

CATALYST COMPONENT AND CATALYST FOR ADDITION POLYMERIZATION,
AND PROCESS FOR PRODUCING ADDITION POLYMER

BACKGROUND OF THE INVENTION

5 Field of the Invention

The present invention relates to a catalyst component for addition polymerization, a catalyst for addition polymerization, and a process for producing an addition polymer. More particularly, the present invention relates to a catalyst component for addition polymerization composed of a transition metal compound having two transition metal atoms in one molecule, a catalyst for addition polymerization prepared by using the same, and a process for producing an addition polymer using this catalyst for addition polymerization.

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Description of Related Arts

With respect to processes for producing an addition polymer such as an olefin polymer and the like using a transition metal compound which forms a single site catalyst such as a metallocene complex and the like, many reports have been reported. For example, JP60-245604A discloses a process for producing a copolymer of ethylene with an α -olefin using a metallocene complex and half metallocene complex.

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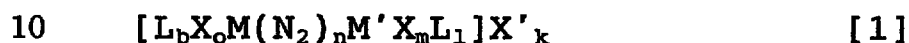
SUMMARY OF THE INVENTION

A catalyst for addition polymerization used in the production of an addition polymer is more efficient when activity thereof is higher, and therefore, a catalyst for addition

polymerization of high activity is required.

An object of the present invention is to provide a catalyst component for addition polymerization useful for preparing a catalyst for addition polymerization having a high activity,
 5 a catalyst for addition polymerization having a high activity, and an efficient process for producing an addition polymer.

Namely, the present invention relates to a catalyst component for addition polymerization composed of a transition metal compound (A) of the following general formula [1]:



(wherein, M and M' each independently represent a transition metal atom of III to X group in the Periodic Table of Elements; X each independently represents a hydrogen atom, halogen atom, alkyl group, aralkyl group, aryl group, substituted silyl group,
 15 alkoxy group, aralkyloxy group, aryloxy group, di-substituted amino group, azido group, cyano group, isothiocyanate group or group having a cyclopentadiene type anion skeleton; L represents a group which bonds to M or M' by a lone-paired electron or a π electron; X' represents a counter anion; k, l, m, o and
 20 p each independently represent an integer of 0 to 5; and n represents an integer of 1 to 3.).

Further, the present invention relates to a catalyst for addition polymerization obtained by a process comprising bringing the above-mentioned transition metal compound (A) into
 25 contact with an organoaluminum compound selected from the group consisting of the following (B1), and an aluminoxane selected from the group consisting of the following (B2) and (B3) and/or a boron compound selected from the group consisting of the following

(C), or with an aluminoxane selected from the group consisting of the following (B2) and (B3) and/or the following (C):

(B1) organoaluