbol R^3 is preferably a halogen atom, a halogenated
 alkyl group, a halogenated aryl group, a cyano group, a cyanated
 alkyl group, a cyanated aryl group, a nitro group, a nitrated

alkyl group, a nitrated aryl group or an ester group. More preferred is a halogen atom, a halogenated alkyl group or a halogenated aryl group. Much more preferred is a fluorine atom, a fluoromethyl group, a difluoromethyl group, a trifluoromethyl group, a 2,2,2-trifluoroethyl group, a 2,2,3,3,3-pentafluoropropyl group, a 2,2,2-trifluoro-1-trifluoromethylethyl group, a 1,1-bis(trifluoromethyl)-2,2,2-trifluoroethyl group, a 2-fluorophenyl group, a 3-fluorophenyl group, a 4-fluorophenyl group, a 2,4-difluorophenyl group, a 2,6-difluorophenyl group, a 3,4-difluorophenyl group, a 3,5-difluorophenyl group, a 2,4,6-trifluorophenyl group, a 2,3,5,6-tetrafluorophenyl group, a pentafluorophenyl group, a 2,3,5,6-tetrafluoro-4-trifluoromethylphenyl group, a 2,3,5,6-tetrafluoro-4-pentafluorophenylphenyl group, a perfluoro-1-naphthyl group, a perfluoro-2-naphthyl group, a chloromethyl group, a dichloromethyl group, a trichloromethyl group, a 2,2,2-trichloroethyl group, a 2,2,3,3,3-pentachloropropyl group, a 2,2,2-trichloro-1-trichloromethylethyl group, a 1,1-bis(trichloromethyl)-2,2,2-trichloroethyl group, a 4-chlorophenyl group, a 2,6-dichlorophenyl group, a 3,5-dichlorophenyl group, a 2,4,6-trichlorophenyl group or a pentachlorophenyl group. Particularly preferred is a fluorine atom, a fluoroalkyl group or a fluoroaryl group, and the most preferred is a fluorine atom.

The symbol R^4 is a hydrogen atom or a hydrocarbon group. The hydrocarbon group as R^4 is preferably an alkyl group, an aryl group or an aralkyl group, and examples thereof are a hydrogen atom and those similar to the hydrocarbon groups explained as L^1 in the above-mentioned general formula [1]. Preferred as R^4 is a hydrogen atom, an alkyl group or an aryl group, more preferred is a hydrogen atom or an alkyl group, and particularly preferred is a hydrogen atom.

In the above general formula [3], R^2 is a hydrocarbon group or a halogenated hydrocarbon group. The hydrocarbon group in R^2 is preferably an alkyl group, an aryl group or an aralkyl group, and the same hydrocarbon group as explained in the general formula [1] as L^1 is used. As the halogenated hydrocarbon group as R^2 , a halogenated alkyl group, a halogenated aryl group and a (halogenated alkyl)aryl group are enumerated, and those enumerated as specific examples of the electron-withdrawing group as R^1 in the general formula [2] are used as the halogenated alkyl group, the halogenated aryl group and the (halogenated alkyl)aryl group.

In the above general formula [3], R^2 is preferably a halogenated hydrocarbon group, and more preferably a fluorinated hydrocarbon group.

Specific examples of the compound (a) having a zinc atom as M^1 are dialkylzincs such as dimethylzinc, diethylzinc, dipropylzinc, di-n-butylzinc, di-isobutylzinc and di-n-hexylzinc; diarylzincs such as diphenylzinc,

dinaphthylzinc and bis(pentafluorophenyl)zinc; dialkenylzincs such as diallylzinc; bis(cyclopentadienyl)zinc; alkylzinc halides such as methylzinc chloride, ethylzinc chloride, propylzinc chloride, n-butylzinc chloride, isobutylzinc chloride, n-hexylzinc chloride, methylzinc bromide, ethylzinc bromide, propylzinc bromide, n-butylzinc bromide, isobutylzinc bromide, n-hexylzinc bromide, methylzinc iodide, ethylzinc iodide, propylzinc iodide, n-butylzinc iodide, isobutylzinc iodide and n-hexylzinc iodide; and zinc halides such as zinc fluoride, zinc chloride, zinc bromide and zinc iodide.

Dialkylzincs are preferred as the compound (a). More preferred is dimethylzinc, diethylzinc, dipropylzinc, di-n-butylzinc, di-isobutylzinc or di-n-hexylzinc, and particularly preferred is dimethylzinc or diethylzinc.

Specific examples of the compound (b) other than that having a group represented by the general formula [4] as R^1 are amines such as di(fluoromethyl)amine, di(chloromethyl)amine, di(bromomethyl)amine, di(iodomethyl)amine, bis(difluoromethyl)amine, bis(dichloromethyl)amine, bis(dibromomethyl)amine, bis(diiodomethyl)amine, bis(trifluoromethyl)amine, bis(trichloromethyl)amine, bis(tribromomethyl)amine, bis(triiodomethyl)amine, bis(2,2,2-trifluoroethyl)amine, bis(2,2,2-trichloroethyl)amine, bis(2,2,2-tribromoethyl)amine, bis(2,2,2-triiodoethyl)amine, bis(2,2,3,3,3-pentafluoropropyl)amine, bis(2,2,3,3,3-pentachloropropyl)amine, bis(2,2,3,3,3-pentabromopropyl)amine,

bis(2,2,3,3,3-pentaiodopropyl)amine,
 bis(2,2,2-trifluoro-1-trifluoromethylethyl)amine,
 bis(2,2,2-trichloro-1-trichloromethylethyl)amine,
 bis(2,2,2-tribromo-1-tribromomethylethyl)amine,
 5 bis(2,2,2-triiodo-1-triiodomethylethyl)amine,
 bis(1,1-bis(trifluoromethyl)-2,2,2-trifluoroethyl)amine,
 bis(1,1-bis(trichloromethyl)-2,2,2-trichloroethyl)amine,
 bis(1,1-bis(tribromomethyl)-2,2,2-tribromoethyl)amine,
 bis(1,1-bis(triiodomethyl)-2,2,2-triiodoethyl)amine,
 10 bis(2-fluorophenyl)amine, bis(3-fluorophenyl)amine,
 bis(4-fluorophenyl)amine, bis(2-chlorophenyl)amine,
 bis(3-chlorophenyl)amine, bis(4-chlorophenyl)amine,
 bis(2-bromophenyl)amine, bis(3-bromophenyl)amine,
 bis(4-bromophenyl)amine, bis(2-iodophenyl)amine,
 15 bis(3-iodophenyl)amine, bis(4-iodophenyl)amine,
 bis(2,6-difluorophenyl)amine, bis(3,5-difluorophenyl)amine,
 bis(2,6-dichlorophenyl)amine, bis(3,5-dichlorophenyl)amine,
 bis(2,6-dibromophenyl)amine, bis(3,5-dibromophenyl)amine,
 bis(2,6-diiodophenyl)amine, bis(3,5-diiodophenyl)amine,
 20 bis(2,4,6-trifluorophenyl)amine,
 bis(2,4,6-trichlorophenyl)amine,
 bis(2,4,6-tribromophenyl)amine,
 bis(2,4,6-triiodophenyl)amine, bis(pentafluorophenyl)amine,
 bis(pentachlorophenyl)amine, bis(pentabromophenyl)amine,
 25 bis(pentaiodophenyl)amine,
 bis(2-(trifluoromethyl)phenyl)amine,
 bis(3-(trifluoromethyl)phenyl)amine,

bis(4-(trifluoromethyl)phenyl)amine,

bis(2,6-di(trifluoromethyl)phenyl)amine,

bis(3,5-di(trifluoromethyl)phenyl)amine,

bis(2,4,6-tri(trifluoromethyl)phenyl)amine,

5 bis(2-cyanophenyl)amine, (3-cyanophenyl)amine,

bis(4-cyanophenyl)amine, bis(2-nitrophenyl)amine,

bis(3-nitrophenyl)amine and bis(4-nitrophenyl)amine. In

addition, phosphine compounds having a phosphorus atom in place
of the nitrogen atom are also exemplified. Such phosphine

10 compounds are those expressed by rewriting the term amine in
the above-mentioned specific examples with the term phosphine.

Further specific examples of compound (b) are alcohols such
as fluoromethanol, chloromethanol, bromomethanol, iodomethanol,
difluoromethanol, dichloromethanol, dibromomethanol,

15 diiodomethanol, trifluoromethanol, trichloromethanol,

tribromomethanol, triiodomethanol, 2,2,2-trifluoroethanol,

2,2,2-trichloroethanol, 2,2,2-tribromoethanol,

2,2,2-triiodoethanol, 2,2,3,3,3-pentafluoropropanol,

2,2,3,3,3-pentachloropropanol, 2,2,3,3,3-pentabromopropanol,

20 2,2,3,3,3-pentaiodopropanol,

2,2,2-trifluoro-1-trifluoromethylethanol,

2,2,2-trichloro-1-trichloromethylethanol,

2,2,2-tribromo-1-tribromomethylethanol,

2,2,2-triiodo-1-triiodomethylethanol,

25 1,1-bis(trifluoromethyl)-2,2,2-trifluoroethanol,

1,1-bis(trichloromethyl)-2,2,2-trichloroethanol,

1,1-bis(tribromomethyl)-2,2,2-tribromoethanol and

1,1-bis(triiodomethyl)-2,2,2-triiodoethanol. In addition, thiol compounds having a sulfur atom in place of the oxygen atom are also exemplified. Such thiol compounds are those expressed by rewriting the terms methanol, ethanol and propanol in the
5 above-mentioned specific examples with the terms methanethiol, ethanethiol and propanethiol, respectively.

Still further specific examples of the compound (b) are phenols such as 2-fluorophenol, 3-fluorophenol, 4-fluorophenol, 2,4-difluorophenol, 2,6-difluorophenol, 3,4-difluorophenol,
10 3,5-difluorophenol, 2,4,6-trifluorophenol, 2,3,5,6-tetrafluorophenol, pentafluorophenol, 2,3,5,6-tetrafluoro-4-trifluoromethylphenol, 2,3,5,6-tetrafluoro-4-pentafluorophenol, perfluoro-1-naphthol, perfluoro-2-naphthol, 2-chlorophenol,
15 3-chlorophenol, 4-chlorophenol, 2,4-dichlorophenol, 2,6-dichlorophenol, 3,4-dichlorophenol, 3,5-dichlorophenol, 2,4,6-trichlorophenol, 2,3,5,6-tetrachlorophenol, pentachlorophenol, 2,3,5,6-tetrachloro-4-trichloromethylphenol,
20 2,3,5,6-tetrachloro-4-pentachlorophenol, perchloro-1-naphthol, perchloro-2-naphthol, 2-bromophenol, 3-bromophenol, 4-bromophenol, 2,4-dibromophenol, 2,6-dibromophenol, 3,4-dibromophenol, 3,5-dibromophenol, 2,4,6-tribromophenol, 2,3,5,6-tetrabromophenol,
25 pentabromophenol, 2,3,5,6-tetrabromo-4-tribromomethylphenol, 2,3,5,6-tetrabromo-4-pentabromophenol, perbromo-1-naphthol, perbromo-2-naphthol, 2-iodophenol, 3-iodophenol, 4-iodophenol,

2,4-diiodophenol, 2,6-diiodophenol, 3,4-diiodophenol,
 3,5-diiodophenol, 2,4,6-triiodophenol,
 2,3,5,6-tetraiodophenol, pentaiodophenol,
 2,3,5,6-tetraiodo-4-triiodomethylphenol,
 5 2,3,5,6-tetraiodo-4-pentaiodophenol, periodo-1-naphthol,
 periodo-2-naphthol, 2-(trifluoromethyl)phenol,
 3-(trifluoromethyl)phenol, 4-(trifluoromethyl)phenol,
 2,6-bis(trifluoromethyl)phenol,
 3,5-bis(trifluoromethyl)phenol,
 10 2,4,6-tris(trifluoromethyl)phenol, 2-cyanophenol,
 3-cyanophenol, 4-cyanophenol, 2-nitrophenol, 3-nitrophenol and
 4-nitrophenol. In addition, thiophenol compounds having a sulfur
 atom in place of the oxygen atom are also exemplified. Such
 thiophenol compounds are those expressed by rewriting the term
 15 phenol in the above-mentioned specific examples with the term
 thiophenol.

As the compound (b), preferred are amines such as
 bis(trifluoromethyl)amine, bis(2,2,2-trifluoromethyl)amine,
 bis(2,2,3,3,3-pentafluoropropyl)amine,
 20 bis(2,2,2-trifluoro-1-trifluoromethylethyl)amine,
 bis(1,1-bis(trifluoromethyl)-2,2,2-trifluoroethyl)amine or
 bis(pentafluorophenyl)amine, alcohols such as
 trifluoromethanol, 2,2,2-trifluoroethanol,
 2,2,3,3,3-pentafluoropropanol,
 25 2,2,2-trifluoro-1-trifluoromethylethanol or
 1,1-bis(trifluoromethyl)-2,2,2-trifluoroethanol, and phenols
 such as 2-fluorophenol, 3-fluorophenol, 4-fluorophenol,

2,6-difluorophenol, 3,5-difluorophenol,
2,4,6-trifluorophenol, pentafluorophenol,
2-(trifluoromethyl)phenol, 3-(trifluoromethyl)phenol,
4-(trifluoromethyl)phenol, 2,6-bis(trifluoromethyl)phenol,
5 3,5-bis(trifluoromethyl)phenol or
2,4,6-tris(trifluoromethyl)phenol.

More preferred as the compound (b) are
bis(trifluoromethyl)amine, bis(pentafluorophenyl) amine,
trifluoromethanol, 2,2,2-trifluoro-1-trifluoromethylethanol,
10 1,1-bis(trifluoromethyl)-2,2,2-trifluoroethanol,
2-fluorophenol, 3-fluorophenol, 4-fluorophenol,
2,6-difluorophenol, 3,5-difluorophenol,
2,4,6-trifluorophenol, pentafluorophenol,
4-(trifluoromethyl)phenol, 2,6-bis(trifluoromethyl)phenol
15 and 2,4,6-tris(trifluoromethyl)phenol. Much more preferred are
3,5-difluorophenol, pentafluorophenol and
1,1-bis(trifluoromethyl)-2,2,2-trifluoroethanol.

Specific examples of the compound (b) having a group
represented by the above general formula [4] as R¹ are phenols
20 such as 3,4,5-trifluorophenol,
3,4,5-tris(trifluoromethyl)phenol,
3,4,5-tris(pentafluorophenyl)phenol,
3,5-difluoro-4-pentafluorophenylphenol,
4,5,6,7,8-pentafluoro-2-naphthol, 3,4,5-trichlorophenol,
25 3,4,5-tris(trichloromethyl)phenol,
3,4,5-tris(pentachlorophenyl)phenol,
3,5-dichloro-4-pentachlorophenylphenol,

4,5,6,7,8-pentachloro-2-naphthol, 3,4,5-tribromophenol,
 3,4,5-tris(tribromomethyl)phenol,
 3,4,5-tris(pentabromophenyl)phenol,
 3,5-dibromo-4-pentabromophenylphenol,
 5 4,5,6,7,8-pentabromo-2-naphthol, 3,4,5-triiodophenol,
 3,4,5-tris(triiodomethyl)phenol,
 3,4,5-tris(pentaiodophenyl)phenol,
 3,5-diiodo-4-pentaiodophenylphenol,
 4,5,6,7,8-pentaio-2-naphthol, 3,5-difluoro-4-nitrophenol,
 10 3,5-dichloro-4-nitrophenol, 3,5-dibromo-4-nitrophenol,
 3,5-diiodo-4-nitrophenol, 3,5-difluoro-4-cyanophenol,
 3,5-dichloro-4-cyanophenol, 3,5-dibromo-4-cyanophenol and
 3,5-diiodo-4-cyanoophenol. In addition, thiophenol compounds
 having a sulfur atom in place of the oxygen atom are also
 15 exemplified. Such thiophenol compounds are those expressed by
 rewriting the term phenol in the above-mentioned specific
 examples with the term thiophenol. Furthermore, there are also
 exemplified phenol compounds or thiophenol compounds, whose
 hydrogen atoms at their 2- and 6-positions (in the case of naphthol,
 20 hydrogen atoms at 1- and 3-positions) are substituted with a
 methyl group, an ethyl group, a n-propyl group, an isopropyl
 group, a n-butyl group, an isobutyl group or a tert-butyl group.

As amines, there are enumerated

bis(3,4,5-trifluorophenyl)amine,
 25 bis(3,4,5-tris(trifluoromethyl)phenyl)amine,
 bis(3,4,5-tris(pentafluorophenyl)phenyl)amine,
 bis(3,5-difluoro-4-pentafluorophenylphenyl)amine,

bis(4,5,6,7,8-pentafluoro-2-naphthyl)amine,
 bis(3,4,5-trichlorophenyl)amine,
 bis(3,4,5-tris(trichloromethyl)phenyl)amine,
 bis(3,4,5-tris(pentachlorophenyl)phenyl)amine,
 5 bis(3,5-dichloro-4-pentachlorophenylphenyl)amine,
 bis(4,5,6,7,8-pentachloro-2-naphthyl)amine,
 bis(3,4,5-tribromophenyl)amine,
 bis(3,4,5-tris(tribromomethyl)phenyl)amine,
 bis(3,4,5-tris(pentabromophenyl)phenyl)amine,
 10 bis(3,5-dibromo-4-pentabromophenylphenyl)amine,
 bis(4,5,6,7,8-pentabromo-2-naphthyl)amine,
 bis(3,4,5-triiodophenyl)amine,
 bis(3,4,5-tris(triiodomethyl)phenyl)amine,
 bis(3,4,5-tris(pentaiodophenyl)phenyl)amine,
 15 bis(3,5-diiodo-4-pentaiodophenylphenyl)amine and
 bis(4,5,6,7,8-pentaiodo-2-naphthyl)amine. In addition,
 phosphine compounds having a phosphorus atom in place of the
 nitrogen atom are also exemplified. Such phosphine compounds
 are those expressed by rewriting the term amine in the
 20 above-mentioned specific examples with the term phosphine.
 Furthermore, there are also exemplified amine compounds or
 phosphine compounds, whose hydrogen atoms at their 2- and
 6-positions (in the case of naphthol, hydrogen atoms at 1- and
 3-positions) are substituted with a methyl group, an ethyl group,
 25 an-n-propyl group, an isopropyl group, an-n-butyl group, an isobutyl
 group or a tert-butyl group.

Preferred as the compound (b) are phenols such as

3,4,5-trifluorophenol, 3,4,5-tris(trifluoromethyl)phenol,
3,4,5-tris(pentafluorophenyl)phenol,
3,5-difluoro-4-pentafluorophenylphenol and
4,5,6,7,8-pentafluoro-2-naphthol, and amines such as
5 bis(3,4,5-trifluorophenyl)amine,
bis(3,4,5-tris(trifluoromethyl)phenyl)amine,
bis(3,4,5-tris(pentafluorophenyl)phenyl)amine,
bis(3,5-difluoro-4-pentafluorophenylphenyl)amine and
bis(4,5,6,7,8-pentafluoro-2-naphthyl)amine.

10 More preferred as the compound (b) are
3,4,5-trifluorophenol, 4,5,6,7,8-pentafluoro-2-naphthol,
bis(3,4,5-trifluorophenyl)amine and
bis(4,5,6,7,8-pentafluoro-2-naphthyl)amine. Particularly
preferred is 3,4,5-trifluorophenol.

15 Specific examples of the compounds (c) are water, hydrogen
sulfide, an alkylamine, an arylamine, an aralkylamine, a
halogenated alkylamine, a halogenated arylamine, and a
(halogenated alkyl)arylamine. More preferred are water,
hydrogen sulfide, methylamine, ethylamine, n-propylamine,
20 isopropylamine, n-butylamine, sec-butylamine,
tert-butylamine, isobutylamine, n-pentylamine,
neopentylamine, isopentylamine, n-hexylamine, n-octylamine,
n-decylamine, n-dodecylamine, n-pentadecylamine,
n-eicosylamine, allylamine, cyclopentadienylamine, aniline,
25 2-tolylamine, 3-tolylamine, 4-tolylamine, 2,3-xylylamine,
2,4-xylylamine, 2,5-xylylamine, 2,6-xylylamine,
3,4-xylylamine, 3,5-xylylamine, 2,3,4-trimethylaniline,

(n-butylphenyl)methylamine, (sec-butylphenyl)methylamine,
 (tert-butylphenyl)methylamine,
 (n-pentylphenyl)methylamine, (neopentylphenyl)methylamine,
 (n-hexylphenyl)methylamine, (n-octylphenyl)methylamine,
 5 (n-decylphenyl)methylamine,
 (n-tetradecylphenyl)methylamine, naphthylmethylamine,
 anthracenylmethylamine, fluoromethylamine,
 chloromethylamine, bromomethylamine, iodomethylamine,
 difluoromethylamine, dichloromethylamine,
 10 dibromomethylamine, diiodomethylamine,
 trifluoromethylamine, trichloromethylamine,
 tribromomethylamine, triiodomethylamine,
 2,2,2-trifluoroethylamine, 2,2,2-trichloroethylamine,
 2,2,2-tribromoethylamine, 2,2,2-triiodoethylamine,
 15 2,2,3,3,3-pentafluoropropylamine,
 2,2,3,3,3-pentachloropropylamine,
 2,2,3,3,3-pentabromopropylamine,
 2,2,3,3,3-pentaiodopropylamine,
 2,2,2-trifluoro-1-trifluoromethylethylamine,
 20 2,2,2-trichloro-1-trichloromethylethylamine,
 2,2,2-tribromo-1-tribromomethylethylamine,
 2,2,2-triiodo-1-triiodomethylethylamine,
 1,1-bis(trifluoromethyl)-2,2,2-trifluoroethylamine,
 1,1-bis(trichloromethyl)-2,2,2-trichloroethylamine,
 25 1,1-bis(tribromomethyl)-2,2,2-tribromoethylamine,
 1,1-bis(triiodomethyl)-2,2,2-triiodoethylamine,
 perfluoropropylamine, perchloropropylamine,

	perbromopropylamine,	perfluorobutylamine,
	perchlorobutylamine,	perbromobutylamine,
	perfluoropentylamine,	perchloropentylamine,
	perbromopentylamine,	perfluorohexylamine,
5	perchlorohexylamine,	perbromohexylamine,
	perfluorooctylamine,	perchlorooctylamine,
	perbromooctylamine,	perfluorododecylamine,
	perchlorododecylamine,	perbromododecylamine,
	perfluoropentadecylamine,	perchloropentadecylamine,
10	perbromopentadecylamine,	perfluoroeicosylamine,
	perchloroeicosylamine,	perbromoeicosylamine,
	2-fluoroaniline,	3-fluoroaniline,
	2-chloroaniline,	3-chloroaniline,
	2-bromoaniline,	3-bromoaniline,
15	2-iodoaniline,	3-iodoaniline,
	2,6-difluoroaniline,	3,5-difluoroaniline,
	2,6-dichloroaniline,	3,5-dichloroaniline,
	2,6-dibromoaniline,	3,5-dibromoaniline,
	2,6-diiodoaniline,	3,5-diiodoaniline,
	3,5-diiodoaniline,	2,4,6-trifluoroaniline,
20	2,4,6-trichloroaniline,	2,4,6-tribromoaniline,
	2,4,6-triiodoaniline,	3,4,5-trifluoroaniline,
	3,4,5-trichloroaniline,	3,4,5-tribromoaniline,
	3,4,5-triiodoaniline,	pentafluoroaniline,
	pentachloroaniline,	pentabromoaniline,
	pentaiodoaniline,	
25	2-(trifluoromethyl)aniline,	3-(trifluoromethyl)aniline,
	4-(trifluoromethyl)aniline,	
	2,6-di(trifluoromethyl)aniline,	

3,5-di(trifluoromethyl)aniline,
2,4,6-tri(trifluoromethyl)aniline and
3,4,5-tri(trifluoromethyl)aniline.

Preferred as the compound (c) are water, hydrogen sulfide,
5 methylamine, ethylamine, n-propylamine, isopropylamine,
n-butylamine, sec-butylamine, tert-butylamine,
isobutylamine, n-octylamine, aniline, 2,6-xylylamine,
2,4,6-trimethylaniline, naphthylamine, anthracenylamine,
benzylamine, trifluoromethylamine, pentafluoroethylamine,
10 perfluoroproylamine, perfluorobutylamine,
perfluoropentylamine, perfluorohexylamine,
perfluorooctylamine, perfluorododecylamine,
perfluoropentadecylamine, perfluoroeicosylamine,
2-fluoroaniline, 3-fluoroaniline, 4-fluoroaniline,
15 2,6-difluoroaniline, 3,5-difluoroaniline,
2,4,6-trifluoroaniline, 3,4,5-trifluoroaniline,
pentafluoroaniline, 2-(trifluoromethyl)aniline,
3-(trifluoromethyl)aniline, 4-(trifluoromethyl)aniline,
2,6-bis(trifluoromethyl)aniline,
20 3,5-bis(trifluoromethyl)aniline and
2,4,6-tris(trifluoromethyl)aniline. Particularly preferred
are water, trifluoromethylamine, perfluorobutylamine,
perfluorooctylamine, perfluoropentadecylamine,
2-fluoroaniline, 3-fluoroaniline, 4-fluoroaniline,
25 2,6-difluoroaniline, 3,5-difluoroaniline,
2,4,6-trifluoroaniline, pentafluoroaniline,
2-(trifluoromethyl)aniline, 3-(trifluoromethyl)aniline,

4-(trifluoromethyl)aniline,
2,6-bis(trifluoromethyl)aniline,
3,5-bis(trifluoromethyl)aniline,
2,4,6-tris(trifluoromethyl)aniline and
5 3,4,5-tris(trifluoromethyl)aniline. The most preferred are
water and pentafluoroaniline.

As the particle (d), that which is generally used as a carrier
is preferably used. A porous substance having a uniform
particle diameter is preferred and an inorganic substance or
10 an organic polymer is preferably used. An inorganic substance
is more preferably used.

From a viewpoint of particle size distribution of a polymer
obtained, the particle (d) has a geometric standard deviation
of preferably not more than 2.5, more preferably not more than
15 2.0, and much more preferably not more than 1.7, wherein the
geometric standard deviation is based on a volume of a particle
diameter of the particle (d).

Examples of the inorganic substance used as the particle
(d) are inorganic oxides. It is also permitted to use clay and
20 clay mineral. These inorganic substances may be used in a
mixture.

Specific examples of the inorganic oxide include
SiO₂(silica), Al₂O₃, MgO, ZrO₂, TiO₂, B₂O₃, CaO, ZnO, BaO, ThO₂
and mixtures thereof such as SiO₂(silica)-MgO,
25 SiO₂(silica)-Al₂O₃, SiO₂(silica)-TiO₂, SiO₂(silica)-V₂O₅,
SiO₂(silica)-Cr₂O₃ and SiO₂(silica)-TiO₂-MgO. Among these
inorganic oxides, SiO₂(silica) and/or Al₂O₃ are/is preferred,

and SiO_2 (silica) is particularly preferred. Incidentally, the above inorganic oxide may contain a small amount of carbonates, sulfates, nitrates and oxide components, such as Na_2CO_3 , K_2CO_3 , CaCO_3 , MgCO_3 , Na_2SO_4 , $\text{Al}_2(\text{SO}_4)_3$, BaSO_4 , KNO_3 , $\text{Mg}(\text{NO}_3)_2$, $\text{Al}(\text{NO}_3)_3$,
5 Na_2O , K_2O and Li_2O .

Examples of the clay or clay mineral are kaolin, bentonite, Kibushi clay, gaerome clay, allophane, hisingerite, pyrophyllite, talc, a mica group, a montmorillonite group, vermiculite, a chlorite group, palygorskite, kaolinite, nacrite, dickite and
10 halloysite.

Among these, smectite, montmorillonite, hectorite, raponite and saponite are preferred, and montmorillonite and hectorite are more preferred.

Among these inorganic substances, inorganic oxides are
15 suitably used.

It is preferable that these inorganic substances are dried to contain substantially no water, and those dried by heat-treatment are preferred. The heat-treatment of inorganic substances, whose water content cannot be visually confirmed,
20 is usually carried out at a temperature of from 100 to 1500°C, preferably from 100 to 1000°C, and more preferably from 200 to 800°C. The heating time is not particularly limited, and preferably from 10 minutes to 50 hours, more preferably from 1 hour to 30 hours. Further, during heating, there can be applied
25 a method of passing a dried inert gas (e.g. nitrogen or argon) at a fixed flow rate, or a method of evacuating. The method is not limited thereto.

Further, it is permitted to use a modified inorganic oxide obtained by substituting an active hydrogen of a surface hydroxide group of the inorganic oxide with a various kind of a substituent. In this case, a preferable substituent is a silyl group. More specifically, there are enumerated inorganic oxides treated with a halogenated alkylsilane compound such as trimethylchlorosilane, tert-butyldimethylchlorosilane and dimethyldichlorosilane, and an alkylidisilazane compound such as 1,1,1,3,3,3-hexamethyldisilazane.

An average particle diameter of the inorganic substance is preferably from 5 to 1000 μm , more preferably from 10 to 500 μm , and more preferably from 10 to 100 μm . A pore volume is preferably not less than 0.1 ml/g, and more preferably 0.3 to 10ml/g. The specific surface area is preferably from 10 to 1000 m^2/g , and more preferably from 100 to 500 m^2/g .

The organic polymer usable as the particle (d) may be any organic polymer, and it is permitted to use a mixture of more than one organic polymer. As the organic polymer, a polymer having an active hydrogen-carrying functional group or a functional group of a non-proton-donating Lewis base is preferred.

The active hydrogen-carrying functional group is not particularly limited as far as it has active hydrogen. Examples thereof are a primary amino group, a secondary amino group, an imino group, an amide group, a hydrazide group, an amidino group, a hydroxy group, a hydroperoxy group, a carboxyl group, a formyl group, a carbamoyl group, a sulfonic acid group, a

sulfinic acid group, a sulfenic acid group, a thiol group, a thioformyl group, a pyrrolyl group, an imidazolyl group, a piperidyl group, an indazolyl group and a carbazolyl group. Among them, preferred is a primary amino group, a secondary amino group, an imino group, an amide group, an imide group, a hydroxy group, a formyl group, a carboxyl group, a sulfonic acid group or a thiol group. Particularly preferred is a primary amino group, a secondary amino group, an amide group or a hydroxy group. These groups may be substituted with a halogen atom or a hydrocarbon group having 1 to 20 carbon atoms.

The functional group of a non-proton-donating Lewis base is not particularly limited as far as it is a functional group having a Lewis base portion containing no active hydrogen atom. Specific examples thereof are a pyridyl group, an N-substituted imidazolyl group, an N-substituted indazolyl group, a nitrile group, an azido group, an N-substituted imino group, an N,N-substituted amino group, an N,N-substituted aminoxy group, an N,N,N-substituted hydrazino group, a nitroso group, a nitro group, a nitroxy group, a furyl group, a carbonyl group, a thiocarbonyl group, an alkoxy group, an alkyloxycarbonyl group, an N,N-substituted carbamoyl group, a thioalkoxy group, a substituted sulfinyl group, a substituted sulfonyl group and a substituted sulfonic acid group. Preferred is a heterocyclic group, and more preferred is an aromatic heterocyclic group having an oxygen atom and/or a nitrogen atom in the ring. Particularly preferred are a pyridyl group, an N-substituted imidazolyl group and an N-substituted indazolyl group, and the

most preferred is a pyridyl group. These groups may be substituted with a halogen atom or a hydrocarbon group having 1 to 20 carbon atoms.

5 An amount of the active hydrogen-carrying functional group or the functional group of a non-proton-donating Lewis base is not particularly limited. It is preferably from 0.01 to 50 mmol/g, and more preferably from 0.1 to 20 mmol/g, in terms of a molar amount of the functional group per g of the organic polymer.

10 The functional group-carrying organic polymer can be obtained, for example, by homopolymerizing a monomer having the active hydrogen-carrying functional group or the functional group of a non-proton-donating Lewis base and more than one polymerizable unsaturated group, or copolymerizing said monomer and another monomer having a polymerizable unsaturated group. At this time, it is preferred to additionally
15 copolymerize a crosslinking polymerizable monomer having two or more polymerizable unsaturated groups.

Examples of the monomer having the active hydrogen-carrying functional group or the functional group of
20 a non-proton-donating Lewis base and more than one polymerizable unsaturated group are monomers having the above-mentioned active hydrogen-carrying functional group and more than one polymerizable unsaturated group, and monomers having the above-mentioned functional group having a Lewis base portion
25 containing no active hydrogen and more than one polymerizable unsaturated group. Examples of the polymerizable unsaturated group are an alkenyl groups such as a vinyl group and an allyl

group, and an alkynyl group such as an ethyne group.

Examples of the monomer having the active hydrogen-carrying functional group and more than one polymerizable unsaturated group are a vinyl group-containing primary amine, a vinyl
5 group-containing secondary amine, a vinyl group-containing amide compound and a vinyl group-containing hydroxy compound. Specific examples thereof are N-(1-ethenyl)amine, N-(2-propenyl)amine, N-(1-ethenyl)-N-methylamine, N-(2-propenyl)-N-methylamine, 1-ethenylamide,
10 2-propenylamide, N-methyl-(1-ethenyl)amide, N-methyl-(2-propenyl)amide, vinyl alcohol and 2-propen-1-ol and 3-buten-1-ol.

Specific examples of the monomer having the functional group having a Lewis base portion containing no active hydrogen and
15 more than one polymerizable unsaturated group are vinylpyridine, vinyl(N-substituted)imidazole and vinyl(N-substituted)indazole.

The other monomers having a polymerizable unsaturated group include ethylene, α -olefins and aromatic vinyl compounds, and
20 specific examples thereof are ethylene, propylene, 1-butene, 1-hexene, 4-methyl-1-pentene and styrene. Preferred is ethylene or styrene. These monomers may be used in a combination of two or more.

As specific examples of the crosslinking polymerizable
25 monomer having two or more polymerizable unsaturated groups, divinylbenzene can be given.

An average particle diameter of the organic polymer is

preferably from 5 to 1000 μm , and more preferably from 10 to 500 μm . A pore volume is preferably not less than 0.1 ml/g, and more preferably 0.3 to 10 ml/g. A specific surface area is preferably from 10 to 1000 m^2/g , and more preferably from 50 to 500 m^2/g .

It is preferable that these organic polymers are dried to contain substantially no water, and those dried by heat-treatment are preferred. The heat-treatment of organic polymers, whose water content cannot be visually confirmed, is usually carried out at a temperature of from 30 to 400°C, preferably from 50 to 200°C, and more preferably from 70 to 150°C. The heating time is not particularly limited, and preferably from 30 minutes to 50 hours, more preferably from 1 hour to 30 hours. Further, during heating, there can be used a method of passing a dried inert gas (e.g. nitrogen or argon) at a fixed flow rate, or a method of evacuating. The method is not limited thereto.

The modified particle in accordance with the present invention is a modified particle obtained by contacting the above-mentioned (a), (b), (c) and (d) with one another. An order for contacting (a), (b), (c) and (d) is not particularly limited, and the following orders are enumerated.

- <1> A contact product obtained by contacting a contact product between (a) and (b) with (c) is contacted with (d).
- <2> A contact product obtained by contacting a contact product between (a) and (b) with (d) is contacted with (c).
- <3> A contact product obtained by contacting a contact product

between (a) and (c) with (b) is contacted with (d).

<4> A contact product obtained by contacting a contact product between (a) and (c) with (d) is contacted with (b).

5 <5> A contact product obtained by contacting a contact product between (a) and (d) with (b) is contacted with (c).

<6> A contact product obtained by contacting a contact product between (a) and (d) with (c) is contacted with (b).

<7> A contact product obtained by contacting a contact product between (b) and (c) with (a) is contacted with (d).

10 <8> A contact product obtained by contacting a contact product between (b) and (c) with (d) is contacted with (a).

<9> A contact product obtained by contacting a contact product between (b) and (d) with (a) is contacted with (c).

15 <10> A contact product obtained by contacting a contact product between (b) and (d) with (c) is contacted with (a).

<11> A contact product obtained by contacting a contact product between (c) and (d) with (a) is contacted with (b).

<12> A contact product obtained by contacting a contact product between (c) and (d) with (b) is contacted with (a).

20 As the contact order, preferred is the above-mentioned <1>, <2>, <3>, <11> or <12>.

It is recommendable to carry out the contact treatment under an inert gas atmosphere. A treating temperature is usually from -100 to 300°C, and preferably from -80 to 200°C. A treating
25 time is usually from 1 minute to 200 hours, and preferably from 10 minutes to 100 hours. It is permitted to carry out such a treatment using a solvent or directly treat the compounds without

use of any solvent.

As the solvent, those inert to either respective components to be contacted or the contact products obtained by the contact when used are usually used. In the case where respective
5 components are contacted step by step as described above, there is a case where a certain solvent reacts with, for example, the above-mentioned component (a), but the solvent does not react with a contact product obtained by contacting the component (a) with another component. In such a case, the solvent can be used
10 in a contact operation wherein said contact product is regarded as one component. There are exemplified the following solvents, which should be appropriately distinguished in the applications in such a manner. Examples of solvents usable are non-polar solvents such as aliphatic hydrocarbon solvents and aromatic
15 hydrocarbon solvents and polar solvents such as halide solvents, ether solvents, alcohol solvents, phenol solvents, carbonyl solvents, phosphoric acid derivatives, nitrile solvents, nitro compounds, amine solvents and sulfur compounds. Specific examples thereof are aliphatic hydrocarbon solvents such as
20 butane, pentane, hexane, heptane, octane, 2,2,4-trimethylpentane and cyclohexane, aromatic hydrocarbon solvents such as benzene, toluene and xylene, halide solvents such as dichloromethane, difluoromethane, chloroform, 1,2-dichloroethane, 1,2-dibromoethane,
25 1,1,2-trichloro-1,2,2-trifluoroethane, tetrachloroethylene, chlorobenzene, bromobenzene and o-dichlorobenzene, ether

solvents such as dimethyl ether, diethyl ether, diisopropyl ether, di-n-butyl ether, methyl-tert-butyl ether, anisole, 1,4-dioxane, 1,2-dimethoxyethane, bis(2-methoxyethyl) ether, tetrahydrofuran and tetrahydropyran, alcohol solvents such as
5 methanol, ethanol, 1-propanol, 2-propanol, 1-butanol, 2-butanol, 2-methyl-1-propanol, 3-methyl-1-butanol, cyclohexanol, benzyl alcohol, ethylene glycol, propylene glycol, 2-methoxy ethanol, 2-ethoxy ethanol, diethylene glycol, triethylene glycol and glycerol, phenol solvents such as phenol
10 and p-cresol, carbonyl solvents such as acetone, ethyl methyl ketone, cyclohexanone, acetic anhydride, ethyl acetate, butyl acetate, ethylene carbonate, propylene carbonate, N,N-dimethylformamide, N,N-dimethylacetamide and N-methyl-2-pyrrolidone, phosphoric acid derivatives such as
15 hexamethylphosphate triamide and triethyl phosphate, nitrile solvents such as acetonitrile, propionitrile, succinonitrile and benzonitrile, nitro compounds such as nitromethane and nitrobenzene, amine solvents such as pyridine, piperidine and morpholine, and sulfur compounds such as dimethylsulfoxide and
20 sulfolane.

In the case where a contact product (e) obtained by contacting (a), (b) and (c) with one another is contacted with the particle (d), namely in the above-mentioned <1>, <3> and <7>, the above-mentioned aliphatic hydrocarbon solvents,
25 aromatic hydrocarbon solvents or ether solvents are preferred as a solvent (s1) used for the production of the contact product

(e).

Whereas, a polar solvent is preferred as a solvent (s2) used for contacting the contact product (e) and the particle (d) with each other. There is known an E_T^N value as an index
5 indicating a polarity of a solvent (C. Reichardt, "Solvents and Solvents Effects in Organic Chemistry", 2nd ed., VCH Verlag (1988)). Particularly preferred is a solvent satisfying $0.8 \geq E_T^N \geq 0.1$. Examples of such a solvent are dichloromethane, dichlorodifluoromethane, chloroform, 1,2-dichloroethane,
10 1,2-dibromoethane, 1,1,2-trichloro-1,2,2-trifluoroethane, tetrachloroethylene, chlorobenzene, bromobenzene, o-dichlorobenzene, dimethyl ether, diethyl ether, diisopropyl ether, di-n-butyl ether, methyl tert-butyl ether, anisole, 1,4-dioxane, 1,2-dimethoxyethane, bis(2-methoxyethyl) ether,
15 tetrahydrofuran, tetrahydropyran, methanol, ethanol, 1-propanol, 2-propanol, 1-butanol, 2-butanol, 2-methyl-1-propanol, 3-methyl-1-butanol, cyclohexanol, benzyl alcohol, ethylene glycol, propylene glycol, 2-methoxy ethanol, 2-ethoxy ethanol, diethylene glycol, triethylene glycol,
20 acetone, ethyl methyl ketone, cyclohexanone, acetic anhydride, ethyl acetate, butyl acetate, ethylene carbonate, propylene carbonate, N,N-dimethylformamide, N,N-dimethylacetamide, N-methyl-2-pyrrolidone, hexamethylphosphate triamide, triethyl phosphate, acetonitrile, propionitrile,
25 succinonitrile, benzonitrile, nitromethane, nitrobenzene, ethylenediamine, pyridine, piperidine, morpholine,

dimethylsulfoxide and sulfolane.

More preferred as the solvent (s2) are dimethyl ether, diethyl ether, diisopropyl ether, di-n-butyl ether, methyl tert-butyl ether, anisole, 1,4-dioxane, 1,2-dimethoxyethane, bis(2-methoxyethyl) ether, tetrahydrofuran, tetrahydropyran, methanol, ethanol, 1-propanol, 2-propanol, 1-butanol, 2-butanol, 2-methyl-1-propanol, 3-methyl-1-butanol, cyclohexanol, benzyl alcohol, ethylene glycol, propylene glycol, 2-methoxy ethanol, 2-ethoxy ethanol, diethylene glycol and triethylene glycol. Particularly preferred are di-n-butyl ether, methyl tert-butyl ether, 1,4-dioxane, tetrahydrofuran, methanol, ethanol, 1-propanol, 2-propanol, 1-butanol, 2-butanol, 2-methyl-1-propanol, 3-methyl-1-butanol and cyclohexanol. The most preferred are tetrahydrofuran, methanol, ethanol, 1-propanol and 2-propanol.

As the above-mentioned solvent (s2), it is permitted to use a mixed solvent of these polar solvents and hydrocarbon solvents. As the hydrocarbon solvents, the above-exemplified aliphatic hydrocarbon solvents and aromatic hydrocarbon solvents are used. Specific examples of the mixed solvents of the polar solvents and the hydrocarbon solvents are a hexane/methanol mixed solvent, a hexane/ethanol mixed solvent, a hexane/1-propanol mixed solvent, a hexane/2-propanol mixed solvent, a heptane/methanol mixed solvent, a heptane/ethanol mixed solvent, a heptane/1-propanol mixed solvent, a heptane/2-propanol mixed solvent, a toluene/methanol mixed

solvent, a toluene/ethanol mixed solvent, a toluene/1-propanol mixed solvent, a toluene/2-propanol mixed solvent, a xylene/methanol mixed solvent, a xylene/ethanol mixed solvent, a xylene/1-propanol mixed solvent and a xylene/2-propanol mixed solvent. Preferable are a hexane/methanol mixed solvent, a hexane/ethanol mixed solvent, a heptane/methanol mixed solvent, a heptane/ethanol mixed solvent, a toluene/methanol mixed solvent, a toluene/ethanol mixed solvent, a xylene/methanol mixed solvent and a xylene/ethanol mixed solvent. More preferred are a hexane/methanol mixed solvent, a hexane/ethanol mixed solvent, a toluene/methanol mixed solvent and a toluene/ethanol mixed solvent. The most preferred is a toluene/ethanol mixed solvent.

A proportion of ethanol in the toluene/ethanol mixed solvent is preferably from 10 to 50% by volume, and more preferably from 15 to 30% by volume.

In the process wherein the contact product (e) obtained by contacting (a), (b) and (c) with one another is contacted with the particle (d), namely in the above-mentioned <1>, <3> and <7>, it is permitted to use the hydrocarbon solvent as either the solvent (s1) or the solvent (s2). In this regard, it is preferred that a time interval from completion of the contact of (a), (b) and (c) to the contact between the obtained contact product (e) and the particle (d) is shorter. The time interval is preferably from 0 to 5 hours, more preferably from 0 to 3 hours, and the most preferably from 0 to 1 hour. A temperature

at which the contact product (e) and the particle (d) are contacted with each other is usually from -100 to 40°C, preferably from -20 to 20°C, and the most preferably from -10 to 10°C.

In the above-mentioned <2>, <5>, <6>, <8>, <9>, <10>, <11>
5 and <12>, either the above-mentioned non-polar solvents or polar solvents can be used. The non-polar solvents are more preferable. Because, it seems that the contact product between (a) and (c) and the contact product obtained by contacting the contact product between (a) and (b) with (c) generally exhibit a low
10 solubility to the non-polar solvent, and therefore, when (d) exists in the reaction system at the time when such contact products are produced, the produced contact products are more steadily precipitated on the surface of (d) rather than exist in the non-polar solvent, so that the contact products are more
15 easily subjected to fixation.

Amounts of respective compounds (a), (b) and (c) mentioned above are not particularly limited. However, it is preferred that y and z substantially satisfy the following expression (1), provided that a molar ratio of the amounts of respective compounds
20 is expressed by (a): (b): (c) = 1: y: z.

$$|m - y - 2z| \leq 1 \quad (1)$$

(In the above expression (1), m is a valence of M¹.)

In the above expression (1), y is preferably a number of from 0.01 to 1.99, more preferably from 0.10 to 1.80, much more
25 preferably from 0.20 to 1.50, and the most preferably from 0.30 to 1.00. A similarly preferable range of z in the above expression

(1) can be determined by m, y and the above expression (1).

In a practical contact treatment of the compounds, it is usual that the amounts used delicately fluctuate even if the compounds are to be used so as to perfectly satisfy the above
5 expression (1), or taking some amounts of unreacted compounds into consideration, the amounts to be used are somewhat increased or decreased. The phrase "substantially satisfy the expression (1)" does not mean that the expression (1) is perfectly satisfied, but it is included to intent to obtain an object product, which
10 will be obtained when contacting respective compounds in a molar ratio satisfying the expression (1).

In preparing the modified particle in accordance with the present invention, it is permitted to appropriately determine a proportion of (d) to (a), provided that a typical metal atom
15 originated from (a) and contained in a particle obtained by contacting (a) and (d) with each other is controlled so as to be preferably not less than 0.1 mmol, and more preferably from 0.5 to 20 mmol, in terms of a mol number of the typical metal atom contained in 1 g of the obtained particle.

20 After completion of the contact treatment as mentioned above, it is recommendable to apply heating, so that the reaction further proceeds. In heating, it is preferred to use a solvent having a higher boiling point to make the heating temperature much higher. In this regard, it is permitted to substitute the
25 solvent used for the contact treatment with another solvent having a higher boiling point.

As a result of such a contact treatment, at least one of the starting (a), (b), (c) and/or (d) may remain as an unreacted material in the modified particle in accordance with the present invention. However, when it is applied to polymerization
5 accompanied by the formation of addition polymer particles, it is recommendable to apply a washing treatment for removing the unreacted material in advance. A solvent used therefor may be the same as or different from that used at the time of contacting.

After completion of the contact treatment and the washing
10 treatment, the solvent is removed from the product by distillation, and the resulting product is dried under reduced pressure preferably at a temperature of not lower than 25°C for 1 to 24 hours, more preferably at a temperature of from 40 to 200°C for 1 to 24 hours, much more preferably at a temperature
15 of from 60 to 200°C for 1 to 24 hours, particularly preferably at a temperature of from 60 to 160°C for 2 to 18 hours, and the most preferably at a temperature of from 80 to 160°C for 4 to 18 hours.

A specific example of a process for producing the modified
20 particle in accordance with the present invention is explained in more detail as follows, wherein M¹ is a zinc atom, the compound (b) is pentafluorophenol, the compound (c) is water and the particle (d) is silica. A hexane solution of diethylzinc is added to tetrahydrofuran as a solvent, the mixture is cooled to 3°C,
25 pentafluorophenol in an amount equimolar to the diethylzinc is dropped thereto, the resulting mixture is stirred at ambient

temperature for 10 minutes to 24 hours, thereafter water in an amount of 0.5 mol per mol of diethylzinc is further added, and the resulting mixture is stirred at ambient temperature for 10 minutes to 24 hours. Thereafter, the solvent is removed by
5 distillation, and the residue is dried under reduced pressure at 120°C for 8 hours. To a solid component obtained through the above-mentioned operations, tetrahydrofuran and silica are added, and the mixture is stirred at 40°C for 2 hours. The resulting solid component is washed with tetrahydrofuran, and
10 dried under reduced pressure at 120°C for 8 hours. Thus, the modified particle in accordance with the present invention can be obtained. Among the modified particles obtained in such a manner, a modified particle which is obtained using a compound (b) having a halogenated hydrocarbon group as R¹ in the general
15 formula [2], and which satisfies the following expressions [5] and [6] or [5] and [7], is preferred from a viewpoint of polymerization activity.

$$N/M > 0.9 \quad [5]$$

(In the expression, N is a substance amount of a halogen atom
20 contained in the modified particle, and M is a substance amount of a typical metal atom M¹ contained in the modified particle.)

$$A/B \geq 0.1 \quad [6]$$

(In the expression, A is an integral intensity of halo showing a peak at Bragg angle (2θ) 33° to 37° in a diffraction intensity
25 profile obtained by an extended X-ray measurement, and B is an integral intensity of halo showing a peak at Bragg angle (2

θ) 18° to 22° in said diffraction intensity profile.)

$$D/C \geq 0.5 \quad [7]$$

(In the expression, C is a peak intensity of the maximum peak present within a range of 1 to 2 nm in a radial distribution function obtained by measuring the modified particle according to an X-ray absorption fine structure analysis, and D is a peak intensity of the maximum peak present within a range of 2.5 to 3.5 nm in said radial distribution function, provided that the radial distribution function is obtained in a manner such that the modified particle is measured according to an X-ray absorption fine structure analysis (XAFS) to obtain X-ray absorption spectra, from which extended X-ray absorption fine structure (EXAFS) spectra of the typical metal atom M^1 in a K absorption edge are obtained and processed according to a Fourier transform.)

A detailed explanation is directed to [5], [6] and [7] as follows.

In the above expression [5], M is a substance amount of a typical metal atom M^1 of the Group 1, 2, 12, 14 or 15. The M can be found by various means generally used in an analytical chemistry (for example, "Encyclopedia of Experimental Chemistry Analysis 4th edition", edited by Japan Chemical Society, issued by Maruzen Co., Ltd. pages 2 and 3). Among these means, a gravimetric method by chemical analysis and a spectroscopy are preferred. Because of easy operations, a spectroscopy is more preferred. Further, of these, an atomic absorption

spectroscopy (AAS analysis) and an inductively coupled plasma-emission spectroscopy (ICP analysis) are particularly preferred for the analysis of trace metal atom.

In the above expression [5], N is a substance amount of
5 a halogen atom contained in the modified particle. The substance
amount of a halogen atom can be found according to a gravimetric
method, an absorptiometry or an ion electrode method as disclosed
in "Encyclopedia of Experimental Chemistry 15 Analysis 4th
edition", edited by Japan Chemical Society, issued by Maruzen
10 Co., Ltd. pages 218 to 231.

In the modified particle in accordance with the present
invention, as clear from the above expression [5], the halogen
atom is so many that the ratio of the substance amount of the
halogen atom to the typical metal atom M^1 contained in the modified
15 particle exceeds 0.9. The ratio (N/M) is preferably within a
range exceeding 0.90.

The extended X-ray measurement in the present
invention is carried out under the following conditions using,
for example, vertical rotating anode X-ray generator ultra X18
20 manufactured by Rigaku International Corporation, as an X-ray
diffraction apparatus.

Load: 50kV-100 mA

Focus: line

Scan mode: $2\theta / \theta$ (2° /min.)

25 Measured angle range: 5° to 70°

Slit: divergence slit 1°

receiving slit 0.15 mm

scatter slit 1°

Detector: scintillation counter

Monochromater: graphite

5 The extended X-ray measurement in the present invention is carried out usually under a condition that a sample to be measured is not exposed to atmosphere.

 In a diffraction intensity profile obtained by the extended X-ray measurement, an integral intensity A of halo showing a
10 peak at Bragg angle (2θ) 33° to 37° and an integral intensity B of halo showing a peak at Bragg angle (2θ) 18° to 22° are determined, and its ratio A/B is calculated.

 It is conceivable that the halo showing a peak at Bragg angle (2θ) 33° to 37° is a halo based on the contact product
15 of the above-mentioned (a), (b) and (c), and the halo showing a peak at Bragg angle (2θ) 18° to 22° is that based on the particle (d). It is preferred that the above-mentioned integral intensity ratio A/B is above a level, and thereby, the catalyst for addition polymerization obtained using the modified particle
20 increases in its activity. In the present invention, the above-mentioned integral intensity ratio A/B is not less than 0.1, preferably from 0.15 to 1, more preferably from 0.2 to 0.8, and particularly preferably from 0.25 to 0.7.

 The above-mentioned expression [7] is an expression formed
25 by the use of an extended X-ray absorption fine structure (EXAFS) analysis. The following explanation is directed to a general

X-ray absorption fine structure (XAFS) analysis. A full detail of the principles and analysis methods is given in literature such as "The Spectroscopical Society of Japan Sokuteihou Series 26 , X-ray absorption fine structure, edited by Yasuo Udagawa (1993)".

When a substance is placed on a line of X-ray beams, an X-ray absorbance of the substance can be calculated from both an intensity of an X-ray irradiated to the substance (incoming X-ray intensity: I_0) and an intensity of an X-ray transmitted through the substance (transmitted X-ray intensity: I_t). In the process of measuring X-ray absorption spectra by varying X-ray energy while monitoring increase and decrease of the X-ray absorbance, it can be observed that the X-ray absorbance suddenly rises at certain X-ray energy (absorption edge). The X-ray energy of the absorption edge corresponds to energy required at the time when an inner shell electron of an atom absorbing X-rays (absorption atom) jumps out as a photoelectron. In an atom, there are various kinds of inner shells such as K shell, L shell and M shell, and there are absorption edges corresponding to respective inner shell electrons. The K shell electron exists in an innermost part, and a corresponding absorption edge is called a K absorption edge. In the X-ray absorption spectra, a fine vibration structure appearing at a region of energy higher than said absorption edge by about 30 to 1000 eV is said to be an extended X-ray absorption fine structure (EXAFS). Such a fine vibration structure appearing on the spectra is caused as a result

of an interference effect between a wave of the photoelectron discharged from the absorption atom owing to absorption of X-rays and a wave of the returning photoelectron owing to scatter caused by a neighboring atom (scattering atom). Accordingly, as a result
5 of detailed analysis, there can be obtained an information about a local structure in the vicinity of the absorption atom.

A background is deducted from the X-ray absorption spectra, and supposing that the absorption atom is isolated, an absorption coefficient having no vibration structure is further deducted
10 therefrom, thereby obtaining EXAFS spectra. The EXAFS spectra are processed according to a Fourier transform in an appropriate region to obtain a radial distribution function focusing on the absorption atom. By examining the radial distribution function in detail, it is possible to obtain an information about a distance
15 between the absorption atom and the scatter atom and another information about a number of the scatter atom from a position of the maximum value in said function and its intensity, respectively, and as a result, there can be clarified a noticeable structural information in the vicinity of the absorption atom.

20 A peak intensity of the maximum peak present within a range of from 1 to 2 nm in the radial distribution function is taken as C, and a peak intensity of the maximum peak present within a range of from 2.5 to 3.5 nm therein is taken as D, and its ratio D/C is calculated. It is conceivable that the peak present
25 within a range of from 1 to 2 nm in the radial distribution function is based on the existence of an atom ① bound to the typical

metal atom M^1 , and the peak present within a range of from 2.5 to 3.5 nm in the radial distribution function is based on the existence of an atom ② bound to the atom ① or coordinated to the atom ①. The peak intensity D of the peak based on the existence of the atom ② increases with increase in numbers of the atom ② present in the vicinity of the typical metal atom M^1 , and this may demonstrate that a structure containing the typical metal atom M^1 , the atom ① and the atom ② becomes a more orderly structure. It is preferred that the ratio of these peak intensity, D/C, is above a level, and thereby, the catalyst for addition polymerization obtained using the modified particle increases in its activity. In the present invention, the above-mentioned peak intensity ratio D/C is not less than 0.5, preferably from 0.55 to 1.2, and more preferably from 0.6 to 0.9.

Usually, such an XAFS measurement is carried out under a condition that a sample to be measured is not exposed to atmosphere.

While, the modified particle meeting the above-mentioned conditions satisfies the expression (1), and further satisfies,

$$2 \leq z/y < 3 \quad (2),$$

wherein y is preferably a number of from 0.20 to 0.42, more preferably a number of from 0.30 to 0.41, and much more preferably a number of from 0.35 to 0.40. The ratio of y to z (z/y) is preferably a number of from 2.2 to 2.8.

Additionally, in the present invention, a typical metal compound component, which is selected from the contact products

(e) obtained by contacting the above-mentioned (a), the above-mentioned (b) and the above-mentioned (c) with one another, and which is insoluble in the following solvent (f) is contacted with the above-mentioned particle (d), thereby obtaining a modified particle. The thus obtained modified particle is also preferred from a viewpoint of polymerization activity.

Solvent (f): a mixed solvent or tetrahydrofuran, wherein the mixed solvent is a mixture of tetrahydrofuran and hexane and satisfies the following expression [8],

$$0.5 \geq V_H / (V_T + V_H) \quad [8]$$

wherein V_T is a volume of tetrahydrofuran, and V_H is a volume of hexane.

It is permitted to use those commercially available as hexane and tetrahydrofuran stated here. It is recommendable to use a supernatant liquid of a mixture of said commercially available product and a dehydrating agent such as silica gel and molecular sieve, or a product put on the market as a dehydrated product (for example, that manufactured by Kanto Kagaku).

The contact product (e) is prepared by contacting the above-mentioned component (a), component (b) and component (c) with one another. An order for contacting these components is not particularly limited, and exemplified as follows.

<13> A contact product between (a) and (b) is contacted with (c).

<14> A contact product between (a) and (c) is contacted with (b).

<15> A contact product between (b) and (c) is contacted with (a).

When (d) participates therein, the above orders correspond to the above-mentioned <1>, <3> and <7>, respectively.

5 As the contacting order, preferred is the above-mentioned <13> or <15>.

The conditions of the contact treatment including treating temperature, treating atmosphere, treating time, presence or absence of a solvent and kind thereof, and using ratios of (a),
10 (b) and (c), are the same as those mentioned above.

Such a typical metal compound component in the present invention can be obtained by contacting the above-mentioned (a), (b) and (c) in the above-mentioned solvent (f) to precipitate an insoluble product.

15 In the production process, an amount of the solvent (f) (VT + VH) to (a) is usually from 0.5 to 5 liters, preferably from 0.7 to 3 liters, and more preferably from 1.0 to 1.5 liters per substance amount (mol) of (a). When the above-mentioned compounds (a), (b) and (c) are used in respective solutions,
20 solvents used therefor may be the same or different from one another in their compositions, provided that the solvent in the treatment liquid obtained after completion of contacting all compounds satisfies the above-mentioned expression [5].

Further, the above-mentioned typical metal compound
25 component can be obtained in a manner such that the above-mentioned (a), (b) and (c) are contacted with one another

in various kinds of the solvent as already mentioned to obtain the contact product (e), the solvent used is removed, successively the contact product (e) is added to the above-mentioned solvent (f), and then a component soluble to
5 the above solvent (f) is removed by filtration or the like.

In this production process, an amount of the solvent (f) (VT + VH) to (e) is usually from 5 to 100 milliliters, preferably from 7 to 50 milliliters, and more preferably from 10 to 20 milliliters per g of the contact product (e).

10 As the typical metal compound component, among the contact products (e), preferred from a viewpoint of higher activity is that insoluble to the following solvent (f1) but soluble to the following solvent (f2).

Solvent (f1): a mixed solvent of tetrahydrofuran and hexane
15 satisfying the following expression [9-1],

$$0.3 \leqslant VH/(VT + VH) \leqslant 0.5 \quad [9-1]$$

wherein VT is a volume of tetrahydrofuran, and VH is a volume of hexane.

Solvent (f2): a mixed solvent or tetrahydrofuran, wherein
20 the mixed solvent is a mixture of tetrahydrofuran and hexane and satisfies the following expression [9-2],

$$0.3 > VH/(VT + VH) \quad [9-2]$$

wherein VT is a volume of tetrahydrofuran, and VH is a volume of hexane.

25 Such a preferable typical metal compound component can be obtained in a manner such that the above-mentioned (a), (b)

and (c) are contacted with one another in various kinds of the solvent as already mentioned to obtain the contact product (e), the solvent used is removed, successively the contact product (e) is added to the above-mentioned solvent (f2), then a component insoluble to the above solvent (f2) is removed by filtration or the like, thereafter a component obtained by removing the above solvent (f2) is added to the above-mentioned solvent (f1), and then a component soluble to the above solvent (f1) is removed by filtration or the like. Here, as an operation carried out after removing the component insoluble to the above-mentioned solvent (f2) by filtration or the like, an operation comprising the steps of adding a fixed amount of hexane to the solution (such as filtrate obtained by filtration), from which the component insoluble to the above-mentioned solvent (f2) has been removed, until the above expression [9-2] is satisfied, and obtaining an insoluble product precipitated by filtration or the like is simple and therefor applicable as a suitable process. In this production process, an amount of the solvent (f1) to the contact product (e) is usually from 2 to 80 milliliters, preferably from 5 to 40 milliliters, and more preferably from 10 to 12 milliliters per g of the contact product (e). An amount of the solvent (f2) is usually from 2 to 80 milliliters, preferably from 5 to 40 milliliters, and more preferably from 9 to 11 milliliters per g of the contact product (e).

The parameter in the present invention, $VH/(VT + VH)$, is an index indicating a polarity of the mixed solvent of hexane

and tetrahydrofuran. With respect to the E_T^N values of hexane and tetrahydrofuran, E_T^N (hexane) = 0.009, and E_T^N (tetrahydrofuran) = 0.207, and therefore, the value of $VH/(VT + VH)$ increases with decrease in the polarity of the mixed solvent.

5 In the present invention, it is permitted to replace the solvent (f), the solvent (f1) and the solvent (f2) used for the above-mentioned processes for producing the typical metal compound component of the present invention with a single solvent or a mixed solvent, which is different from such solvents but
10 has a polarity equal to their E_T^N .

By contacting the above-mentioned typical metal compound component and the above-mentioned particle (d) with each other, the modified particle can be obtained. Conditions of the contact treatment are the same as those in the above-mentioned contact
15 treatment of the contact product (e) and the particle (d). By using the thus obtained modified particle, a catalyst having a higher polymerization activity can be obtained.

While, the above-mentioned typical metal compound component is also useful as a catalyst component for addition
20 polymerization (particularly as a catalyst component for olefin polymerization). As the catalyst for addition polymerization obtained using such a typical metal compound component, there are exemplified a catalyst for addition polymerization obtained by contacting the above-mentioned typical metal compound
25 component and a transition metal compound (B) of the Groups 3 to 11 or lanthanoid series with each other, and a catalyst for

addition polymerization obtained by contacting the above-mentioned typical metal compound component, a transition metal compound (B) of the Groups 3 to 11 or lanthanoide series and an organoaluminum compound (C) with one another. Of these,
5 the latter is higher in activity and more preferred.

The modified particle in accordance with the present invention is useful as a carrier for supporting the catalyst component for addition polymerization comprising the transition metal compound capable of forming a single site catalyst, which
10 is suitably used for polymerization accompanied by the formation of addition polymer particles. In addition, the modified particle in accordance with the present invention is useful as the catalyst component for addition polymerization (particularly the catalyst component for olefin polymerization). As the catalyst
15 component for addition polymerization in accordance with the present invention, there are exemplified a catalyst for addition polymerization obtained by contacting the above-mentioned modified particle (A) and a transition metal compound (B) of the Groups 3 to 11 or lanthanoide series with each other, and
20 a catalyst for addition polymerization obtained by contacting the above-mentioned modified particle (A), a transition metal compound (B) of the Groups 3 to 11 or lanthanoide series and an organoaluminum compound (C) with one another. Of these, the latter is higher in activity and more preferred.

25 (B) Transition metal compound of the Groups 3 to 11 or Lanthanoide Series

As the transition metal compound (B) of the Groups 3 to 11 or lanthanoide series used for the catalyst for addition polymerization in accordance with the present invention, a transition metal compound capable of forming a single site catalyst is used. It is not particularly limited as far as it is a transition metal compound of the Groups 3 to 11 or lanthanoide series, and exhibits addition polymerization activity when used in combination with the above-mentioned modified particle (A) (and additionally organoaluminum compound (C)) as a co-catalyst component for activation use. Incidentally, the single site catalyst stated here is conceptually distinguished from a conventional type solid catalyst, and includes not only a narrow sense single site catalyst giving an addition polymer of a narrow molecular weight distribution or a narrow composition distribution in the case of copolymer, but also a catalyst giving an addition polymer of a broad molecular weight distribution or a broad composition distribution in the case of a copolymer, as far as the catalyst is obtained in a preparation manner similar to that of the narrow sense single site catalyst.

As such a transition metal compound (B), preferred is a transition metal compound represented by the following general formula [10] or its μ -oxo type transition metal compound dimer.



(In the formula, M^2 is a transition metal atom of the Groups 3 to 11 or lanthanoide series, L^2 is a cyclopentadienyl type anion skeleton-carrying group or a hetero atom-containing group, more than one L^2 may be linked directly or through a residual

group containing a carbon atom, a silicone atom, a nitrogen atom, an oxygen atom, a sulfur atom or a phosphorus atom, X is a halogen atom, a hydrocarbon group (excluding the cyclopentadienyl type anion skeleton-carrying group) or $-OR^{11}$ (R^{11} is a hydrocarbon group or a halogenated hydrocarbon group, and more than one R^{11} may be the same or different from each other), a is a number satisfying $0 < a \leq 8$, and b is a number satisfying $0 < b \leq 8$.)

In the general formula [10], M^2 is a transition metal atom of the Groups 3 to 11 of the periodic table (IUPAC 1989) or lanthanoide series. Examples thereof are a scandium atom, an yttrium atom, a titanium atom, a zirconium atom, a hafnium atom, a vanadium atom, a niobium atom, a tantalum atom, a chromium atom, an iron atom, a ruthenium atom, a cobalt atom, a rhodium atom, a nickel atom, a palladium atom, a samarium atom and an ytterbium atom. Preferred as M^2 in the general formula [10] is a titanium atom, a zirconium atom, a hafnium atom, a vanadium atom, a chromium atom, an iron atom, a cobalt atom or a nickel atom, and particularly preferred is a titanium atom, a zirconium atom or a hafnium atom.

In the general formula [10], L^2 is a cyclopentadienyl type anion skeleton-carrying group or a hetero atom-containing group, and more than one L^2 may be the same or different from one another. Further, more than one L^2 may be linked directly or through a residual group containing a carbon atom, a silicone atom, a nitrogen atom, an oxygen atom, a sulfur atom or a phosphorus atom.

Examples of the cyclopentadienyl type anion

skeleton-carrying group as L^2 are η^5 -(substituted) cyclopentadienyl groups, η^5 -(substituted)indenyl groups and η^5 -(substituted) fluorenyl groups. Specific examples thereof are an η^5 -cyclopentadienyl group, an η^5 -methylcyclopentadienyl group, an η^5 -ethylcyclopentadienyl group, an η^5 -n-butylcyclopentadienyl group, an η^5 -tert-butylcyclopentadienyl group, an η^5 -1,2-dimethylcyclopentadienyl group, an η^5 -1,3-dimethylcyclopentadienyl group, an η^5 -1-methyl-2-ethylcyclopentadienyl group, an η^5 -1-methyl-3-ethylcyclopentadienyl group, an η^5 -1-tert-butyl-2-methylcyclopentadienyl group, an η^5 -1-tert-butyl-3-methylcyclopentadienyl group, an η^5 -1-methyl-2-isopropylcyclopentadienyl group, an η^5 -1-methyl-3-isopropylcyclopentadienyl group, an η^5 -1-methyl-2-n-butylcyclopentadienyl group, an η^5 -1-methyl-3-n-butylcyclopentadienyl group, an η^5 -1,2,3-trimethylcyclopentadienyl group, an η^5 -1,2,4-trimethylcyclopentadienyl group, an η^5 -tetramethylcyclopentadienyl group, an η^5 -pentamethylcyclopentadienyl group, an η^5 -indenyl group, an η^5 -4,5,6,7-tetrahydroindenyl group, an η^5 -2-methylindenyl group, an η^5 -3-methylindenyl group, an η^5 -4-methylindenyl group, an η^5 -5-methylindenyl group, an η^5 -6-methylindenyl group, an η^5 -7-methylindenyl group, an η^5 -2-tert-butylindenyl group, an η^5 -3-tert-butylindenyl group, an η^5 -4-tert-butylindenyl group, an η^5 -5-tert-butylindenyl group,

an η^5 -6-tert-butylindenyl group, an η^5 -7-tert-butylindenyl group, an η^5 -2,3-dimethylindenyl group, an η^5 -4,7-dimethylindenyl group, an η^5 -2,4,7-trimethylindenyl group, an η^5 -2-methyl-4-isopropylindenyl group, an η^5 -4,5-benzindenyl group, an η^5 -2-methyl-4,5-benzindenyl group, an η^5 -4-phenylindenyl group, an η^5 -2-methyl-5-phenylindenyl group, an η^5 -2-methyl-4-phenylindenyl group, an η^5 -2-methyl-4-naphthylindenyl group, an η^5 -fluorenyl group, an η^5 -2,7-dimethylfluorenyl group, an η^5 -2,7-di-tert-butylfluorenyl group and their substitution compounds.

Incidentally, in the present specification, " η^5 -" in the name of the transition metal compound may be omitted in some cases.

The hetero atom in the hetero atom-containing group includes an oxygen atom, a sulfur atom, a nitrogen atom and a phosphorus atom. Examples of such a group are preferably an alkoxy group, an aryloxy group, a thioalkoxy group, a thioaryloxy group, an alkylamino group, an arylamino group, an alkylphosphino group, an arylphosphino group, a chelating ligand, and an aromatic or aliphatic heterocyclic group having an oxygen atom, a sulfur atom, a nitrogen atom and/or a phosphorus atom in the ring.

Specific examples of the hetero atom-containing group are a methoxy group, an ethoxy group, a propoxy group, a butoxy group, a phenoxy group, a 2-methylphenoxy group, a 2,6-dimethylphenoxy group, a 2,4,6-trimethylphenoxy group, a 2-ethylphenoxy group, a 4-n-propylphenoxy group, a 2-isopropylphenoxy group, a

2,6-diisopropylphenoxy group, a 4-sec-butylphenoxy group, a
 4-tert-butylphenoxy group, a 2,6-di-sec-butylphenoxy group, a
 2-tert-butyl-4-methylphenoxy group, a
 2,6-di-tert-butylphenoxy group, a 4-methoxyphenoxy group, a
 5 2,6-dimethoxyphenoxy group, a 3,5-dimethoxyphenoxy group, a
 2-chlorophenoxy group, a 4-nitrosophenoxy group, a
 4-nitrophenoxy group, a 2-aminophenoxy group, a 3-aminophenoxy
 group, a 4-aminothiophenoxy group, a 2,3,6-trichlorophenoxy
 group, a 2,4,6-trifluorophenoxy group, a thiomethoxy group, a
 10 dimethylamino group, a diethylamino group, a dipropylamino group,
 a diphenylamino group, an isopropylamino group, a
 tert-butylamino group, a pyrrolyl group, a dimethylphosphino
 group, a 2-(2-oxy-1-propyl)phenoxy group, catechol, resorcinol,
 4-isopropylcatechol, 3-methoxycatechol, a
 15 1,8-dihydroxynaphthyl group, a 1,2-dihydroxynaphthyl group, a
 2,2'-biphenyldiol group, a 1,1'-bi-2-naphthol group, a
 2,2'-dihydroxy-6,6'-dimethylbiphenyl group, a
 4,4',6,6'-tetra-tert-butyl-2,2'-methylenediphenoxy group and
 a 4,4',6,6'-tetramethyl-2,2'-isobutylidenediphenoxy group.

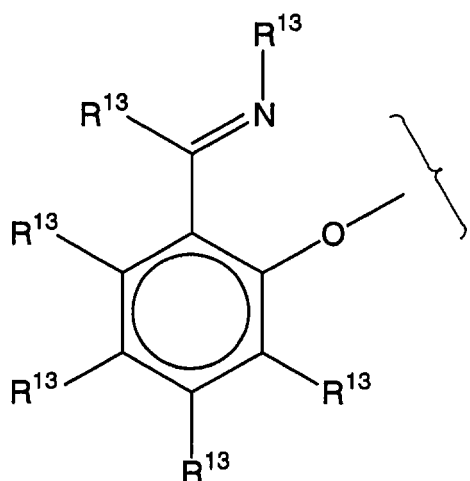
20 Further, as the above-mentioned hetero atom-containing
 group, there is exemplified a group represented by the following
 general formula [11].



(In the formula, respective R^{12} s are independently of one another
 25 a hydrogen atom, a halogen atom or a hydrocarbon group, and may
 be the same or different from one another, and two or more thereof
 may be bound with one another and may form a ring.)

Specific examples of R^{12} in the above general formula [11] are a hydrogen atom, a fluorine atom, a chlorine atom, a bromine atom, an iodine atom, a methyl group, an ethyl group, a n-propyl group, an isopropyl group, a n-butyl group, a tert-butyl group, a cyclopropyl group, a cyclobutyl group, a cycloheptyl group, a cyclohexyl group, a phenyl group, a 1-naphthyl group, a 2-naphthyl group and a benzyl group, but are not limited thereto.

Furthermore, as the hetero atom-containing group, there is also exemplified a group represented by the following formula [12].



[12]

(In the formula, respective R^{13} s are independently of one another a hydrogen atom, a halogen atom, a hydrocarbon group, a halogenated hydrocarbon group, a hydrocarbyloxy group, a silyl group or an amino group, and may be the same or different from one another, and two or more thereof may be bound with one another and may form a ring.)

Specific examples of R^{13} in the general formula [12] are

a hydrogen atom, a fluorine atom, a chlorine atom, a bromine atom, an iodine atom, a phenyl group, a 1-naphthyl group, a 2-naphthyl group, a tert-butyl group, a 2,6-dimethylphenyl group, a 2-fluorenyl group, a 2-methylphenyl group, a
5 4-trifluoromethylphenyl group, a 4-methoxyphenyl group, a 4-pyridyl group, a cyclohexyl group, a 2-isopropylphenyl group, a benzyl group, a methyl group, a triethylsilyl group, a diphenylmethylsilyl group, a 1-methyl-1-phenylethyl group, a 1,1-dimethylpropyl group, a 2-chlorophenyl group and a
10 pentafluorophenyl group, but are not limited thereto.

The above-mentioned chelating ligand means a ligand having more than one coordinating position. Specific examples thereof are acetylacetonate, diimine, oxazoline, bisoxazoline, terpyridine, acylhydrazone, diethylenetriamine,
15 triethylenetetramine, porphyrin, crown ether and cryptate.

Specific examples of the above-mentioned heterocyclic group are a pyridyl group, an N-substituted imidazolyl group and an N-substituted indazolyl group. Preferable is a pyridyl group.

20 Two cyclopentadienyl type anion skeleton-carrying groups, the cyclopentadienyl type anion skeleton-carrying group and the hetero atom-containing group, or two hetero atom-containing groups may be bound with each other directly or through a residual group containing a carbon atom, a silicone atom, a nitrogen atom,
25 an oxygen atom, a sulfur atom or a phosphorus atom, respectively. As such a residual group, preferred is a divalent residual group in which an atom binding two L²s is a carbon atom, a silicone

atom, a nitrogen atom, an oxygen atom, a sulfur atom and/or a phosphorus atom. More preferred is a divalent residual group in which an atom binding two L²s is a carbon atom, a silicone atom, a nitrogen atom, an oxygen atom, a sulfur atom and/or a phosphorus atom, provided that a minimum atom number of the atom binding two L²s is not more than 3 (a case where the atom binding two L²s is single is included). Specific examples thereof are alkylene groups such as a methylene group, an ethylene group and a propylene group, substituted alkylene groups such as a dimethylmethylene group (an isopropylidene group) and a diphenylmethylene group, a silylene group, a substituted silylene group such as a dimethylsilylene group, a diethylsilylene group, a diphenylsilylene group, a tetramethyldisilylene group and a dimethoxysilylene group, and a hetero atom such as a nitrogen atom, an oxygen atom, a sulfur atom and a phosphorus atom. Particularly preferred are a methylene group, an ethylene group, a dimethylmethylene group (an isopropylidene group), a diphenylmethylene group, a dimethylsilylene group, a diethylsilylene group, a diphenylsilylene group and a dimethoxysilylene group.

In the general formula [10], X is a halogen atom, a hydrocarbon group or OR¹¹. Particularly, OR¹¹ is preferable because of high polymerization activity.

Specific examples of the halogen atom are a fluorine atom, a chlorine atom, a bromine atom and an iodine atom. The hydrocarbon group stated here does not include the cyclopentadiene type anion skeleton-carrying group. Examples of the hydrocarbon group

stated here are an alkyl group, an aralkyl group, an aryl group and an alkenyl group. Preferred are an alkyl group having 1 to 20 carbon atoms, an aralkyl group having 7 to 20 carbon atoms, an aryl group having 6 to 20 carbon atoms and an alkenyl group
5 having 3 to 20 carbon atoms.

Examples of the alkyl group having 1 to 20 carbon atoms are a methyl group, an ethyl group, a n-propyl group, an isopropyl group, a n-butyl group, a sec-butyl group, a tert-butyl group, an isobutyl group, a n-pentyl group, a neopentyl group, an amyl
10 group, a n-hexyl group, a n-octyl group, a n-decyl group, a n-dodecyl group, a n-pentadecyl group and a n-eicosyl group. More preferred are a methyl group, an ethyl group, an isopropyl group, a tert-butyl group, an isobutyl group and an amyl group.

All of these alkyl groups may be substituted with a halogen
15 atom such as a fluorine atom, a chlorine atom, a bromine atom or an iodine atom. Examples of the alkyl group having 1 to 10 carbon atoms which is substituted with the halogen atom are a fluoromethyl group, a trifluoromethyl group, a chloromethyl group, a trichloromethyl group, a fluoroethyl group, a
20 pentafluoroethyl group, a perfluoropropyl group, a perfluorobutyl group, a perfluorohexyl group, a perfluorooctyl group, a perchloropropyl group, a perchlorobutyl group and a perbromopropyl group.

Further, all of these alkyl groups may be partially
25 substituted with an alkoxy group such as a methoxy group and an ethoxy group, an aryloxy group such as a phenoxy group or an aralkyloxy group such as a benzyloxy group.

Examples of the aralkyl group having 7 to 20 carbons are a benzyl group, a (2-methylphenyl)methyl group, a (3-methylphenyl)methyl group, a (4-methylphenyl)methyl group, a (2,3-dimethylphenyl)methyl group, a

5 (2,4-dimethylphenyl)methyl group, a (2,5-dimethylphenyl)methyl group, a (2,6-dimethylphenyl)methyl group, a (3,4-dimethylphenyl)methyl group, a (3,5-dimethylphenyl)methyl group, a

10 (2,3,4-trimethylphenyl)methyl group, a (2,3,5-trimethylphenyl)methyl group, a (2,3,6-trimethylphenyl)methyl group, a (3,4,5-trimethylphenyl)methyl group, a (2,4,6-trimethylphenyl)methyl group, a

15 (2,3,4,5-tetramethylphenyl)methyl group, a (2,3,4,6-tetramethylphenyl)methyl group, a (2,3,5,6-tetramethylphenyl)methyl group, a (pentamethylphenyl)methyl group, an (ethylphenyl)methyl group, a (n-propylphenyl)methyl group, an (isopropylphenyl)methyl

20 group, a (n-butylphenyl)methyl group, a (sec-butylphenyl)methyl group, a (tert-butylphenyl)methyl group, a (n-pentylphenyl)methyl group, a (neopentylphenyl)methyl group, a (n-hexylphenyl)methyl group, a (n-octylphenyl)methyl group, (n-decylphenyl)methyl group, a

25 (n-dodecylphenyl)methyl group, a naphthylmethyl group and an anthracenylmethyl group. More preferred is a benzyl group.

All of these aralkyl groups may be partially substituted

with a halogen atom such as a fluorine atom, a chlorine atom, a bromine atom or an iodine atom, an alkoxy group such as a methoxy group or an ethoxy group, an aryloxy group such as a phenoxy group or an aralkyloxy group such as a benzyloxy group.

5 Examples of the aryl group having 6 to 20 carbon atoms are a phenyl group, a 2-tolyl group, a 3-tolyl group, a 4-tolyl group, a 2,3-xylyl group, a 2,4-xylyl group, a 2,5-xylyl group, a 2,6-xylyl group, a 3,4-xylyl group, a 3,5-xylyl group, a 2,3,4-trimethylphenyl group, a 2,3,5-trimethylphenyl group, a 10 2,3,6-trimethylphenyl group, a 2,4,6-trimethylphenyl group, a 3,4,5-trimethylphenyl group, a 2,3,4,5-tetramethylphenyl group, a 2,3,4,6-tetramethylphenyl group, a 2,3,5,6-tetramethylphenyl group, a pentamethylphenyl group, an ethylphenyl group, a n-propylphenyl group, an isopropylphenyl group, a n-butylphenyl 15 group, a sec-butylphenyl group, a tert-butylphenyl group, a n-pentylphenyl group, a neopentylphenyl group, a n-hexylphenyl group, a n-octylphenyl group, a n-decylphenyl group, a n-dodecylphenyl group, a n-tetradecylphenyl group, a naphthyl group and an anthracenyl group. More preferred is a phenyl group.

20 All of these aryl groups may be partially substituted with a halogen atom such as a fluorine atom, a chlorine atom, a bromine atom or an iodine atom, an alkoxy group such as a methoxy group or an ethoxy group, an aryloxy group such as a phenoxy group or an aralkyloxy group such as a benzyloxy group.

25 Examples of the alkenyl group having 3 to 20 carbon atoms are an allyl group, a methallyl group, a crotyl group and a 1,3-diphenyl-2-propenyl group. More preferred are an allyl group

and a methallyl group.

When X is a halogen atom or a hydrocarbon group, preferred X are a chlorine atom, a methyl group, an ethyl group, a n-propyl group, an isopropyl group, a n-butyl group and a benzyl group.

5 In the general formula [10], a is a number satisfying $0 < a \leq 8$, b is a number satisfying $0 < b \leq 8$, and both are appropriately selected depending upon the valence of M^2 .

While, R^{11} in the OR^{11} is a hydrocarbon group or a halogenated hydrocarbon group. Examples of the hydrocarbon group are an alkyl group, an aralkyl group and an aryl group. Preferred are an alkyl group having 1 to 20 carbon atoms, an aralkyl group having 7 to 20 carbon atoms and an aryl group having 6 to 20 carbon atoms. Specific examples thereof are the same as those mentioned above. Examples of the halogenated hydrocarbon group are a halogenated alkyl group, a halogenated aryl group and a halogenated aralkyl group. Preferred are a halogenated alkyl group having 1 to 20 carbon atoms, a halogenated aryl group having 6 to 20 carbon atoms and a halogenated aralkyl group having 7 to 20 carbon atoms.

Specific examples of the halogenated alkyl group are a fluoromethyl group, a difluoromethyl group, a trifluoromethyl group, a chloromethyl group, a dichloromethyl group, a trichloromethyl group, a bromomethyl group, a dibromomethyl group, a tribromomethyl group, an iodomethyl group, a diiodomethyl group, a triiodomethyl group, a fluoroethyl group, a difluoroethyl group, a trifluoroethyl group, a tetrafluoroethyl group, a pentafluoroethyl group, a chloroethyl group, a dichloroethyl group, a trichloroethyl group, a

tetrachloroethyl group, a pentachloroethyl group, a bromoethyl group, a dibromoethyl group, a tribromoethyl group, a tetrabromoethyl group, a pentabromoethyl group, a perfluoropropyl group, a perfluorobutyl group, a
5 perfluoropentyl group, a perfluorohexyl group, a perfluorooctyl group, a perfluorododecyl group, a perfluoropentadecyl group, a perfluoroeicosyl group, a perchloropropyl group, a perchlorobutyl group, a perchloropentyl group, a perchlorohexyl group, a perchlorooctyl group, a perchlorododecyl group, a
10 perchloropentadecyl group, a perchloroeicosyl group, a perbromopropyl group, a perbromobutyl group, a perbromopentyl group, a perbromohexyl group, a perbromooctyl group, a perbromododecyl group, a perbromopentadecyl group and a perbromoeicosyl group.

15 Specific examples of the halogenated aryl group are a 2-fluorophenyl group, a 3-fluorophenyl group, a 4-fluorophenyl group, a 2,4-difluorophenyl group, a 2,6-difluorophenyl group, a 3,4-difluorophenyl group, a 3,5-difluorophenyl, a 3,4,5-trifluorophenyl group, a 2,4,6-trifluorophenyl group, a
20 2,3,5,6-tetrafluorophenyl group, a pentafluorophenyl group, a 2,3,5,6-tetrafluoro-4-trifluoromethylphenyl group, a 2,3,5,6-tetrafluoro-4-pentafluorophenylphenyl group, a perfluoro-1-naphthyl group, a perfluoro-2-naphthyl group, a 2-chlorophenyl group, a 3-chlorophenyl group, a 4-chlorophenyl
25 group, a 2,4-dichlorophenyl group, a 2,6-dichlorophenyl group, a 3,4-dichlorophenyl group, a 3,5-dichlorororophenyl, a 2,4,6-trichlorororophenyl group, a 2,3,5,6-tetrachlorophenyl

group, a pentachlorophenyl group, a
 2,3,5,6-tetrachloro-4-trichloromethylphenyl group, a
 2,3,5,6-tetrachloro-4-pentachlorophenylphenyl group, a
 perchloro-1-naphthyl group, a perchloro-2-naphthyl group, a
 5 2-bromophenyl group, a 3-bromophenyl group, a 4-bromophenyl
 group, a 2,4-dibromophenyl group, a 2,6-dibromophenyl group,
 a 3,4-dibromophenyl group, a 3,5-dibromophenyl, a
 2,4,6-tribromophenyl group, a 2,3,5,6-tetrabromophenyl group,
 a pentabromophenyl group, a
 10 2,3,5,6-tetrabromo-4-tribromomethylphenyl group, a
 2,3,5,6-tetrabromo-4-pentabromophenylphenyl group, a
 perbromo-1-naphthyl group, a perbromo-2-naphthyl group, a
 2-iodophenyl group, a 3-iodophenyl group, a 4-iodophenyl group,
 a 2,4-diiodophenyl group, a 2,6-diiodophenyl group, a
 15 3,4-diiodophenyl group, a 3,5-diiodophenyl, a
 2,4,6-triiodophenyl group, a 2,3,5,6-tetraiodophenyl group, a
 pentaiodophenyl group, a
 2,3,5,6-tetraiodo-4-triiodomethylphenyl group, a
 2,3,5,6-tetraiodo-4-pentaiodophenylphenyl group, a
 20 periodo-1-naphthyl group and a periodo-2-naphthyl group.

Examples of the halogenated aralkyl group are a
 tris(pentafluorophenyl)methyl group and
 bis(trifluoromethyl)phenylmethyl group.

As R^{11} , particularly preferred are a methyl group, an ethyl
 25 group, an isopropyl group, a tert-butyl group, an isobutyl group,
 a trifluoromethyl group, a perfluorobutyl group, a phenyl group,
 a 2,6-di(tert-butyl)phenyl group, a pentafluorophenyl group,

a 2,3,5,6-tetrafluoro-4-pentafluorophenylphenyl group and a benzyl group. As R¹¹, particularly preferred are a methyl group, a phenyl group, a 2,6-di(tert-butyl)phenyl group, a pentafluorophenyl group and a benzyl group, and the most
5 preferred is a phenyl group.

In the transition metal compound represented by the general formula [10], specific examples of those having a titanium atom, a zirconium atom or a hafnium metal as the transition metal atom are bis(cyclopentadienyl)titanium dichloride,
10 bis(methylcyclopentadienyl)titanium dichloride,
bis(ethylcyclopentadienyl)titanium dichloride,
bis(n-butylcyclopentadienyl)titanium dichloride,
bis(tert-butylcyclopentadienyl)titanium dichloride,
bis(1,2-dimethylcyclopentadienyl)titanium dichloride,
15 bis(1,3-dimethylcyclopentadienyl)titanium dichloride,
bis(1-methyl-2-ethylcyclopentadienyl)titanium dichloride,
bis(1-methyl-3-ethylcyclopentadienyl)titanium dichloride,
bis(1-methyl-2-n-butylcyclopentadienyl)titanium dichloride,
bis(1-methyl-3-n-butylcyclopentadienyl)titanium dichloride,
20 bis(1-methyl-2-isopropylcyclopentadienyl)titanium
dichloride,
bis(1-methyl-3-isopropylcyclopentadienyl)titanium
dichloride,
bis(1-tert-butyl-2-methylcyclopentadienyl)titanium
25 dichloride,
bis(1-tert-butyl-3-methylcyclopentadienyl)titanium
dichloride, bis(1,2,3-trimethylcyclopentadienyl)titanium

dichloride, bis(1,2,4-trimethylcyclopentadienyl)titanium
 dichloride, bis(tetramethylcyclopentadienyl)titanium
 dichloride, bis(pentamethylcyclopentadienyl)titanium
 dichloride,
 5 bis(indenyl)titanium dichloride,
 bis(4,5,6,7-tetrahydroindenyl)titanium dichloride,
 bis(fluorenyl)titanium dichloride,
 bis(2-phenylindenyl)titanium dichloride,
 bis[2-(bis-3,5-trifluoromethylphenyl)indenyl]titanium
 10 dichloride, bis[2-(4-tert-butylphenyl)indenyl]titanium
 dichloride, bis[2-(4-trifluoromethylphenyl)indenyl]titanium
 dichloride, bis[2-(4-methylphenyl)indenyl]titanium dichloride,
 bis[2-(3,5-dimethylphenyl)indenyl]titanium dichloride,
 bis[2-(pentafluorophenyl)indenyl]titanium dichloride,
 15 cyclopentadienyl(pentamethylcyclopentadienyl)titanium
 dichloride, cyclopentadienyl(indenyl)titanium dichloride,
 cyclopentadienyl(fluorenyl)titanium dichloride,
 indenyl(fluorenyl)titanium dichloride,
 pentamethylcyclopentadienyl(indenyl)titanium dichloride,
 20 pentamethylcyclopentadienyl(fluorenyl)titanium dichloride,
 cyclopentadienyl(2-phenylindenyl)titanium dichloride,
 pentamethylcyclopentadienyl(2-phenylindenyl)titanium
 dichloride, dimethylsilylenebis(cyclopentadienyl)titanium
 dichloride,
 25 dimethylsilylenebis(2-methylcyclopentadienyl)titanium
 dichloride, dimethylsilylenebis(3-methylcyclopentadienyl)
 titanium dichloride,

dimethylsilylenebis(2-n-butylcyclopentadienyl)titanium
 dichloride, dimethylsilylenebis(3-n-butylcyclopentadienyl)
 titanium dichloride,
 dimethylsilylenebis(2,3-dimethylcyclopentadienyl)titanium
 5 dichloride,
 dimethylsilylenebis(2,4-dimethylcyclopentadienyl) titanium
 dichloride,
 dimethylsilylenebis(2,5-dimethylcyclopentadienyl)titanium
 dichloride,
 10 dimethylsilylenebis(3,4-dimethylcyclopentadienyl) titanium
 dichloride,
 dimethylsilylenebis(2,3-ethylmethylcyclopentadienyl)titanium
 m dichloride,
 dimethylsilylenebis(2,4-ethylmethylcyclopentadienyl)titanium
 15 m dichloride,
 dimethylsilylenebis(2,5-ethylmethylcyclopentadienyl)titanium
 m dichloride,
 dimethylsilylenebis(3,5-ethylmethylcyclopentadienyl)titanium
 m dichloride,
 20 dimethylsilylenebis(2,3,4-trimethylcyclopentadienyl)titanium
 m dichloride,
 dimethylsilylenebis(2,3,5-trimethylcyclopentadienyl)titanium
 m dichloride,
 dimethylsilylenebis(tetramethylcyclopentadienyl)titanium
 25 dichloride, dimethylsilylenebis(indenyl) titanium dichloride,
 dimethylsilylenebis(2-methylindenyl)titanium dichloride,
 dimethylsilylenebis(2-tert-butylindenyl)titanium dichloride,

dimethylsilylenebis(2,3-dimethylindenyl)titanium dichloride,
 dimethylsilylenebis(2,4,7-trimethylindenyl)titanium
 dichloride,
 dimethylsilylenebis(2-methyl-4-isopropylindenyl)titanium
 5 dichloride, dimethylsilylenebis(4,5-benzindenyl)titanium
 dichloride,
 dimethylsilylenebis(2-methyl-4,5-benzindenyl)titanium
 dichloride, dimethylsilylenebis(2-phenylindenyl)titanium
 dichloride, dimethylsilylenebis(4-phenylindenyl)titanium
 10 dichloride,
 dimethylsilylenebis(2-methyl-4-phenylindenyl)titanium
 dichloride,
 dimethylsilylenebis(2-methyl-5-phenylindenyl)titanium
 dichloride,
 15 dimethylsilylenebis(2-methyl-4-naphthylindenyl)titanium
 dichloride,
 dimethylsilylenebis(4,5,6,7-tetrahydroindenyl)titanium
 dichloride,
 dimethylsilylene(cyclopentadienyl)(indenyl)titanium
 20 dichloride, dimethylsilylene(methylcyclopentadienyl)
 (indenyl)titanium dichloride,
 dimethylsilylene(n-butylcyclopentadienyl)(indenyl)titanium
 dichloride, dimethylsilylene(tetramethylcyclopentadienyl)
 (indenyl)titanium dichloride, dimethylsilylene
 25 (cyclopentadienyl)(fluorenyl)titanium dichloride,
 dimethylsilylene(methylcyclopentadienyl)(fluorenyl)titanium
 dichloride,

dimethylsilylene(n-butylcyclopentadienyl)(fluorenyl)titanium
 dichloride, dimethylsilylene(tetramethylcyclopentadienyl)
 (indenyl)titanium dichloride,
 dimethylsilylene(indenyl)(fluorenyl)titanium dichloride,
 5 dimethylsilylenebis(fluorenyl)titanium dichloride,
 dimethylsilylene(cyclopentadienyl)(tetramethylcyclopentadie
 nyl)titanium dichloride,
 dimethylsilylene(tetramethylcyclopentadienyl)
 (fluorenyl)titanium dichloride, cyclopentadienyltitanium
 10 trichloride, pentamethylcyclopentadienyltitanium trichloride,
 cyclopentadienyl(dimethylamido)titanium dichloride,
 cyclopentadienyl(phenoxy)titanium dichloride,
 cyclopentadienyl(2,6-dimethylphenyl)titanium dichloride,
 cyclopentadienyl(2,6-diisopropylphenyl)titanium dichloride,
 15 cyclopentadienyl(2,6-di-tert-butylphenyl)titanium
 dichloride,
 pentamethylcyclopentadienyl(2,6-dimethylphenyl)titanium
 dichloride,
 pentamethylcyclopentadienyl(2,6-diisopropylphenyl)titanium
 20 dichloride,
 pentamethylcyclopentadienyl(2,6-tert-butylphenyl)titanium
 dichloride, indenyl(2,6-diisopropylphenyl)titanium
 dichloride, fluorenyl(2,6-diisopropylphenyl)titanium
 dichloride,
 25 dimethylsilylene(cyclopentadienyl)(2-phenoxy)titanium
 dichloride,
 dimethylsilylene(cyclopentadienyl)(3-methyl-2-phenoxy)titan

ium dichloride,
 dimethylsilylene(cyclopentadienyl)(3,5-dimethyl-2-phenoxy)t
 itanium dichloride,
 dimethylsilylene(cyclopentadienyl)(3-tert-butyl-2-phenoxy)t
 5 itanium dichloride,
 dimethylsilylene(cyclopentadienyl)(3-tert-butyl-5-methyl-2-
 phenoxy)titanium dichloride,
 dimethylsilylene(cyclopentadienyl)(3,5-di-tert-butyl-2-phen
 oxy)titanium dichloride,
 10 dimethylsilylene(cyclopentadienyl)(5-methyl-3-phenyl-2-phen
 oxy)titanium dichloride,
 dimethylsilylene(cyclopentadienyl)(3-tert-butyldimethylsily
 l-5-methyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(cyclopentadienyl)(5-methyl-3-trimethylsily
 15 l-2-phenoxy)titanium dichloride,
 dimethylsilylene(cyclopentadienyl)(3-tert-butyl-5-methoxy-2
 -phenoxy)titanium dichloride,
 dimethylsilylene(cyclopentadienyl)(3-tert-butyl-5-chloro-2-
 phenoxy)titanium dichloride,
 20 dimethylsilylene(cyclopentadienyl)(3,5-diamyl-2-phenoxy)tit
 anium dichloride,
 dimethylsilylene(cyclopentadienyl)(3-phenyl-2-phenoxy)titan
 ium dichloride,
 dimethylsilylene(cyclopentadienyl)(1-naphthox-2-yl)titanium
 25 dichloride,
 dimethylsilylene(methylcyclopentadienyl)(2-phenoxy)titanium
 dichloride,

dimethylsilylene(methylcyclopentadienyl)(3-methyl-2-phenoxy
)titanium dichloride,
 dimethylsilylene(methylcyclopentadienyl)(3,5-dimethyl-2-phe
 noxy)titanium dichloride,
 5 dimethylsilylene(methylcyclopentadienyl)(3-tert-butyl-2-phe
 noxy)titanium dichloride,
 dimethylsilylene(methylcyclopentadienyl)(3-tert-butyl-5-met
 hyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(methylcyclopentadienyl)(3,5-di-tert-butyl-
 10 2-phenoxy)titanium dichloride,
 dimethylsilylene(methylcyclopentadienyl)(5-methyl-3-phenyl-
 2-phenoxy)titanium dichloride,
 dimethylsilylene(methylcyclopentadienyl)(3-tert-butyl dimeth
 ylsilyl-5-methyl-2-phenoxy)titanium dichloride,
 15 dimethylsilylene(methylcyclopentadienyl)(5-methyl-3-trimeth
 ylsilyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(methylcyclopentadienyl)(3-tert-butyl-5-met
 hox-2-phenoxy)titanium dichloride,
 dimethylsilylene(methylcyclopentadienyl)(3-tert-butyl-5-chl
 20 oro-2-phenoxy)titanium dichloride,
 dimethylsilylene(methylcyclopentadienyl)(3,5-diamyl-2-pheno
 xy)titanium dichloride,
 dimethylsilylene(methylcyclopentadienyl)(3-phenyl-2-phenoxy
)titanium dichloride,
 25 dimethylsilylene(methylcyclopentadienyl)(1-naphthox-2-yl)ti
 tanium dichloride,
 dimethylsilylene(n-butylcyclopentadienyl)(2-phenoxy)titanium

m dichloride,
 dimethylsilylene(n-butylcyclopentadienyl)(3-methyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(n-butylcyclopentadienyl)(3,5-dimethyl-2-phenoxy)titanium dichloride,
 5 dimethylsilylene(n-butylcyclopentadienyl)(3-tert-butyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(n-butylcyclopentadienyl)(3-tert-butyl-5-methyl-2-phenoxy)titanium dichloride,
 10 dimethylsilylene(n-butylcyclopentadienyl)(3,5-di-tert-butyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(n-butylcyclopentadienyl)(5-methyl-3-phenyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(n-butylcyclopentadienyl)(3-tert-butyl-5-methylsilyl-5-methyl-2-phenoxy)titanium dichloride,
 15 dimethylsilylene(n-butylcyclopentadienyl)(5-methyl-3-trimethylsilyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(n-butylcyclopentadienyl)(3-tert-butyl-5-methoxy-2-phenoxy)titanium dichloride,
 20 dimethylsilylene(n-butylcyclopentadienyl)(3-tert-butyl-5-chloro-2-phenoxy)titanium dichloride,
 dimethylsilylene(n-butylcyclopentadienyl)(3,5-diamyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(n-butylcyclopentadienyl)(3-phenyl-2-phenoxy)titanium dichloride, dimethylsilylene(n-butylcyclopentadienyl)(1-naphthox-2-yl)titanium dichloride,
 25 dimethylsilylene(tert-butylcyclopentadienyl)(2-phenoxy)titanium dichloride,

nium dichloride,
 dimethylsilylene(tert-butylcyclopentadienyl)(3-methyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(tert-butylcyclopentadienyl)(3,5-dimethyl-2
 5 -phenoxy)titanium dichloride,
 dimethylsilylene(tert-butylcyclopentadienyl)(3-tert-butyl-2
 -phenoxy)titanium dichloride,
 dimethylsilylene(tert-butylcyclopentadienyl)(3-tert-butyl-5
 -methyl-2-phenoxy)titanium dichloride,
 10 dimethylsilylene(tert-butylcyclopentadienyl)(3,5-di-tert-bu
 tyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(tert-butylcyclopentadienyl)(5-methyl-3-phe
 nyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(tert-butylcyclopentadienyl)(3-tert-butyl-di
 15 methylsilyl-5-methyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(tert-butylcyclopentadienyl)(5-methyl-3-tri
 methylsilyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(tert-butylcyclopentadienyl)(3-tert-butyl-5
 -methoxy-2-phenoxy)titanium dichloride,
 20 dimethylsilylene(tert-butylcyclopentadienyl)(3-tert-butyl-5
 -chloro-2-phenoxy)titanium dichloride,
 dimethylsilylene(tert-butylcyclopentadienyl)(3,5-diamyl-2-p
 henoxo)titanium dichloride,
 dimethylsilylene(tert-butylcyclopentadienyl)(3-phenyl-2-phenoxy)titanium dichloride,
 25 noxy)titanium dichloride,
 dimethylsilylene(tert-butylcyclopentadienyl)(1-naphthoxy-2-yl)
 1)titanium dichloride,

dimethylsilylene(tetramethylcyclopentadienyl)(2-phenoxy)titanium dichloride,
 dimethylsilylene(tetramethylcyclopentadienyl)(3-methyl-2-phenoxy)titanium dichloride,
 5 dimethylsilylene(tetramethylcyclopentadienyl)(3,5-dimethyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(tetramethylcyclopentadienyl)(3-tert-butyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(tetramethylcyclopentadienyl)(3-tert-butyl-5-methyl-2-phenoxy)titanium dichloride,
 10 dimethylsilylene(tetramethylcyclopentadienyl)(3,5-di-tert-butyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(tetramethylcyclopentadienyl)(5-methyl-3-phenyl-2-phenoxy)titanium dichloride,
 15 dimethylsilylene(tetramethylcyclopentadienyl)(3-tert-butyl-5-methyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(tetramethylcyclopentadienyl)(5-methyl-3-tert-butyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(tetramethylcyclopentadienyl)(3-tert-butyl-5-methoxy-2-phenoxy)titanium dichloride,
 20 dimethylsilylene(tetramethylcyclopentadienyl)(3-tert-butyl-5-chloro-2-phenoxy)titanium dichloride,
 dimethylsilylene(tetramethylcyclopentadienyl)(3,5-diamyl-2-phenoxy)titanium dichloride,
 25 dimethylsilylene(tetramethylcyclopentadienyl)(3-phenyl-2-phenoxy)titanium dichloride,
 dimethylsilylene(tetramethylcyclopentadienyl)(1-naphthoxy-2

-yl)titanium dichloride,
dimethylsilylene(trimethylsilylcyclopentadienyl)(2-phenoxy)
titanium dichloride,
dimethylsilylene(trimethylsilylcyclopentadienyl)(3-methyl-2
5 -phenoxy)titanium dichloride,
dimethylsilylene(trimethylsilylcyclopentadienyl)(3,5-dimeth
yl-2-phenoxy)titanium dichloride,
dimethylsilylene(trimethylsilylcyclopentadienyl)(3-tert-but
yl-2-phenoxy)titanium dichloride,
10 dimethylsilylene(trimethylsilylcyclopentadienyl)(3-tert-but
yl-5-methyl-2-phenoxy)titanium dichloride,
dimethylsilylene(trimethylsilylcyclopentadienyl)(3,5-di-ter
t-butyl-2-phenoxy)titanium dichloride,
dimethylsilylene(trimethylsilylcyclopentadienyl)(5-methyl-3
15 -phenyl-2-phenoxy)titanium dichloride,
dimethylsilylene(trimethylsilylcyclopentadienyl)(3-tert-but
yldimethylsilyl-5-methyl-2-phenoxy)titanium dichloride,
dimethylsilylene(trimethylsilylcyclopentadienyl)(5-methyl-3
-3-trimethylsilyl-2-phenoxy)titanium dichloride,
20 dimethylsilylene(trimethylsilylcyclopentadienyl)(3-tert-but
yl-5-methoxy-2-phenoxy)titanium dichloride,
dimethylsilylene(trimethylsilylcyclopentadienyl)(3-tert-but
yl-5-chloro-2-phenoxy)titanium dichloride,
dimethylsilylene(trimethylsilylcyclopentadienyl)(3,5-diamyl
25 -2-phenoxy)titanium dichloride,
dimethylsilylene(trimethylsilylcyclopentadienyl)(3-phenyl-2
-phenoxy)titanium dichloride,

dimethylsilylene(trimethylsilylcyclopentadienyl)(1-naphthox
 -2-yl)titanium dichloride,
 dimethylsilylene(indenyl)(2-phenoxy)titanium dichloride,
 dimethylsilylene(indenyl)(3-methyl-2-phenoxy)titanium
 5 dichloride,
 dimethylsilylene(indenyl)(3,5-dimethyl-2-phenoxy)titanium
 dichloride,
 dimethylsilylene(indenyl)(3-tert-butyl-2-phenoxy)titanium
 dichloride,
 10 dimethylsilylene(indenyl)(3-tert-butyl-5-methyl-2-phenoxy)t
 itanium dichloride,
 dimethylsilylene(indenyl)(3,5-di-tert-butyl-2-phenoxy)titan
 ium dichloride,
 dimethylsilylene(indenyl)(5-methyl-3-phenyl-2-phenoxy)titan
 15 ium dichloride,
 dimethylsilylene(indenyl)(3-tert-butyl dimethylsilyl-5-methy
 l-2-phenoxy)titanium dichloride,
 dimethylsilylene(indenyl)(5-methyl-3-trimethylsilyl-2-pheno
 xy)titanium dichloride,
 20 dimethylsilylene(indenyl)(3-tert-butyl-5-methoxy-2-phenoxy)
 titanium dichloride,
 dimethylsilylene(indenyl)(3-tert-butyl-5-chloro-2-phenoxy)t
 itanium dichloride,
 dimethylsilylene(indenyl)(3,5-diamyl-2-phenoxy)titanium
 25 dichloride,
 dimethylsilylene(indenyl)(3-phenyl-2-phenoxy)titanium
 dichloride,

dimethylsilylene(indenyl)(1-naphthox-2-yl)titanium
 dichloride, dimethylsilylene(fluorenyl)(2-phenoxy)titanium
 dichloride,
 dimethylsilylene(fluorenyl)(3-methyl-2-phenoxy)titanium
 5 dichloride,
 dimethylsilylene(fluorenyl)(3,5-dimethyl-2-phenoxy)titanium
 dichloride,
 dimethylsilylene(fluorenyl)(3-tert-butyl-2-phenoxy)titanium
 dichloride,
 10 dimethylsilylene(fluorenyl)(3-tert-butyl-5-methyl-2-phenoxy
)titanium dichloride,
 dimethylsilylene(fluorenyl)(3,5-di-tert-butyl-2-phenoxy)tit
 anium dichloride,
 dimethylsilylene(fluorenyl)(5-methyl-3-phenyl-2-phenoxy)tit
 15 anium dichloride,
 dimethylsilylene(fluorenyl)(3-tert-butyl-5-methoxy-2-phenoxy
)titanium dichloride,
 dimethylsilylene(fluorenyl)(3-tert-butyl-5-chloro-2-phenoxy
)titanium dichloride,
 dimethylsilylene(fluorenyl)(3,5-diamyl-2-phenoxy)titanium
 20 dichloride,
 dimethylsilylene(fluorenyl)(3-phenyl-2-phenoxy)titanium
 dichloride,

dimethylsilylene(fluorenyl)(1-naphthox-2-yl)titanium
 dichloride,
 (tert-butylamido)tetramethylcyclopentadienyl-1,2-ethanediyl
 titanium dichloride,
 5 (methylamido)tetramethylcyclopentadienyl-1,2-ethanediyltita
 nium dichloride,
 (ethylamido)tetramethylcyclopentadienyl-1,2-ethanediyltitan
 ium dichloride,
 (tert-butylamido)tetramethylcyclopentadienyl-dimethylsilane
 10 titanium dichloride,
 (benzylamido)tetramethylcyclopentadienyl-dimethylsilane
 titanium dichloride,
 (phenylphosphido)tetramethylcyclopentadienyl-dimethylsilane
 titanium dichloride, (tert-butylamido)indenyl-1,2-ethanediyl
 15 titanium dichloride,
 (tert-butylamido)tetrahydroindenyl-1,2-ethanediyl titanium
 dichloride, (tert-butylamido)fluorenyl-1,2-ethanediyl
 titanium dichloride, (tert-butylamido)indenyl dimethylsilane
 titanium dichloride,
 20 (tert-butylamido)tetrahydroindenyl dimethylsilane titanium
 dichloride, (tert-butylamido)fluorenyl dimethylsilane
 titanium dichloride,
 (dimethylaminomethyl)tetramethylcyclopentadienyl
 titanium(III) dichloride,
 25 (dimethylaminoethyl)tetramethylcyclopentadienyl titanium(III)
 dichloride,
 (dimethylaminopropyl)tetramethylcyclopentadienyl

titanium(III) dichloride,
 (N-pyrrolidinyloethyl)tetramethylcyclopentadienyl titanium
 dichloride, (B-dimethylaminoborabenzene)cyclopentadienyl
 titanium dichloride,
 5 cyclopentadienyl(9-mesitylboraanthracenyl)titanium
 dichloride,
 2,2'-thiobis[4-methyl-6-tert-butylphenoxy]titanium
 dichloride,
 2,2'-thiobis[4-methyl-6-(1-methylethyl)phenoxy]titanium
 10 dichloride, 2,2'-thiobis[4,6-dimethylphenoxy]titanium
 dichloride,
 2,2'-thiobis(4-methyl-6-tert-butylphenoxy)titanium
 dichloride,
 2,2'-methylenebis(4-methyl-6-tert-butylphenoxy)titanium
 15 dichloride,
 2,2'-ethylenebis(4-methyl-6-tert-butylphenoxy)titanium
 dichloride,
 2,2'-sulfinylbis(4-methyl-6-tert-butylphenoxy)titanium
 dichloride,
 20 2,2'-(4,4',6,6'-tetra-tert-butyl-1,1'-biphenoxy)titanium
 dichloride, (di-tert-butyl-1,3-propanediamido)titanium
 dichloride, (dicyclohexyl-1,3-propanediamido)titanium
 dichloride,
 [bis(trimethylsilyl)-1,3-propanediamido]titanium dichloride,
 25 [bis(tert-butyl-dimethylsilyl)-1,3-propanediamido]titanium
 dichloride,
 [bis(2,6-dimethylphenyl)-1,3-propanediamido]titanium

dichloride,
 [bis(2,6-diisopropylphenyl)-1,3-propanediamido]titanium
 dichloride,
 [bis(2,6-di-tert-butylphenyl)-1,3-propanediamido]titanium
 5 dichloride,
 [bis(triisopropylsilyl)naphthalenediamido]titanium
 dichloride, [bis(trimethylsilyl)naphthalenediamido]titanium
 dichloride,
 [bis(tert-butyl dimethylsilyl)naphthalenediamido]titanium
 10 dichloride,
 [hydrotris(3,5-dimethylpyrazolyl)borate]titanium
 trichloride,
 [hydrotris(3,5-diethylpyrazolyl)borate]titanium trichloride,
 [hydrotris(3,5-di-tert-butylpyrazolyl)borate]titanium
 15 trichloride, [tris(3,5-dimethylpyrazolyl)methyl]titanium
 trichloride, [tris(3,5-diethylpyrazolyl)methyl]titanium
 trichloride,
 [tris(3,5-di-tert-butylpyrazolyl)methyl]titanium
 trichloride, and those formed by replacing the titanium of these
 20 compounds with zirconium or hafnium, those formed by replacing
 the 2-phenoxy of these compounds with (3-phenyl-2-phenoxy),
 (3-trimethylsilyl-2-phenoxy) or
 (3-tert-butyl dimethylsilyl-2-phenoxy), those formed by
 replacing the dimethylsilylene of these compounds with methylene,
 25 ethylene, dimethylmethylene (isoprpylidene),
 diphenylmethylene, diethylsilylene, diphenylsilylene or
 dimethoxysilylene, those formed by replacing the dichloride of

these compounds with difluoride, dibromide, diiodide, dimethyl, diethyl, diisopropyl, diphenyl, dibenzyl, dimethoxide, diethoxide, di-n-propoxide, diisopropoxide, di-n-butoxide, diisobutoxide, di-tert-butoxide, diphenoxide, di(pentafluorophenoxide) or di(2,6-di-tert-butylphenoxide), and those formed by replacing the trichloride with trifluoride, tribromide, triiodide, trimethyl, triethyl, triisopropyl, triphenyl, tribenzyl, trimethoxide, triethoxide, tri-n-propoxide, triisopropoxide, tri-n-butoxide, triisobutoxide, tri-tert-butoxide, triphenoxide, tri(pentafluorophenoxide) or tri(2,6-di-tert-butylphenoxide).

In the transition metal compounds represented by the general formula [10], specific examples of a compound having a nickel atom as a transition metal atom are

2,2'-methylenebis[(4R)-4-phenyl-5,5'-dimethyloxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-phenyl-5,5'-diethyloxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-phenyl-5,5'-di-n-propyloxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-phenyl-5,5'-diisopropyloxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-phenyl-5,5'-dicyclohexyloxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-phenyl-5,5'-dimethoxyoxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-phenyl-5,5'-diethoxyoxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-phenyl-5,5'-diphenyloxazoline]nickel dibromide,

5 2,2'-methylenebis[(4R)-4-methyl-5,5-di-(2-methylphenyl)oxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-methyl-5,5-di-(3-methylphenyl)oxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-methyl-5,5-di-(4-methylphenyl)oxazoline]nickel dibromide,

10 2,2'-methylenebis[(4R)-4-methyl-5,5-di-(2-methoxyphenyl)oxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-methyl-5,5-di-(3-methoxyphenyl)oxazoline]nickel dibromide,

15 2,2'-methylenebis[(4R)-4-methyl-5,5-di-(4-methoxyphenyl)oxazoline]nickel dibromide,

2,2'-methylenebis[spiro{(4R)-4-methyloxazoline-5,1'-cyclobutane}]nickel dibromide,

2,2'-methylenebis[spiro{(4R)-4-methyloxazoline-5,1'-cyclopentane}]nickel dibromide,

20 2,2'-methylenebis[spiro{(4R)-4-methyloxazoline-5,1'-cyclohexane}]nickel dibromide,

2,2'-methylenebis[spiro{(4R)-4-methyloxazoline-5,1'-cycloheptane}]nickel dibromide,

25 2,2'-methylenebis[(4R)-4-isopropyl-5,5-dimethyloxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-isopropyl-5,5-diethyloxazoline]nickel dibromide,

kel dibromide,
 2,2'-methylenebis[(4R)-4-isopropyl-5,5-di-n-propyloxazoline
]nickel dibromide,
 2,2'-methylenebis[(4R)-4-isopropyl-5,5-diisopropyloxazoline
 5]nickel dibromide,
 2,2'-methylenebis[(4R)-4-isopropyl-5,5-dicyclohexyloxazolin
 e]nickel dibromide,
 2,2'-methylenebis[(4R)-4-isopropyl-5,5-diphenyloxazoline]ni
 ckel dibromide,
 10 2,2'-methylenebis[(4R)-4-isopropyl-5,5-di-(2-methylphenyl)o
 xazoline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-isopropyl-5,5-di-(3-methylphenyl)o
 xazoline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-isopropyl-5,5-di-(4-methylphenyl)o
 15 xazoline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-isopropyl-5,5-di-(2-methoxyphenyl)
 oxazoline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-isopropyl-5,5-di-(3-methoxyphenyl)
 oxazoline]nickel dibromide,
 20 2,2'-methylenebis[(4R)-4-isopropyl-5,5-di-(4-methoxyphenyl)
 oxazoline]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-isopropyloxazoline-5,1'-cycl
 obutane}]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-isopropyloxazoline-5,1'-cycl
 25 opentane}]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-isopropyloxazoline-5,1'-cycl
 ohexane}]nickel dibromide,

2,2'-methylenebis[spiro{(4R)-4-isopropylloxazoline-5,1'-cycloheptane}]nickel dibromide,

2,2'-methylenebis[(4R)-4-isobutyl-5,5-dimethylloxazoline]nickel dibromide,

5 2,2'-methylenebis[(4R)-4-isobutyl-5,5-diethylloxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-isobutyl-5,5-di-n-propylloxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-isobutyl-5,5-diisopropylloxazoline]nickel dibromide,

10 2,2'-methylenebis[(4R)-4-isobutyl-5,5-dicyclohexylloxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-isobutyl-5,5-diphenylloxazoline]nickel dibromide,

15 2,2'-methylenebis[(4R)-4-isobutyl-5,5-di-(2-methylphenyl)loxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-isobutyl-5,5-di-(3-methylphenyl)loxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-isobutyl-5,5-di-(4-methylphenyl)loxazoline]nickel dibromide,

20 2,2'-methylenebis[(4R)-4-isobutyl-5,5-di-(2-methoxyphenyl)loxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-isobutyl-5,5-di-(3-methoxyphenyl)loxazoline]nickel dibromide,

25 2,2'-methylenebis[(4R)-4-isobutyl-5,5-di-(4-methoxyphenyl)loxazoline]nickel dibromide,

2,2'-methylenebis[spiro{(4R)-4-isobutylloxazoline-5,1'-cyclo

butane}]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-isobutyloxazoline-5,1'-cyclo
 pentane}]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-isobutyloxazoline-5,1'-cyclo
 5 hexane}]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-isobutyloxazoline-5,1'-cyclo
 heptane}]nickel dibromide,
 2,2'-methylenebis[(4R)-4-tert-butyl-5,5-dimethyloxazoline]n
 ickel dibromide,
 10 2,2'-methylenebis[(4R)-4-tert-butyl-5,5-diethyloxazoline]ni
 ckel dibromide,
 2,2'-methylenebis[(4R)-4-tert-butyl-5,5-di-n-propyloxazol
 ine]nickel dibromide,
 2,2'-methylenebis[(4R)-4-tert-butyl-5,5-diisopropyloxazol
 ine]nickel dibromide,
 15 2,2'-methylenebis[(4R)-4-tert-butyl-5,5-diphenyloxazoline]n
 ickel dibromide,
 2,2'-methylenebis[(4R)-4-tert-butyl-5,5-dicyclohexyloxazoli
 ne]nickel dibromide,
 20 2,2'-methylenebis[(4R)-4-tert-butyl-5,5-di-(2-methylphenyl)
 oxazoline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-tert-butyl-5,5-di-(3-methylphenyl)
 oxazoline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-tert-butyl-5,5-di-(4-methylphenyl)
 25 oxazoline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-tert-butyl-5,5-di-(2-methoxyphenyl
)oxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-tert-butyl-5,5-di-(3-methoxyphenyl)
)oxazoline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-tert-butyl-5,5-di-(4-methoxyphenyl)
)oxazoline]nickel dibromide,
 5 2,2'-methylenebis[spiro{(4R)-4-tert-butyloxazoline-5,1'-cyc
 lobutane}]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-tert-butyloxazoline-5,1'-cyc
 lopentane}]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-tert-butyloxazoline-5,1'-cyc
 10 lohexane}]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-tert-butyloxazoline-5,1'-cyc
 loheptane}]nickel dibromide,
 2,2'-methylenebis[(4R)-4-phenyl-5,5-dimethyloxazoline]nicke
 1 dibromide,
 15 2,2'-methylenebis[(4R)-4-phenyl-5,5-diethyloxazoline]nickel
 dibromide,
 2,2'-methylenebis[(4R)-4-phenyl-5,5-di-n-propyloxazoline]ni
 ckel dibromide,
 2,2'-methylenebis[(4R)-4-phenyl-5,5-diisopropyloxazoline]ni
 20 ckel dibromide,
 2,2'-methylenebis[(4R)-4-phenyl-5,5-dicyclohexyloxazoline]n
 ickel dibromide,
 2,2'-methylenebis[(4R)-4-phenyl-5,5-diphenyloxazoline]nicke
 1 dibromide,
 25 2,2'-methylenebis[(4R)-4-phenyl-5,5-di-(2-methylphenyl)oxaz
 oline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-phenyl-5,5-di-(3-methylphenyl)oxaz

oline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-phenyl-5,5-di-(4-methylphenyl)oxaz
 oline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-phenyl-5,5-di-(2-methoxyphenyl)oxa
 5 zoline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-phenyl-5,5-di-(3-methoxyphenyl)oxa
 zoline]nickel dibromide,
 2,2'-methylenebis[(4R)-4-phenyl-5,5-di-(4-methoxyphenyl)oxa
 zoline]nickel dibromide,
 10 2,2'-methylenebis[spiro{(4R)-4-phenyloxazoline-5,1'-cyclobu
 tane}]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-phenyloxazoline-5,1'-cyclope
 ntane}]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-phenyloxazoline-5,1'-cyclohe
 15 xane}]nickel dibromide,
 2,2'-methylenebis[spiro{(4R)-4-phenyloxazoline-5,1'-cyclohe
 ptane}]nickel dibromide,
 2,2'-methylenebis[(4R)-4-benzyl-5,5-dimethyloxazoline]nicke
 l dibromide,
 20 2,2'-methylenebis[(4R)-4-benzyl-5,5-diethyloxazoline]nickel
 dibromide,
 2,2'-methylenebis[(4R)-4-benzyl-5,5-di-n-propyloxazoline]ni
 ckel dibromide,
 2,2'-methylenebis[(4R)-4-benzyl-5,5-di-isopropyloxazoline]n
 25 ickel dibromide,
 2,2'-methylenebis[(4R)-4-benzyl-5,5-dicyclohexyloxazoline]n
 ickel dibromide,

2,2'-methylenebis[(4R)-4-benzyl-5,5-diphenyl]oxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-benzyl-5,5-di-(2-methylphenyl)oxazoline]nickel dibromide,

5 2,2'-methylenebis[(4R)-4-benzyl-5,5-di-(3-methylphenyl)oxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-benzyl-5,5-di-(4-methylphenyl)oxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-benzyl-5,5-di-(2-methoxyphenyl)oxazoline]nickel dibromide,

10 2,2'-methylenebis[(4R)-4-benzyl-5,5-di-(3-methoxyphenyl)oxazoline]nickel dibromide,

2,2'-methylenebis[(4R)-4-benzyl-5,5-di-(4-methoxyphenyl)oxazoline]nickel dibromide,

15 2,2'-methylenebis[spiro{(4R)-4-benzyloxazoline-5,1'-cyclobutane}]nickel dibromide,

2,2'-methylenebis[spiro{(4R)-4-benzyloxazoline-5,1'-cyclopentane}]nickel dibromide,

2,2'-methylenebis[spiro{(4R)-4-benzyloxazoline-5,1'-cyclohexane}]nickel dibromide,

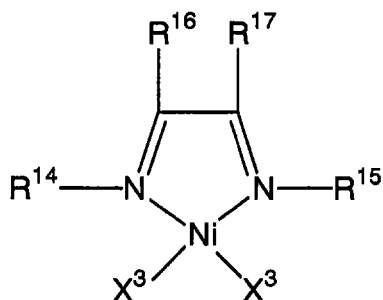
20 2,2'-methylenebis[spiro{(4R)-4-benzyloxazoline-5,1'-cycloheptane}]nickel dibromide, and antidopes of the above-mentioned compounds. Further, there are exemplified those formed by reversing a steric configuration of the asymmetric carbon on

25 one hand oxazoline ring of the above-mentioned bisoxazoline compound, and those formed by replacing the dibromide of said compounds with difluoride, dichloride, diiodide, dimethyl,

diethyl, diisopropyl, diphenyl, dibenzyl, dimethoxide,
 diethoxide, di-n-propoxide, diisopropoxide, di-n-butoxide,
 diisobutoxide, di-tert-butoxide, diphenoxide,
 di(pentafluorophenoxide) and di(2,6-di-tert-butylphenoxide).

5 Further, specific examples of the nickel compound are
 [hydrotris(3,5-dimethylpyrazolyl)borate]nickel chloride,
 [hydrotris(3,5-diethylpyrazolyl)borate]nickel chloride,
 [hydrotris(3,5-di-tert-butylpyrazolyl)borate]nickel
 chloride, and those formed by replacing the chloride in those
 10 compounds with bromide, iodide, methyl, ethyl, allyl, methallyl,
 methoxide, ethoxide, n-propoxide, isopropoxide, n-butoxide,
 isobutoxide, tert-butoxide, phenoxide, pentafluorophenoxide
 and 2,6-di-tert-butylphenoxide.

As the nickel compound, there are enumerated compounds
 15 represented by the following structural formula.



(In the formula, R¹⁴ and R¹⁵ are each a 2,6-diisopropylphenyl
 20 group, R¹⁶ and R¹⁷ are each a hydrogen atom or a methyl group,
 or R¹⁴ and R¹⁵ may be bound to form an acenaphthene group, and
 X³ is a fluorine atom, a chlorine atom, a bromine atom, an iodine
 atom, a methyl group, an ethyl group, a n-propyl group, an

isopropyl group, a n-butyl group, a phenyl group, a benzyl group, a methoxy group, an ethoxy group, a n-propoxy group, an isopropoxy group, a n-butoxy group, an isobutoxy group or a phenoxy group.)

Further, there are exemplified those formed by replacing
5 the nickel atom in the above-mentioned nickel compounds with palladium, cobalt, rhodium and ruthenium.

In the transition metal compound represented by the general formula [10], specific examples of a compound having an iron atom as a transition metal atom are

10 2,6-bis-[1-(2,6-dimethylphenylimino)ethyl]pyridineiron dichloride, 2,6-bis-[1-(2,6-diisopropylphenylimino)ethyl]pyridineiron dichloride, 2,6-bis-[1-(2-tert-butyl-phenylimino)ethyl]pyridineiron dichloride, and those formed by replacing the dichloride thereof
15 with difluoride, dibromide, diiodide, dimethyl, diethyl, dimethoxide, diethoxide, di-n-propoxide, diisopropoxide, di-n-butoxide, diisobutoxide, di-tert-butoxide, diphenoxide, di(pentafluorophenoxide) and di(2,6-di-tert-butylphenoxide).

Further, specific examples of the iron compound are
20 [hydrotris(3,5-dimethylpyrazolyl)borate]iron chloride, [hydrotris(3,5-diethylpyrazolyl)borate]iron chloride, [hydrotris(3,5-di-tert-butylpyrazolyl)borate]iron chloride and those formed by replacing the chloride in these compounds with fluoride, bromide, iodide, methyl, ethyl, allyl, methallyl,
25 methoxide, ethoxide, n-propoxide, isopropoxide, n-butoxide, isobutoxide, tert-butoxide, phenoxide, pentafluorophenoxide and 2,6-di-tert-butylphenoxide.

Furthermore, there are similarly exemplified those formed by replacing the iron in the above-mentioned iron compounds with cobalt and nickel.

Moreover, specific examples of a μ -oxo type transition metal compound of the transition metal compounds represented by the general formula [10] are μ -oxobis[isopropylidene(cyclopentadienyl)(2-phenoxy)titanium chloride], μ -oxobis[isopropylidene(cyclopentadienyl)(3-tert-butyl-5-methyl-2-phenoxy)titanium chloride], μ -oxobis[isopropylidene(methylcyclopentadienyl)(2-phenoxy)titanium chloride], μ -oxobis[isopropylidene(methylcyclopentadienyl)(3-tert-butyl-5-methyl-2-phenoxy)titanium chloride], μ -oxobis[isopropylidene(tetramethylcyclopentadienyl)(2-phenoxy)titanium chloride], μ -oxobis[isopropylidene(tetramethylcyclopentadienyl)(3-tert-butyl-5-methyl-2-phenoxy)titanium chloride], μ -oxobis[dimethylsilylene(cyclopentadienyl)(2-phenoxy)titanium chloride], μ -oxobis[dimethylsilylene(cyclopentadienyl)(3-tert-butyl-5-methyl-2-phenoxy)titanium chloride], μ -oxobis[dimethylsilylene(methylcyclopentadienyl)(2-phenoxy)titanium chloride], μ -oxobis[dimethylsilylene(methylcyclopentadienyl)(3-tert-butyl-5-methyl-2-phenoxy)titanium chloride], μ -oxobis[dimethylsilylene(tetramethylcyclopentadienyl)(2-phenoxy)titanium chloride] and μ -oxobis[dimethylsilylene(tetramethylcyclopentadienyl)(3-tert-butyl-5-methyl-2-phenoxy)titanium chloride].

xy)titanium chloride]. In addition, there are exemplified those formed by replacing the chloride in said compounds with fluoride, bromide, iodide, methyl, ethyl, isopropyl, phenyl benzyl, methoxide, ethoxide, n-propoxide, isopropoxide, n-butoxide, 5 isobutoxide, tert-butoxide, phenoxide, pentafluorophenoxide and 2,6-di-tert-butylphenoxide.

Besides the transition metal compound represented by the general formula [10] and its μ -oxo type transition metal compound, whose examples are as mentioned above, examples of a compound 10 used as the transition metal compound (B) are as follows. Examples of a compound having a nickel atom as the transition metal atom are nickel chloride, nickel bromide, nickel iodide, nickel sulfate, nickel nitrate, nickel perchlorate, nickel acetate, nickel trifluoroacetate, nickel cyanide, nickel oxalate, 15 acetylacetonatonickel, bis(allyl)nickel, bis(1,5-cyclooctadiene)nickel, dichloro(1,5-cyclooctadiene)nickel, dichlorobis(acetonitrile)nickel, dichlorobis(benzonitrile)nickel, 20 carbonyltris(triphenylphosphine)nickel, dichlorobis(triethylphosphine)nickel, diacetobis(triphenylphosphine)nickel, tetrakis(triphenylphosphine)nickel, dichloro[1,2-bis(diphenylphosphino)ethane]nickel, 25 bis[1,2-bis(diphenylphosphino)ethane]nickel, dichloro[1,3-bis(diphenylphosphino)propane]nickel, bis[1,3-bis(diphenylphosphino)propane]nickel,

tetraaminenickel nitrate, tetraaxis(acetonitrile)nickel tetrafluoroborate and nickelpthalocianine.

Similarly, examples of a compound having a vanadium atom as the transition metal atom are vanadium acetylacetonate,
5 vanadium tetrachloride and vanadium oxytrichloride.

Example of a compound having a samarium atom as the transition metal atom is bis(pentamethylcyclopentadienyl) samarium methyltetrahydrofuran.

Example of a compound having an ytterbium atom as the
10 transition metal atom is bis(pentamethylcyclopentadienyl) ytterbium methyltetrahydrofuran.

These transition metal compounds may be used singly or in a combination of two or more thereof.

Among those transition metal compounds exemplified above,
15 those represented by the general formula [10] are preferred as the transition metal compound (B) used in the present invention. In particular, the transition metal compounds having the 4 Group atom as M^2 in the above general formula [10] are preferred, and those having at least one cyclopentadiene type anion
20 skeleton-carrying group as L^2 in the general formula [10] are particularly preferred.

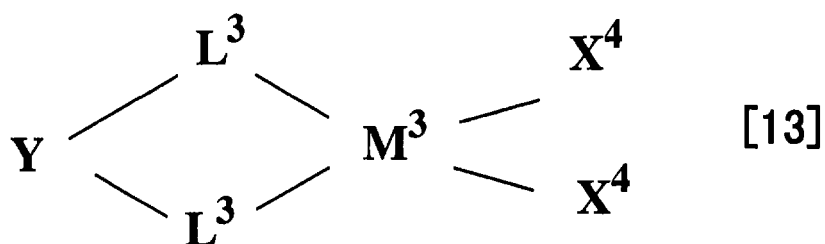
Further, in the case where X is OR^{11} , the 4 Group transition metal compounds having two cyclopentadiene type anion skeleton-carrying groups as L^2 , which groups are linked through
25 a residual group containing a carbon atom, a silicon atom, a nitrogen atom, an oxygen atom, a sulfur atom or a phosphorus atom, are preferred from a viewpoint of an improvement of activity.

The 4 Group transition metal compounds having two η ⁵-(substituted)indenyl groups as L², which groups are linked through a methylene group, an ethylene group, a dimethylmethylenegroup, a dimethylsilylene group, a diethylsilylene group or a diphenylsilylene group, are more preferred from a viewpoint of an improvement of activity. A racemic ethylenebis(1-indenyl)zirconium diphenoxide is particularly preferred from a viewpoint of an improvement of activity.

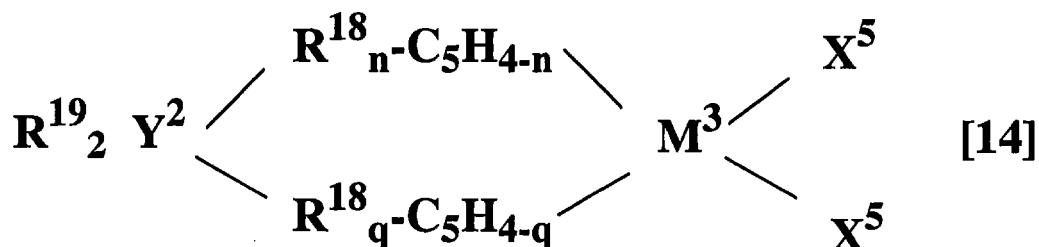
The transition metal compound having OR¹¹ as X in the above general formula [10] can be obtained by preparing an R¹¹OH corresponding to the OR¹¹, reacting the same with a strong base such as an organolithium compound, and thereafter reacting the obtained compound with a transition metal compound having a chlorine atom in place of OR¹¹ in the above general formula [10].

Further, a transition metal compound having a stereoregular polymerization ability of an α -olefin can be selected from the transition metal compound (B) represented by the formula [10]. Thereby, a stereoregular polymerization of an α -olefin can be accomplished.

As the transition metal compound having a stereoregular polymerization ability, preferred is a transition metal compound represented by the following general formula [13] or a transition metal compound represented by the following general formula [14].



(In the formula, M^3 is a transition metal atom of the Groups
 3 to 11 or lanthanoid series, L^3 is an η^5 -indenyl group or a
 5 substituted η^5 -indenyl group, and both of L^3 s may be the same
 or different from each other, Y is a bridging group for linking
 both of L^3 s, and both of X^4 s are independently of each other
 a hydrogen atom, a halogen atom, an alkyl group, an aralkyl group,
 an aryl group, a substituted silyl group, an alkoxy group, an
 10 aralkyloxy group, an aryloxy group or a heterocyclic group.)



(In the formula, M^3 is a transition metal atom of the Groups
 15 3 to 11 or lanthanoid series, Y^2 is a silicon atom, a germanium
 atom or a tin atom, the $(R^{18}_n-C_5H_{4-n})$ and $(R^{18}_q-C_5H_{4-q})$ are
 independently of each other a substituted η
 5-cyclopentadienyl group, n and q are independently of each
 other an integer of 1 to 3, and both of R^{18} s may be the same
 20 or different from each other and are independently of each

other a halogen atom, an alkyl group, an aralkyl group, an aryl group, a substituted silyl group, an alkoxy group, an aralkyloxy group, an aryloxy group or a heterocyclic group, provided that the position and/or kind of each R^{18} in each substituted η^5 -cyclopentadienyl group are (is) selected, so that a symmetry plane containing M^3 does not exist, R^{19} and X^5 are independently of each other a hydrogen atom, a halogen atom, an alkyl group, an aralkyl group, an aryl group, a substituted silyl group, an alkoxy group, an aralkyloxy group, an aryloxy group or a heterocyclic group, and R^{19} and X^5 may be all the same or different from each other.)

The transition metal atom represented by M^3 in the above general formulas [13] and [14] is defined as the same as M^2 in the above-mentioned general formula [10].

In the general formula [13], L^3 is an η^5 -indenyl group or a substituted η^5 -indenyl group, and both of L^3 s may be the same or different from each other. Specific examples of L^3 are an η^5 -indenyl group, an η^5 -methylindenyl group, an η^5 -dimethylindenyl group, an η^5 -n-propylindenyl group, an η^5 -isopropylindenyl group, an η^5 -n-butylindenyl group, an η^5 -tert-butylindenyl group, an η^5 -phenylindenyl group, an η^5 -methylphenylindenyl group, an η^5 -naphthylindenyl group, an η^5 -trimethylsilylindenyl group and an η^5 -tetrahydroindenyl group.

In the above-mentioned general formula [13], Y is a bridging group for linking both of L^3 s. As the bridging group, there is given a divalent bridging group containing an atom of

the Group 14 of the periodic table of elements (IUPAC, inorganic chemical nomenclature, revised edition, 1989). Preferred is a divalent bridging group containing a carbon atom, a silicon atom, a germanium atom or a tin atom. More preferred is a divalent
5 bridging group, whose atom connected with both of L^3 's is a carbon atom, a silicon atom, a germanium atom and/or a tin atom. Much more preferred is a divalent bridging group, whose atom connected with both of L^3 's is a carbon atom, a silicon atom, a germanium atom and/or a tin atom, and whose minimum atom number between
10 atoms connected with both of L^3 's is not more than 3 (the case where the atom connected with both of L^3 's is single is included). Specific examples thereof are a methylene group, an ethylene group, a propylene group, a dimethylmethylene group (an isopropylidene group), a diphenylmethylene group, a
15 tetramethylethylene group, a silylene group, a dimethylsilylene group, a diethylsilylene group, a diphenylsilylene group, a tetramethyldisilylene group and a dimethoxysilylene group. Particularly preferred are a methylene group, an ethylene group, a dimethylmethylene group (an isopropylidene group), a
20 dimethylsilylene group, a diethylsilylene group and a diphenylsilylene group.

In the above-mentioned general formula [13], both of X^4 's are independently of each other a hydrogen atom, a halogen atom, an alkyl group, an aralkyl group, an aryl group, a substituted
25 silyl group, an alkoxy group, an aralkyloxy group, an aryloxy group or a heterocyclic group.

Among these, examples of a halogen atom, an alkyl group,

an aralkyl group and an aryl group are similar to those of X in the above general formula [10]. While, examples of a substituted silyl group, an alkoxy group, an aralkyloxy group, an aryloxy group and a heterocyclic group are as follows.

5 The substituted silyl group means a hydrocarbon group-substituted silyl group. Here, as the hydrocarbon group, there are enumerated an alkyl group having 1 to 10 carbon atoms such as a methyl group, an ethyl group, a n-propyl group, an isopropyl group, a n-butyl group, a sec-butyl group, a tert-butyl
10 group, an isobutyl group, a n-pentyl group, a n-hexyl group and a cyclohexyl group, and an aryl group such as a phenyl group. As the substituted silyl group having 1 to 20 carbon atoms, there are exemplified a mono-substituted silyl group having 1 to 20 carbon atoms such as a methylsilyl group, an ethylsilyl group
15 and a phenylsilyl group, a di-substituted silyl group having 2 to 20 carbon atoms such as a dimethylsilyl group, a diethylsilyl group and a diphenylsilyl group, and a tri-substituted silyl group having 3 to 20 carbon atoms such as a trimethylsilyl group, a triethylsilyl group, a tri-n-propylsilyl group, a
20 triisopropylsilyl group, a tri-n-butylsilyl group, a tri-sec-butylsilyl group, a tri-tert-butylsilyl group, a triisobutylsilyl group, a tert-butyl dimethylsilyl group, a tri-n-pentylsilyl group, a tri-n-hexylsilyl group, a tricyclohexylsilyl group and a triphenylsilyl group. Preferred
25 are a trimethylsilyl group, a tert-butyl dimethylsilyl group and a triphenylsilyl group. Any of the hydrocarbon group appended to these substituted silyl groups may be partially substituted

with a halogen atom such as a fluorine atom, a chlorine atom, a bromine atom or an iodine atom, an alkoxy group such as a methoxy group or an ethoxy group, an aryloxy group such as a phenoxy group, and an aralkyloxy group such as a benzyloxy group.

5 As the alkoxy group, preferred are those having 1 to 20 carbon atoms. Examples thereof are a methoxy group, an ethoxy group, a n-propoxy group, an isopropoxy group, a n-butoxy group, a sec-butoxy group, a tert-butoxy group, a n-pentoxy group, a neopentoxy group, a n-hexoxy group, a n-octoxy group, a
10 n-dodecoxy group, a n-pentadecoxy group and an eicosoxy group, and more preferred are a methoxy group, an ethoxy group, an isopropoxy group and a tert-butoxy group. Any of these alkoxy groups may be partially substituted with a halogen atom such as a fluorine atom, a chlorine atom, a bromine atom or an iodine
15 atom, an alkoxy group such as a methoxy group or an ethoxy group, an aryloxy group such as a phenoxy group, and an aralkyloxy group such as a benzyloxy group.

As the aralkyloxy group, preferred are those having 7 to 20 carbon atoms. Examples thereof are a benzyloxy group, a
20 (2-methylphenyl)methoxy group, a (3-methylphenyl)methoxy group, a (4-methylphenyl)methoxy group, a (2,3-dimethylphenyl)methoxy group, a (2,4-dimethylphenyl)methoxy group, a (2,5-dimethylphenyl)methoxy group, a (2,6-dimethylphenyl)methoxy group, a
25 (3,4-dimethylphenyl)methoxy group, a (3,5-dimethylphenyl)methoxy group, a (2,3,4-trimethylphenyl)methoxy group, a

(2,3,5-trimethylphenyl)methoxy group, a
(2,3,6-trimethylphenyl)methoxy group, a
(2,4,5-trimethylphenyl)methoxy group, a
(2,4,6-trimethylphenyl)methoxy group, a
5 (3,4,5-trimethylphenyl)methoxy group, a
(2,3,4,5-tetramethylphenyl)methoxy group, a
(2,3,4,6-tetramethylphenyl)methoxy group, a
(2,3,5,6-tetramethylphenyl)methoxy group, a
(pentamethylphenyl)methoxy group, an (ethylphenyl)methoxy
10 group, a (n-propylphenyl)methoxy group, an
(isopropylphenyl)methoxy group, a (n-butylphenyl)methoxy group,
a (sec-butylphenyl)methoxy group, a (tert-butylphenyl)methoxy
group, a (n-hexylphenyl)methoxy group, a
(n-octylphenyl)methoxy group, a (n-decylphenyl)methoxy group,
15 a naphthylmethoxy group and an anthracenylmethoxy group. More
preferred is a benzyloxy group.

Any of these aralkyloxy groups may be partially substituted
with a halogen atom such as a fluorine atom, a chlorine atom,
a bromine atom or an iodine atom, an alkoxy group such as a methoxy
20 group or an ethoxy group, an aryloxy group such as a phenoxy
group, and an aralkyloxy group such as a benzyloxy group.

As the aryloxy group, preferred are those having 6 to 20
carbon atoms. Examples thereof are a phenoxy group, a
2-methylphenoxy group, a 3-methylphenoxy group, a
25 4-methylphenoxy group, a 2,3-dimethylphenoxy group, a
2,4-dimethylphenoxy group, a 2,5-dimethylphenoxy group, a
2,6-dimethylphenoxy group, a 3,4-dimethylphenoxy group, a

3,5-dimethylphenoxy group, a 2-tert-butyl-3-methylphenoxy
 group, a 2-tert-butyl-4-methylphenoxy group, a
 2-tert-butyl-5-methylphenoxy group, a
 2-tert-butyl-6-methylphenoxy group, a 2,3,4-trimethylphenoxy
 5 group, a 2,3,5-trimethylphenoxy group, a
 2,3,6-trimethylphenoxy group, a 2,4,5-trimethylphenoxy group,
 a 2,4,6-trimethylphenoxy group, a
 2-tert-butyl-3,4-dimethylphenoxy group, a
 2-tert-butyl-3,5-dimethylphenoxy group, a
 10 2-tert-butyl-3,6-dimethylphenoxy group, a
 2,6-di-tert-butyl-3-methylphenoxy group, a
 2-tert-butyl-4,5-dimethylphenoxy group, a
 2,6-di-tert-butyl-4-methylphenoxy group, a
 3,4,5-trimethylphenoxy group, a 2,3,4,5-tetramethylphenoxy
 15 group, a 2-tert-butyl-3,4,5-trimethylphenoxy group, a
 2,3,4,6-tetramethylphenoxy group, a
 2-tert-butyl-3,4,6-trimethylphenoxy group, a
 2,6-di-tert-butyl-3,4-dimethylphenoxy group, a
 2,3,5,6-tetramethylphenoxy group, a
 20 2-tert-butyl-3,5,6-trimethylphenoxy group, a
 2,6-di-tert-butyl-3,5-dimethylphenoxy group, a
 pentamethylphenoxy group, an ethylphenoxy group, a
 n-propylphenoxy group, an isopropylphenoxy group, a
 n-butylphenoxy group, a sec-butylphenoxy group, a
 25 tert-butylphenoxy group, a n-hexylphenoxy group, a
 n-octylphenoxy group, a n-decylphenoxy group, a
 n-tetradecylphenoxy group, a naphthoxy group and an anthracenoxy

group.

Any of these aryloxy groups may be partially substituted with a halogen atom such as a fluorine atom, a chlorine atom, a bromine atom or an iodine atom, an alkoxy group such as a methoxy group or an ethoxy group, an aryloxy group such as a phenoxy group, and an aralkyloxy group such as a benzyloxy group.

The heterocyclic ring group is a group having a heterocyclic ring, preferably a 4 to 8 membered heterocyclic ring, and more preferably a 4 to 8 membered aromatic heterocyclic ring. A hetero atom contained in the heterocyclic ring is preferably a nitrogen atom, an oxygen atom or a sulfur atom. Examples of such a heterocyclic ring group are an indolyl group, a furyl group, a thienyl group, a pyridyl group, a piperidyl group, a quinolyl group and an isoquinolyl group. A more preferred example of the heterocyclic ring group is a furyl group.

Any of these heterocyclic ring groups may be partially substituted with a halogen atom such as a fluorine atom, a chlorine atom, a bromine atom or an iodine atom, an alkoxy group such as a methoxy group or an ethoxy group, an aryloxy group such as a phenoxy group, and an aralkyloxy group such as a benzyloxy group.

A compound in a category of the transitional metal compound having the stereoregular polymerization ability can be selected, for example, from the above-exemplified compounds as a specific example, preferred are those represented by the general formula [13]. For example, those having a di-substituted silicon atom as the bridging group Y in the general formula [13] are preferred.

Specific examples thereof are

dimethylsilylenebis(2-methylindenyl)zirconium dichloride,
dimethylsilylenebis(2-n-propylindenyl)zirconium dichloride,
dimethylsilylenebis(2-n-butylindenyl)zirconium dichloride,
5 dimethylsilylenebis(2-n-hexylindenyl)zirconium dichloride,
dimethylsilylenebis(2-isopropylindenyl)zirconium dichloride,
dimethylsilylenebis(2-isobutylindenyl)zirconium dichloride,
dimethylsilylenebis(2-tert-butylindenyl)zirconium
dichloride, dimethylsilylenebis(2-phenylindenyl)zirconium
10 dichloride,
dimethylsilylenebis(2-trimethylsilylindenyl)zirconium
dichloride,
dimethylsilylenebis(2,4-dimethylindenyl)zirconium
dichloride,
15 dimethylsilylenebis(2,4,7-trimethylindenyl)zirconium
dichloride,
dimethylsilylenebis(2-methyl-4-isopropylindenyl) zirconium
dichloride,
dimethylsilylenebis(2-methyl-4-phenylindenyl)zirconium
20 dichloride,
dimethylsilylenebis(2-methyl-4-naphtylindenyl)zirconium
dichloride,
dimethylsilylenebis(2-methyl-4,5-benzoindenyl)zirconium
dichloride,
25 dimethylsilylenebis(4,5,6,7-tetrahydroindenyl)zirconium
dichloride, diphenylsilylenebis(2-methylindenyl)zirconium
dichloride, diphenylsilylenebis(2-n-propylindenyl)zirconium

dichloride, diphenylsilylenebis(2-n-butylindenyl)zirconium
 dichloride, diphenylsilylenebis(2-n-hexylindenyl)zirconium
 dichloride,
 diphenylsilylenebis(2-isopropylindenyl)zirconium dichloride,
 5 diphenylsilylenebis(2-isobutylindenyl)zirconium dichloride,
 diphenylsilylenebis(2-tert-butylindenyl)zirconium
 dichloride, diphenylsilylenebis(2-phenylindenyl)zirconium
 dichloride,
 diphenylsilylenebis(2-trimethylsilylindenyl)zirconium
 10 dichloride,
 diphenylsilylenebis(2,4-dimethylindenyl)zirconium
 dichloride,
 diphenylsilylenebis(2,4,7-trimethylindenyl)zirconium
 dichloride,
 15 diphenylsilylenebis(2-methyl-4-isopropylindenyl) zirconium
 dichloride,
 diphenylsilylenebis(2-methyl-4-phenylindenyl)zirconium
 dichloride,
 diphenylsilylenebis(2-methyl-4-naphtylindenyl)zirconium
 20 dichloride,
 diphenylsilylenebis(2-methyl-4,5-benzoindenyl)zirconium
 dichloride and
 diphenylsilylenebis(4,5,6,7-tetrahydroindenyl)zirconium
 dichloride.

25 In the general formula [14], ($R^{18}_n-C_5H_{4-n}$) and ($R^{18}_q-C_5H_{4-q}$)
 are each a substituted η^5 -cyclopentadienyl group, and n and
 q are each an integer from 1 to 3. Both of R^{18} s may be the same

or different from each other, and are each a halogen atom, an alkyl group, an aralkyl group, an aryl group, a substituted silyl group, an alkoxy group, an aralkyloxy group, an aryloxy group or a heterocyclic group, provided that the position and/or kind
5 of the R^{18} in the substituted η^5 -cyclopentadienyl group are (is) selected, so that a symmetry plane including M^3 does not exist.

The halogen atom and the alkyl, aralkyl, aryl, substituted silyl, alkoxy, aralkyloxy, aryloxy and heterocyclic groups stated here are as explained in the general formula [13]. As
10 R^{18} , preferred is an alkyl group or a heterocyclic ring group.

Any of n and q is an integer of from 1 to 3.

As examples of the $(R^{18}_n-C_5H_{4-n})$ and $(R^{18}_q-C_5H_{4-q})$ having R^{18} , in which substituted η^5 -cyclopentadienyl groups the position and/or kind of the R^{18} are (is) selected, so that a symmetry
15 plane including M^3 does not exist, there are enumerated a 2-methylcyclopentadienyl group, a 2-ethylcyclopentadienyl group, a 2-n-propylcyclopentadienyl group, a 2-isopropylcyclopentadienyl group, a 2-n-butylcyclopentadienyl group, a 2-isobutylcyclopentadienyl
20 group, a 2-tert-butylcyclopentadienyl group, a 2-n-hexylcyclopentadienyl group, a 3-methylcyclopentadienyl group, a 3-ethylcyclopentadienyl group, a 3-n-propylcyclopentadienyl group, a 3-isopropylcyclopentadienyl group, a 3-n-butylcyclopentadienyl group, a 3-isobutylcyclopentadienyl
25 group, a 3-tert-butylcyclopentadienyl group, a 3-n-hexylcyclopentadienyl group, a

2,3-dimethylcyclopentadienyl group, a
 2,3-diethylcyclopentadienyl group, a
 2,3-di-n-propylcyclopentadienyl group, a
 2,3-diisopropylcyclopentadienyl group, a
 5 2,3-di-n-butylcyclopentadienyl group, a
 2,3-diisobutylcyclopentadienyl group, a
 2,3-di-tert-butylcyclopentadienyl group, a
 2,3-di-n-hexylcyclopentadienyl group, a
 2,4-dimethylcyclopentadienyl group, a
 10 2,4-diethylcyclopentadienyl group, a
 2,4-di-n-propylcyclopentadienyl group, a
 2,4-diisopropylcyclopentadienyl group, a
 2,4-di-n-butylcyclopentadienyl group, a
 2,4-diisobutylcyclopentadienyl group, a
 15 2,4-di-tert-butylcyclopentadienyl group, a
 2,4-di-n-hexylcyclopentadienyl group, a
 3,5-dimethylcyclopentadienyl group, a
 3,5-diethylcyclopentadienyl group, a
 3,5-di-n-propylcyclopentadienyl group, a
 20 3,5-diisopropylcyclopentadienyl group, a
 3,5-di-n-butylcyclopentadienyl group, a
 3,5-diisobutylcyclopentadienyl group, a
 3,5-di-tert-butylcyclopentadienyl group, a
 3,5-di-n-hexylcyclopentadienyl group,
 25 a 2-ethyl-3-methylcyclopentadienyl group, a
 2-methyl-3-ethylcyclopentadienyl group, a
 2-methyl-3-n-propylcyclopentadienyl group, a

2-methyl-3-isopropylcyclopentadienyl group, a
 2-ethyl-3-isopropylcyclopentadienyl group, a
 2-methyl-3-n-butylcyclopentadienyl group, a
 2-methyl-3-isobutylcyclopentadienyl group, a
 5 2-methyl-3-tert-butylcyclopentadienyl group, a
 2-methyl-3-n-hexylcyclopentadienyl group, a
 2-ethyl-4-methylcyclopentadienyl group, a
 2-methyl-4-ethylcyclopentadienyl group, a
 2-methyl-4-n-propylcyclopentadienyl group, a
 10 2-methyl-4-isopropylcyclopentadienyl group, a
 2-ethyl-4-isopropylcyclopentadienyl group, a
 2-methyl-4-n-butylcyclopentadienyl group, a
 2-methyl-4-isobutylcyclopentadienyl group, a
 2-methyl-4-tert-butylcyclopentadienyl group, a
 15 2-methyl-4-n-hexylcyclopentadienyl group, a
 3-ethyl-5-methylcyclopentadienyl group, a
 3-methyl-5-ethylcyclopentadienyl group, a
 3-methyl-5-n-propylcyclopentadienyl group, a
 3-methyl-5-isopropylcyclopentadienyl group, a
 20 3-ethyl-5-isopropylcyclopentadienyl group, a
 3-methyl-5-n-butylcyclopentadienyl group, a
 3-methyl-5-isobutylcyclopentadienyl group, a
 3-methyl-5-tert-butylcyclopentadienyl group, a
 3-methyl-5-n-hexylcyclopentadienyl group, a
 25 2,3,5-trimethylcyclopentadienyl group, a
 2,4,5-trimethylcyclopentadienyl group, a
 2-(2-furyl)-3,5-dimethylcyclopentadienyl group and a

2-(2-furyl)-4,5-dimethylcyclopentadienyl group.

In the above general formula [14], Y^2 is a carbon atom, a silicon atom, a germanium atom or a tin atom. Preferred is a carbon atom or a silicon atom.

5 In the above general formula [14], R^{19} and X^5 are independently of each other a hydrogen atom, a halogen atom, an alkyl group, an aralkyl group, an aryl group, a substituted silyl group, an alkoxy group, an aralkyloxy group, an aryloxy group or a heterocyclic group, and R^{19} and X^5 may be the same
10 or different from each other. The halogen atom and the alkyl, aralkyl, aryl, substituted silyl, alkoxy, aralkyloxy, aryloxy and heterocyclic groups stated here are as already explained as R^{18} .

As R^{19} , preferred is a hydrogen atom or an alkyl group.

15 As X^5 , preferred is a halogen atom, an alkyl group, an aralkyl group or an alkoxy group. Examples thereof are similar to those of X in the general formula [10].

As specific examples of the transition metal compound represented by the general formula [14], those corresponding
20 to the general formula [14] are selected from examples of the transition metal compound represented by the above general formula [10]. Of these, preferred are

dimethylsilylene(cyclopentadienyl)(3-methylcyclopentadienyl)
zirconium dichloride,

25 dimethylsilylene(cyclopentadienyl)(3-tert-butylcyclopentadienyl)zirconium dichloride,

dimethylsilylene(cyclopentadienyl)(2,4-dimethylcyclopentadi

enyl)zirconium dichloride,
 dimethylsilylene(cyclopentadienyl)(2,3,5-trimethylcyclopent
 adienyl)zirconium dichloride,
 dimethylsilylene(3-methylcyclopentadienyl)(4-methylcyclopen
 5 tadienyl)zirconium dichloride,
 dimethylsilylene(3-tert-butylcyclopentadienyl)(4-tert-butyl
 cyclopentadienyl)zirconium dichloride,
 dimethylsilylene(2,4-dimethylcyclopentadienyl)(3,5-dimethyl
 cyclopentadienyl)zirconium dichloride,
 10 dimethylsilylene(2,3,5-trimethylcyclopentadienyl)(2,4,5-tri
 methylcyclopentadienyl)zirconium dichloride,
 methylene(cyclopentadienyl)(3-methylcyclopentadienyl)zircon
 ium dichloride, methylene(cyclopentadienyl)
 (3-tert-butylcyclopentadienyl)zirconium dichloride,
 15 methylene(cyclopentadienyl)
 (2,4-dimethylcyclopentadienyl)zirconium dichloride,
 methylene(cyclopentadienyl)(2,3,5-trimethylcyclopentadienyl
)zirconium dichloride, methylene(3-methylcyclopentadienyl)
 (4-methylcyclopentadienyl)zirconium dichloride,
 20 methylene(3-tert-butylcyclopentadienyl)(4-tert-butylcyclope
 ntadienyl)zirconium dichloride,
 methylene(2,4-dimethylcyclopentadienyl)(3,5-dimethylcyclope
 ntadienyl)zirconium dichloride and
 methylene(2,3,5-trimethylcyclopentadienyl)(2,4,5-trimethylc
 25 ylopentadienyl)zirconium dichloride.

In addition, there are exemplified compounds formed by
 replacing the zirconium atom in the above-mentioned zirconium

compounds with a titanium atom or a hafnium atom.

(C) Organoaluminum compound

As the organoaluminum compound of the component(C) used for the catalyst for addition polymerization in accordance with the present invention, a known organoaluminum compound can be used. Preferred is an organoaluminum compound represented by the following general formula [15].



(In the formula, R^{20} is a hydrocarbon group and all of R^{20} may be the same or different from one another, Y^3 is a hydrogen atom, a halogen atom, an alkoxy group, a aralkyloxy group, or an aryloxy group, all of Y^3 may be the same or different from one another, and c is a number satisfying $0 < c \leq 3$.)

In the general formula [15] representing the organoaluminum compound, as R^{20} , preferred is a hydrocarbon group having 1 to 24 carbon atoms, and more preferred is an alkyl group having 1 to 24 carbon atoms. Specific examples thereof are a methyl group, an ethyl group, a n-propyl group, a n-butyl group, an isobutyl group, a n-hexyl group, a 2-methylhexyl group and a n-octyl group. Preferred are an ethyl group, a n-butyl group, an isobutyl group, a n-hexyl group and a n-octyl group.

When Y^3 is a halogen atom, specific examples thereof are a fluorine atom, a chlorine atom, a bromine atom and an iodine atom, and preferred is a chlorine atom.

When Y^3 is an alkoxy group, preferred is an alkoxy group having 1 to 24 carbon atoms. Specific examples thereof are a methoxy group, an ethoxy group, a n-propoxy group, an isopropoxy

group, a n-butoxy group, a sec-butoxy group, a tert-butoxy group, a n-pentoxy group, a neopentoxy group, a n-hexoxy group, a n-octoxy group, a n-dodecoxy group, a n-pentadecoxy group and a n-eicoxy group, and preferred are a methoxy group, an ethoxy group and a tert-butoxy group.

When Y^3 is an aryloxy group, preferred is an aryloxy having 6 to 24 carbon atoms. Specific examples thereof are a phenoxy group, a 2-methylphenoxy group, a 3-methylphenoxy group, a 4-methylphenoxy group, a 2,3-dimethylphenoxy group, a 2,4-dimethylphenoxy group, a 2,5-dimethylphenoxy group, a 2,6-dimethylphenoxy group, a 3,4-dimethylphenoxy group, a 3,5-dimethylphenoxy group, a 2,3,4-trimethylphenoxy group, a 2,3,5-trimethylphenoxy group, a 2,3,6-trimethylphenoxy group, a 2,4,5-trimethylphenoxy group, a 2,4,6-trimethylphenoxy group, a 3,4,5-trimethylphenoxy group, a 2,3,4,5-tetramethylphenoxy group, a 2,3,4,6-tetramethylphenoxy group, a 2,3,5,6-tetramethylphenoxy group, a pentamethylphenoxy group, an ethylphenoxy group, a n-propylphenoxy group, an isopropylphenoxy group, a n-butylphenoxy group, a sec-butylphenoxy group, a tert-butylphenoxy group, a n-hexylphenoxy group, a n-octylphenoxy group, a n-decylphenoxy group, a n-tetradecylphenoxy group, a naphthoxy group and an anthracenoxy group.

When Y^3 is an aralkyloxy group, preferred is an aralkyloxy having 7 to 24 carbon atoms. Specific examples thereof are a benzyloxy group, a (2-methylphenyl)methoxy group, a (3-methylphenyl)methoxy group, a (4-methylphenyl)methoxy group,

a (2,3-dimethylphenyl)methoxy group, a
 (2,4-dimethylphenyl)methoxy group, a
 (2,5-dimethylphenyl)methoxy group, a
 (2,6-dimethylphenyl)methoxy group, a
 5 (3,4-dimethylphenyl)methoxy group, a
 (3,5-dimethylphenyl)methoxy group, a
 (2,3,4-trimethylphenyl)methoxy group, a
 (2,3,5-trimethylphenyl)methoxy group, a
 (2,3,6-trimethylphenyl)methoxy group, a
 10 (2,4,5-trimethylphenyl)methoxy group, a
 (2,4,6-trimethylphenyl)methoxy group, a
 (3,4,5-trimethylphenyl)methoxy group, a
 (2,3,4,5-tetramethylphenyl)methoxy group, a
 (2,3,5,6-tetramethylphenyl)methoxy group, a
 15 (pentamethylphenyl)methoxy group, an (ethylphenyl)methoxy
 group, a (n-propylphenyl)methoxy group, an
 (isopropylphenyl)methoxy group, a (n-butylphenyl)methoxy group,
 a (sec-butylphenyl)methoxy group, a (tert-butylphenyl)methoxy
 group, a (n-hexylphenyl)methoxy group, a
 20 (n-octylphenyl)methoxy group, a (n-decylphenyl)methoxy group,
 a (n-tetradecylphenyl)methoxy group, a naphthylmethoxy group
 and an anthracenylmethoxy group. Preferred is a benzyloxy group.

Specific examples of the organoaluminum compound
 represented by the general formula [15] are trialkylaluminums
 25 such as trimethylaluminum, triethylaluminum,
 tri-n-propylaluminum, tri-n-butylaluminum,
 triisobutylaluminum, tri-n-hexylaluminum and

tri-n-octylaluminum; dialkylaluminum chlorides such as dimethylaluminum chloride, diethylaluminum chloride, di-n-propylaluminum chloride, di-n-butylaluminum chloride, diisobutylaluminum chloride and di-n-hexylaluminum chloride;
5 alkylaluminum dichlorides such as methylaluminum dichloride, ethylaluminum dichloride, n-propylaluminum dichloride, n-butylaluminum dichloride, isobutylaluminum dichloride and n-hexylaluminum dichloride; dialkylaluminum hydrides such as dimethylaluminum hydride, diethylaluminum hydride,
10 di-n-propylaluminum hydride, di-n-butylaluminum hydride, diisobutylaluminum hydride and di-n-hexylaluminum hydride; trialkoxyaluminums such as trimethoxyaluminum, triethoxyaluminum and tri(tert-butoxy)aluminum; alkylalkoxyaluminums such as methy(dimethoxy)aluminum,
15 methyl(diethoxy)aluminum and methyl(di-tert-butoxy)aluminum; dialkylalkoxyaluminums such as dimethy(methoxy)aluminum, dimethyl(ethoxy)aluminum and dimethyl(tert-butoxy)aluminum; triaryloxyaluminums such as triphenoxyaluminum, tris(2,6-diisopropylphenoxy)aluminum and
20 tris(2,6-diphenylphenoxy)aluminum; alkyl diaryloxyaluminums such as methyl(diphenoxy)aluminum, methylbis(2,6-diisopropylphenoxy)aluminum and methylbis(2,6-diphenylphenoxy)aluminum; and dialkylaryloxyaluminums such as dimethyl(phenoxy)aluminum,
25 dimethyl (2,6-diisopropylphenoxy)aluminum and dimethyl (2,6-diphenylphenoxy)aluminum.

Among these, preferred are trialkylaluminums, more

preferred are trimethylaluminum, triethylaluminum, tri-n-butylaluminum, triisobutylaluminum, tri-n-hexylaluminum and tri-n-octylaluminum, and particularly preferred are triisobutylaluminum and tri-n-octylaluminum.

5 These organoaluminum compounds may be used singly or in combination of two or more.

 The component (B) is used in an amount of usually from 1×10^{-6} to 1×10^{-3} mol, and preferably from 5×10^{-6} to 5×10^{-4} mol per g of the component (A). The component (C) is used in
10 an amount of preferably from 0.01 to 10,000, more preferably from 0.1 to 5,000, and the most preferably from 1 to 2,000 in terms of a molar ratio (C)/(B), namely in terms of the aluminum atom in the organoaluminum compound of the compound (c) to the transition metal atom in the transition metal compound of the
15 component (B).

 As the catalyst for addition polymerization in accordance with the present invention, a reaction product obtained in advance by contacting the component (A) and the component (B) and optionally the component (C) with one another may be used,
20 or these components may be used in a manner of separately feeding in a polymerization reaction apparatus. When the components (A), (B) and (C) are used, optional two components selected therefrom may be contacted in advance with each other, followed by contact of the remaining one component.

25 When the stereoregular polymerization of α -olefin is carried out, it is recommendable to use the catalyst obtained in advance by contacting the component (A), the component (B)

and the component (C).

The contact may be conducted in a reactor for catalyst preparation use different from a polymerization reactor, or in a polymerization reactor, in which neither a monomer nor a comonomer is introduced. The contact may be conducted by supplying the component (A), the component (B) and the component (C) at the same time, or one after another in an arbitrary order in the reactor for catalyst preparation use or the polymerization reactor. Conditions of contacting the component (A), the component (B) and the component (C) are not particularly limited, and a contact temperature is preferably from -5 to 90°C, and more preferably from 5 to 40°C.

As a method for supplying the catalyst to a polymerization vessel, there are usually applied a method comprising supplying the catalyst under a water-free condition using an inert gas such as nitrogen or argon, or hydrogen or ethylene, and a method comprising supplying the catalyst in a slurry state obtained by dilution in a solvent. Respective catalyst components may be separately supplied, or before supplying, arbitrarily selected components may be contacted in advance in an arbitrary order.

A method for supplying respective catalyst components to the catalyst preparation reactor or the polymerization reactor is also not particularly limited. There can be applied a method comprising supplying respective components in each solid state, or in a solution, suspension or slurry state obtained using a hydrocarbon solvent, provided that any ingredient capable of

de-activating the catalyst components such as water and oxygen has been sufficiently eliminated from the hydrocarbon solvent. Examples of the solvent are aliphatic hydrocarbon solvents such as butane, pentane, heptane and octane, aromatic hydrocarbon solvents such as benzene and toluene and halogenated hydrocarbons such as methylene chloride. Preferred are aliphatic hydrocarbons and aromatic hydrocarbons.

When the catalyst components are supplied in their solution, suspension or slurry state, a concentration of the component (A) is usually from 0.01 to 1000 g/l, and preferably from 0.1 to 500 g/l. A concentration of the component (C) is usually from 0.0001 to 100 mol/l, and preferably from 0.01 to 10 mol/l in terms of Al atom. A concentration of the component (B) is usually from 0.0001 to 1000 mmol/l, and preferably from 0.01 to 50 mmol/l in terms of the transition metal atom.

The polymerization process is not particularly limited, and it is permitted to apply a gas phase polymerization carried out in a gaseous monomer or a solution polymerization or slurry polymerization using a solvent. A solvent used for the solution polymerization or the slurry polymerization includes an aliphatic hydrocarbon solvent such as butane, hexane, pentane, heptane and octane, an aromatic hydrocarbon solvent such as benzene and toluene, and a halogenated hydrocarbon solvent such as methylene chloride. Alternatively, it is possible to use an olefin itself as a solvent (bulk polymerization). As the polymerization process, any of a batch polymerization and a continuous polymerization

can be applied. It is permitted to carry out the polymerization dividing into two or more steps different from one another in their reaction conditions. A polymerization time can be appropriately determined according to a kind of the desired
5 olefin polymer and a reaction apparatus, and it is permitted to adopt a period of time of from 1 minute to 20 hours.

The present invention can be particularly suitably applied to a polymerization accompanied by the formation of addition polymer particles (for example, a slurry
10 polymerization, a gas phase polymerization and a bulk polymerization).

The slurry polymerization may be carried out in a conventional slurry polymerization manner and condition, but it is not limited thereto. According to a preferred slurry
15 polymerization, a monomer (and a comonomer), a material to be supplied, a diluent and the like are, if desired continuously added, and a continuous type reactor is used, so that a produced polymer is continuously or at least periodically taken out. Further, there can be applied a means
20 using a loop reactor or a means using a stirring reactor. In addition, more than one stirring reactor, which may be different in a reactor or in a reaction condition, may be combined in series, in parallel or in series and parallel.

As the diluent, there can be used an inert diluent
25 (medium) such as paraffin, cycloparaffin and aromatic hydrocarbons. A temperature of the polymerization reactor or a reaction zone is usually from about 0 to about 150°C,

and preferably from 30 to 100°C. A pressure can be varied within a range of from about 0.1 to about 10 MPa, and preferably from 0.5 to 5 MPa. It is permitted to apply a pressure, under which a monomer and a comonomer are contacted with each other while maintaining the catalyst in a suspended state, and the medium and at least a part of the monomer and comonomer in a solution phase. Accordingly, the medium, temperature and pressure may be selected so as to produce the addition polymer in a solid particle form and recover it in that form.

A molecular weight of the addition polymer can be controlled by known manners, for example, by controlling a temperature of a reaction zone or introducing hydrogen.

The catalyst components and a monomer (and a comonomer) can be fed into a reactor or a reaction zone in an arbitrary order according to a conventional manner. For example, the catalyst components and a monomer (and a comonomer) are fed into a reaction zone at the same time or one after the other. If desired, the catalyst components and a monomer (and a comonomer) can be subjected to pre-contact in an inert atmosphere before their contact.

The gas phase polymerization may be carried out in a conventional gas phase polymerization manner and condition, but it is not limited thereto. As a gas phase polymerization apparatus, a fluidized bed reactor, preferably a fluidized bed reactor having an expansion can be used. A reaction apparatus equipped with a stirring blade in its reaction zone may also be used without any problem.

Respective components can be fed into a polymerization zone in a water-free state using an inert gas such as nitrogen or argon, or hydrogen or ethylene, or in feeding, respective components can be dissolved or diluted in a solvent to form
5 a solution or a slurry. The catalyst components may be fed independently, or in feeding, arbitrarily selected catalyst components may be contacted in advance with one another in an arbitrary order.

With respect to polymerization conditions, a
10 temperature is lower than that at which the polymer is melted, preferably from 0 to 150°C, and particularly preferably from 30 to 100°C. Furthermore, for the purpose of controlling the melt flow of a final product, it is permitted to add hydrogen as a molecular weight regulator. In addition, it is permitted
15 that an inert gas coexists in the mixed gas during polymerization.

In the present invention, a pre-polymerization as mentioned below may be carried out before carrying out the above-mentioned polymerization (real polymerization).

20 The pre-polymerization can be carried out by feeding a small amount of an olefin in the presence of the modified particle (A) and the transition metal compound (B), or further in the presence of the organoaluminum compound (A). It is recommendable to carry out in a slurry state. Examples of
25 a solvent used for formation of the slurry are inert hydrocarbons such as propane, butane, isobutane, pentane, isopentane, hexane, heptane, octane, cyclohexane, benzene

and toluene. Further, in forming the slurry, it is permitted to use a liquefied olefin in place of a part or the whole of the inert hydrocarbon.

5 In the pre-polymerization, an amount of the organoaluminum compound (C) can be determined within a wide range of from 0.5 to 700 mol per mol of the transition metal compound (B). Preferable is from 0.8 to 500 mol per mol thereof, and particularly preferable is from 1 to 200 mol per mol thereof.

10 An amount of the olefin to be subjected to the pre-polymerization is usually from 0.01 to 1000 g per g of the modified particle, preferably from 0.05 to 500 g per g thereof, and particularly preferably from 0.1 to 200 g per g thereof.

15 In the pre-polymerization, a slurry concentration is preferably from 0.1 to 50 g-above-mentioned modified particle/l-solvent, and particularly preferably from 0.5 to 20 g-the above-mentioned modified particle/l-solvent. A pre-polymerization temperature is preferably from -20 to 20 100°C, and particularly preferably from 0 to 80°C. A partial pressure of the olefin in the gas phase during the pre-polymerization is preferably from 0.001 to 2 MPa, and particularly preferably from 0.01 to 1 MPa, excepting an olefin which is liquid under the pressure and temperature
25 conditions of pre-polymerization. Further, a pre-polymerization time is not particularly limited, and usually from 2 minutes to 15 hours.

In carrying out the pre-polymerization, as a process for supplying the above-mentioned modified particle (A), transition metal compound (B), organoaluminum compound (C) and olefin, there can be applied a process comprising the
5 step of contacting the above-mentioned modified particle (A) and the transition metal compound (B), and if desired, the organoaluminum compound (C), and thereafter supplying the olefin, a process comprising the step of contacting the above-mentioned modified particle (A), the transition metal
10 compound (B) and the olefin, and thereafter supplying the organoaluminum compound (C), and a process comprising the step of contacting the organoaluminum compound (C) and the transition metal compound (B) in the presence of the olefin, and thereafter supplying the above-mentioned modified
15 particle (A). At the time when the above-mentioned modified particle (A) and the organoaluminum compound (C) are contacted with each other, it is recommendable that the olefin coexists in advance. Further, as a supply method of the olefin, any of a method comprising supplying the olefin in order while
20 keeping the pre-determined pressure of the polymerization zone, and a method comprising supplying all the pre-determined amount of the olefin in the first place can be applied. It is also permitted to add a chain transfer agent such as hydrogen to regulate a molecular weight of the polymer obtained.

25 In the present invention, that obtained by the pre-polymerization as mentioned above can be used as a catalyst component or a catalyst. The pre-polymerized

catalyst component in accordance with the present invention includes a pre-polymerized catalyst component for addition polymerization obtained by pre-polymerizing an olefin in the presence of the catalyst for addition polymerization obtained by contacting the above-mentioned modified particle (A) and the transition metal compound of the Groups 3 to 11 or lanthanoide series with each other, and a pre-polymerized catalyst component for addition polymerization obtained by pre-polymerizing an olefin in the presence of the catalyst for addition polymerization obtained by contacting the above-mentioned modified particle (A), the transition metal compound of the Groups 3 to 11 or lanthanoide series and the organoaluminum compound (C) with one another. The pre-polymerized catalyst in accordance with the present invention includes a catalyst for addition polymerization obtained by pre-polymerizing an olefin in the presence of the catalyst for addition polymerization obtained by contacting the above-mentioned modified particle (A) and the transition metal compound of the Groups 3 to 11 or lanthanoide series with each other, and a catalyst for addition polymerization obtained by pre-polymerizing an olefin in the presence of the catalyst for addition polymerization obtained by contacting the above-mentioned modified particle (A), the transition metal compound of the Groups 3 to 11 or lanthanoide series and the organoaluminum compound (C) with one another. The catalyst obtained using said pre-polymerized catalyst component for addition polymerization in accordance with the

present invention includes a catalyst for addition polymerization obtained by contacting said pre-polymerized catalyst component for addition polymerization and the organoaluminum compound (C) with each other.

5 Examples of the monomer used in the polymerization are olefins having 2 to 20 carbon atoms, diolefins, cyclic olefins, alkenyl aromatic hydrocarbons and polar monomers. Two or more kinds of the monomer may be used at the same time. It is permitted to use two or more monomers at the same time.

10 Specific examples thereof are olefins such as ethylene, propylene, 1-butene, 1-pentene, 4-methyl-1-pentene, 5-methyl-1-hexene, 1-hexene, 1-heptene, 1-octene, 1-nonene and 1-decene; diolefins such as 1,5-hexadiene, 1,4-hexadiene, 1,4-pentadiene, 1,7-octadiene, 1,8-nonadiene, 15 1,9-decadiene, 4-methyl-1,4-hexadiene, 5-methyl-1,4-hexadiene, 7-methyl-1,6-octadiene, 5-ethylidene-2-norbornene, dicyclopentadiene, 5-vinyl-2-norbornene, norbornadiene, 5-methylene-2-norbornene, 1,5-cyclooctadiene, 20 5,8-endomethylenehexahydronaphthalene, 1,3-butadiene, isoprene, 1,3-hexadiene, 1,3-octadiene, 1,3-cyclooctadiene and 1,3-cyclohexadiene; cyclic olefins such as norbornene, 5-methyl-2-norbornene, 5-ethyl-2-norbornene, 5-butyl-2-norbornene, 25 5-phenyl-2-norbornene, 5-benzyl-2-norbornene, tetracyclododecene, tricyclodecene, tricycloundecene, pentacyclopentadecene, pentacyclohexadecene,

methacrylate, n-propylmethacrylate, isopropylmethacrylate, n-butylmethacrylate and isobutylmethacrylate, unsaturated dicarboxylic acids such as maleic acid and itaconic acid, vinyl esters such as vinyl acetate, vinyl propionate, vinyl caproate, vinyl caprate, vinyl laurate, vinyl stearate and vinyl trifluoroacetate, and unsaturated carboxylic acid glycidyl esters such as glycidyl acrylate, glycidyl methacrylate and monoglycidyl itaconate.

The present invention can be applied to homopolymerization or copolymerization of these monomers. Examples of copolymer-constituting monomers are ethylene-propylene, ethylene-1-butene, ethylene-1-hexene, ethylene-1-octene, propylene-1-butene, ethylene-propylene-butene and ethylene-propylene-1-hexene, but the present invention should not be limited thereto.

Among those mentioned above, α -olefins having 3 to 20 carbon atoms can be used as the monomer for polymerization carried out using the above-mentioned stereoregular catalyst.

Further, copolymerization between said α -olefin and a comonomer copolymerizable with said α -olefin can be suitably carried out. Examples of the comonomer copolymerizable with said α -olefin are ethylene and diolefin compounds as mentioned above.

The stereoregular polymerization is suitable particularly for the production of an isotactic stereoregular α -olefin polymer, more particularly for the production of

an isotactic stereoregular propylene polymer.

Specific examples of the isotactic stereoregular propylene polymer are a homopolymer of propylene; a random copolymer of propylene with a comonomer such as ethylene
5 and/or an α -olefin having 4 to 12 carbon atoms in an amount controlled as not to lose crystallinity; and a block copolymer obtained by performing homopolymerization of propylene or copolymerization of propylene with ethylene or an α -olefin having 4 to 12 carbon atoms (referred to as the first part
10 polymerization), and then performing one step or multi-step polymerization of an α -olefin having 3 to 12 carbon atoms with ethylene (referred to as the latter part polymerization). The amount controlled as not to lose crystallinity for the random copolymer varies depending upon a kind of the comonomer.
15 For example, in the case of ethylene, an amount of the repeating unit derived from ethylene in the copolymer is usually not more than 10% by weight, and in the case of other α -olefins such as 1-butene, an amount of the repeating unit derived from the α -olefin in the copolymer is usually not more than
20 30% by weight, and preferably not more than 10% by weight. In the first part polymerization for the block copolymer, for example, a polymerization amount of ethylene is usually not more than 10% by weight, preferably not more than 3% by weight, and more preferably not more than 0.5% by weight,
25 and a polymerization amount of the α -olefin is usually not more than 15% by weight, and preferably not more than 10% by weight. In the latter part polymerization, a polymerization

amount of ethylene is usually from 20 to 80% by weight, and preferably from 30 to 50% by weight.

Specific examples of a combination of a copolymer-constituting monomer are propylene-ethylene, propylene-1-butene, propylene-1-hexene and propylene-1-octene, but the present invention should not be limited thereto.

As an index of the isotactic stereoregularity, an isotactic pentad proportion is used.

It means a proportion of a propylene monomer unit in the middle of a pentad unit isotactic chain in a crystalline polypropylene molecule chain, which is measured using the method disclosed in "Macromolecules", by A. Zambelli et al., 6, 925 (1973), namely, using ^{13}C -NMR. The pentad unit isotactic chain can be rephrased by a chain formed through meso linkage of five propylene monomer units in succession. Attribution of the NMR absorption peak is based on "Macromolecules", 8, 687 (1975), issued after that. As the case may be, the isotactic pentad proportion is abbreviated as "mmmm". An upper limit of a theoretical mmmm value is 1.000.

When the stereoregular polymerization of the α -olefin is carried out, particularly when that of propylene is carried out, the obtained stereoregular α -olefin polymer is preferably that having mmmm of not less than 0.900 (more preferably not less than 0.940, and much more preferably not less than 0.955).

Example

The present invention is illustrated in more detail with reference to Examples and Comparative Examples, but the present invention is not limited thereto. Measurement values of respective items in Examples were obtained by the following methods.

(1) A content of a repeating unit derived from the α -olefin in the polymer was obtained as follows. Using an infrared spectrophotometer (FT-IR 7300, manufactured by Japan Spectroscopic Co., Ltd.), characteristic absorption of ethylene and the α -olefin were measured to prepare a calibration curve, and the content was obtained using the calibration curve and expressed in terms of a short chain branch number (SCB) per 1000 carbon atoms.

(2) Intrinsic viscosity = $[\eta]$: measured in a tetralin solution at 135°C using Ubbelohde viscometer. (Unit: dl/g)

(3) Melt flow rate = MFR: a melt flow rate value (Unit: g/10 min.) measured according to the method prescribed in JIS K7210-1995, under conditions of 190°C and a load of 21.18 N (2.16 kg).

(4) Swelling ratio = SR: a value obtained by dividing a strand diameter measured at the time of the MFR measurement by an inner diameter of a die, namely 2.095 mm.

(5) Melt flow rate ratio = MFRR: a value obtained by dividing a melt flow rate value obtained according to the method prescribed in JIS K7210-1995, under conditions of 190°C and a load of 211.82 N (21.60 kg) by the melt flow rate (MFR) value obtained under the

load of 21.18 N (2.16 kg).

In the above-mentioned melt flow rate measurement, a polymer blended in advance with 1000 ppm of an antioxidant was used.

5 (6) Molecular weight and molecular weight distribution

It was measured under the conditions described below according to gel permeation chromatography (GPC). A calibration curve was prepared using a standard polystyrene. The molecular weight distribution was represented by a ratio
10 (Mw/Mn) of a weight average molecular weight (Mw) to a number average molecular weight (Mn).

Measurement machine: 150C type, manufactured by Millipore Waters Co.

Column: TSK-GEL GMH-HT 7.5 × 600 × 2 columns

15 Measurement temperature: 140°C

Solvent: ortho-dichlorobenzene

Measurement concentration: 5mg/5ml

Provisos that; in Examples 6 to 12 and 17, the measurement was carried out using Column: TSK-GEL GMH6-HT 7.5 × 300 ×
20 3 columns, and in Examples 67 to 69, the measurement was carried out under the following column, measurement temperature and measurement concentration conditions.

Column: Shodex AT806-M/S

Measurement temperature: 145°C set up

25 Measurement concentration: 8mg/8ml o-dichlorobenzene

(7) Elementary analysis

Zn: A sample was put in a sulfuric acid aqueous solution

(1 mol/l), a supersonic wave was applied thereto to extract a metal component, and the obtained liquid portion was quantitatively determined according to an IPC emission spectrochemical analysis.

5 F: A sample was burned in a flask filled with oxygen, thereby producing a combustion gas, the gas was absorbed in a sodium hydroxide aqueous solution (10%), and the resulting solution was quantitatively determined according to an ion selective electrode method.

10 (8) Film formation of polymer

To 0.030 kg of the polymer powder obtained by polymerization, IRGANOX 1076 (manufactured by Ciba Specialty Chemicals Co., Ltd.) as an antioxidant in an amount corresponding to 1500 ppm and calcium stearate as a
15 neutralizing agent in an amount corresponding to 500 ppm were blended, and the blend was processed under the following molding conditions according to a tubular film process, thereby obtaining a film having a thickness of 15 μ m. A degree of production of fish eyes was examined to determine
20 appearance of the film.

Extruder: single screw 20 mm ψ , screw rotating speed: 16 rpm, die: 25 mm ψ , rip width: 0.7 mm, drawing speed: 5.5 m/min. and temperature: 170°C

(9) Extended X-ray measurement

25 A sample and a cell attached with a polyimide film (KAPTON, manufactured by DU PONT-TORAY CO., LTD.) were placed in a nitrogen bag, and after purging with nitrogen, the sample

was introduced in the cell. The cell sealed hermetically was taken out and mounted to an X-ray measurement apparatus. The extended X-ray measurement was carried out as follows.

X-ray diffraction apparatus: vertical rotating anode
5 X-ray generator UltraX18 manufactured by Rigaku International Corporation

Load: 50kV-100 mA

Focus: line

Scan mode: $2\theta/\theta$ (2° /min.)

10 Measured angle range: 5° to 70°

Slit: divergence slit 1°

receiving slit 0.15 mm

scatter slit 1°

Detector: scintillation counter

15 Monochromater: graphite

(10) Measurement method of XAFS

① Preparation of measurement sample

(i) A center portion of a polyester sheet was cut off, and a polyimide film (KAPTON, manufactured by DU PONT-TORAY
20 CO.,LTD.) was attached thereto.

(ii) Another polyester sheet (PET) whose center portion had been similarly cut off was placed thereon.

(iii) A sample was spread over the cut off portion.

(iv) The other polyester sheet (PET) whose center portion
25 had been cut off and attached with a polyimide film (KAPTON, manufactured by DU PONT-TORAY CO.,LTD.) was placed thereon,

thereby obtaining a measurement sample.

Among the above-mentioned operations, (iii) and (iv) were carried out under a nitrogen atmosphere.

② Measurement

5 Measurement of K absorption edge XAFS on a Zn atom in the sample was carried out using a measurement apparatus of XAFS, with the beam line 10B (BL-10B) of the Photon Factory in the National Laboratory for High-Energy Physics. Using an Si (311) channel cut analyzing crystal, an incoming X-ray intensity (I_0) and a transmitted X-ray intensity (I_t) were measured at ambient
10 temperature using a 17 cm ion chamber with use of nitrogen as a detection gas and a 31 cm ion chamber with use of a mixed gas of 50% nitrogen and 50% argon as a detection gas, respectively. At this time, measuring regions, intervals of measuring points
15 and elapsed time per measuring point were set as follows.

 In the region between 9160 eV and 9610 eV of an incoming X-ray energy (E), the elapsed time was 1 second at each interval of 6.43 eV (70 measuring points).

 In the region between 9610 eV and 9760 eV of an incoming
20 X-ray energy (E), the elapsed time was 1 second at each interval of 1 eV (150 measuring points).

 In the region between 9760 eV and 10160 eV of an incoming X-ray energy (E), the elapsed time was 2 seconds at each interval of 2.5 eV (160 measuring points).

25 In the region between 10160 eV and 10760 eV of an incoming X-ray energy (E), the elapsed time was 2 seconds at each interval

of 6 eV (101 measuring points). Incidentally, calibration of the energy was conducted by adjusting an angle of the analyzing crystal to 24.934° at an energy value position (8980.3 eV) of pre-edge peak appeared in the X-ray absorption near edge structure (XANES) spectra on a K absorption edge of a metal copper.

Determination method of EXAFS

The K absorption edge energy E_0 on the Zn atom in the sample was found to be 9663 eV. Incidentally, E_0 means an energy value at which a first order differential coefficient thereof becomes an absolute maximum in the XANES spectra.

A background determined by applying Victoreen's expression ($A\lambda^3 - B\lambda^4 + C$; λ is a wave length of the incoming X-ray, and A, B and C are an optional constant) to an absorption coefficient of an energy region lower than the absorption edge according to the least-squares method was deducted, and successively an absorbance of a lone atom was estimated according to a weighed Cubic Spline method. Thereby, EXAFS function $\chi(k)$ was extracted. Incidentally, k means a photoelectron wave as defined by $0.5123 \times (E - E_0)^{1/2}$, wherein a unit of k is $\text{\AA}^{-1}$.

Finally, the k^3 -weighed EXAFS function $k^3 \chi(k)$ was processed according to a Fourier transform within a range of k of from $2.5 \text{ \AA}^{-1}$ to $15.7 \text{ \AA}^{-1}$, thereby obtaining a radial distribution function, provided that Hanning function and $0.05 \text{ \AA}^{-1}$ were used as a window function of the Fourier transform and the window width, respectively.

(11) ^{13}C -NMR measurement

A solution of 200 mg of the polymer in 3 ml of a mixed solvent of o-dichlorobenzene/deuterated o-dichlorobenzene (o-dichlorobenzene/deuterated o-dichlorobenzene = 4/1 (volume ratio)) was subjected to measurement using JNM-EX270,
5 manufactured by JEOL.

(12) Bulk density

The bulk density was measured according to JISK-6721-1966.

(13) Melt tension (MT): Using 5 g of a sample, a molten resin strand was extruded through an orifice having a diameter
10 of 2.09 mm ϕ and a length of 8 mm at 190°C with a piston having an extrusion speed of 5.5 mm/min. and wound into a roller having a diameter of 50 mm while increasing a rotating speed at a rate of 40 rpm/min., wherein a tension immediately before cutting of the strand was measured to determine the melt tension (MT)
15 (unit = cN).

[Example 1]

(1) Production of contact product (e)

In a 5 l four necked flask purged with nitrogen, 1.48 l of tetrahydrofuran and 1.47 l (2.93 mol) of a hexane solution
20 of diethylzinc (1.99 mol/l) were introduced and cooled to -50°C. To the mixture was dropped a solution prepared by dissolving 269.5 g (1.46 mol) of pentafluorophenol in 500 ml of tetrahydrofuran over 25 minutes. After completion of dropping, the mixture was gradually heated to raise the
25 temperature to ambient temperature and stirred for 3.2 hours. Thereafter, the mixture was heated to 40°C and stirred for

35 minutes. The temperature was lowered to 20°C using an ice bath, and a solution prepared by dissolving 39.85 g (2.21 mol) of H₂O in 200 ml of tetrahydrofuran was dropped thereto over 1.3 hours. As a result, the mixture was separated into
5 a yellow transparent liquid product and a yellow gel product. The mixture was stirred for 1 hour at 52°C, and thereafter allowed to stand for 12 hours at ambient temperature. The yellow transparent liquid product and the yellow gel product were separated from each other and placed into different
10 flasks purged with nitrogen, respectively. Both were treated to remove a volatile component by distillation and dried for 12 hours at 120°C and for 8 hours at 130°C. Thereafter, 72% by weight of a solid product obtained from the yellow transparent liquid product and all of a solid product obtained
15 from the yellow gel product were combined and introduced into a 5 l four necked flask purged with nitrogen, and 3 l of tetrahydrofuran was added thereto. The mixture was heated to 50°C, and cooled to ambient temperature and allowed to stand for 3 days. Under reduced pressure, a volatile component
20 was removed by distillation, and thereafter, the residue was dried under reduced pressure for 11 hours at 120°C. As a result, 408 g of a solid product was obtained.

(2) Production of component (A)

In a 5 l four necked flask purged with nitrogen, 408 g
25 of the solid product obtained in the above Example 1 (1) and 3 l of tetrahydrofuran were introduced and stirred. To the mixture, 288.4 g of silica (Sylopol 948, manufactured by Davison Co.,

Ltd.; average particle diameter = $61\ \mu\text{m}$; pore volume = $1.61\ \text{ml/g}$; specific surface area = $296\ \text{m}^2/\text{g}$), which had been heat-treated at 300°C under a nitrogen stream, was added. The mixture was heated to 40°C , stirred for 2 hours and thereafter allowed to stand, thereby precipitating a solid component. At the time when an interface between a layer of the precipitated solid component and an upper layer of a slurry portion appeared, the upper slurry portion was removed. For completing washing, 3 l of tetrahydrofuran was added to the residue, and the mixture was stirred, and thereafter allowed to stand, thereby precipitating a solid component, and then the upper slurry portion was removed similarly at the time when the interface appeared. Said washing operation was repeated 5 times. Further, 3 l of hexane was added thereto, and the mixture was stirred, and thereafter allowed to stand, thereby precipitating a solid component, and then the upper slurry portion was removed similarly at the time when the interface appeared. Using a glass filter, the liquid component was removed, and thereafter, the residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 422 g of a component (A). An elementary analysis revealed $\text{Zn} = 2.4\ \text{mmol/g}$, and $\text{F} = 3.8\ \text{mmol/g}$.

(3) Pre-polymerization

A 0.4 l-volume autoclave equipped with a stirrer was made vacuum, hydrogen was fed therein to obtain a pressure of 0.003 MPa, 90 g of butane was fed therein as a solvent, and then a temperature of the autoclave was raised to 50°C . Further, ethylene

was fed therein to obtain a partial pressure of 0.2 MPa, and after stabilizing the system, 1.2 mmol of triisobutylaluminum, 45 μ mol of racemic ethylenebis(1-indenyl)zirconium dichloride, and successively 0.66 g of the component (A) obtained in the above Example 1 (2) were introduced therein to initiate polymerization. During the polymerization, a mixed gas of ethylene and hydrogen (hydrogen concentration = 0.27 mol%) was continuously supplied while adjusting so as to keep the gas phase pressure in the autoclave to 0.65 MPa. After 240 minutes, ethylene, butane and hydrogen gas were released, and a solid remaining in the autoclave was dried for 30 minutes at ambient temperature under reduced pressure. As a result, a polyethylene-pre-polymerized catalyst component for addition polymerization was obtained in a yield of 57 g per g of the component (A). Intrinsic viscosity $[\eta]$ and MFR of said polyethylene were 1.65 dl/g and 2.4 g/10 min., respectively.

(4) Real polymerization

A 5 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.018 MPa, 1066 g of butane and 200 ml of 1-hexene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 1.06 mol%. To the resultant

was added 1.5 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 1.13 g of the pre-polymerized catalyst component for addition polymerization obtained in the above Example 1 (3) was added thereto. Polymerization was continued for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.76 mol%) so as to keep the whole pressure constant. As a result, 90 g of an olefin polymer having a good particle property was obtained. Polymerization activity per the solid catalyst component (the component (A) obtained in the above Example 1 (2)) was found to be 4500 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 15.5, MFR = 1.02, MFRR = 86, SR = 1.36 and MT = 3.7.

(5) Formation of film

Using the ethylene/1-hexene copolymer obtained in the above (4), the formation of film was conducted, thereby obtaining a film having a little fish eye and a good appearance.

[Example 2]

(1) Pre-polymerization

In a 210 l-volume autoclave equipped with a stirrer, which had been purged with nitrogen, 68 l of butane containing isobutylaluminum in a concentration of 3.7 mmol/l and 25 l of hydrogen of ordinary temperature and pressure were fed, and then a temperature of the autoclave was raised to 40°C. Further, ethylene was fed therein to make a gas phase pressure in the autoclave 0.6 MPa, and after stabilizing the system, 89.3 mmol

of triisobutylaluminum, 12.8 mmol of racemic ethylenebis(1-indenyl)zirconium dichloride, and successively 170 g of the component (A) obtained in the above Example 1 (2) were introduced therein to initiate polymerization. During the
5 polymerization, 3.0 kg/hour of ethylene and 7.5 l/hour of hydrogen of ordinary temperature and pressure were supplied while keeping the temperature to 50°C, and polymerization was continued for 2.8 hours. After completion of the polymerization, ethylene, butane and hydrogen gas were released, and a remaining solid
10 was dried in vacuo at ambient temperature under a nitrogen stream, thereby obtaining a polyethylene-pre-polymerized catalyst component for addition polymerization in a yield of 47 g per g of the component (A). Intrinsic viscosity $[\eta]$ and MFR of said polyethylene were 1.14 dl/g and 5.15 g/10 min., respectively.

15 (2) Real polymerization

Using the pre-polymerized catalyst component for addition polymerization obtained in the above (1) and a continuous type fluidized gas phase polymerization apparatus, copolymerization of ethylene and 1-hexene was carried out at 70°C under the whole
20 pressure of 2.0 MPa. The pre-polymerized catalyst component for addition polymerization obtained in the above (1) and triisobutylaluminum were continuously supplied in rates of 192 g/hour and 40 mmol/hour, respectively, while continuously feeding ethylene, 1-hexene, hydrogen, nitrogen and butane so
25 as to keep the gas composition constant. As a result, an ethylene/1-hexene copolymer having a good particle property was

obtained in 15.4 kg/hour. The obtained ethylene/1-hexene copolymer showed, SCB = 21.4, MFR = 3.00, MFRR = 80, SR = 1.56 and MT = 2.7.

(3) Formation of film

5 Using the ethylene/1-hexene copolymer obtained in the above (2), the formation of film was conducted, thereby obtaining a film having a little fish eye and a good appearance.

[Comparative Example 1]

(1)

10 To a 5 l four necked flask purged with nitrogen, a stirrer, a dropping funnel and a thermometer were equipped. In the flask, 382 g of silica (Sylopol 948, manufactured by Davison Co., Ltd.; average particle diameter = $64\ \mu\text{m}$; pore volume = 1.62 ml/g; specific surface area = $312\ \text{m}^2/\text{g}$), which had been heat-treated
15 at 300°C under a nitrogen stream, was get. To the silica, 3.3 l of toluene was added to form a slurry, and the slurry was cooled to 5°C using an ice bath, and 375 ml of a toluene solution of trimethylaluminum having a concentration of 2.1 mmol/ml was gradually dropped thereto. At this moment, generation of a gas
20 was observed. The mixture was stirred for 30 minutes at 5°C , and stirred for 2 hours at 80°C , and thereafter, a supernatant was removed. The remaining solid product was washed 4 times with each 3.3 l of toluene. Thereafter, 3.3 l of toluene was again added thereto to form a slurry.

25 (2) The slurry obtained in the above Comparative Example 1 (1) was cooled to 5°C using an ice bath, and thereafter 420 ml of

a toluene solution of pentafluorophenol adjusted to a concentration of 2 mmol/ml was gradually dropped thereto. At this moment, generation of a gas was observed. The mixture was stirred for 30 minutes at 5°C, and stirred for 2 hours at 80°C, and thereafter, a supernatant was removed by filtration. The remaining solid product was washed 4 times with each 3.3 l of toluene, and washed 2 times with each 3.3 l of hexane. Thereafter, the solid product was dried under reduced pressure to obtain 434 g of a solid product having flowability. Elementary analysis revealed, Al = 1.2 mmol/g and F = 4.6 mmol/g.

(3) Polymerization A 5 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.007 MPa, 1145 g of butane and 55 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.37 mol% and 1-butene = 1.55 mol%. To the resultant was added 1.5 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Next, 0.5 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml was introduced therein, and successively, 36 mg of the solid product obtained in the above Comparative Example 1 (2) was added therein as a solid catalyst

component. Polymerization was continued for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.22 mol%) so as to keep the whole pressure constant. As a result, 135 g of an olefin polymer was obtained. Polymerization activity per the zirconium atom was found to be 1.4×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 3720 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 12.4, MFR = 2.55, MFRR = 48, and SR = 1.35.

10 (4) Formation of film

Using the ethylene/1-hexene copolymer obtained in the above (3), the formation of film was conducted, thereby obtaining a film having many fish eyes and a poor appearance.

[Example 3]

15 (1) Production of contact product (e)

In a 300 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 20 ml (20 mmol) of a hexane solution of diethylzinc (1.02 mol/l) and 100 ml of tetrahydrofuran were introduced, cooled to -78°C and stirred. To the mixture was dropped 4.4 ml (5.1 mmol) of a hexane solution of pentafluorophenol (1.17 mol/l). After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2 hours. Thereafter, 321 μ l (17.8 mmol) of H₂O was dropped thereto. After completion of dropping, the mixture was stirred for 14 hours at ambient temperature. Under reduced pressure,

a volatile component was removed by distillation, and thereafter, the resulting solid was dried under reduced pressure for 8 hours at 120°C. As a result, a solid product was obtained.

5 (2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 0.73 g of silica (Sylopol 948, manufactured by Davison Co., Ltd.; average particle diameter = 55 μ m; pore volume = 1.66 ml/g; specific surface area = 309 m²/g), which had been heat-treated at 300°C under a nitrogen stream and 40 ml of tetrahydrofuran were introduced and stirred. To the mixture, 1.01 g of the solid product obtained in the above Example 3 (1) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed using a glass filter, and the solid component was washed 4 times with each 20 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining a component (A).

(3) Polymerization

20 A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.012 MPa, 700 g of butane was fed therein, 50 g of 1-butene was fed therein, and then a temperature was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized.

As a result of gas chromatography, a gas composition in the system was found to be hydrogen = 0.477 mol%, and 1-butene = 2.29 mol%. To there, 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml was added. Successively,
5 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml was added thereto, and further 15.3 mg of the component (A) obtained in the above Example 3 (2) was added thereto. Polymerization was carried out for 60 minutes
10 at 70°C while feeding a mixed gas of ethylene and hydrogen (hydrogen concentration = 0.199 mol%) so as to keep the whole pressure constant. As a result, 46.5 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 9.3×10^7 g/mol-Zr/hour, and
15 polymerization activity per the solid catalyst component was found to be 3040 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 14.9, Mw = 169000, Mw/Mn = 10.2, MFR = 0.070, MFRR = 192 and SR = 1.05.

[Example 4]

20 (1) Production of contact product (e) In a 300 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 20 ml (20 mmol) of a hexane solution of diethylzinc (1.02 mol/l) and 100 ml of tetrahydrofuran were introduced, cooled to -78°C and stirred.
25 To the mixture was dropped 8.7 ml (10 mmol) of a hexane solution of pentafluorophenol (1.17 mol/l). After completion of

dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2 hours. Thereafter, 275 μ l (15.3 mmol) of H₂O was dropped thereto. After completion of dropping, the mixture was stirred all
5 night at ambient temperature. Under reduced pressure, a volatile component was removed by distillation, and thereafter, the resulting solid was dried under reduced pressure for 8 hours at 120°C. As a result, a solid product was obtained.

10 (2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 0.93 g of the same lot heat-treated silica as that used in Example 3 (2) and 40 ml of tetrahydrofuran were introduced and stirred. To
15 the mixture, 1.34 g of the solid product obtained in the above Example 4 (1) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed using a glass filter, and the solid component was washed 4 times with each 20 ml of tetrahydrofuran and once with 20 ml of hexane.
20 The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining a component (A). Elementary analysis revealed Zn = 2.6 mmol/g, and F = 3.9 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that
25 21.0 mg of the component (A) obtained in Example 4 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.494

mol% and 1-butene = 2.74 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.192 mol%. As a result, 70.2 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.4×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 3340 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 14.8, Mw = 143000, Mw/Mn = 5.28, MFR = 0.158, MFRR = 138 and SR = 1.06.

10 [Example 5]

(1) Production of contact product (e)

In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 10 ml (10 mmol) of a hexane solution of diethylzinc (1.02 mol/l) and 15 50 ml of tetrahydrofuran were introduced, cooled to -78°C and stirred. To the mixture was dropped 6.9 ml (10 mmol) of a hexane solution of pentafluorophenol (1.48 mol/l). After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 20 2 hours. Thereafter, 93.7 mg (5.20 mmol) of H_2O was dropped thereto. After completion of dropping, the mixture was stirred for 20 hours at ambient temperature. Under reduced pressure, a volatile component was removed by distillation, and thereafter, the resulting solid was dried under reduced 25 pressure for 8 hours at 120°C . As a result, 2.34 g of a solid product was obtained.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 0.38 g of the same lot heat-treated silica as that used in Example 3 (2) and 40 ml of tetrahydrofuran were introduced and stirred. To the mixture, 0.877 g of the solid product obtained in the above Example 5 (1) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed using a glass filter, and the solid component was washed 4 times with each 20 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining a component (A).

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 12.6 mg of the component (A) obtained in Example 5 (2) was used as a solid catalyst component, and a gas composition in the system found by gas chromatography was composed of hydrogen = 0.530 mol% and 1-butene = 2.38 mol%. As a result, 6.2 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.2×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 492 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 14.0, Mw = 112000, Mw/Mn = 6.93, MFR = 0.428, MFRR = 84.1 and SR = 1.30.

25 [Example 6]

(1) Production of contact product (e)

In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) and 40 ml of tetrahydrofuran were introduced, cooled to 3°C and stirred. To the mixture was dropped 8.6 ml (20 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.33 mol/l) at a rate of 17.9 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 125 minutes. Thereafter, 1.26 ml (69.9 mmol) of H₂O was dropped thereto at 20°C at a rate of 0.54 ml/hour. After completion of dropping, the mixture was stirred for 65 minutes at ambient temperature and thereafter allowed to stand for 11.3 hours. Under reduced pressure, a volatile component was removed by distillation, and thereafter, the resulting solid was dried under reduced pressure for 8 hours at 120°C. As a result, a solid product was obtained.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with nitrogen, 3.21 g of the solid product obtained in the above Example 6 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the mixture, 3.02 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the mixture was allowed to stand, thereby precipitating a solid component. At the time when an interface between a layer of the precipitated solid component

and an upper layer of a slurry portion appeared, the upper slurry portion was removed. Thereafter, filtration was carried out using a filter. For completing washing, 30 ml of tetrahydrofuran was added to the remainder, and the mixture was stirred, and thereafter allowed to stand, thereby precipitating a solid component, and then the upper slurry portion was removed similarly at the time when the interface appeared. Thereafter, filtration was carried out using a filter. The above washing operation was repeated 5 times. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 4.16 g of a component (A). Elementary analysis revealed Zn = 2.6 mmol/g, and F = 2.2 mmol/g. Accordingly, F/Zn = 0.85 (molar ratio). As a result of extended X-ray measurement, A/B = 0.212. As a result of EXAFS, peak intensity C = 13.1, peak intensity D = 5.74, and peak intensity ratio D/C = 0.44.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 16.3 mg of the component (A) obtained in the above Example 6 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.466 mol% and 1-butene = 2.82 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.187 mol%.

As a result, 20.5 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 4.1×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was

found to be 1260 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 12.7, Mw = 123000, Mw/Mn = 4.47, MFR = 0.167, MFRR = 108 and SR = 1.11.

[Example 7]

5 (1) Production of contact product (e)

In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) and 40 ml of tetrahydrofuran were introduced, cooled to 3°C and
10 stirred. To the mixture was dropped 17.2 ml (40 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.33 mol/l) at a rate of 17.9 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2.6 hours. Thereafter,
15 1.08 ml (60.0 mmol) of H₂O was dropped thereto at 20°C at a rate of 0.54 ml/hour. After completion of dropping, the mixture was stirred for 3.2 hours at ambient temperature and thereafter allowed to stand for 11 hours. Under reduced
20 pressure, a volatile component was removed by distillation, and thereafter, the resulting solid was dried under reduced pressure for 8 hours at 120°C, thereby obtaining a solid product.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under
25 reduced pressure and thereafter purged with nitrogen, 4.34 g of the solid product obtained in the above Example 7 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the

mixture, 2.87 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed, and thereafter the remaining solid component was washed 5 times
5 with each 30 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 3.21 g of a component (A). Elementary analysis revealed Zn = 1.5 mmol/g and F = 2.5 mmol/g. Accordingly, F/Zn = 1.7 (molar ratio). As a result of extended X-ray measurement, A/B = 0.034. As a result
10 of EXAFS, peak intensity C = 13.2, peak intensity D = 4.76, and peak intensity ratio D/C = 0.36.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 15.1 mg of the component (A) obtained in the above Example 7
15 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.441 mol% and 1-butene = 2.69 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.187 mol%.

As a result, 25.7 g of an olefin polymer having a good
20 particle property was obtained. Polymerization activity per zirconium atom was found to be 5.1×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 1700 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 14.6, Mw = 124000,
25 Mw/Mn = 4.32, MFR = 0.166, MFRR = 116 and SR = 1.12.

[Example 8]

(1) Production of contact product (e)

In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) and 40 ml of tetrahydrofuran were introduced, cooled to 3°C and stirred. To the mixture was dropped 34.3 ml (80 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.33 mol/l) at a rate of 17.9 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2.7 hours. Thereafter, 0.72 ml (40 mmol) of H₂O was dropped thereto at 20°C at a rate of 0.54 ml/hour. After completion of dropping, the mixture was stirred for 2.4 hours at ambient temperature and thereafter allowed to stand for 12 hours. Under reduced pressure, a volatile component was removed by distillation, and thereafter, the resulting solid was dried under reduced pressure for 8 hours at 120°C, thereby obtaining a solid product.

(2) Production of component (A) In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with nitrogen, 6.47 g of the solid product obtained in the above Example 8 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the mixture, 3.05 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the mixture was allowed to stand, thereby precipitating a solid component. At the time when an interface between a layer of the precipitated

solid component and an upper layer of a slurry portion appeared, the upper slurry portion was removed. Thereafter, filtration was carried out using a filter. For completing washing, 30 ml of tetrahydrofuran was added to the residue, and the mixture
5 was stirred, and thereafter allowed to stand, thereby precipitating a solid component, and then the upper slurry portion was removed similarly at the time when the interface appeared. Thereafter, filtration was carried out using a filter. The above washing operation was repeated 2 times. (Operated by
10 dip tube. Ogane) Thereafter, the liquid component was removed using a filter, and the resulting solid component was washed 2 times with each 30 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 3.35 g of a component (A). Elementary analysis revealed Zn =
15 1.2 mmol/g, and F = 2.4 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 15.5 mg of the component (A) obtained in the above Example 8 (2) was used as a solid catalyst component, a gas composition
20 in the system found by gas chromatography was composed of hydrogen = 0.475 mol% and 1-butene = 2.71 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.187 mol%.

As a result, 36.7 g of an olefin polymer having a good particle property was obtained. Polymerization activity per
25 zirconium atom was found to be 7.3×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was

found to be 2370 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 13.5, Mw = 122000, Mw/Mn = 4.06, MFR = 0.170, MFRR = 106 and SR = 1.15.

[Example 9]

5 (1) Production of contact product (e)

In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) and 40 ml of tetrahydrofuran were introduced, cooled to 3°C and
10 stirred. To the mixture was dropped 51.3 ml (120 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.34 mol/l) at a rate of 17.9 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2.4 hours. Thereafter,
15 0.36 ml (20 mmol) of H₂O was dropped thereto at 20°C at a rate of 0.54 ml/hour. After completion of dropping, the mixture was stirred for 3.4 hours at ambient temperature and thereafter allowed to stand for 13.5 hours. Under reduced pressure, a volatile component was removed by distillation,
20 and thereafter, the resulting solid was dried under reduced pressure for 8 hours at 120°C, thereby obtaining 26.2 g of a solid product.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under
25 reduced pressure and thereafter purged with argon, 9.28 g of the solid product obtained in the above Example 9 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the mixture,

2.97 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed, and thereafter the remaining solid component was washed 5 times with each 30 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining a component (A). Elementary analysis revealed Zn = 1.0 mmol/g and F = 2.8 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 15.7 mg of the component (A) obtained in the above Example 9 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.469 mol% and 1-butene = 2.48 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.192 mol%.

As a result, 12.9 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 2.6×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 822 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 7.91, Mw = 99200, Mw/Mn = 2.81, MFR = 0.582, MFRR = 48.4 and SR = 1.39.

[Example 10]

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 20.1 mg of the component (A) obtained in the above Example 1

(2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.480 mol% and 1-butene = 3.11 mol%, and the polymerization time was 15 minutes.

5 As a result, 20.0 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.6×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 4000 g/g-solid catalyst component/hour. Further,
10 the obtained olefin polymer showed, SCB = 14.7, Mw = 150000, Mw/Mn = 3.77, MFR = 0.06, MFRR = 233 and SR = 1.05.

[Example 11]

(1) Production of contact product (e)

15 In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) and 40 ml of tetrahydrofuran were introduced, cooled to 3°C and stirred. To the mixture was dropped 13.7 ml (31.9 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.33 mol/l)
20 at a rate of 17.9 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2.2 hours. Thereafter, 1.01 ml (56.1 mmol) of H₂O was dropped thereto at 20°C at a rate of 0.54 ml/hour. After completion of dropping, the
25 mixture was allowed to stand for 19.4 hours. Under reduced pressure, a volatile component was removed by distillation,

and thereafter, the resulting solid was dried under reduced pressure for 8 hours at 120°C, thereby obtaining a solid product.

(2) Production of component (A)

5 In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 4.05 g of the solid product obtained in the above Example 11 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the mixture, 2.79 g of the same lot heat-treated silica as that used in Example 10 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed, and thereafter the remaining solid component was washed 5 times with each 30 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining a component 15 (A). Elementary analysis revealed Zn = 1.5 mmol/g and F = 2.0 mmol/g.

[0269]

(3) Polymerization

 Polymerization of Example 3 (3) was repeated, except that 20 17.3 mg of the component (A) obtained in the above Example 11 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.479 mol% and 1-butene = 2.83 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.185 mol%.

25 As a result, 9.8 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium

atom was found to be 2.0×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 566 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 8.64, Mw = 90800, Mw/Mn = 2.83, MFR = 0.302, 5 MFRR = 119 and SR = 1.35.

[Example 12]

(1) Production of contact product (e)

In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80 10 mmol) of a hexane solution of diethylzinc (2 mol/l) and 40 ml of tetrahydrofuran were introduced, cooled to 3°C and stirred. To the mixture was dropped 6.9 ml (16.1 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.33 mol/l) at a rate of 17.9 ml/hour. After completion of dropping, the 15 mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2 hours. Thereafter, 1.44 ml (79.9 mmol) of H₂O was dropped thereto at 20°C at a rate of 0.54 ml/hour. After completion of dropping, the mixture was allowed to stand for 16.8 hours. Under reduced pressure, 20 a volatile component was removed by distillation, and thereafter, the resulting solid was dried under reduced pressure for 8 hours at 120°C, thereby obtaining a solid product.

(2) Production of component (A)

25 In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 3.02 g of the solid product obtained in the above Example 12 (1) and 30

ml of tetrahydrofuran were introduced and stirred. To the mixture, 2.92 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed, and then
5 the remaining solid component was washed 5 times with each 30 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining a component (A). Elementary analysis revealed Zn = 3.1 mmol/g and F = 2.2 mmol/g.

(3) Polymerization

10 Polymerization of Example 3 (3) was repeated, except that 12.9 mg of the component (A) obtained in the above Example 12 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.450 mol% and 1-butene = 2.72 mol%, and a hydrogen concentration
15 of the ethylene/hydrogen mixed gas was 0.185 mol%.

As a result, 12.6 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 2.5×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was
20 found to be 980 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 8.46, Mw = 106000, Mw/Mn = 3.45, MFR = 0.346, MFRR = 64.0 and SR = 1.28.

[Example 13]

(1) Production of contact product (e)

25 In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80

mmol) of a hexane solution of diethylzinc (2 mol/l) and 40 ml of tetrahydrofuran were introduced, cooled to 3°C and stirred. To the mixture, 1.01 ml (56.1 mmol) of H₂O was dropped thereto at a rate of 0.54 ml/hour. After completion of dropping, 5 the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2 hours, and further stirred at 40°C for 1.5 hours. After cooling to 25°C, 13.7 ml (31.9 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.33 mol/l) was dropped thereto at a rate 10 of 17.9 ml/hour. After completion of dropping, the mixture was stirred at 40°C for 2 hours and allowed to stand for 60.5 hours. Under reduced pressure, a volatile component was removed by distillation, and thereafter, the resulting solid was dried under reduced pressure for 8 hours at 120°C, thereby 15 obtaining a solid product.

(2) Production of component (A) In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 3.80 g of the solid product obtained in the above Example 13 (1) and 30 ml of tetrahydrofuran were introduced and 20 stirred. To the mixture, 3.07 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed, and thereafter the remaining solid component was washed 5 times with each 30 ml of tetrahydrofuran. 25 The residue was dried for 10 hours at 120°C under reduced pressure, thereby obtaining a component (A). Elementary analysis revealed Zn = 1.4 mmol/g and F = 1.9 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 16.6 mg of the component (A) obtained in the above Example 13 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.457 mol% and 1-butene = 2.61 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.193 mol%.

As a result, 8.2 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.6×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 494 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 6.37, Mw = 84700, Mw/Mn = 2.82, MFR = 1.35, MFRR = 38.0 and SR = 1.51.

15 [Example 14]

(1) Production of contact product (e)

In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) and 40 ml of tetrahydrofuran were introduced, cooled to 3°C and stirred. To the mixture, 1.15 ml (63.8 mmol) of H₂O was dropped thereto at a rate of 0.54 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2 hours, and further stirred at 40°C for 1.8 hours. After cooling to 24°C, 6.9 ml (16.1 mmol) of a tetrahydrofuran solution of

pentafluorophenol (2.33 mol/l) was dropped thereto at a rate of 17.9 ml/hour. After completion of dropping, the mixture was stirred at 40°C for 2 hours and allowed to stand for 15.3 hours. Under reduced pressure, a volatile component was removed by distillation, and thereafter, the resulting solid was dried under reduced pressure for 8 hours at 120°C, thereby obtaining a solid product.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with nitrogen, 2.92 g of the solid product obtained in the above Example 14 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the mixture, 2.84 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed, and thereafter the remaining solid component was washed 5 times with each 30 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 3.16 g of a component (A). Elementary analysis revealed Zn = 1.5 mmol/g and F = 1.2 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 16.7 mg of the component (A) obtained in the above Example 14 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.516 mol% and 1-butene = 2.49 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.198 mol%.

As a result, 5.5 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.1×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 329 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 4.35, Mw = 69600, Mw/Mn = 2.30, MFR = 2.00, MFRR = 30.6 and SR = 1.64.

[Example 15]

(1) Production of contact product (e)

10 In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) and 40 ml of tetrahydrofuran were introduced, cooled to 3°C and stirred. To the mixture, 1.26 ml (69.9 mmol) of H₂O was dropped
15 thereto at a rate of 0.54 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2.2 hours, and further stirred at 40°C for 1.5 hours. After cooling to 25°C, 8.6 ml (20.0 mmol) of a tetrahydrofuran solution of
20 pentafluorophenol (2.33 mol/l) was dropped thereto at a rate of 17.2 ml/hour. After completion of dropping, the mixture was stirred at 40°C for 2 hours and allowed to stand for 84 hours. Under reduced pressure, a volatile component was removed by distillation, and thereafter, the resulting solid
25 was dried under reduced pressure for 8 hours at 120°C, thereby obtaining a solid product.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 3.08 g of the solid product obtained in the above Example 15 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the mixture,
5 2.90 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed, and thereafter the remaining solid component was washed 5 times with each 30 ml of tetrahydrofuran. The residue was dried for 8.5
10 hours at 120°C under reduced pressure, thereby obtaining a component (A). Elementary analysis revealed Zn = 2.4 mmol/g and F = 3.1 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that
15 16.6 mg of the component (A) obtained in the above Example 15 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.560 mol% and 1-butene = 2.80 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.193 mol%.

20 As a result, 12.6 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 2.5×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 759 g/g-solid catalyst component/hour. Further, the
25 obtained olefin polymer showed, SCB = 12.1, Mw = 108000, Mw/Mn = 4.76, MFR = 0.415, MFRR = 74.4 and SR = 1.22.

[Example 16]

(1) Production of contact product (e)

In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) and 40 ml of tetrahydrofuran were introduced, cooled to 3°C and stirred. To the mixture, 1.44 ml (79.9 mmol) of H₂O was dropped thereto at a rate of 0.54 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2 hours, and further stirred at 40°C for 1.5 hours. After cooling to 26°C, 6.9 ml (16.1 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.33 mol/l) was dropped thereto at a rate of 17.9 ml/hour. After completion of dropping, the mixture was stirred at 40°C for 2 hours and allowed to stand for 10 hours. Under reduced pressure, a volatile component was removed by distillation, and thereafter, the resulting solid was dried under reduced pressure for 8 hours at 120°C, thereby obtaining a solid product.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with nitrogen, 3.04 g of the solid product obtained in the above Example 16 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the mixture, 3.07 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed,

and thereafter the remaining solid component was washed 5 times with each 30 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 3.15 g of a component (A). Elementary analysis revealed Zn = 3.5 mmol/g and F = 2.8 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 18.4 mg of the component (A) obtained in the above Example 16 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.473 mol% and 1-butene = 2.48 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.198 mol%.

As a result, 15.7 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 3.1×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 853 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 11.7, Mw = 108000, Mw/Mn = 4.69, MFR = 1.04, MFRR = 31.9 and SR = 1.23.

20 [Example 17]

(1) Production of contact product (e)

In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) and 53.6 ml of tetrahydrofuran were introduced, cooled to -53°C and stirred. To the mixture was dropped 17.2 ml (40.1 mmol) of

a tetrahydrofuran solution of pentafluorophenol (2.33 mol/l) at a rate of 41.3 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 2 hours, and further stirred at 40°C for 0.5 hour. After cooling to 17°C, 1.08 ml (60.0 mmol) of H₂O was dropped thereto at a rate of 0.81 ml/hour. After completion of dropping, the mixture was gradually heated to ambient temperature, stirred for 2 hours, further stirred at 50°C for 1 hour, and thereafter allowed to stand at ambient temperature for 12 hours. As a result, a content of the flask was separated into a yellow transparent liquid product and a brown gel product. Then, 72% by volume of the yellow transparent liquid product and the whole of the brown gel product were separated from each other and placed into different flasks purged with nitrogen, respectively. Both were treated to remove a volatile component by distillation and dried for 11 hours at 120°C. Thereafter, the solid product originated from the yellow transparent liquid product was mixed with 80 ml of tetrahydrofuran to be dissolved, and the resulting solution was added to the solid product originated from the brown gel product. The mixture was stirred at ambient temperature for 48 hours. Thereby, a content of the flask turned into an orange turbid product and a small amount of a powdery product having a skin color. Under reduced pressure, a volatile component was removed by distillation, and thereafter, the residue was dried under reduced pressure for 8 hours at 120°C, thereby obtaining a solid product.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with nitrogen, 4.25 g of the solid product obtained in the above Example 17 (1) and 5 30 ml of tetrahydrofuran were introduced and stirred. To the mixture, 2.89 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the mixture was allowed to stand, thereby precipitating a solid component. At the time when an 10 interface between a layer of the precipitated solid component and an upper layer of a slurry portion appeared, the upper slurry portion was removed. Thereafter, filtration was carried out using a filter. For completing washing, 30 ml of tetrahydrofuran was added to the residue, and the mixture was stirred, and thereafter 15 allowed to stand, thereby precipitating a solid component, and then the upper slurry portion was removed similarly at the time when the interface appeared. Thereafter, filtration was carried out using a filter. The above washing operation was repeated 5 times. The residue was dried for 8 hours at 120°C under reduced 20 pressure, thereby obtaining 3.75 g of a component (A). Elementary analysis revealed Zn = 2.1 mmol/g, and F = 2.7 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 17.6 mg of the component (A) obtained in the above Example 17 25 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen

= 0.516 mol% and 1-butene = 2.89 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.195 mol%.

As a result, 60.4 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.2×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 3430 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 16.0, Mw = 112000, Mw/Mn = 4.11, MFR = 0.463, MFRR = 69.7 and SR = 1.21.

10 [Example 18]

(1) Production of contact product (e)

In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) and 53.6 ml of tetrahydrofuran were introduced, cooled to -60°C and stirred. To the mixture was dropped 17.2 ml (40.2 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.34 mol/l) at a rate of 41.3 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 3.2 hours, and further stirred at 40°C for 0.5 hour. After cooling to 20°C , 1.08 ml (60.0 mmol) of H_2O was dropped thereto at a rate of 0.81 ml/hour. After completion of dropping, the mixture was gradually heated to ambient temperature, stirred for 2.3 hours, further stirred at 50°C for 1 hour, and thereafter allowed to stand all night at ambient temperature. As a result, a content of the flask

was separated into a yellow transparent liquid product and a brown gel product. Then, the whole amount of the yellow transparent liquid product and the whole amount of the brown gel product were separated from each other and placed into different flasks purged with nitrogen, respectively. Both were treated to remove a volatile component by distillation and dried for 8 hours at 120°C. Thereafter, the solid product originated from the yellow transparent liquid product was mixed with 80 ml of tetrahydrofuran to be dissolved, and the resulting solution was added to the solid product originated from the brown gel product. The mixture was stirred at ambient temperature for 67 hours. Thereby, a content of the flask turned into an orange turbid product and a small amount of a powdery product having a cream color. Under reduced pressure, a volatile component was removed by distillation, and thereafter, the residue was dried under reduced pressure for 8 hours at 120°C, thereby obtaining 13.8 g of a solid product.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 4.44 g of the solid product obtained in the above Example 18 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the mixture, 3.24 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Then, after removing the liquid component, the solid component was washed 5 times with each 30 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure,

thereby obtaining a component (A). Elementary analysis revealed Zn = 1.8 mmol/g and F = 3.1 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that
5 16.0 mg of the component (A) obtained in the above Example 18 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.533 mol% and 1-butene = 2.66 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.184 mol%.

10 As a result, 31.3 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 6.3×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 1960 g/g-solid catalyst component/hour. Further,
15 the obtained olefin polymer showed, SCB = 12.9, MFR = 0.296, MFRR = 76.0 and SR = 1.23.

[Example 19]

(1) Production of contact product (e)

In a 200 ml four necked flask, which had been dried under
20 reduced pressure and thereafter purged with argon, 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) and 53.6 ml of tetrahydrofuran were introduced, cooled to -65°C and stirred. To the mixture was dropped 17.2 ml (40.2 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.34 mol/l)
25 at a rate of 41.3 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to

ambient temperature and stirred for 3.2 hours, and further stirred at 40°C for 0.5 hour. After cooling to 20°C, 1.08 ml (60.0 mmol) of H₂O was dropped thereto at a rate of 0.81 ml/hour. After completion of dropping, the mixture was gradually heated to ambient temperature, stirred for 2.3 hours, further stirred at 50°C for 1 hour, and thereafter allowed to stand all night at ambient temperature. As a result, a content of the flask was separated into a yellow transparent liquid product and an orange oily product. Then, 28% by volume of the yellow transparent liquid product was taken out of the flask, and a mixture of the yellow transparent liquid product and the orange oily product was treated to remove a volatile component by distillation and dried under reduced pressure for 8 hours at 120°C. Thereafter, the solid product was mixed with 80 ml of tetrahydrofuran to be dissolved, and the resulting solution was stirred at ambient temperature for 67 hours. Thereby, a content of the flask turned into an orange transparent liquid product and a small amount of a cream-colored powdery product. Under reduced pressure, a volatile component was removed by distillation, and thereafter, the residue was dried under reduced pressure for 8 hours at 120°C. As a result, 10.5 g of a solid product was obtained.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 4.30 g of the solid product obtained in the above Example 19 (1) and 30

ml of tetrahydrofuran were introduced and stirred. To the mixture, 2.96 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the liquid component was removed, and then
5 the solid component was washed 5 times with each 30 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining a component (A). Elementary analysis revealed Zn = 2.4 mmol/g, and F = 3.3 mmol/g.

(3) Polymerization

10 Polymerization of Example 3 (3) was repeated, except that 11.4 mg of the component (A) obtained in the above Example 19 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.541 mol% and 1-butene = 2.73 mol%, and a hydrogen concentration
15 of the ethylene/hydrogen mixed gas was 0.184 mol%.

As a result, 61.0 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.2×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was
20 found to be 5350 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 15.9, MFR = 0.866, MFRR = 62.5 and SR = 1.25.

[Example 20]

(1) Production of contact product (e)

25 In a 200 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 40 ml (80

mmol) of a hexane solution of diethylzinc (2 mol/l) and 53.6 ml of tetrahydrofuran were introduced, cooled to -68°C and stirred. To the mixture was dropped 17.2 ml (40.2 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.34 mol/l) at a rate of 41.3 ml/hour. After completion of dropping, the mixture was gradually heated to raise the temperature to ambient temperature and stirred for 3.2 hours, and further stirred at 40°C for 0.5 hour. After cooling to 20°C , 1.08 ml (60.0 mmol) of H_2O was dropped thereto at a rate of 0.81 ml/hour. After completion of dropping, the mixture was gradually heated to ambient temperature, stirred for 2 hours, further stirred at 50°C for 1 hour, and thereafter allowed to stand all night at ambient temperature. As a result, a content of the flask was separated into a yellow transparent liquid product and an orange oily product. Then, 28% by volume of the yellow transparent liquid product was taken out of the flask, and a mixture of the yellow transparent liquid product and the orange oily product was treated to remove a volatile component by distillation, and dried under reduced pressure for 8 hours at 120°C . As a result, 10.6 g of a solid product was obtained.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with argon, 4.40 g of the solid product obtained in the above Example 20 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the mixture, 2.81 g of the same lot heat-treated silica as that used in Example 3 (2) was added. The mixture was heated to 40°C and stirred for

2 hours. Thereafter, the liquid component was removed, and then the solid component was washed 5 times with each 30 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining a component (A).

5 (3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 14.7 mg of the component (A) obtained in the above Example 20 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen
10 = 0.564 mol% and 1-butene = 2.57 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.193 mol%.

As a result, 53.6 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.1×10^8 g/mol-Zr/hour, and
15 polymerization activity per the solid catalyst component was found to be 3650 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 15.7, MFR = 0.396, MFRR = 86.6 and SR = 1.21.

[Example 21]

20 (1) Production of contact product (e)

In a 5 l four necked flask purged with nitrogen, 1.28 l of tetrahydrofuran and 1.28 l (2.55 mol) of a hexane solution of diethylzinc (1.99 mol/l) were introduced and cooled to -50°C. To the mixture was dropped a solution prepared by
25 dissolving 233.6 g (1.27 mol) of pentafluorophenol in 427 ml of tetrahydrofuran over 25 minutes. After completion of

dropping, the mixture was gradually heated to ambient temperature, and 500 ml of tetrahydrofuran was further added thereto. A solution prepared by dissolving 34.69 g (1.93 mol) of H₂O in 50 ml of tetrahydrofuran was dropped thereto at
5 ambient temperature. After completion of dropping, the mixture was stirred all night. Under reduced pressure, a volatile component was removed by distillation and the residue was dried under reduced pressure for 8 hours at 120°C and further for 8 hours at 130°C. As a result, 403 g of a solid
10 product was obtained.

(2) Production of component (A) (i)

In a 5 l four necked flask purged with nitrogen, 201.7 g of the solid product obtained in the above Example 21 (1) and 3 l of tetrahydrofuran were introduced and stirred. To the mixture,
15 136.1 g of the same lot heat-treated silica as in that used in Example 1 (2) was added. The mixture was heated to 40°C and stirred for 2 hours.

(3) Production of component (A) (ii)

In a 5 l four necked flask purged with nitrogen, 201.7
20 g of the solid product obtained in the above Example 21 (1) and 3.1 l of tetrahydrofuran were introduced and stirred. To the mixture, 140.4 g of the same lot heat-treated silica as that used in Example 1 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the solid product was allowed
25 to precipitate, and a slurry portion of the upper layer was removed. To the residue was added the whole amount of that obtained in the above Example 21 (2) in a slurry state. The mixture was stirred,

thereafter, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. For completing washing, 3 l of tetrahydrofuran was added thereto, the mixture was stirred, the solid product was allowed to precipitate, and
5 a slurry portion of the upper layer was removed. The above-mentioned washing operation was repeated 5 times. Further, 3 l of tetrahydrofuran was added thereto, the mixture was stirred, the solid product was allowed to precipitate, and a slurry portion of the upper layer was removed. After removing the liquid
10 component using a glass filter, the residue was dried for 8 hours at 120°C under reduced pressure. As a result, 372 g of a component (A) was obtained. Elementary analysis revealed, Zn = 1.8 mmol/g and F = 2.9 mmol/g.

(3) Polymerization

15 Polymerization of Example 3 (3) was repeated, except that 18.4 mg of the component (A) obtained in the above Example 21 (3) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.515 mol% and 1-butene = 2.01 mol%, and a hydrogen concentration
20 of the ethylene/hydrogen mixed gas was 0.195 mol%.

As a result, 54.5 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.1×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was
25 found to be 2960 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 13.9, Mw = 110000,

Mw/Mn = 4.35, MFR = 0.194, MFRR = 117 and SR = 1.18.

[Example 22]

(1) Polymerization

Polymerization of Example 3 (3) was repeated, except that
5 15.3 mg of the component (A) obtained in the above Example 21
(3) was used as a solid catalyst component, the amount of the
toluene solution of racemic ethylenebis(1-indenyl)zirconium
dichloride adjusted to a concentration of 2 μ mol/ml was changed
to 1.25 ml, a gas composition in the system found by gas
10 chromatography was composed of hydrogen = 0.513 mol% and 1-butene
= 3.03 mol%, and a hydrogen concentration of the
ethylene/hydrogen mixed gas was 0.197 mol%.

As a result, 72.9 g of an olefin polymer was obtained.
Polymerization activity per zirconium atom was found to be 2.9
15 $\times 10^7$ g/mol-Zr/hour, and polymerization activity per the solid
catalyst component was found to be 4770 g/g-solid catalyst
component/hour. Further, the obtained olefin polymer showed,
SCB = 13.4, Mw = 120000, Mw/Mn = 4.26, MFR = 0.326, MFRR = 93.8
and SR = 1.16.

20 [Example 23]

(1) Polymerization

Polymerization of Example 3 (3) was repeated, except that
14.6 mg of the component (A) obtained in the above Example 21
(3) was used as a solid catalyst component, the amount of the
25 toluene solution of racemic ethylenebis(1-indenyl)zirconium
dichloride adjusted to a concentration of 2 μ mol/ml was changed

to 0.05 ml, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.498 mol% and 1-butene = 3.02 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.196 mol%.

5 As a result, 8.2 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 8.2×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 560 g/g-solid catalyst component/hour. Further, the obtained olefin
10 polymer showed, SCB = 15.7, $M_w = 110000$, $M_w/M_n = 3.62$, MFR = 0.649, MFRR = 58.7 and SR = 1.26.

[Example 24]

(1) Polymerization

Polymerization of Example 3 (3) was repeated, except that
15 15.0 mg of the component (A) obtained in the above Example 21 (3) was used as a solid catalyst component, the amount of the heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml was changed to 4.5 ml, a gas composition in the system found by gas chromatography was composed of hydrogen
20 = 0.509 mol% and 1-butene = 2.84 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.196 mol%.

As a result, 31.7 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 6.3×10^7 g/mol-Zr/hour, and
25 polymerization activity per the solid catalyst component was found to be 2110 g/g-solid catalyst component/hour. Further,

the obtained olefin polymer showed, SCB = 15.0, Mw = 124000, Mw/Mn = 3.79, MFR = 0.183, MFRR = 125 and SR = 1.13.

[Example 25]

(1) Polymerization

5 Polymerization of Example 3 (3) was repeated, except that 16.0 mg of the component (A) obtained in the above Example 21 (3) was used as a solid catalyst component, the amount of 1-butene was changed to 30 g, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.500 mol% and
10 1-butene = 1.60 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.25 mol%.

As a result, 40.0 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 8.0×10^7 g/mol-Zr/hour, and
15 polymerization activity per the solid catalyst component was found to be 2500 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 21.0, Mw = 120000, Mw/Mn = 3.61, MFR = 0.150, MFRR = 134 and SR = 1.20.

[Example 26]

20 (1) Polymerization

Polymerization of Example 3 (3) was repeated, except that 16.1 mg of the component (A) obtained in the above Example 1 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen
25 = 0.468 mol% and 1-butene = 2.62 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.189 mol%.

As a result, 83.4 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.7×10^6 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 5180 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 15.6, MFR = 0.917, MFRR = 50.6 and SR = 1.32.

[Example 27]

(1) Production of contact product (e)

10 In a 5 l four necked flask purged with nitrogen, 2 l of tetrahydrofuran and 1.35 l (2.7 mol) of a hexane solution of diethylzinc (2 mol/l) were introduced and cooled to -50°C . To the mixture was dropped a solution prepared by dissolving 248.7 g (1.35 mol) of pentafluorophenol in 390 ml of
15 tetrahydrofuran over 25 minutes. After completion of dropping, the mixture was gradually heated to ambient temperature and stirred for 2.9 hours, and further stirred at 40°C for 1 hour. After cooling to 21°C using an ice bath, 37.3 g (2.1 mol) of H_2O was dropped thereto over 1.4 hours. After completion
20 of dropping, the mixture was stirred at ambient temperature for 2 hours, and further stirred at 50°C for 1 hour. As a result, a content was separated into a yellow transparent liquid product and a yellow gel product. Thereafter, the mixture was cooled to ambient temperature and allowed to stand
25 all night. Then, 72% by weight of the yellow transparent liquid product and the whole amount of the yellow gel product were separated from each other and placed into different flasks

purged with nitrogen, respectively. Both were treated to remove a volatile component by distillation and dried under reduced pressure for 8 hours at 120°C. Thereafter, both the solid product originated from the yellow transparent liquid product and the solid product originated from the yellow gel product were introduced in a 5 l four necked flask purged with nitrogen, and 3 l of tetrahydrofuran was added thereto. The mixture was allowed to stand for 3 days. Under reduced pressure, a volatile component was removed by distillation, and the residue was dried under reduced pressure for 8 hours at 120°C. As a result, 376 g of a solid product was obtained.

(2) Production of component (A)

In a 5 l four necked flask purged with nitrogen, 376 g of the solid product obtained in the above Example 27 (1) and 3 l of tetrahydrofuran were introduced and stirred. To the mixture, 272.3 g of the same lot heat-treated silica as that used in Example 1 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. For the completion of washing, 3 l of tetrahydrofuran was added thereto, the mixture was stirred, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. The above-mentioned washing operation was repeated 5 times. After removing the liquid component using a glass filter, the residue was dried for 9 hours at 120°C under reduced pressure, thereby obtaining 419 g of a component (A). Elementary analysis revealed Zn = 2.4 mmol/g, and F = 3.6 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 16.1 mg of the component (A) obtained in the above Example 27 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.470 mol% and 1-butene = 2.72 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.188 mol%.

As a result, 78.5 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.6×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 4880 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 15.8, MFR = 0.783, MFRR = 66.1 and SR = 1.44.

15 [Example 28]

(1) Production of contact product (e)

In a 5 l four necked flask purged with nitrogen, 2 l of tetrahydrofuran and 1.35 l (2.7 mol) of a hexane solution of diethylzinc (2 mol/l) were introduced and cooled to -50°C. To the mixture was dropped a solution prepared by dissolving 251.5 g (1.37 mol) of pentafluorophenol in 390 ml of tetrahydrofuran over 25 minutes. After completion of dropping, the mixture was gradually heated to ambient temperature and stirred for 3 hours, and thereafter further stirred at 45°C for 1 hour. After cooling to 20°C using an ice bath, 37.72 g (2.09 mol) of H₂O was dropped thereto over 1.4 hours. As

a result, a content was separated into a yellow transparent liquid product and a yellow gel product. After completion of dropping, the mixture was stirred for 2 hours, and further stirred at 40°C for 1 hour. The mixture was allowed to stand
5 all night at ambient temperature. Then, 72% by weight of the yellow transparent liquid product and the whole amount of the yellow gel product were separated from each other and placed into different flasks purged with nitrogen, respectively. Both were treated to remove a volatile component
10 by distillation and dried under reduced pressure for 8 hours at 120°C. Thereafter, the solid product originated from the yellow transparent liquid product was dissolved in 3 l of tetrahydrofuran, and the resulting solution was transferred into a 5 l flask, in which the solid product originated from
15 the yellow gel product had been placed. The mixture was allowed to stand for 69 hours at ambient temperature, and thereafter dried under reduced pressure for 8 hours at 120°C. As a result, 374 g of a solid product was obtained.

(2) Production of component (A)

20 In a 5 l four necked flask purged with nitrogen, 374 g of the solid product obtained in the above Example 28 (1) and 3 l of tetrahydrofuran were introduced and stirred. To the mixture, 282 g of the same lot heat-treated silica as that used in Example
1 (2) was added. The mixture was heated to 40°C and stirred for
25 2 hours. Thereafter, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. For the completion of washing, 3 l of tetrahydrofuran was added

thereto, the mixture was stirred, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. The above-mentioned washing operation was repeated 5 times. After removing the liquid component using a glass filter, the residue
5 was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 452 g of a component (A). Elementary analysis revealed Zn = 2.8 mmol/g, and F = 3.6 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that
10 14.0 mg of the component (A) obtained in the above Example 28 (2) was used as a solid catalyst component, and a gas composition in the system found by gas chromatography was composed of hydrogen = 0.571 mol% and 1-butene = 2.69 mol%.

As a result, 75.8 g of an olefin polymer having a good
15 particle property was obtained. Polymerization activity per zirconium atom was found to be 1.5×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 5410 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 16.1, MFR = 0.965,
20 MFRR = 66.2 and SR = 1.38.

[Example 29]

(1)

In a 5 l four necked flask purged with nitrogen, 2 l of tetrahydrofuran and 1.35 l (2.7 mol) of a hexane solution
25 of diethylzinc (2 mol/l) were introduced and cooled to -58°C. To the mixture was dropped a solution prepared by dissolving

251.5 g (1.37 mol) of pentafluorophenol in 390 ml of tetrahydrofuran over 25 minutes. After completion of dropping, the mixture was gradually heated to ambient temperature and stirred for 3 hours, and thereafter further stirred at 38°C for 1 hour. After cooling to 18°C using an ice bath, 37.74 g (2.09 mol) of H₂O was dropped thereto over 1.4 hours. As a result, a content in the flask turned to a yellowish white slurry. After completion of dropping, the mixture was stirred for 2 hours, and further stirred at 40°C for 1 hour. The mixture was allowed to stand all night at ambient temperature. Then, 72% by weight of the liquid product and the whole amount of the solid product were separated from each other and placed into different flasks purged with nitrogen, respectively. Both were treated to remove a volatile component by distillation and dried under reduced pressure for 8 hours at 120°C. Thereafter, the solid product originated from the liquid product was dissolved in 3 l of tetrahydrofuran, and the resulting solution was transferred in a 5 l flask, in which the solid product originated from the solid product had been placed. The mixture was allowed to stand for 69 hours at ambient temperature, and dried under reduced pressure for 8 hours at 120°C. As a result, 387 g of a solid product was obtained.

(2) Production of component (A)

In a 5 l four necked flask purged with nitrogen, 387 g of the solid product obtained in the above Example 29 (1) and 3 l of tetrahydrofuran were introduced and stirred. To the mixture,

281 g of the same lot heat-treated silica as that used in Example 1 (2) was added. The mixture was heated to 40°C and stirred for 2 hours. Thereafter, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed.

5 For the completion of washing, 3 l of tetrahydrofuran was added thereto, the mixture was stirred, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. The above-mentioned washing operation was repeated 5 times. After removing the liquid component using a glass filter, the residue
10 was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 403 g of a component (A). Elementary analysis revealed Zn = 2.3 mmol/g, and F = 3.2 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that
15 14.3 mg of the component (A) obtained in the above Example 29 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.523 mol% and 1-butene = 2.78 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.183 mol%.

20 As a result, 51.2 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.0×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 3580 g/g-solid catalyst component/hour. Further,
25 the obtained olefin polymer showed, SCB = 15.8, MFR = 0.600, MFRR = 92.0 and SR = 1.26.

[Example 30]

(1) Production of contact product (e)

In a 1 l four necked flask purged with nitrogen, 238 ml of tetrahydrofuran and 180.5 ml (361 mmol) of a hexane solution of diethylzinc (2 mol/l) were introduced and cooled to -65°C. To the mixture was dropped 81.7 ml (181 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.21 mol/l) over 25 minutes. After completion of dropping, the mixture was gradually heated to ambient temperature and stirred for 3.2 hours, and thereafter further stirred at 40°C for 1 hour. After cooling to 20°C using an ice bath, 4.9 ml (272 mmol) of H₂O was dropped thereto over 1.3 hours. As a result, a content in the flask turned to a yellowish white slurry. After completion of dropping, the mixture was stirred for 2 hours, and further stirred at 50°C for 1 hour. The mixture was allowed to stand all night at ambient temperature. Thereafter, the liquid product and the solid product were separated from each other and placed into different flasks purged with nitrogen, respectively. Both were treated to remove a volatile component by distillation and dried under reduced pressure for 8 hours at 120°C. As a result, 48.2 g of a solid product originated from the liquid product and 16.5 g of a solid product originated from the solid product were obtained.

(2) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 4.41 g of the solid product originated from the solid product obtained in the above Example 30 (1), 2.90 g of the same lot heat-treated

silica as that used in Example 1 (2) and 30 ml of tetrahydrofuran were introduced and stirred for 2 hours at 40°C. The solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. For the completion of washing, 30
5 ml of tetrahydrofuran was added thereto, the mixture was stirred, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. The above-mentioned washing operation was repeated 5 times. After removing the liquid component using a glass filter, the residue was dried for 8 hours
10 at 120°C under reduced pressure, thereby obtaining 5.19 g of a component (A). Elementary analysis revealed Zn = 1.5 mmol/g, and F = 4.4 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that
15 13.5 mg of the component (A) obtained in the above Example 30 (2) was used as a solid catalyst component, a gas composition in the system found by gas chromatography was composed of hydrogen = 0.493 mol% and 1-butene = 2.75 mol%, and a hydrogen concentration of the ethylene/hydrogen mixed gas was 0.188 mol%.

20 As a result, 106.2 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 2.1×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 7870 g/g-solid catalyst component/hour. Further,
25 the obtained olefin polymer showed, SCB = 14.3, MFR = 0.550, MFRR = 64.0 and SR = 1.20.

[Example 31]

(1) Production of contact product (e)

In a 300 ml four necked flask purged with nitrogen, 106 ml of tetrahydrofuran and 40 ml (80 mmol) of a hexane solution
5 of diethylzinc (2 mol/l) were introduced and cooled to 4°C. To the mixture was dropped 14.5 ml (32 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.21 mol/l) over 50 minutes. After completion of dropping, the mixture was further stirred for 1 hour at 5°C. After cooling to 20°C using an ice bath,
10 7.3 ml (72.3 mmol) of a tetrahydrofuran solution of H₂O (9.9 mol/l) was dropped thereto over 1 hour. After completion of dropping, the mixture was stirred for 2 hours, and allowed to stand all night at ambient temperature. Thereafter, the mixture was stirred under reflux condition (63°C) for 8 hours.
15 As a result, 146 ml of a yellow transparent solution was obtained.

(2) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 2.93 g of the same lot heat-treated silica as that used in Example
20 1 (2) and 47 ml of the yellow transparent liquid obtained in the above Example 31 (1) were introduced and stirred for 2 hours at 40°C. After removing the liquid component using a glass filter, the residue was washed 5 times with each 30 ml of tetrahydrofuran, and dried for 8 hours at 120°C under reduced pressure, thereby
25 obtaining 3.89 g of a component (A). Elementary analysis revealed Zn = 2.0 mmol/g, and F = 2.0 mmol/g.

(3) Polymerization

Polymerization of Example 3 (3) was repeated, except that 14.6 mg of the component (A) obtained in the above Example 31 (2) was used as a solid catalyst component, and a gas composition in the system found by gas chromatography was composed of hydrogen
5 = 0.561 mol% and 1-butene = 2.70 mol%.

As a result, 65.1 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.3×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was
10 found to be 4460 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 13.8, MFR = 0.14, MFRR = 138 and SR = 1.10.

[Example 32]

(1) Production of contact product (e)

15 In a 200 ml four necked flask purged with nitrogen, 40 ml of tetrahydrofuran and 40 ml (80 mmol) of a hexane solution of diethylzinc (2.0 mol/l) were introduced and cooled to -5°C. To the mixture was dropped 34.3 ml (80 mmol) of a tetrahydrofuran solution of 3,5-difluorophenol (2.33 mol/l) over 120 minutes.
20 After completion of dropping, the mixture was stirred for 90 minutes at 5°C and further stirred for 70 minutes at 40°C. After cooling to 20°C using an ice bath, 0.72 ml (40 mmol) of H₂O was dropped thereto over 80 minutes. After completion of dropping, the mixture was stirred for 60 minutes at 20°C,
25 and further stirred for 120 minutes at 40°C. As a result, a white yellow slurry product was obtained. It was allowed

to stand all night at ambient temperature. Thereafter, a volatile component was removed by distillation and the residue was dried for 8 hours at 120°C under reduced pressure. As a result, 14.7 g of a yellow solid product was obtained.

5 (2) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 5.33 g of the yellow solid product obtained in the above Example 32 (1), 30 ml of tetrahydrofuran, and 3.03 g of the same lot heat-treated silica as that used in Example 1 (2) were introduced and stirred for 2 hours at 40°C. After removing the liquid component using a glass filter, the residue was washed 5 times with each 30 ml of tetrahydrofuran, and thereafter dried for 8 hours at 120°C under reduced pressure, thereby obtaining 3.00 g of a component (A). Elementary analysis revealed Zn = 1.1 mmol/g, and F = 1.3 mmol/g.

(3) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.012 MPa, 700 g of butane and 50 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.50 mol% and 1-butene = 2.62 mol%. To the resultant was added 0.9 ml of a heptane solution

of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 19.0 mg of the component (A) obtained in the above Example 32 (2) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.25 mol%) so as to keep the whole pressure constant. As a result, 7 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.4×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 370 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, $M_w = 8.44 \times 10^4$, $M_w/M_n = 2.16$, and $[\eta] = 1.35$.

[Example 33]

(1) Production of contact product (e)

In a 200 ml four necked flask purged with nitrogen, 40 ml of tetrahydrofuran and 40 ml (80 mmol) of a hexane solution of diethylzinc (2.0 mol/l) were introduced and cooled to -5°C. To the mixture was dropped 34.3 ml (80 mmol) of a tetrahydrofuran solution of perfluoro-tert-butanol (2.33 mol/l) over 120 minutes. After completion of dropping, the mixture was stirred for 90 minutes at 5°C and further stirred for 70 minutes at 40°C. After cooling to 20°C using an ice bath, 0.72 ml (40 mmol) of H₂O was dropped thereto over 80 minutes. After completion of dropping, the mixture was stirred for 60 minutes at 20°C,

and further stirred for 120 minutes at 40°C. As a result, a white yellow solution was obtained. It was allowed to stand all night at ambient temperature. Thereafter, a volatile component was removed by distillation and the residue was
5 dried for 8 hours at 120°C under reduced pressure. As a result, 26.35 g of a yellow solid product was obtained.

(2) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 8.29 g of the yellow solid product obtained in the above Example 33
10 (1), 30 ml of tetrahydrofuran, and 3.09 g of the same lot heat-treated silica as that used in Example 1 (2) were introduced and stirred for 2 hours at 40°C. After removing the liquid component using a glass filter, the residue was washed 5 times with each 30 ml of tetrahydrofuran, and thereafter dried for
15 8 hours at 120°C under reduced pressure, thereby obtaining 4.39 g of a component (A). Elementary analysis revealed Zn = 2.8 mmol/g, and F = 3.4 mmol/g.

(3) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had
20 been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.012 MPa, 700 g of butane and 50 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain
25 a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition

in the system was found to be hydrogen = 0.51 mol% and 1-butene = 2.47 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.25 ml of a toluene solution of racemic
5 ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 20.0 mg of the component (A) obtained in the above Example 33 (2) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed
10 gas (hydrogen = 0.25 mol%) so as to keep the whole pressure constant. As a result, 13 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 2.6×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 650
15 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 14.3, MFR = 0.96, MFRR = 44.5, SR = 1.40, $M_w = 8.98 \times 10^4$, $M_w/M_n = 2.65$, and $[\eta] = 1.33$.

[Example 34]

(1) Production of contact product (e)

20 In a 200 ml four necked flask purged with nitrogen, 40 ml of tetrahydrofuran and 40 ml (80 mmol) of a hexane solution of diethylzinc (2.0 mol/l) were introduced and cooled to -5°C. To the mixture was dropped 34.3 ml (80 mmol) of a tetrahydrofuran solution of 1,1,1,3,3,3-hexafluoro-2-propanol (2.33 mol/l)
25 over 120 minutes. After completion of dropping, the mixture was stirred for 90 minutes at 5°C and further stirred for

70 minutes at 40°C. After cooling to 20°C using an ice bath, 0.72 ml (40 mmol) of H₂O was dropped thereto over 80 minutes. After completion of dropping, the mixture was stirred for 60 minutes at 20°C, and further stirred for 120 minutes at 40°C. As a result, a white yellow solution was obtained. It was allowed to stand all night at ambient temperature. Thereafter, a volatile component was removed by distillation and the residue was dried for 8 hours at 120°C under reduced pressure. As a result, 15.47 g of a yellow solid product was obtained.

(2) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 6.53 g of the yellow solid product obtained in the above Example 34 (1), 30 ml of tetrahydrofuran, and 3.11 g of the same lot heat-treated silica as that used in Example 1 (2) were introduced and stirred for 2 hours at 40°C. After removing the liquid component using a glass filter, the residue was washed 5 times with each 30 ml of tetrahydrofuran, and thereafter dried for 8 hours at 120°C under reduced pressure, thereby obtaining 4.44 g of a component (A). Elementary analysis revealed Zn = 2.9 mmol/g, and F = 3.0 mmol/g.

(3) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.012 MPa, 700 g of butane and 50 g of 1-butene were fed therein, and then a temperature of the autoclave was

raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.51 mol% and 1-butene = 2.35 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μmol/ml, and then 16.0 mg of the component (A) obtained in the above Example 34 (2) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.25 mol%) so as to keep the whole pressure constant. As a result, 8 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.6×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 500 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, MFR = 0.77, MFRR = 48.4, SR = 1.38, Mw = 8.28×10^4 , Mw/Mn = 2.46, and $[\eta] = 1.32$.

[Example 35]

(1) Production of contact product (e)

In a 2 l four necked flask purged with nitrogen, 535 ml of tetrahydrofuran and 361 ml (722 mol) of a hexane solution of diethylzinc (2 mol/l) were introduced and cooled to -53°C. To the mixture was dropped a solution prepared by dissolving

68.13 g (370 mmol) of pentafluorophenol in 104 ml of tetrahydrofuran over 20 minutes. After completion of dropping, the mixture was gradually heated to ambient temperature and stirred for 1 hour, and further stirred at 47°C for 1 hour. After
5 cooling to 20°C using an ice bath, 10.5 g (583 mmol) of H₂O was dropped thereto over 100 minutes. As a result, a content was separated into a yellow transparent liquid product and a white crystal. The mixture was stirred at ambient temperature for 2 hours, and further stirred at 50°C for 1 hour. Thereby, the content
10 turned into a yellow transparent liquid product and a yellow oil product. Then, 71% by volume of the yellow transparent liquid product and the whole amount of the yellow gel product were separated from each other and placed into different flasks purged with nitrogen, respectively. Both were treated to remove a
15 volatile component by distillation and dried under reduced pressure for 8 hours at 120°C. Thereafter, the solid product originated from the yellow transparent liquid product was dissolved in 803 ml of tetrahydrofuran, and the resulting solution was combined with the solid product originated from the yellow
20 oil product. The mixture was allowed to stand for 59.5 hours. Thereby, the content turned into a yellow transparent liquid product and a small amount of a white powder. After removing a volatile component by distillation, the residue was dried under reduced pressure for 8 hours at 120°C. As a result, 124 g of
25 a solid product was obtained. Elementary analysis revealed Zn = 6.0 mmol/g, and F = 13 mmol/g.

(2) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with nitrogen, 4.30 g of the solid product obtained in the above Example 35 (1), 30 ml of tetrahydrofuran, and 2.95 g of silica (Sylopol 948, manufactured by Davison Co., Ltd.; average particle diameter = 52.0 μm ; pore volume = 1.58 ml/g; specific surface area = 302 m^2/g), which had been heat-treated at 300°C under a nitrogen stream, were introduced and stirred for 2 hours at 40°C. After removing the liquid component, the residue was washed 5 times with each 30 ml of tetrahydrofuran, and thereafter dried for 8 hours at 120°C under reduced pressure, thereby obtaining 4.35 g of a component (A). Elementary analysis revealed Zn = 2.6 mmol/g, and F = 3.5 mmol/g.

(3) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.012 MPa, 700 g of butane and 50 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.481 mol% and 1-butene = 2.92 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml.

Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 16.3 mg of the component (A) obtained in the above Example 35 (2) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.181 mol%) so as to keep the whole pressure constant. As a result, 54.1 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.1×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 3320 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 16.0, Mw = 108000, Mw/Mn = 4.40, MFR = 0.846, MFRR = 58.9, and SR = 1.31.

[Example 36]

(1) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with nitrogen, 4.41 g of the solid product obtained in the above Example 35 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the mixture, 3.02 g of silica (Sylopol 948, manufactured by Davison Co., Ltd.; average particle diameter = 45.6 μ m; pore volume = 1.63 ml/g; specific surface area = 527 m²/g), which had been heat-treated at 300°C under a nitrogen stream, was added and the resulting mixture was stirred for 2 hours at 40°C. After removing the liquid component, the solid component was washed

5 times with each 30 ml of tetrahydrofuran, and thereafter dried for 8 hours at 120°C under reduced pressure, thereby obtaining 3.98 g of a component (A). Elementary analysis revealed Zn = 2.6 mmol/g, and F = 4.2 mmol/g.

5 (2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.012 MPa, 700 g of butane and 50 g of 1-butene
10 were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.452 mol% and 1-butene
15 = 2.82 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 13.3 mg of the component
20 (A) obtained in the above Example 36 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.194 mol%) so as to keep the whole pressure constant. As a result, 46.5 g of an olefin polymer having a good
25 particle property was obtained. Polymerization activity per zirconium atom was found to be 9.3×10^7 g/mol-Zr/hour, and

polymerization activity per the solid catalyst component was found to be 3500 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 14.1, Mw = 118000, Mw/Mn = 7.44, MFR = 0.180, MFRR = 144, and SR = 1.13.

5 [Example 37]

(1) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with nitrogen, 4.45 g of the solid product obtained in the above Example 35 (1) and
10 30 ml of tetrahydrofuran were introduced and stirred. To the mixture was added 3.09 g of an organic polymer carrier, an acrylonitrile-amine cross-linking product (KA-892, manufactured by Sumitomo Chemical Co., Ltd.; specific surface area = 34.4 m²/g; amine-functional group content = 7 mmol/g).
15 The resulting mixture was heated to 40°C, and stirred for 2 hours. After removing the liquid component, the solid component was washed 5 times with each 30 ml of tetrahydrofuran, and thereafter dried for 8 hours at 120°C under reduced pressure, thereby obtaining a component (A). Elementary analysis revealed Zn =
20 0.58 mmol/g, and F = 1.1 mmol/g.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a
25 partial pressure of 0.012 MPa, 700 g of butane and 50 g of 1-butene were fed therein, and then a temperature of the autoclave was

raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.528 mol% and 1-butene = 3.00 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 13.8 mg of the component (A) obtained in the above Example 37 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.184 mol%) so as to keep the whole pressure constant. As a result, 11.4 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 2.3×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 826 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 5.39, Mw = 80900, Mw/Mn = 2.40, MFR = 1.41, MFRR = 34.7, and SR = 1.47.

[Example 38]

(1) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with nitrogen, 4.34 g of the solid product obtained in the above Example 35 (1) and 30 ml of tetrahydrofuran were introduced and stirred. To the

mixture was added 3.07 g of an organic polymer carrier, a 4-vinylpyridine polymer (4VP, manufactured by Kojie Chemical Co., Ltd.; specific surface area = 35.8 m²/g; pyridine-functional group content = 3.6 mmol/g). The resulting mixture was heated
5 to 40°C, and stirred for 2 hours. After removing the liquid component, the solid component was washed 5 times with each 30 ml of tetrahydrofuran, and thereafter dried for 8 hours at 120°C under reduced pressure, thereby obtaining a component (A). Elementary analysis revealed Zn = 1.4 mmol/g, and F = 3.3 mmol/g.

10 (2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.012 MPa, 700 g of butane and 50 g of 1-butene
15 were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.577 mol% and 1-butene
20 = 3.00 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μmol/ml, and then 11.4 mg of the component
25 (A) obtained in the above Example 38 (1) as a solid catalyst component were added thereto. Polymerization was carried out

for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.184 mol%) so as to keep the whole pressure constant. As a result, 13.8 g of an olefin polymer having a good particle property was obtained. Polymerization activity per
5 zirconium atom was found to be 2.8×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 1210 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 6.17, Mw = 87200, Mw/Mn = 2.44, MFR = 0.679, MFRR = 54.8, and SR = 1.48.

10 [Example 39]

(1) Production of component (A)

In a 50 ml four necked flask, which had been dried under reduced pressure and thereafter purged with nitrogen, 4.37 g of the solid product obtained in the above Example 35 (1) and
15 30 ml of tetrahydrofuran were introduced and stirred. To the mixture was added 3.07 g of an organic polymer carrier, a styrene-divinylbenzene copolymer (CS-1, manufactured by Sumitomo Chemical Co., Ltd.; divinylbenzene unit content = 39.5% by weight; average particle diameter = 33.0 μ m; pore volume
20 = 1.05 ml/g). The resulting mixture was heated to 40°C, and stirred for 2 hours. After removing tetrahydrofuran by distillation under reduced pressure, 40 ml of tetrahydrofuran was again added to the residue, and the mixture was heated to 40°C. Thereafter, the liquid component was removed using a glass filter, and the
25 resulting solid component was washed 5 times with each 30 ml of tetrahydrofuran. The residue was dried for 8 hours at 120°C

under reduced pressure, thereby obtaining a component (A). Elementary analysis revealed $\text{Zn} = 0.077 \text{ mmol/g}$, and $\text{F} = 1.3 \text{ mmol/g}$.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had
5 been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.012 MPa, 700 g of butane and 50 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C . Thereafter, ethylene was fed therein to obtain
10 a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.54 mol% and 1-butene = 2.78 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml.
15 Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of $2 \mu\text{mol/ml}$, and then 15.3 mg of the component (A) obtained in the above Example 39 (1) as a solid catalyst component were added thereto. Polymerization was carried out
20 for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.184 mol%) so as to keep the whole pressure constant. As a result, 8.3 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be $1.7 \times 10^7 \text{ g/mol-Zr/hour}$, and
25 polymerization activity per the solid catalyst component was found to be 542 g/g-solid catalyst component/hour. Further, the

obtained olefin polymer showed, SCB = 7.20, MFR = 2.54, MFRR = 28.7, SR = 1.61.

[Example 40]

(1) Production of contact product (e)

5 In a 300 ml four necked flask purged with nitrogen, 100 ml of toluene and 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) were introduced and cooled to -5°C. To the mixture was dropped 11.6 ml (32.1 mmol) of a toluene solution of pentafluorophenol (2.77 mol/l) over 40 minutes. After
10 completion of dropping, the mixture was gradually heated to 20°C. Thereafter, 1.58 ml (87.7 mmol) of H₂O was dropped thereto over 60 minutes at 20°C. After completion of dropping, the mixture was stirred for 6 hours under reflux condition (91°C). As a result, 140 ml of a slurry of a contact product (e) was obtained. According
15 to calculations based on the amounts of reagents used, a slurry concentration was found to be 0.57 mmol (in terms of Zn atom)/ml (slurry).

(2) Production of component (A)

 In a 50 ml four necked flask purged with nitrogen, 45 ml
20 (25.7 mmol-Zn) of the slurry of the contact product (e) obtained in the above Example 40 (1) was introduced. After removing the solvent using a glass filter, 40 ml of methyl-tert-butyl ether and 2.99 g of silica (Sylopol 948, manufactured by Davison Co., Ltd.; average particle diameter = 59 μm; pore volume = 1.61 ml/g;
25 specific surface area = 306 m²/g), which had been heat-treated at 300°C under a nitrogen stream, were introduced, heated to

40°C and stirred for 2 hours. After removing the liquid component using a glass filter, the residue was washed 4 times with each 30 ml of toluene and washed once with 30 ml of hexane, and thereafter dried for 8 hours at 120°C under reduced pressure, thereby obtaining 5.98 g of a component (A). Elementary analysis revealed Zn = 3.2 mmol/g, and F = 5.1 mmol/g.

(3) Polymerization A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.017 MPa, 695 g of butane and 55 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.759 mol% and 1-butene = 2.62 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μmol/ml, and then 8.1 mg of the component (A) obtained in the above Example 40 (2) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70 °C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.203 mol%) so as to keep the whole pressure constant. As a result, 28.6 g of an olefin polymer having a good particle property was obtained.

Polymerization activity per zirconium atom was found to be 1.9×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 3530 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed,
5 SCB = 14.3, MFR = 1.90, MFRR = 43.4 and SR = 1.32.

[Example 41]

(1) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 45 ml (25.7 mmol-Zn) of the slurry of the contact product (e) obtained
10 in the above Example 40 (1) was introduced. After removing the solvent using a glass filter, 40 ml of ethanol was introduced therein. Thus, a proportion of ethanol was 100% by volume. Then, 3.10 g of the same lot heat-treated silica as that used in Example 40 (2) was introduced therein. The mixture was heated to 40°C
15 and stirred for 2 hours. After removing the liquid component using a glass filter, the residue was washed 4 times with each 30 ml of toluene and washed once with 30 ml of hexane, and thereafter dried for 8 hours at 120°C under reduced pressure, thereby obtaining 3.82 g of a component (A). Elementary analysis
20 revealed Zn = 2.4 mmol/g, and F = 2.5 mmol/g.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a
25 partial pressure of 0.017 MPa, 695 g of butane and 55 g of 1-butene were fed therein, and then a temperature of the autoclave was

raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.752 mol% and 1-butene = 2.62 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μmol/ml, and then 8.4 mg of the component (A) obtained in the above Example 41 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.203 mol%) so as to keep the whole pressure constant. As a result, 11.9 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 7.9×10^6 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 1420 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 14.6, MFR = 2.03, MFRR = 48.8 and SR = 1.53.

[Example 42]

(1) Production of contact product (e)

In a 1000 ml four necked flask purged with nitrogen, 300 ml of toluene and 120 ml (240 mmol) of a hexane solution of diethylzinc (2 mol/l) were introduced and cooled to -5°C. To the mixture was dropped 34.8 ml (96.3 mmol) of a toluene solution

of pentafluorophenol (2.77 mol/l) over 40 minutes. After completion of dropping, the mixture was gradually heated to 20°C. Thereafter, 4.74 ml (263 mmol) of H₂O was dropped thereto over 60 minutes at 20°C. After completion of the drop, the mixture
5 was stirred for 6 hours under reflux condition (91°C). As a result, 420 ml of a slurry of a contact product (e) was obtained. According to calculations based on the amounts of reagents used, a slurry concentration was found to be 0.57 mmol (in terms of Zn atom)/ml (slurry).

10 (2) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 45 ml (25.7 mmol-Zn) of the slurry of the contact product (e) obtained in the above Example 42 (1) was introduced. After removing the solvent using a glass filter, 2 ml of ethanol and 38 ml of toluene
15 were introduced therein. Thus, a proportion of ethanol was 5% by volume. Then, 3.03 g of the same lot heat-treated silica as that used in Example 40 (3) was introduced therein. The mixture was heated to 40°C and stirred for 2 hours. The resulting solid component was allowed to precipitate, and a slurry portion of
20 the upper layer was removed. For completing washing, 30 ml of toluene was added to the residue, the mixture was stirred, thereafter, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. The above-mentioned washing operation was repeated 4 times. A similar
25 washing operation was repeated 2 times using 30 ml of hexane. After removing the liquid component using a glass filter, the

residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 3.51 g of a component (A). Elementary analysis revealed Zn = 2.4 mmol/g, and F = 3.8 mmol/g.

(3) Polymerization

5 A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.017 MPa, 695 g of butane and 55 g of 1-butene were fed therein, and then a temperature of the autoclave was
10 raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.713 mol% and 1-butene = 2.55 mol%. To the resultant was added 0.9 ml of a heptane solution
15 of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 8.0 mg of the component (A) obtained in the above Example 42 (2) as a solid catalyst
20 component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.200 mol%) so as to keep the whole pressure constant. As a result, 28.7 g of an olefin polymer having a good particle property was obtained. Polymerization activity per
25 zirconium atom was found to be 1.9×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was

found to be 3590 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 13.2, MFR = 0.780, MFRR = 58.6 and SR = 1.25.

[Example 43]

5 (1) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 45 ml (25.7 mmol-Zn) of the slurry of the contact product (e) obtained in the above Example 42 (1) was introduced. After removing the solvent using a glass filter, 8 ml of ethanol and 32 ml of toluene
10 were introduced therein. Thus, a proportion of ethanol was 20% by volume. Then, 2.65 g of the same lot heat-treated silica as that used in Example 40 (3) was introduced therein. The mixture was heated to 40°C and stirred for 2 hours. The resulting solid component was allowed to precipitate, and a slurry portion of
15 the upper layer was removed. For completing washing, 30 ml of toluene was added to the residue, the mixture was stirred, thereafter, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. The above-mentioned washing operation was repeated 4 times. A similar
20 washing operation was carried out 1 time using 30 ml of hexane. After removing the liquid component using a glass filter, the residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 4.17 g of a component (A). Elementary analysis revealed Zn = 3.7 mmol/g, and F = 3.8 mmol/g.

25 (2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had

been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.017 MPa, 695 g of butane and 55 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.693 mol% and 1-butene = 2.79 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 9.3 mg of the component (A) obtained in the above Example 43 (2) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.200 mol%) so as to keep the whole pressure constant. As a result, 60.6 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 4.0×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 6520 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 17.1, MFR = 4.61, MFRR = 41.2 and SR = 1.42.

[Example 44]

(1) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 45 ml (25.7 mmol-Zn) of the slurry of the contact product (e) obtained in the above Example 42 (1) was introduced. After removing the solvent using a glass filter, 16 ml of ethanol and 24 ml of toluene were introduced therein. Thus, a proportion of ethanol was 40% by volume. Then, 3.15 g of the same lot heat-treated silica as that used in Example 40 (3) was introduced therein. The mixture was heated to 40°C and stirred for 2 hours. The resulting solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. For completing washing, 30 ml of toluene was added to the residue, the mixture was stirred, thereafter, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. The above-mentioned washing operation was repeated 4 times. A similar washing operation was carried out 1 time using 30 ml of hexane. After removing the liquid component using a glass filter, the residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 4.98 g of a component (A). Elementary analysis revealed Zn = 3.8 mmol/g, and F = 3.7 mmol/g.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.017 MPa, 695 g of butane and 55 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain

a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.680 mol% and 1-butene = 2.87 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 9.2 mg of the component (A) obtained in the above Example 44 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.200 mol%) so as to keep the whole pressure constant. As a result, 44.9 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 3.0×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 4880 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 12.5, MFR = 1.23, MFRR = 54.6 and SR = 1.41.

[Example 45]

(1) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 20 ml of toluene and 12.8 ml (25.6 mmol) of a hexane solution of diethylzinc (2 mol/l) were introduced and cooled to 2°C. To the mixture was dropped 3.7 ml (10.2 mmol) of a toluene solution of pentafluorophenol (2.77 mol/l) over 40 minutes. After

completion of dropping, the mixture was stirred at 2°C for 20 minutes. Thereafter, 0.51 ml (28.3 mmol) of H₂O was dropped thereto over 60 minutes at 2°C. Immediately thereafter, 2.98 g of the same lot heat-treated silica as that used in Example 5 40 (2) was introduced therein at 2°C. The mixture was stirred at 2°C for 1 hour, thereafter heated to 60°C and stirred for 2 hours. The resulting solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. For completing washing, 30 ml of toluene was added to the residue, the mixture 10 was stirred, thereafter, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. The above-mentioned washing operation was repeated 4 times. A similar washing operation was carried out 1 time using 30 ml of hexane. After removing the liquid component using a glass 15 filter, the residue was dried at ambient temperature under reduced pressure, thereby obtaining 5.88 g of a component (A). Elementary analysis revealed Zn = 3.5 mmol/g, and F = 6.3 mmol/g.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had 20 been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.017 MPa, 695 g of butane and 55 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain 25 a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition

in the system was found to be hydrogen = 0.707 mol% and 1-butene = 2.82 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic
5 ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 8.8 mg of the component (A) obtained in the above Example 45 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed
10 gas (hydrogen = 0.200 mol%) so as to keep the whole pressure constant. As a result, 94.9 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 6.3×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was
15 found to be 10800 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 15.8, MFR = 3.15, MFRR = 41.9 and SR = 1.38.

[Example 46]

(1) Production of component (A)

20 In a 50 ml four necked flask purged with nitrogen, 20 ml of toluene and 12.8 ml (25.6 mmol) of a hexane solution of diethylzinc (2 mol/l) were introduced and cooled to 5°C. To the mixture was dropped 3.7 ml (10.2 mmol) of a toluene solution of pentafluorophenol (2.77 mol/l) over 40 minutes. After
25 completion of dropping, the mixture was stirred at 5°C for 20 minutes. Thereafter, 0.51 ml (28.3 mmol) of H₂O was dropped

thereto over 60 minutes at 5°C. After completion of dropping, the mixture was stirred at 5°C for 1 hour, and then 2.93 g of the same lot heat-treated silica as that used in Example 40 (2) was introduced therein at 2°C. The mixture was stirred at 2°C
5 for 1 hour, thereafter heated to 40°C and stirred for 2 hours. The resulting solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. For completion of washing, 30 ml of toluene was added to the residue, the mixture was stirred, thereafter, the solid component was allowed to
10 precipitate, and a slurry portion of the upper layer was removed. The above-mentioned washing operation was repeated 4 times. A similar washing operation was carried out 1 time using 30 ml of hexane. After removing the liquid component using a glass filter, the residue was dried at ambient temperature under
15 reduced pressure, thereby obtaining 5.84 g of a component (A). Elementary analysis revealed Zn = 3.4 mmol/g, and F = 5.3 mmol/g.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with
20 argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.017 MPa, 695 g of butane and 55 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized.
25 As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.748 mol% and 1-butene

= 2.59 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 8.8 mg of the component (A) obtained in the above Example 46 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.200 mol%) so as to keep the whole pressure constant. As a result, 57.4 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 3.8×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 6520 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 15.4, MFR = 0.453, MFRR = 40.2 and SR = 1.16.

[Example 47]

(1) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 20 ml of toluene and 12.8 ml (25.6 mmol) of a hexane solution of diethylzinc (2 mol/l) were introduced and cooled to 5°C. To the mixture was dropped 3.7 ml (10.2 mmol) of a toluene solution of pentafluorophenol (2.77 mol/l) over 40 minutes. After completion of dropping, the mixture was stirred at 5°C for 20 minutes. Thereafter, 0.51 ml (28.3 mmol) of H₂O was dropped thereto over 60 minutes at 5°C. After completion of dropping,

the mixture was stirred at 5°C for 3 hours, and then 3.01 g of the same lot heat-treated silica as that used in Example 40 (2) was introduced therein at 4°C. The mixture was stirred at 4°C for 1 hour, thereafter heated to 60°C and stirred for 2 hours.

5 The resulting solid component was allowed to precipitate, and a slurry portion of the upper layer was removed. For completion of washing, 30 ml of toluene was added to the residue, the mixture was stirred, thereafter, the solid component was allowed to precipitate, and a slurry portion of the upper layer was removed.

10 The above-mentioned washing operation was repeated 4 times. A similar washing operation was carried out 1 time using 30 ml of hexane. After removing the liquid component using a glass filter, the residue was dried at ambient temperature under reduced pressure, thereby obtaining 4.31 g of a component (A).

15 Elementary analysis revealed Zn = 2.4 mmol/g, and F = 3.9 mmol/g.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a

20 partial pressure of 0.017 MPa, 695 g of butane and 55 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition

25 in the system was found to be hydrogen = 0.624 mol% and 1-butene = 2.67 mol%. To the resultant was added 0.9 ml of a heptane solution

of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 8.3 mg of the component
5 (A) obtained in the above Example 47 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.200 mol%) so as to keep the whole pressure constant. As a result, 42.6 g of an olefin polymer having a good
10 particle property was obtained. Polymerization activity per zirconium atom was found to be 2.8×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 5130 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 14.3, MFR = 0.135,
15 MFRR = 141 and SR = 1.12.

[Example 48]

(1) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 3.00 g of the same lot heat-treated silica as that used in Example
20 40 (2), 30 ml of toluene and 0.149 g (8.25 mmol) of H₂O were introduced. The mixture was stirred at ambient temperature for 30 minutes, during which H₂O existing in toluene in the form of a liquid drop visually disappeared, thereby forming a uniform slurry (it seems that the silica absorbed it). To the mixture
25 was added 1.8 ml (3.6 mmol) of a toluene solution of pentafluorophenol (2.0 mol/l), and the resulting mixture was

cooled to 5°C using an ice bath. Then, 4.5 ml (9.0 mmol) of a hexane solution of diethylzinc (2 mol/l) was dropped thereto over 1 hour. After completion of dropping, the mixture was stirred at 5°C for 1 hour, heated to 40°C, stirred at 40°C for 1 hour, 5 further heated to 60°C and stirred at 60°C for 1 hour. After filtration using a filter, the residue was washed 3 times with 30 ml of toluene at 40°C and 2 times with 30 ml of hexane at ambient temperature with use of a filter. Successively, the remaining solid component was dried at 40°C for 1 hour and at 10 120°C for 8 hours under reduced pressure, thereby obtaining a component (A). Elementary analysis of the component (A) dried at 120°C revealed Zn = 2.1 mmol/g, and F = 3.4 mmol/g.

(2) Polymerization A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and 15 thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.017 MPa, 695 g of butane and 55 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the 20 system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.738 mol% and 1-butene = 2.93 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene 25 solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 7.3 mg of

the component (A) obtained in the above Example 48 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70 °C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.203 mol%) so as to
5 keep the whole pressure constant. As a result, 44.5 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 3.0×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 6100 g/g-solid catalyst
10 component/hour. Further, the obtained olefin polymer showed, SCB = 13.0, MFR = 1.32, MFRR = 42.7 and SR = 1.44.

[Example 49]

(1) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 2.99
15 g of the same lot heat-treated silica as that used in Example 40 (2), 30 ml of hexane and 0.149 g (8.29 mmol) of H₂O were introduced. The mixture was stirred at ambient temperature for 30 minutes, during which H₂O existing in toluene in the form of a liquid drop visually disappeared, thereby forming a uniform
20 slurry (it seems that the silica absorbed it). To the mixture was added 1.8 ml (3.6 mmol) of a toluene solution of pentafluorophenol (2.0 mol/l), and the resulting mixture was cooled to 5°C using an ice bath. Then, 4.5 ml (9.0 mmol) of a hexane solution of diethylzinc (2 mol/l) was dropped thereto
25 over 1 hour. After completion of dropping, the mixture was stirred at 5°C for 1 hour, heated to 40°C, and stirred at 40°C for 2

hours. After filtration using a filter, the residue was washed 3 times with 30 ml of toluene at 40°C and 2 times with 30 ml of hexane at ambient temperature with use of a filter. Successively, the remaining solid component was dried at 40°C
5 for 1 hour and at 120°C for 8 hours under reduced pressure, thereby obtaining a component (A). Elementary analysis of the component (A) dried at 120°C revealed Zn = 2.1 mmol/g, and F = 4.0 mmol/g.
(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had
10 been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.017 MPa, 695 g of butane and 55 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain
15 a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.789 mol% and 1-butene = 2.78 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml.
20 Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 7.8 mg of the component (A) obtained in the above Example 49 (1) as a solid catalyst component were added thereto. Polymerization was carried out
25 for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.200 mol%) so as to keep the whole pressure

constant. As a result, 53.9 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 3.6×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 6910 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 15.0, MFR = 3.00, MFRR = 42.0 and SR = 1.48.

[Example 50]

(1) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 2.98 g of the same lot heat-treated silica as that used in Example 40 (2), 30 ml of tetrahydrofuran, 5.4 ml (10.8 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.0 mol/l) and 0.458 g (25.4 mmol) of H₂O were introduced and cooled to 5°C using an ice bath. Then, 13.4 ml (27 mmol) of a hexane solution of diethylzinc (2 mol/l) was dropped thereto over 1 hour. After completion of dropping, the mixture was stirred at 5°C for 1 hour, heated to 40°C, and stirred at 40°C for 2 hours. The resulting solid component was allowed to precipitate, a slurry portion of the upper layer was removed, and further the liquid component was removed using a glass filter. As a washing operation, 30 ml of tetrahydrofuran was added thereto, stirring was carried out until the temperature reached 40°C, thereafter the solid component was allowed to precipitate, a slurry portion of the upper layer was removed, and the remaining liquid component was removed using a glass filter. Said washing operation was repeated

5 times. Successively, the remaining solid component was dried under reduced pressure at 120°C for 8 hours, thereby obtaining 4.92 g of a component (A). Elementary analysis revealed Zn = 3.5 mmol/g, and F = 4.4 mmol/g.

5 (2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.012 MPa, 700 g of butane and 50 g of 1-butene
10 were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.574 mol% and 1-butene
15 = 2.40 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 13.4 mg of the component
20 (A) obtained in the above Example 50 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.193 mol%) so as to keep the whole pressure constant. As a result, 42.1 g of an olefin polymer having a good
25 particle property was obtained. Polymerization activity per zirconium atom was found to be 8.4×10^7 g/mol-Zr/hour, and

polymerization activity per the solid catalyst component was found to be 3140 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 17.8, MFR = 22.0 and SR = 1.54.

5 [Example 51]

(1) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 3.05 g of the same lot heat-treated silica as that used in Example 40 (2), 30 ml of hexane and 0.081 g (4.5 mmol) of H₂O were introduced.

10 The mixture was stirred at ambient temperature for 30 minutes, during which H₂O existing in hexane in the form of a liquid drop visually disappeared, thereby forming a uniform slurry (it seems that the silica absorbed it). After cooling to 5°C using an ice bath, 4.48 ml (9 mmol) of a hexane solution of diethylzinc (2.01

15 mol/l) was dropped over 1 hour. After completion of dropping, the mixture was stirred at 5°C for 30 minutes, heated to 60°C, and stirred at 60°C for 1 hour. Successively, After cooling to 5°C using an ice bath, 4.5 ml (9.0 mmol) of a toluene solution of pentafluorophenol (2.00 mol/l) was dropped thereto over 1

20 hour. After completion of dropping, the mixture was stirred at 5°C for 30 minutes, heated to 60°C, and stirred at 60°C for 1 hour. After filtration using a filter, the residue was washed with 30 ml of hexane at 60°C 4 times using a filter. Successively, the remaining solid component was dried at 40°C for 1 hour and

25 at 120°C for 8 hours under reduced pressure, thereby obtaining a component (A). Elementary analysis of the component (A)

revealed Zn = 1.8 mmol/g, and F = 6.4 mmol/g.

[0270]

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had
5 been dried under reduced pressure and thereafter purged with
argon, was made vacuum, hydrogen was fed therein to obtain a
partial pressure of 0.015 MPa, 695 g of butane and 55 g of 1-butene
were fed therein, and then a temperature of the autoclave was
raised to 70°C. Thereafter, ethylene was fed therein to obtain
10 a partial pressure of 1.6 MPa, and the system was stabilized.
As a result of a gas chromatography analysis, a gas composition
in the system was found to be hydrogen = 0.802 mol% and 1-butene
= 2.32 mol%. To the resultant was added 0.9 ml of a heptane solution
of triisobutylaluminum adjusted to a concentration of 1 mmol/ml.
15 Successively, 0.75 ml of a toluene solution of racemic
ethylenebis(1-indenyl)zirconium dichloride adjusted to a
concentration of 2 μ mol/ml, and then 10.3 mg of the component
(A) obtained in the above Example 51 (1) as a solid catalyst
component were added thereto. Polymerization was carried out
20 for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed
gas (hydrogen = 0.212 mol%) so as to keep the whole pressure
constant. As a result, 58 g of an olefin polymer having a good
particle property was obtained. Polymerization activity per
zirconium atom was found to be 3.9×10^7 g/mol-Zr/hour, and
25 polymerization activity per the solid catalyst component was
found to be 5630 g/g-solid catalyst component/hour. Further,

the obtained olefin polymer showed, SCB = 11.7, MFR = 1.54, MFRR = 43.6 and SR = 1.40.

[Example 52]

(1) Production of component (A)

5 In a 5 l four necked flask purged with nitrogen, 1.5 l of hexane and 1.49 l (3.0 mol) of a hexane solution of diethylzinc (2.01 mol/l) were introduced and cooled to 5°C. To the mixture, a solution prepared by dissolving 198.8 g (1.07 mol) of pentafluorophenol in a mixed solvent of 342 ml of hexane and
10 194 ml of toluene was dropped over 85 minutes. After completion of dropping, the mixture was stirred for 1 hour at 5°C, thereafter heated to 40°C, stirred for 1 hour, and thereafter allowed to stand over night at ambient temperature. After cooling to 20°C using an ice bath, 359 g of the same lot heat-treated silica
15 as that used in Example 40 (2) was introduced thereto, and then 0.3 l of hexane was poured therein so as to wash away the silica adhering to a wall of the flask. After stirring the mixture for 5 minutes, 44.2 g (2.46 mol) of H₂O was added thereto over 1 hour. After completion of the dropping, the mixture was stirred
20 at 20°C for 1 hour, at 40°C for 2 hours and at 60°C for 2 hours. Thereafter, the mixture was allowed to stand, thereby precipitating a solid component. At the time when an interface between a layer of the precipitated solid component and an upper layer of a slurry portion appeared, the upper slurry portion
25 was removed. Thereafter, the remaining liquid component was removed by filtration. For completing washing, 3 l of hexane

was added to the residue, and the mixture was stirred, and thereafter allowed to stand, thereby precipitating a solid component, and then the upper slurry portion was removed similarly at the time when the interface appeared. Thereafter, the remaining liquid component was removed by filtration. The above washing operation was repeated 3 times. Thereafter, the solid component was dried for 4 hours at 60°C under reduced pressure, thereby obtaining 814 g of a component (A). Elementary analysis revealed Zn = 3.7 mmol/g, and F = 6.3 mmol/g.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.015 MPa, 690 g of butane and 60 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.76 mol% and 1-butene = 2.42 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 9.4 mg of the component (A) obtained in the above Example 52 (1) as a solid catalyst component were added thereto. Polymerization was carried out

for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.23 mol%) so as to keep the whole pressure constant. As a result, 83 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 5.5×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 8830 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 15.2, MFR = 0.09, MFRR = 127 and SR = 1.05. [Example 53]

10 (1) Production of component (A)

In a 5 l four necked flask purged with nitrogen, 1.5 l of hexane and 1.49 l (3.0 mol) of a hexane solution of diethylzinc (2.01 mol/l) were introduced and cooled to 5°C. To the mixture, a solution prepared by dissolving 277.1 g (1.51 mol) of pentafluorophenol in 585 ml of toluene was dropped over 100 minutes. After completion of dropping, the mixture was stirred for 1 hour at 5°C, thereafter heated to 40°C, and stirred for 1 hour. After cooling to 5°C using an ice bath, 355 g of the same lot heat-treated silica as that used in Example 40 (2) was introduced thereto, and then 0.3 l of toluene was poured therein so as to wash away the silica adhering to a wall of the flask. After stirring the mixture for 5 minutes, 41.0 g (2.28 mol) of H₂O was added thereto over 1 hour. After completion of the dropping, the mixture was stirred at 5°C for 2.5 hours, and at 80°C for 25 2 hours. Thereafter, the mixture was allowed to stand, thereby precipitating a solid component. At the time when an interface

between a layer of the precipitated solid component and an upper layer of a slurry portion appeared, the upper slurry portion was removed. The remaining liquid component was removed by filtration. Thereafter, 3 l of toluene was added thereto, and
5 the mixture was allowed to stand overnight at ambient temperature. After that, washing of the solid component was repeated 4 times with each 3 l of toluene at 80°C and 2 times with each 3 l of hexane at ambient temperature, in a manner such that the mixture was allowed to stand to precipitate a solid component, the upper
10 slurry portion was removed at the time when the interface between a layer of the precipitated solid component and an upper layer of a slurry portion appeared, and then, the remaining liquid component was removed by filtration. Thereafter, the solid component was dried for 2 hours at 60°C under reduced pressure,
15 thereby obtaining 864 g of a component (A). Elementary analysis revealed Zn = 3.5 mmol/g, and F = 6.8 mmol/g.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with
20 argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.015 MPa, 690 g of butane and 60 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized.
25 As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.70 mol% and 1-butene

= 3.01 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 8.7 mg of the component (A) obtained in the above Example 53 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.26 mol%) so as to keep the whole pressure constant. As a result, 85 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 5.7×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 9770 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 16.8.

[Example 54]

(1) Production of component (A)

In a 5 l four necked flask purged with nitrogen, 2.5 l of toluene, 71 ml of tetrahydrofuran and 0.871 l (1.75 mol) of a hexane solution of diethylzinc (2.01 mol/l) were introduced and cooled to 5°C. To the mixture, a solution prepared by dissolving 164.8 g (0.90 mol) of pentafluorophenol in 277 ml of toluene was dropped over 80 minutes. After completion of dropping, the mixture was stirred for 1 hour at 5°C, thereafter heated to 40°C, and stirred for 1 hour. After cooling to 5°C using an ice bath, 350 g of the same lot heat-treated silica

as that used in Example 40 (2) was introduced therein, and then 0.1 l of toluene was poured therein so as to wash away the silica adhering to a wall of the flask. After stirring the mixture for 5 minutes, 23.8 g (1.32 mol) of H₂O was added thereto over 1.5 hours. After completion of the dropping, the mixture was stirred at 5°C for 85 minutes, and at 40°C for 2 hours. Thereafter, the mixture was allowed to stand over night at ambient temperature. Thereafter, the mixture was stirred for 2 hours at 80°C, and then allowed to stand, thereby precipitating a solid component.

At the time when an interface between a layer of the precipitated solid component and an upper layer of a slurry portion appeared, the upper slurry portion was removed. The remaining liquid component was removed by filtration. Thereafter, 3.5 l of toluene was added thereto, and the mixture was stirred for 2 hours at 95°C. After that, washing of the solid component was repeated 4 times with each 3.5 l of toluene at 95°C and 2 times with each 3.5 l of hexane at ambient temperature, in a manner such that the mixture was allowed to stand, thereby precipitating a solid component, the upper slurry portion was removed at the time when the interface between a layer of the precipitated solid component and an upper layer of a slurry portion appeared, and then the remaining liquid component was removed by filtration. Thereafter, the solid component was dried for 4 hours at 120°C under reduced pressure, thereby obtaining 571 g of a component (A). Elementary analysis revealed Zn = 2.6 mmol/g, and F = 4.9 mmol/g.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.015 MPa, 690 g of butane and 60 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.74 mol% and 1-butene = 2.75 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 8.4 mg of the component (A) obtained in the above Example 54 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.26 mol%) so as to keep the whole pressure constant. As a result, 116 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 7.7×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 13800 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 16.9, MFR = 1.70, MFRR = 46.0 and SR = 1.29.

[Example 55]

(1) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 12.5 ml (25 mmol) of a hexane solution of diethylzinc (2.01 mol/l) and 12.5 ml of tetrahydrofuran were introduced and cooled to 5°C. To the mixture, 6.25 ml of a tetrahydrofuran solution of pentafluorophenol adjusted to a concentration of 2 mmol/ml was dropped over 21 minutes. After completion of dropping, the mixture was stirred for 1 hour at 5°C and for 1 hour at 40°C. After cooling to 5°C using an ice bath, 2.98 g of the same lot heat-treated silica as that used in Example 40 (2) was introduced thereto, and then 8 ml of tetrahydrofuran was poured therein so as to wash away the silica adhering to a wall of the flask. After stirring the mixture for 5 minutes, 0.34 g (18.8 mmol) of H₂O was added thereto over 38 minutes. After completion of dropping, the mixture was stirred at 5°C for 2.5 hours, and at 60°C for 2 hours. Thereafter, washing was carried out using a filter 4 times with each 25 ml of toluene at 80°C, and 2 times with 25 ml of hexane at ambient temperature. Thereafter, the solid component was dried for 2 hours at 120°C under reduced pressure, thereby obtaining 4.80 g of a component (A). Elementary analysis revealed Zn = 2.9 mmol/g, and F = 5.3 mmol/g.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.015 MPa, 690 g of butane and 60 g of 1-butene

were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition
5 in the system was found to be hydrogen = 0.74 mol% and 1-butene = 3.19 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a
10 concentration of 2 μmol/ml, and then 8.0 mg of the component (A) obtained in the above Example 55 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.25 mol%) so as to keep the whole pressure constant.
15 As a result, 174 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.2×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 21800 g/g-solid catalyst component/hour. Further, the obtained olefin
20 polymer showed, SCB = 14.7, MFR = 0.89, MFRR = 63.3 and SR = 1.29.

[Example 56]

(1) Production of contact product (e)

In a 200 ml four necked flask purged with nitrogen, 40
25 ml of tetrahydrofuran and 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) were introduced and cooled to 5°C. To

the mixture was dropped 34.3 ml (80 mmol) of a tetrahydrofuran solution of 3,4,5-trifluorophenol (2.33 mol/l) over 120 minutes. After completion of dropping, the mixture was stirred at 5°C for 90 minutes and further at 40°C for 70 minutes. Thereafter, the temperature was lowered to 20°C using an ice bath, 0.72 ml (40 mmol) of H₂O was dropped thereto over 80 minutes. After completion of dropping, the mixture was stirred at 20°C for 60 minutes and further stirred at 40°C for 120 minutes. As a result, the mixture was separated into a transparent liquid product and a yellow gel product. After allowing to stand all night at ambient temperature, a volatile component was removed by distillation, and the residue was dried at 120°C for 8 hours under reduced pressure. As a result, 17.05 g of a yellow solid product was obtained.

(2) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 5.98 g of the yellow solid product obtained in the above Example 56 (1), 30 ml of tetrahydrofuran and 3.15 g of same lot heat-treated silica as that used in Example 1(2) were introduced and stirred at 40°C for 2 hours. After removing the liquid component using a glass filter, the residue was washed 5 times with each 30 ml of tetrahydrofuran and dried for 8 hours at 120°C under reduced pressure, thereby obtaining 4.45 g of a component (A). Elementary analysis revealed Zn = 2.1 mmol/g, and F = 2.4 mmol/g.

(3) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had

been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.009 MPa, 700 g of butane and 50 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.36 mol% and 1-butene = 2.67 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 16.6 mg of the component (A) obtained in the above Example 56 (2) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.25 mol%) so as to keep the whole pressure constant. As a result, 78 g of an olefin polymer was obtained. Polymerization activity per zirconium atom was found to be 1.6×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 4700 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 16.3, MFR = 1.99, MFRR = 50.4, SR = 1.36, $M_w = 9.51 \times 10^4$, $M_w/M_n = 8.99$, and $[\eta] = 1.26$.

[Example 57]

(1) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.012 MPa, 700 g of butane and 50 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.51 mol% and 1-butene = 2.91 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 18.0 mg of the component (A) obtained in the above Example 56 (2) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.25 mol%) so as to keep the whole pressure constant. As a result, 75 g of an olefin polymer was obtained. Polymerization activity per zirconium atom was found to be 1.5×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 4170 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 17.2, MFR = 7.03, MFRR = 39.9, SR = 1.45, $M_w = 6.83 \times 10^4$, $M_w/M_n = 7.32$, and $[\eta] = 1.04$.
[Example 58]

(1) Production of contact product (e)

In a 200 ml four necked flask purged with nitrogen, 40 ml of tetrahydrofuran and 40 ml (80 mmol) of a hexane solution of diethylzinc (2 mol/l) were introduced and cooled to 5°C. To
5 the mixture was dropped 13.7 ml (32 mmol) of a tetrahydrofuran solution of 3,4,5-trifluorophenol (2.33 mol/l) over 45 minutes. After completion of dropping, the mixture was stirred at 5°C for 90 minutes and further at 40°C for 70 minutes. Thereafter, the temperature was lowered to 20°C using an ice bath, 1.30 ml
10 (72 mmol) of H₂O was dropped thereto over 145 minutes. After completion of dropping, the mixture was stirred at 20°C for 60 minutes and further stirred at 40°C for 120 minutes. As a result, the mixture was separated into a transparent liquid product and a yellow gel product. After allowing to stand overnight at ambient
15 temperature, a volatile component was removed by distillation, and the residue was dried at 120°C for 8 hours under reduced pressure. As a result, 10.39 g of a yellow solid product was obtained.

(2) Production of component (A)

20 In a 50 ml four necked flask purged with nitrogen, 3.65 g of the yellow solid product obtained in the above Example 56 (1), 30 ml of tetrahydrofuran and 3.08 g of same lot heat-treated silica as that used in Example 1 (2) were introduced and stirred at 40°C for 2 hours. After removing the liquid component using
25 a glass filter, the residue was washed 5 times with each 30 ml of tetrahydrofuran and dried for 8 hours at 120°C under reduced

pressure, thereby obtaining 4.76 g of a component (A). Elementary analysis revealed Zn = 3.1 mmol/g, and F = 2.4 mmol/g.

(3) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had
5 been dried under reduced pressure and thereafter purged with
argon, was made vacuum, hydrogen was fed therein to obtain a
partial pressure of 0.004 MPa, 690 g of butane and 60 g of 1-butene
were fed therein, and then a temperature of the autoclave was
raised to 70°C. Thereafter, ethylene was fed therein to obtain
10 a partial pressure of 1.6 MPa, and the system was stabilized.
As a result of a gas chromatography analysis, a gas composition
in the system was found to be hydrogen = 0.17 mol% and 1-butene
= 2.85 mol%. To the resultant was added 0.9 ml of a heptane solution
of triisobutylaluminum adjusted to a concentration of 1 mmol/ml.
15 Successively, 0.25 ml of a toluene solution of racemic
ethylenebis(1-indenyl)zirconium dichloride adjusted to a
concentration of 2 μ mol/ml, and then 9.8 mg of the component
(A) obtained in the above Example 58 (2) as a solid catalyst
component were added thereto. Polymerization was carried out
20 for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed
gas (hydrogen = 0.22 mol%) so as to keep the whole pressure constant.
As a result, 87 g of an olefin polymer was obtained. Polymerization
activity per zirconium atom was found to be 1.7×10^8
g/mol-Zr/hour, and polymerization activity per the solid
25 catalyst component was found to be 8880 g/g-solid catalyst
component/hour. Further, the obtained olefin polymer showed,

SCB = 17.6, MFR = 2.62, MFRR = 45.4, SR = 1.36, $M_w = 9.93 \times 10^4$, $M_w/M_n = 4.53$, and $[\eta] = 1.32$.

[Example 59]

(1) Pre-polymerization

5 A 0.4 l-volume autoclave equipped with a stirrer was made vacuum, hydrogen was fed therein to obtain a pressure of 0.002 MPa, 90 g of butane was fed therein as a solvent, and then a temperature of the autoclave was raised to 50°C. Further, ethylene was fed therein to obtain a partial pressure of 0.15 MPa, and
10 the system was stabilized. As a result of gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.02 mol%. Then, 0.5 ml of a hexane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml was introduced therein. Successively, 0.2 ml of a hexane solution
15 of racemic ethylenebis(1-indenyl)zirconium dichloride and triisobutylaluminum adjusted to a zirconium concentration and an aluminum concentration to 150 μ mol/ml and 1.0 mmol/ml, respectively, was added thereto, and then 0.42 g of the component (A) obtained in the above Example 56 (2) was introduced therein
20 to initiate polymerization. During the polymerization, a mixed gas of ethylene and hydrogen (hydrogen concentration = 0.24 mol%) was continuously supplied while adjusting so as to keep the gas phase pressure in the autoclave to 0.68 MPa. After 72 minutes, ethylene, butane and hydrogen gas were released, and a solid
25 remaining in the autoclave was dried for 30 minutes at ambient temperature under reduced pressure. As a result, a

polyethylene-pre-polymerized catalyst component for addition polymerization was obtained in a yield of 51 g per g of the component (A). Intrinsic viscosity $[\eta]$ and MFR of said polyethylene were 1.16 dl/g and 4.2 g/10 min., respectively.

5 (2) Real polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.008 MPa, 700 g of butane and 50 g of 1-butene
10 were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.32 mol% and 1-butene
15 = 2.75 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.427 g of the pre-polymerized catalyst component for addition polymerization obtained in the above Example 59(1) was added thereto. Polymerization was continued for 60 minutes
20 at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.25 mol%) so as to keep the whole pressure constant. As a result, 48 g of an olefin polymer was obtained. Polymerization activity per the solid catalyst component (the component (A) obtained in the above Example 56 (2)) was found to be 5750 g/g-solid
25 catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 17.8, MFR = 22.8 and $[\eta]$ = 0.89.

[Example 60]

(1) Production of solid product

In a 5 l four necked flask purged with nitrogen, 1.65 l of tetrahydrofuran and 1.50 l (2.98 mol) of a hexane solution of diethylzinc (1.99 mol/l) were introduced and cooled to 5°C. To the mixture, a solution prepared by dissolving 223.36 g (1.21 mol) of pentafluorophenol in 370 ml of tetrahydrofuran was dropped over 70 minutes. During the dropping, temperature in the flask was kept from 4 to 6°C. After completion of dropping, the mixture was stirred at that temperature for 1 hour, thereafter, gradually heated to 45°C and stirred for 1 hour. Thereafter, the temperature was lowered to 20°C using an ice bath, and 48.2 g (2.67 mol) of H₂O was dropped thereto over 1.5 hours at that temperature. As a result, the mixture was separated into a yellow transparent liquid product and a yellow gel product. After stirring at 20°C for 1 hour, the mixture was heated to 45°C, stirred for 1 hour at that temperature, and thereafter allowed to stand for 12 hours at ambient temperature. A volatile component was removed by distillation under reduced pressure, and the residue was dried at 120°C for 8 hours under reduced pressure. As a result, 472.6 g of a solid product was obtained.

In a 200 ml four necked flask purged with nitrogen, 19.7 g of the above-mentioned solid product and 139 ml of tetrahydrofuran were introduced and heated to 40°C. After stirring for 1 hour at that temperature, the mixture was cooled to ambient temperature, and allowed to stand for 2 days.

Filtration gave 0.20 g of an insoluble matter. The filtrate was transferred to a 300 ml four necked flask purged with nitrogen, and 48.5 ml of hexane was added to the filtrate, thereby precipitating a solid. Thus, $VH/(VT + VH) = 0.26$. The mixture
5 was heated to 40°C, stirred for 1 hour at that temperature, thereafter cooled to ambient temperature and filtered. The solid separated by filtration was dried at 120°C for 2 hours under reduced pressure. As a result, 1.35 g of a solid product was obtained.

10 [0271]

(2) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 928.4 mg of the solid product obtained in the above Example 60 (1) and 7 ml of tetrahydrofuran were introduced and stirred for 1
15 hour at ambient temperature. To the mixture, 721 mg of the same lot heat-treated silica as that used in Example 1 (2) was added thereto. The resulting mixture was heated to 40°C, stirred for 2 hours, and thereafter filtered. To the residue, 7 ml of tetrahydrofuran was added, and the mixture was stirred at ambient
20 temperature and thereafter filtered. This operation was repeated 5 times. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 834.2 mg of a component (A).

(3) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had
25 been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein below 10°C to

obtain a partial pressure of 0.01 MPa, 700 g of butane and 50 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium dichloride adjusted to a concentration of 2 μ mol/ml, and then 11.9 mg of the component (A) obtained in the above Example 60 (2) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene so as to keep the whole pressure constant. As a result, 63.6 g of an olefin polymer having a good particle property was obtained. Polymerization activity per the solid catalyst component (the component (A) obtained in the above-mentioned (2)) was found to be 5,340 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 13.5, MFR = 0.08, MFRR = 171 and SR = 1.09.

[Example 61]

(1) Production of solid product

To the filtrate obtained in Example 60 (1), 15 ml of hexane was further added to precipitate a solid. Thus, $VH/(VT + VH) = 0.31$. The mixture was heated to 40°C, stirred for 1 hour at that temperature, thereafter cooled to ambient temperature and then filtered. The solid separated by filtration was dried for

2 hours at 120°C under reduced pressure. As a result, 3.71 g of a solid product was obtained.

[0272]

(2) Production of component (A)

5 Example 60 (2) was repeated, except that the solid product was changed to 2.60 g of the solid product obtained in the above Example 61 (1), and the amounts of tetrahydrofuran and the silica were changed to 17.6 ml and 1.82 g, respectively. As a result, 3.36 g of a component (A) was obtained.

10 (3) Polymerization

 Polymerization of Example 60 (3) was repeated, except that the solid catalyst component was changed to 9.3 mg of the component (A) obtained in the above Example 61 (2). As a result, 108.8 g of a polymer having a good particle property was obtained.

15 Polymerization activity per the solid catalyst component was found to be 11,700 g/solid catalyst component-g/hour. The value of $VH/(VT + VH)$ was larger than 0.3, and the activity was found to be high. Further, the olefin polymer obtained showed, $SCB = 14.1$, $MFR = 0.018$, $MFRR = 292$ and $SR = 1.00$.

20 [Example 62]

 (1) Production of solid product

 To the filtrate obtained in Example 61 (1), 30 ml of hexane was further added to precipitate a solid. Thus, $VH/(VT + VH) = 0.40$. The mixture was heated to 40°C, stirred for 1 hour at
25 that temperature, thereafter cooled to ambient temperature and then filtered. The solid separated by filtration was dried for

2 hours at 120°C under reduced pressure. As a result, 4.01 g of a solid product was obtained.

(2) Production of component (A)

5 Example 60 (2) was repeated, except that the solid product was changed to 1.70 g of the solid product obtained in the above Example 62 (1), and the amounts of tetrahydrofuran and the silica were changed to 10.5 ml and 1.29 g, respectively. As a result, 2.22 g of a component (A) was obtained.

(3) Polymerization

10 Polymerization of Example 60 (3) was repeated, except that the solid catalyst component was changed to 10.5 mg of the component (A) obtained in the above Example 62 (2). As a result, 87.8 g of a polymer having a good particle property was obtained. Polymerization activity per the solid catalyst component was
15 found to be 8,360 g/solid catalyst component-g/hour. The value of $VH/(VT + VH)$ was larger than 0.3, and the activity was found to be high. Further, the olefin polymer obtained showed, $SCB = 14.3$, $MFR = 0.04$, $MFRR = 158$ and $SR = 1.04$.

[Reference Example 1]

20 (1) Production of solid product

To the filtrate obtained in Example 62 (1), 60 ml of hexane was further added to precipitate a solid. Thus, $VH/(VT + VH) = 0.52$. The mixture was heated to 40°C, stirred for 1 hour at that temperature, thereafter cooled to ambient temperature and
25 then filtered. The solid separated by filtration was dried for 2 hours at 120°C under reduced pressure. As a result, 2.26 g

of a solid product was obtained.

(2) Production of component (A)

Example 60 (2) was repeated, except that the solid product was changed to 1.63 g of the solid product obtained in the above
5 Reference Example 1(1), and the amount of the silica was changed to 1.27 g. As a result, 1.86 g of a component (A) was obtained.

[0273]

(3) Polymerization

Polymerization of Example 60 (3) was repeated, except that
10 the solid catalyst component was changed to 10.2 mg of the component (A) obtained in the above Reference Example 1 (2). As a result, 40.1 g of a polymer having a good particle property was obtained. Polymerization activity per the solid catalyst component was found to be 3,930 g/solid catalyst component-g/hour.
15 The value of $VH/(VT + VH)$ was larger than 0.5, and the activity was found to be somewhat low. Further, the olefin polymer obtained showed, SCB = 14.1, MFR = 0.08, MFRR = 138 and SR = 1.07.

[Example 63]

(1) Production of contact product (e)

20 In a 5 l four necked flask purged with nitrogen, 1.5 l of tetrahydrofuran and 1.35 l (2.70 mol) of a hexane solution of diethylzinc (2.00 mol/l) were introduced and cooled using an ice bath. To the mixture, a solution prepared by dissolving 199.8 g (1.09 mol) of pentafluorophenol in 340 ml of
25 tetrahydrofuran was added over 65 minutes. After completion of dropping, the mixture was stirred for 1 hour. The temperature

was raised to 45°C over 0.5 hour, and the mixture was stirred for 1 hour at that temperature. The temperature was lowered to 20°C using an ice bath, 45.09 g (2.51 mol) of H₂O was dropped thereto over 1.5 hours. Thereafter, the mixture was stirred at
5 20°C for 1 hour, heated to 45°C and stirred for 1 hour at that temperature. After cooling to ambient temperature, a volatile component was removed by distillation, and the residue was dried at 120°C for 8 hours under reduced pressure. As a result, 425.5 g of a solid product was obtained.

10 (2) Production of component (A)

In the flask of the above (1), 3 l of tetrahydrofuran was introduced and stirred. To the mixture, 327.4 g of the same lot heat-treated silica as that used in Example 1 (2) was introduced. The mixture was heated to 40°C, and stirred for 2 hours. Thereafter,
15 the solid component was allowed to precipitate, and the mixture was filtered. As a washing operation, 3 l of tetrahydrofuran was added thereto, the mixture was stirred, and after allowing the solid component to precipitate, the mixture was filtered. Said washing operation was repeated 5 times. The residue was
20 dried for 8 hours at 120°C under reduced pressure, thereby obtaining 510.0 g of a component (A). Elementary analysis revealed Zn = 2.5 mmol/g, and F = 2.4 mmol/g. Thus, F/Zn = 0.96 (molar ratio). As a result of an extended X-ray measurement, A/B was calculated to be 0.274, provided that A is an integral
25 intensity of halo showing a peak at Bragg angle (2θ) 33° to 37° in a diffraction intensity profile and B is an integral

intensity of halo showing a peak at Bragg angle (2θ) 18° to 22° in said diffraction intensity profile.

(3) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had
5 been dried under reduced pressure and thereafter purged with
argon, was made vacuum, hydrogen was fed therein to obtain a
partial pressure of 0.012 MPa, 700 g of butane and 50 g of 1-butene
were fed therein, and then a temperature of the autoclave was
raised to 70°C . Thereafter, ethylene was fed therein to obtain
10 a partial pressure of 1.6 MPa, and the system was stabilized.
As a result of a gas chromatography analysis, a gas composition
in the system was found to be hydrogen = 0.45 mol%. To the resultant
was added 0.9 ml of a heptane solution of triisobutylaluminum
adjusted to a concentration of 1 mmol/ml. Successively, 14.3
15 mg of the component (A) obtained in the above Example 63 (2)
as a solid catalyst component was added thereto. Polymerization
was carried out for 60 minutes at 70°C while feeding an
ethylene/hydrogen mixed gas (hydrogen = 0.200 vol%) so as to
keep the whole pressure constant. As a result, 77.4 g of an olefin
20 polymer was obtained. Polymerization activity per the solid
catalyst component (the component (A) obtained in the above
Example 63 (2)) was found to be 5,410 g/g-solid catalyst
component/hour. Further, the obtained olefin polymer showed,
SCB = 15.5, MFR = 0.35, MFRR = 118 and SR = 1.15.

25 [Example 64]

(1) Production of contact product (e)

In a 300 ml four necked flask purged with nitrogen, 106 ml of tetrahydrofuran and 40 ml (80.0 mmol) of a hexane solution of diethylzinc (2.00 mol/l) were introduced and cooled using an ice bath. To the mixture, a 14.5 ml (32.0 mmol) of a
5 tetrahydrofuran solution of pentafluorophenol (2.21 mol/l) was added over 50 minutes. After completion of dropping, the mixture was stirred at 5°C for 1 hour. The temperature was raised to 20°C, and 8.9 ml (88.1 mmol) of a tetrahydrofuran solution of H₂O (9.90 mol/l) was dropped thereto over 1 hour. Thereafter,
10 the mixture was stirred for 2 hours, and then allowed to stand at ambient temperature for 15.3 hours. The resulting mixture was refluxed at 63°C and cooled to ambient temperature, thereby obtaining 146 ml of a yellow solution.

(2) Production of component (A)

15 In a 50 ml four necked flask purged with nitrogen, 3.12 g of the same lot heat-treated silica as that used in Example 3 (2) (Sylopol 948, manufactured by Davison Co., Ltd.; average particle diameter = 55 μ m; pore volume = 1.66 ml/g; specific surface area = 309 m²/g) was introduced, and then 50 ml of the
20 yellow solution obtained in the above Example 64 (1) was added thereto. The mixture was heated to 40°C, and stirred for 2 hours. Thereafter, the solid component was allowed to precipitate, and the mixture was filtered. As a washing operation, 30 ml of tetrahydrofuran was added thereto, the mixture was stirred, and
25 after allowing the solid component to precipitate, the mixture was filtered. Said washing operation was repeated 5 times. The

residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 5.60 g of a component (A). Elementary analysis revealed Zn = 3.7 mmol/g, and F = 4.1 mmol/g. Thus, F/Zn = 1.1 (molar ratio). As a result of an extended X-ray measurement, 5 A/B was calculated to be 0.696, provided that A is an integral intensity of halo showing a peak at Bragg angle (2θ) 33° to 37° in a diffraction intensity profile and B is an integral intensity of halo showing a peak at Bragg angle (2θ) 18° to 22° in said diffraction intensity profile. Further, as a result 10 of EXAFS, the peak intensity C of the maximum peak present within a range of 1 to 2 nm in a radial distribution function obtained according to a Fourier transform was 14.2, and the peak intensity D of the maximum peak present within a range of 2.5 to 3.5 nm in said radial distribution function was 10.1, and thus, a peak 15 intensity ratio, D/C = 0.71.

(3) Polymerization

Example 63 (3) was repeated, except that the solid catalyst component was changed to 13.6 mg of the component (A) obtained in Example 64 (2), and a hydrogen concentration of the 20 ethylene/hydrogen mixed gas to keep the whole pressure of the autoclave constant was changed to 0.199 vol%. As a result, 72.3 g of an olefin polymer was obtained. Polymerization activity per the solid catalyst component (the component (A) obtained in the above Example 64 (2)) was found to be 5,320 g/g-solid 25 catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 15.5, MFR = 1.57, MFRR = 58.2 and SR = 1.30.

[Example 65]

(1) Production of contact product (e)

In a 5 l four necked flask purged with nitrogen, 1.5 l of tetrahydrofuran and 1.35 l (2.70 mol) of a hexane solution of diethylzinc (2.00 mol/l) were introduced and cooled using an ice bath. To the mixture, a solution prepared by dissolving 197.3 g (1.07 mol) of pentafluorophenol in 380 ml of tetrahydrofuran was added over 65 minutes. After completion of dropping, the mixture was stirred for 1 hour. The temperature was raised to 45°C over 0.5 hour, and the mixture was stirred for 1 hour at that temperature. After lowering the temperature to 20°C using an ice bath, 44.0 g (2.44 mol) of H₂O was dropped thereto over 1.5 hours. Thereafter, the mixture was stirred for 1 hour at 20°C, thereafter heated to 45°C and then stirred for 1 hour at that temperature. After cooling to ambient temperature, a volatile component was removed by distillation, and the residue was dried for 8 hours at 120°C under reduced pressure. As a result, 423.3 g of a solid product was obtained.

[0274]

(2) Production of component (A)

In the flask of the above Example 65 (1), 3 l of tetrahydrofuran was introduced and stirred, and 324.6 g of the same lot heat-treated silica as that used in Example 1 (2) was introduced. The mixture was heated to 40°C, and stirred for 2 hours. Thereafter, the solid component was allowed to precipitate, and the mixture was filtered. As a washing operation, 3 l of

tetrahydrofuran was added thereto, the mixture was stirred, and after allowing the solid component to precipitate, the mixture was filtered. Said washing operation was repeated 5 times. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 515.6 g of a component (A). Elementary analysis revealed Zn = 2.6 mmol/g, and F = 3.2 mmol/g. Thus, F/Zn = 1.2 (molar ratio). As a result of EXAFS, the peak intensity C of the maximum peak present within a range of 1 to 2 nm in a radial distribution function obtained according to a Fourier transform was 13.3, and the peak intensity D of the maximum peak present within a range of 2.5 to 3.5 nm in said radial distribution function was 8.51, and thus, a peak intensity ratio, D/C = 0.64.

(3) Polymerization

Example 63 (3) was repeated, except that the solid catalyst component was changed to 13.1 mg of the component (A) obtained in Example 65 (2), and a hydrogen concentration of the ethylene/hydrogen mixed gas to keep the whole pressure of the autoclave constant was changed to 0.199 vol%. As a result, 71.5 g of an olefin polymer was obtained. Polymerization activity per the solid catalyst component (the component (A) obtained in the above Example 65 (2)) was found to be 5,460 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 14.3, MFR = 0.13, MFRR = 126 and SR = 1.10.

[Example 66]

(1) Production of contact product (e)

In a 300 ml four necked flask purged with nitrogen, 100

ml of tetrahydrofuran and 39.6 ml (80.0 mmol) of a hexane solution of diethylzinc (2.02 mol/l) were introduced and cooled using an ice bath. To the mixture, a 14.5 ml (32.0 mmol) of a tetrahydrofuran solution of pentafluorophenol (2.21 mol/l) was added over 50 minutes. After completion of dropping, the mixture was stirred at 5°C for 1 hour. The temperature was raised to 20°C, and 9.7 ml (96.0 mmol) of a tetrahydrofuran solution of H₂O (9.90 mol/l) was dropped thereto over 1 hour. Thereafter, the mixture was stirred for 2 hours, and then allowed to stand at ambient temperature for 15.3 hours. The resulting mixture was refluxed at 63°C and cooled to ambient temperature, thereby obtaining a yellowish white slurry.

(2) Production of component (A)

In a 50 ml four necked flask purged with nitrogen, 2.88 g of the same lot heat-treated silica as that used in Example 3 (2) was introduced, and then 50 ml of the yellowish white slurry obtained in the above Example 66 (1) was added thereto. The mixture was heated to 40°C, and stirred for 2 hours. Thereafter, the solid component was allowed to precipitate, and the mixture was filtered. As a washing operation, 30 ml of tetrahydrofuran was added thereto, the mixture was stirred, and after allowing the solid component to precipitate, the mixture was filtered. Said washing operation was repeated 5 times. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 3.00 g of a component (A). Elementary analysis revealed Zn = 2.8 mmol/g, and F = 1.9 mmol/g. Thus, F/Zn = 0.68 (molar ratio).

As a result of an extended X-ray measurement, A/B was calculated to be 0.472. Further, as a result of EXAFS, the peak intensity C was 14.0, and the peak intensity D was 14.4, and thus, a peak intensity ratio, $D/C = 1.03$.

5 (3) Polymerization

Example 63 (3) was repeated, except that the solid catalyst component was changed to 12.4 mg of the component (A) obtained in Example 66 (2), and a hydrogen concentration of the ethylene/hydrogen mixed gas to keep the whole pressure of the autoclave constant was changed to 0.186 vol%. As a result, 29.4 g of an olefin polymer was obtained. Polymerization activity per the solid catalyst component (the component (A) obtained in the above Comparative Example 4 (2)) was found to be 2,370 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 14.7, MFR = 1.63, MFRR = 52.1 and SR = 1.26.

[Example 67]

(1) Polymerization of propylene

20 In a 3 l-volume stainless steel autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, 780 g of propylene was fed and heated to 40°C. In a different vessel, 145.5 mg of the modified particle obtained in the above Example 1 (2) as the catalyst component (A), 1.5 ml of a toluene solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml
25 as the catalyst component (C), and 5 ml of a toluene solution

of racemic dimethylsilylenebis(2-methyl-1-indenyl)zirconium dichloride adjusted to a concentration of $2 \mu\text{mol/ml}$ as the catalyst component (B) had been collectively mixed. The resulting mixture was fed into the above-mentioned autoclave, and
5 polymerization was carried out at 40°C . After 60 minutes, the unreacted monomer was released to stop the polymerization. The resulting polymer was dried at 60°C for 5 hours under reduced pressure, thereby obtaining 282 g of a polypropylene powder.

Accordingly, yield of the polypropylene per mol of the
10 transition metal compound and per hour of the polymerization time (hereinafter referred to as g-PP/mol-Zr·Hr) was 2.7×10^7 , a bulk density of the obtained propylene polymer was 0.44 (g/ml), and stereoregularity thereof mmmm% was 0.941. Polymerization results were as shown in Table 1.

15 [Example 68]

(1) Polymerization of propylene

Example 67 (1) was repeated to carry out the polymerization of propylene, except that the amounts of the modified particle (catalyst component (A)) and the toluene solution as the catalyst
20 component (B) were changed to 155.3 mg and 0.9 ml, respectively, and the polymerization time was changed to 30 minutes. Polymerization results were as shown in Table 1.

[Example 69]

(1) Polymerization of propylene

25 In a 3 l-volume stainless steel autoclave equipped with a stirrer, which had been dried under reduced pressure and

thereafter purged with argon, 500 g of butane was fed and heated to 40°C. In a different vessel, 304.7 mg of the modified particle obtained in the above Example 1 (2) as the catalyst component (A), 1.5 ml of a toluene solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml as the catalyst component (C), and 12 ml of a toluene solution of racemic dimethylsilylenebis(2-methyl-1-indenyl)zirconium dichloride adjusted to a concentration of 2 μmol/ml as the catalyst component (B) had been collectively mixed. The resulting mixture was fed into the above-mentioned autoclave, and polymerization was carried out at 40°C while keeping inner pressure 0.7 MPa using propylene. After 60 minutes, the unreacted monomer was released. The resulting polymer was dried at 60°C for 5 hours under reduced pressure, thereby obtaining 49 g of a polypropylene powder. Polymerization results were as shown in Table 1.

[Reference Example 2]

(1) Polymerization of propylene

In a 3 l-volume stainless steel autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, a toluene slurry prepared by adding 304.4 mg of the modified particle obtained in the above Example 1 (2) as the catalyst component (A) to 10 ml of toluene was introduced, thereafter 780 g of propylene was fed, and a temperature was raised to 40°C. While, 2.0 ml of a toluene solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml as the catalyst component (C) and 8.9 ml of a toluene solution

of dimethylsilylenebis(2-methyl-1-indenyl)zirconium
dichloride adjusted to a concentration of $2\ \mu\text{mol/ml}$ as the
catalyst component (B) had been contacted with each other in
a different stainless steel vessel. The resultant was introduced
5 in the foregoing autoclave, and polymerization was carried out
at 40°C . After 60 minutes, the unreacted monomer was released
to stop the polymerization. The resulting polymer was dried at
 60°C for 5 hours under reduced pressure, thereby obtaining 0.6
g of a polypropylene powder. Polymerization results were as shown
10 in Table 1.

Table 1

	Polymerization activity (g-PP/mol-Zr·hr)	Bulk density (g/ml)	[mmmm]	Mw	Mw/Mn
Example 67	2.7×10^7	0.44	0.942	214,000	1.9
Example 68	5.1×10^8	0.42	0.931	186,000	1.9
Example 69	2.7×10^6	0.41	0.925	173,000	2.0
Reference Example 2	3.4×10^4				

[Example 70]

(1) Production of transition metal compound

5 In a 3 l four necked flask purged with argon, 55.3 g (588 mmol) of phenol and 1200 ml of toluene were introduced and stirred. After cooling to 5°C, 377 ml (588 mmol) of a hexane solution of n-BuLi (1.56 mol/l) was gradually dropped thereto. The content turned to white slurry. After completion of dropping, 700 ml

10 of toluene was added thereto. Then, 110 g (263 mmol) of racemic ethylenebis(1-indenyl)zirconiumdichloride was gradually added thereto, and the mixture was stirred at 5°C for 1 hour. Thereafter, the mixture was stirred for 3 hours under a reflux condition. It was observed that a white solid was precipitated. After

15 allowing the mixture to stand over night, it was observed that a white solid was precipitated downward. Using a syringe, a supernatant yellow transparent liquid was transferred to a flask purged with argon and concentrated, thereby precipitating a yellow solid. Hexane was added thereto, thereby further

20 precipitating the yellow solid. After allowing the mixture to stand in a refrigerator, the yellow solid was separated out,

thereby obtaining 46.8 g of racemic ethylenebis(indenyl)zirconium diphenoxide as a first crystal.

After completion of the reaction, dichloromethane was added to the white solid precipitated and separated. The mixture
5 was filtered using sellalite. The filtrate was concentrated, thereby precipitating a yellow solid. Hexane was added thereto, thereby further precipitating the yellow solid. After allowing the mixture to stand in a refrigerator, the yellow solid was separated out, thereby obtaining 59.3 g of racemic
10 ethylenebis(1-indenyl)zirconium diphenoxide as a second crystal. A total amount combined with the first crystal was 106 g (199 mmol). Yield 76%.

(1) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had
15 been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.018 MPa, 700 g of butane and 50 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain
20 a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.67 mol% and 1-butene = 2.37 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml.
25 Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium diphenoxide adjusted to a

concentration of 2 μ mol/ml, and then 15.6 mg of the component (A) obtained in the above Example 28 (2) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.31 mol%) so as to keep the whole pressure constant. As a result, 75 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.5×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 4810 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 13.2, MFR = 0.43, MFRR = 109, SR = 1.21, $M_w = 1.12 \times 10^5$, $M_w/M_n = 10.4$, and $[\eta] = 1.35$.

[Example 71]

(1) Polymerization A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.028 MPa, 700 g of butane and 50 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.90 mol% and 1-butene = 2.80 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.25 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium

diphenoxide adjusted to a concentration of 2 μ mol/ml, and then 15.8 mg of the component (A) obtained in the above Example 28 (2) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.31 mol%) so as to keep the whole pressure constant. As a result, 67 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 1.3×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 4240 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 16.1, MFR = 1.54, MFRR = 76.9, SR = 1.34, $M_w = 8.90 \times 10^4$, $M_w/M_n = 9.5$, and $[\eta] = 1.16$.

[Example 72]

(1) Production of contact product (e)

In a 5 l four necked flask purged with nitrogen, 1.5 l of tetrahydrofuran and 1.35 l (2.7 mol) of a hexane solution of diethylzinc (2.01 mol/l) were introduced and cooled to 5°C. To the mixture, a solution prepared by dissolving 197.3 g (1.07 mol) of pentafluorophenol in 380 ml of tetrahydrofuran was added over 55 minutes. After completion of dropping, the mixture was stirred at 5°C for 1 hour. Thereafter, the temperature was raised to 45°C, and the mixture was stirred for 1 hour. After lowering the temperature to 20°C using an ice bath, 44.0 g (2.44 mol) of H₂O was dropped thereto over 1.5 hours. As a result, the mixture turned to yellowish white slurry. After completion of dropping,

the mixture was stirred for 1 hour, thereafter heated to 45°C and then stirred for 1 hour. After allowing the mixture to stand over night at ambient temperature, a volatile component was removed by distillation, and the residue was dried for 8 hours
5 at 120°C under reduced pressure. As a result, 423 g of a solid product was obtained.

(2) Production of component (A)

In a 5 l four necked flask purged with nitrogen, 423 g of the solid product obtained in the above Example 72 (1), and
10 3 l of tetrahydrofuran were introduced and stirred. To the mixture, 325 g of the same lot heat-treated silica as that used in Example 1 (2) was introduced. The mixture was heated to 40°C, and stirred for 2 hours. Thereafter, the solid component was allowed to precipitate, and a slurry component of the upper layer was removed.
15 The remaining liquid component was filtered using a glass filter. As a washing operation, 3 l of tetrahydrofuran was added thereto, the mixture was stirred, after allowing the solid component to precipitate, a slurry component of the upper layer was removed, and the remaining liquid component was filtered using a glass
20 filter. Said washing operation was repeated 5 times. The residue was dried for 8 hours at 120°C under reduced pressure, thereby obtaining 516 g of a component (A). Elementary analysis revealed Zn = 2.6 mmol/g, and F = 3.2 mmol/g.

(3) Pre-polymerization

25 A 0.4 l-volume autoclave equipped with a stirrer was made vacuum, hydrogen was fed therein to obtain a pressure of 0.005

MPa at ambient temperature, 90 g of butane was fed therein as a solvent, and then a temperature of the autoclave was raised to 50°C. Further, ethylene was fed therein to obtain a partial pressure of 0.5 MPa, and the system was stabilized. As a result of gas chromatography, a gas composition in the system was found to be hydrogen = 0.12 mol%. Then, 0.2 ml of a hexane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml was introduced therein. Successively, 1 ml of a hexane solution of racemic ethylenebis(1-indenyl)zirconium diphenoxide and triisobutylaluminum adjusted to a zirconium concentration and an aluminum concentration to 50 μ mol/ml and 0.5 mmol/ml, respectively, was added thereto, and then 0.42 g of the component (A) obtained in the above Example 72 (2) was introduced therein to initiate polymerization. During the polymerization, a mixed gas of ethylene and hydrogen (hydrogen concentration = 0.28 mol%) was continuously supplied while adjusting so as to keep the gas phase pressure in the autoclave to 1.0 MPa. After 6.3 hours, ethylene, butane and hydrogen gas were released, and a solid remaining in the autoclave was dried for 30 minutes at ambient temperature under reduced pressure. As a result, a polyethylene-pre-polymerized catalyst component for addition polymerization was obtained in a yield of 102 g per g of the component (A). MFR of said polyethylene was 2.0 g/10 min.

(4) Real polymerization

A 5 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with

argon, was made vacuum, hydrogen was fed at 10°C therein to obtain a partial pressure of 0.018 MPa, 1066 g of butane and 200 ml of 1-hexene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.98 mol%. To the resultant was added 1.5 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 2.17 g of the pre-polymerized catalyst component for addition polymerization obtained in the above Example 72(3) was added thereto. Polymerization was continued for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.69 mol%) so as to keep the whole pressure constant. As a result, 129 g of an olefin polymer having a good particle property was obtained. Polymerization activity per the solid catalyst component (the component (A) obtained in the above Example 72 (2)) was found to be 6100 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, MFR = 3.52, MFRR = 53, and SR = 1.52.

[Example 73]

(1) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.015 MPa, 690 g of butane and 60 g of 1-butene

were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.75 mol% and 1-butene = 2.76 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium diphenoxide adjusted to a concentration of 2 μmol/ml, and then 10.0 mg of the component (A) obtained in the above Example 52 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.23 mol%) so as to keep the whole pressure constant. As a result, 89 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 5.9×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 8900 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 15.7, MFR = 0.14, MFRR = 118 and SR = 1.05.

[Example 74]

(1) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.015 MPa, 690 g of butane and 60 g of 1-butene

were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.67 mol% and 1-butene = 3.29 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium diphenoxide adjusted to a concentration of 2 μmol/ml, and then 8.3 mg of the component (A) obtained in the above Example 54 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.25 mol%) so as to keep the whole pressure constant. As a result, 133 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium atom was found to be 8.9×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 16000 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 17.7, MFR = 2.50, MFRR = 45.9 and SR = 1.33.

[Example 75]

(3) Pre-polymerization

A 0.4 l-volume autoclave equipped with a stirrer was made vacuum, hydrogen was fed therein under pressure at ambient temperature to obtain a pressure of 0.004 MPa, 90 g of butane

was fed therein as a solvent, and then a temperature of the autoclave was raised to 50°C. Further, ethylene was fed therein to obtain a partial pressure of 0.15 MPa, and the system was stabilized. As a result of gas chromatography, a gas composition
5 in the system was found to be hydrogen = 0.15 mol%. Then, 0.2 ml of a hexane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml was introduced therein. Successively,
2 ml of a hexane solution of racemic ethylenebis(1-indenyl)zirconium diphenoxide and
10 triisobutylaluminum adjusted to a zirconium concentration and an aluminum concentration to 150 μ mol/ml and 1.0 mmol/ml, respectively, was added thereto, and then 2.12 g of the component (A) obtained in the above Example 54 (1) was introduced therein to initiate polymerization. During the polymerization, a mixed
15 gas of ethylene and hydrogen (hydrogen concentration = 0.19 mol%) was continuously supplied while adjusting so as to keep the gas phase pressure in the autoclave to 0.8 MPa. After 3.4 hours, ethylene, butane and hydrogen gas were released, and a solid remaining in the autoclave was dried for 30 minutes at ambient
20 temperature under reduced pressure. As a result, a polyethylene-pre-polymerized catalyst component for addition polymerization was obtained in a yield of 20.8 g per g of the component (A).

(4) Real polymerization

25 A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with

argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.015 MPa, 690 g of butane and 60 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.72 mol% and 1-butene = 3.09 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 227 mg of the pre-polymerized catalyst component for addition polymerization obtained in the above Example 75(1) was added thereto. Polymerization was continued for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.26 mol%) so as to keep the whole pressure constant. As a result, 100 g of an olefin polymer having a good particle property was obtained. Polymerization activity per the solid catalyst component (the component (A) obtained in the above Example 54 (1)) was found to be 9170 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 19.3, MFR = 5.37, MFRR = 33.8 and SR = 1.47.

[Example 76]

(1) Production of component (A)

In a 5 l four necked flask purged with nitrogen, 3.0 l of toluene, 43 ml of tetrahydrofuran and 0.522 l (1.05 mol) of a hexane solution of diethylzinc (2.00 mol/l) were introduced and cooled to 5°C. To the mixture, a solution prepared by

dissolving 98.1 g (0.53 mol) of pentafluorophenol in 160 ml of toluene was dropped over 60 minutes. After completion of dropping, the mixture was stirred for 1 hour at 5°C, thereafter heated to 40°C, and stirred for 1 hour. After cooling to 5°C using an ice bath, 350 g of the same lot heat-treated silica as that used in Example 40 (2) was introduced thereto, and then 0.1 l of toluene was poured therein so as to wash away the silica adhering to a wall of the flask. After stirring the mixture for 5 minutes, 14.4 g (0.80 mol) of H₂O was added thereto over 1.5 hours. After completion of the dropping, the mixture was stirred at 5°C for 1.5 hours, and at 40°C for 2 hours. Thereafter, the mixture was allowed to stand over night at ambient temperature. Then, the mixture was stirred for 2 hours at 80°C, and thereafter allowed to stand, thereby precipitating a solid component. At the time when an interface between a layer of the precipitated solid component and an upper layer of a slurry portion appeared, the upper slurry portion was removed. The remaining liquid component was removed by filtration. Thereafter, 3.5 l of toluene was added thereto, and the mixture was stirred for 2 hours at 95°C. After that, washing of the solid component was repeated 4 times with each 3.5 l of toluene at 95°C and 2 times with each 3.5 l of hexane at ambient temperature in a manner such that the mixture was allowed to stand, thereby precipitating a solid component, and the upper slurry portion was removed at the time when the interface between a layer of the precipitated solid component and an upper layer of a slurry portion appeared. Thereafter,

the remaining liquid component was removed by filtration. Thereafter, the solid component was dried for 4 hours at 120°C under reduced pressure, thereby obtaining 516 g of a component (A). Elementary analysis revealed Zn = 2.0 mmol/g and F = 4.3 mmol/g.

(2) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.015 MPa, 690 g of butane and 60 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.72 mol% and 1-butene = 3.21 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium diphenoxide adjusted to a concentration of 2 μ mol/ml, and then 6.2 mg of the component (A) obtained in the above Example 76 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.25 mol%) so as to keep the whole pressure constant. As a result, 81 g of an olefin polymer having a good particle property was obtained. Polymerization activity per zirconium

atom was found to be 5.4×10^7 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 13100 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 16.1, MFR = 0.92, MFRR = 63.0 and SR = 1.23.

[Example 77]

(1) Polymerization

A 3 l-volume autoclave equipped with a stirrer, which had been dried under reduced pressure and thereafter purged with argon, was made vacuum, hydrogen was fed therein to obtain a partial pressure of 0.015 MPa, 690 g of butane and 60 g of 1-butene were fed therein, and then a temperature of the autoclave was raised to 70°C. Thereafter, ethylene was fed therein to obtain a partial pressure of 1.6 MPa, and the system was stabilized. As a result of a gas chromatography analysis, a gas composition in the system was found to be hydrogen = 0.73 mol% and 1-butene = 3.38 mol%. To the resultant was added 0.9 ml of a heptane solution of triisobutylaluminum adjusted to a concentration of 1 mmol/ml. Successively, 0.75 ml of a toluene solution of racemic ethylenebis(1-indenyl)zirconium diphenoxide adjusted to a concentration of 2 μ mol/ml, and then 8.2 mg of the component (A) obtained in the above Example 55 (1) as a solid catalyst component were added thereto. Polymerization was carried out for 60 minutes at 70°C while feeding an ethylene/hydrogen mixed gas (hydrogen = 0.25 mol%) so as to keep the whole pressure constant. As a result, 163 g of an olefin polymer having a good particle

property was obtained. Polymerization activity per zirconium atom was found to be 1.1×10^8 g/mol-Zr/hour, and polymerization activity per the solid catalyst component was found to be 19900 g/g-solid catalyst component/hour. Further, the obtained olefin polymer showed, SCB = 16.2, MFR = 1.37, MFRR = 51.8 and SR = 1.29.

Industrial Applicability

As mentioned above in detail, according to the present invention, there are provided a particle, a carrier and a catalyst component for addition polymerization, which are used for addition polymerization in combination with a transition metal compound capable of forming a single site catalyst, thereby obtaining a film having a good appearance, a process for producing said particle, a pre-polymerized catalyst component for addition polymerization prepared by using said particle, an addition polymerization catalyst prepared by using said particle, and a process for producing an addition polymer comprising using said addition polymerization catalyst. Further, according to the present invention, there are provided a particle, a carrier and a catalyst component for addition polymerization, which are useful for the production of an addition polymerization catalyst, which can afford an addition polymer superior in its shape and particle property when used for polymerization accompanied by the formation of addition polymer particles (for example, slurry polymerization, gas phase polymerization or bulk

polymerization), a process for producing said particle, a pre-polymerized catalyst component for addition polymerization prepared by using said particle, an addition polymerization catalyst prepared by using said particle, and a process for
5 producing an addition polymer superior in its particle property, which comprises using said addition polymerization catalyst for polymerization accompanied by the formation of addition polymer particles (for example, slurry polymerization, gas phase polymerization or bulk polymerization). Still further,
10 according to the present invention, there are provided an addition polymerization catalyst having a high activity and a process for producing an addition polymer efficiently. Furthermore, there is provided a process for producing a stereoregular α -olefin polymer much more efficiently.

15 Incidentally, the single site catalyst mentioned here is conceptually distinguished from a multiple site solid catalyst. The catalyst includes not only a single site catalyst of a restricted sense capable of affording an addition polymer having a narrow molecular weight distribution or an addition polymer
20 having a narrow composition distribution in the case of a copolymer, but also a catalyst capable of affording an addition polymer having a wide molecular weight distribution or an addition polymer having a wide composition distribution in the case of a copolymer, as far as the latter catalyst can be obtained
25 according to a process similar to a process for producing the single site catalyst of the restricted sense.

Claims

1. A modified particle obtained by contacting the following (a), the following (b), the following (c) and a particle (d) with one another,

(a): a compound represented by the following general formula [1],



(b): a compound represented by the following general formula [2],



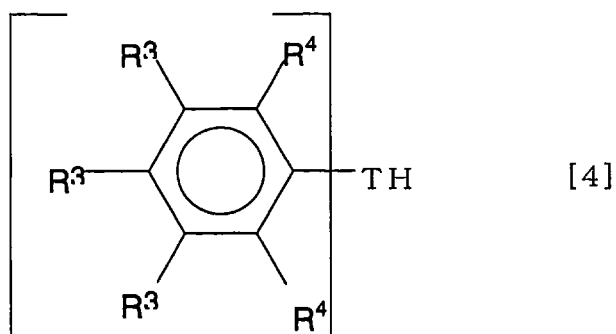
(c): a compound represented by the following general formula [3],



in which general formulas [1] to [3], M^1 is a typical metal atom of the Group 1, 2, 12, 14 or 15 of the periodic table, m is a valence of M^1 , L^1 is a hydrogen atom, a halogen atom or a hydrocarbon group, when more than one L^1 exists, they may be the same or different from one another, R^1 is an electron-withdrawing group or an electron-withdrawing group-containing group, when more than one R^1 exists, they may be the same or different from one another, R^2 is a hydrocarbon group or a halogenated hydrocarbon group, T in each compound is independently of each other a non-metal atom of the Group 15 or 16 of the periodic table, and t is a valence of T in each compound.

2. The modified particle according to Claim 1, wherein

the compound of (b) is a compound represented by the following general formula [4],



5 wherein R³ is an electron-withdrawing group or an electron-withdrawing group-containing group, three R³'s may be the same or different from one another, optional two of R³ may be combined with each other at adjacent positions of the benzene ring to form a condensed ring structure, R⁴ is a hydrogen atom
10 or a hydrocarbon group, and T is as defined above.

3. The modified particle according to Claim 1 or 2, wherein the particle (d) is a porous substance.

4. The modified particle according to any of Claims 1 to 3, wherein the particle (d) is an inorganic substance
15 heat-treated at 100 to 1500°C for 10 minutes to 50 hours.

5. The modified particle according to Claims 4, wherein the particle (d) is silica heat-treated at 200 to 800°C for 1 to 30 hours.

6. The modified particle according to any of Claims
20 1 to 5, which satisfies the following conditions of [5] and [6],

$$N/M > 0.9 \quad [5]$$

wherein N is a substance amount of a halogen atom contained in the modified particle, and M is a substance amount of a typical metal atom M^1 contained in the modified particle, and

$$A/B \geq 0.1 \quad [6]$$

5 wherein A is an integral intensity of halo showing a peak at Bragg angle (2θ) 33° to 37° in a diffraction intensity profile obtained by an extended X-ray measurement, and B is an integral intensity of halo showing a peak at Bragg angle (2θ) 18° to 22° in said diffraction intensity profile.

10 7. The modified particle according to any of Claims 1 to 6, which satisfies the following conditions of [5] and [7],

$$N/M > 0.9 \quad [5]$$

wherein N is a substance amount of a halogen atom contained in the modified particle, and M is a substance amount of a typical
15 metal atom M^1 contained in the modified particle, and

$$D/C \geq 0.5 \quad [7]$$

wherein C is a peak intensity of the maximum peak present within a range of 1 to 2 nm in a radial distribution function obtained by measuring the modified particle according to an X-ray
20 absorption fine structure analysis, and D is a peak intensity of the maximum peak present within a range of 2.5 to 3.5 nm in said radial distribution function, provided that the radial distribution function is obtained in a manner such that the modified particle is measured according to an X-ray absorption
25 fine structure analysis (XAFS) to obtain X-ray absorption spectra, from which extended X-ray absorption fine structure (EXAFS)

spectra of the typical metal atom M^1 in a K absorption edge are found and processed according to a Fourier transform.

8. The modified particle according to any of Claims 1 to 7, which is obtained by contacting (a), (b) and (c) with one another to obtain a contact product (e) and contacting the contact product (e) and the particle (d) with each other.

9. The modified particle according to any of Claims 1 to 8, wherein the contact product (e) is obtained by contacting (a) and (b) with each other and contacting the resulting contact product and (c) with each other, or by contacting (a) and (c) with each other and contacting the resulting contact product and (b) with each other.

10. The modified particle according to any of Claims 1 to 9, which is obtained by contacting a typical metal component and (d) with each other, wherein the typical metal component is selected from the contact product (e) obtained by contacting (a), (b) and (c) with one another, and is insoluble in the following solvent (f),

solvent (f): a mixed solvent or tetrahydrofuran, wherein the mixed solvent is a mixture of tetrahydrofuran and hexane and satisfies the following expression [8],

$$0.5 \geq V_H / (V_T + V_H) \quad [8]$$

wherein V_T is a volume of tetrahydrofuran, and V_H is a volume of hexane.

11. The modified particle according to any of Claims 1 to 10, wherein respective amounts to be used of the compounds

(a), (b) and (c) satisfy the following expressions (1) and (2),

$$|m-y-2z| \leq 1 \quad (1)$$

$$2 \leq z/y < 3 \quad (2)$$

wherein m in the above expression (1) is a valence of M^1 , and
5 y and z in the above expressions (1) and (2) are defined, so
that a molar ratio of the amounts of respective compounds is
expressed by (a): (b): (c) = 1: y: z.

12. A carrier comprising the modified particle
according to any of Claims 1 to 11.

10 13. A catalyst component for addition polymerization,
which comprises the modified particle according to any of Claims
1 to 11.

14. A catalyst for addition polymerization, which is
obtained by contacting the modified particle (A) according to
15 any of Claims 1 to 11 and a transition metal compound (B) of
the Groups 3 to 11 or a lanthanoide series with each other.

15. A catalyst for addition polymerization, which is
obtained by contacting the modified particle (A) according to
any of Claims 1 to 11, a transition metal compound (B) of the
20 Groups 3 to 11 or a lanthanoide series and an organoaluminum
compound (C) with one another.

16. The catalyst for addition polymerization according
to Claim 14 or 15, wherein the transition metal compound (B)
of the Groups 3 to 11 or a lanthanoide series is a transition
25 metal compound having at least one cyclopentadiene type anion
skeleton-carrying group.

17. The catalyst for addition polymerization according to Claim 16, wherein the transition metal compound (B) of the Groups 3 to 11 or a lanthanoide series is a transition metal compound represented by a general formula [10] or its μ -oxo type transition metal compound dimer,



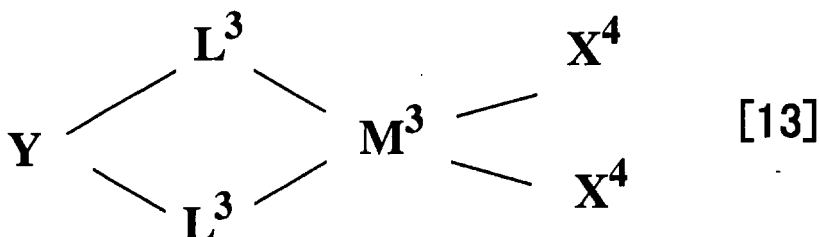
wherein M^2 is a transition metal atom of the Groups 3 to 11 or lanthanoide series, L^2 is a cyclopentadienyl type anion skeleton-carrying group or a hetero atom-containing group, and more than one L^2 may be linked directly or through a residual group containing a carbon atom, a silicone atom, a nitrogen atom, an oxygen atom, a sulfur atom or a phosphorus atom, X is a halogen atom, a hydrocarbon group (excluding the cyclopentadienyl type anion skeleton-carrying group) or $-OR^{11}$, in which R^{11} is a hydrocarbon group or a halogenated hydrocarbon group, and more than one R^{11} may be the same or different from each other, a is a number satisfying $0 < a \leq 8$, and b is a number satisfying $0 < b \leq 8$.

18. The catalyst for addition polymerization according to Claim 17, wherein X is OR^{11} .

19. The catalyst for addition polymerization according to Claim 14 or 15, wherein the transition metal compound (B) of the Groups 3 to 11 or a lanthanoide series is a transition metal compound (B) having a stereoregular polymerization ability of an α -olefin.

20. The catalyst for addition polymerization according to Claim 19, wherein the transition metal compound (B) having

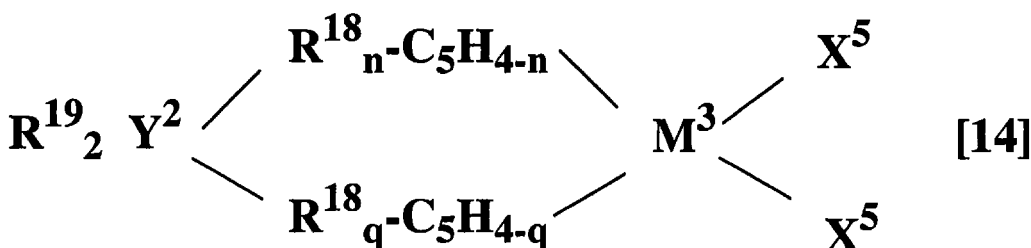
a stereoregular polymerization ability of an α -olefin is a transition metal compound represented by the following general formula [13] or [14],



5

wherein M^3 is a transition metal atom of the Groups 3 to 11 or lanthanoide series, L^3 is an η^5 -indenyl group or a substituted η^5 -indenyl group, and both of L^3 s may be the same or different from each other, Y is a bridging group for linking both of L^3 s, and both of X^4 s are independently of each other a hydrogen atom, a halogen atom, an alkyl group, an aralkyl group, an aryl group, a substituted silyl group, an alkoxy group, an aralkyloxy group, an aryloxy group or a heterocyclic group,

10



15

wherein M^3 is a transition metal atom of the Groups 3 to 11 or lanthanoide series, Y^2 is a silicon atom, a germanium atom or a tin atom, the $(R^{18}_n\text{-C}_5\text{H}_{4-n})$ and $(R^{18}_q\text{-C}_5\text{H}_{4-q})$ are each a substituted η^5 -cyclopentadienyl group, n and q are each an

20

integer of 1 to 3, and both of R^{18} s may be the same or different from each other and are independently of each other a halogen atom, an alkyl group, an aralkyl group, an aryl group, a substituted silyl group, an alkoxy group, an aralkyloxy group, an aryloxy group or a heterocyclic group, the position and/or kind of each R^{18} in each substituted η^5 -cyclopentadienyl group are (is) selected, so that a symmetry plane containing M^3 does not exist, R^{19} and X^5 are independently of each other a hydrogen atom, a halogen atom, an alkyl group, an aralkyl group, an aryl group, a substituted silyl group, an alkoxy group, an aralkyloxy group, an aryloxy group or a heterocyclic group, and R^{19} and X^5 may be all the same or different from each other.

21. A pre-polymerized catalyst component for addition polymerization, which is obtained by pre-polymerizing an olefin in the presence of a catalyst for addition polymerization, wherein the catalyst is obtained by contacting the modified particle (A) according to any of Claims 1 to 11, a transition metal compound (B) of the Groups 3 to 11 or a lanthanoide series and an organoaluminum compound (C) with one another.

22. The pre-polymerized catalyst component for addition polymerization according to Claim 21, wherein the transition metal compound (B) of the Groups 3 to 11 or a lanthanoide series is a transition metal compound having at least one cyclopentadiene type anion skeleton-carrying group.

23. The pre-polymerized catalyst component for addition polymerization according to Claim 21, wherein the

transition metal compound (B) of the Groups 3 to 11 or a lanthanoid series is a transition metal compound represented by a general formula [10] or its μ -oxo type transition metal compound dimer,



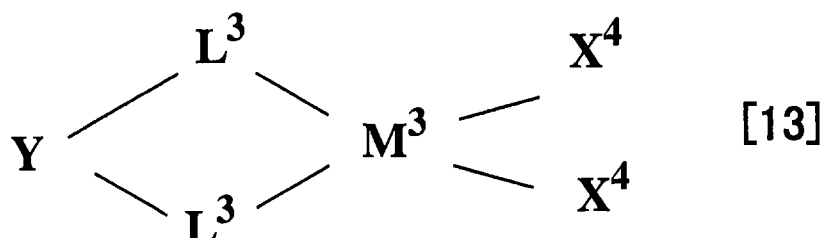
5 wherein M^2 is a transition metal atom of the Groups 3 to 11 or lanthanide series, L^2 is a cyclopentadienyl type anion skeleton-carrying group or a hetero atom-containing group, and more than one L^2 may be linked directly or through a residual group containing a carbon atom, a silicone atom, a nitrogen atom,
10 an oxygen atom, a sulfur atom or a phosphorus atom, X is a halogen atom, a hydrocarbon group (excluding the cyclopentadienyl type anion skeleton-carrying group) or $-OR^{11}$, in which R^{11} is a hydrocarbon group or a halogenated hydrocarbon group, and more than one R^{11} may be the same or different from each other, a
15 is a number satisfying $0 < a \leq 8$, and b is a number satisfying $0 < b \leq 8$.

24. The pre-polymerized catalyst component for addition polymerization according to Claim 22, wherein X is OR^{11} .

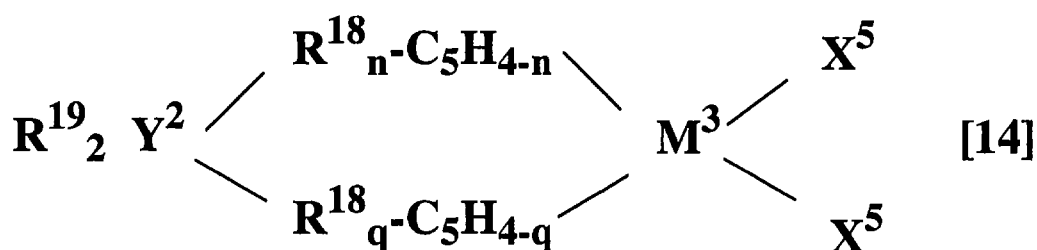
25. The pre-polymerized catalyst component for
20 addition polymerization according to Claim 21, wherein the transition metal compound (B) of the Groups 3 to 11 or a lanthanoid series is a transition metal compound (B) having a stereoregular polymerization ability of an α -olefin.

26. The pre-polymerized catalyst component for
25 addition polymerization according to Claim 25, wherein the transition metal compound (B) having a stereoregular polymerization ability of an α -olefin is a transition metal

compound represented by the following general formula [13] or [14],



5 wherein M^3 is a transition metal atom of the Groups 3 to 11 or
 lanthanide series, L^3 is an η^5 -indenyl group or a substituted
 η^5 -indenyl group, and both of L^3 s may be the same or different
 from each other, Y is a bridging group for linking both of L^3 s,
 10 and both of X^4 s are independently of each other a hydrogen atom,
 a halogen atom, an alkyl group, an aralkyl group, an aryl group,
 a substituted silyl group, an alkoxy group, an aralkyloxy group,
 an aryloxy group or a heterocyclic group,



15 wherein M^3 is a transition metal atom of the Groups 3 to 11
 or lanthanide series, Y^2 is a silicon atom, a germanium atom
 or a tin atom, the $(R^{18}_n-C_5H_{4-n})$ and $(R^{18}_q-C_5H_{4-q})$ are each a
 substituted η^5 -cyclopentadienyl group, n and q are each an
 20 integer of 1 to 3, and both of R^{18} s may be the same or different

from each other and are independently of each other a halogen atom, an alkyl group, an aralkyl group, an aryl group, a substituted silyl group, an alkoxy group, an aralkyloxy group, an aryloxy group or a heterocyclic group, the position and/or
5 kind of each R^{18} in each substituted η^5 -cyclopentadienyl group are (is) selected, so that a symmetry plane containing M^3 does not exist, R^{19} and X^5 are independently of each other a hydrogen atom, a halogen atom, an alkyl group, an aralkyl group, an aryl group, a substituted silyl group, an alkoxy
10 group, an aralkyloxy group, an aryloxy group or a heterocyclic group, and R^{19} and X^5 may be all the same or different from each other.

27. A catalyst for addition polymerization, which is obtained by contacting the pre-polymerized catalyst component
15 for addition polymerization according to any of Claims 21 to 26 and an organoaluminum compound (C) with each other.

28. A pre-polymerized catalyst for addition polymerization, which is obtained by pre-polymerizing an olefin in the presence of a catalyst for addition polymerization,
20 wherein the catalyst is obtained by contacting the modified particle (A) according to any of Claims 1 to 11 and a transition metal compound (B) of the Groups 3 to 11 or a lanthanoid series with each other.

29. A pre-polymerized catalyst for addition
25 polymerization, which is obtained by pre-polymerizing an olefin in the presence of a catalyst for addition polymerization, wherein the catalyst is obtained by contacting the modified

particle (A) according to any of Claims 1 to 11, a transition metal compound (B) of the Groups 3 to 11 or a lanthanoid series and an organoaluminum compound (C) with one another.

30. A process for producing an addition polymer, which
5 comprises polymerizing an addition polymerizable monomer in the presence of the catalyst for addition polymerization according to any of Claims 14 to 17 and 27 to 29.

31. The process for producing an addition polymer according to Claim 30, wherein the addition polymerizable monomer
10 is an α -olefin polymer.

32. The process for producing an addition polymer according to Claim 31, wherein the addition polymerizable monomer is a mixture of ethylene and an α -olefin.

33. A process for producing a stereoregular α -olefin
15 polymer, which comprises polymerizing an α -olefin having 3 to 20 carbon atoms in the presence of the catalyst for addition polymerization according to Claim 19 or 20.

34. A process for producing a stereoregular α -olefin polymer, which comprises polymerizing an α -olefin having 3 to
20 20 carbon atoms in the presence of the catalyst for addition polymerization, wherein the catalyst is obtained by contacting the pre-polymerized catalyst component for addition polymerization according to Claim 25 or 26 and an organoaluminum compound (C) with each other.

25 35. The process for producing a stereoregular α -olefin polymer according to Claim 33 or 34, wherein the α -olefin is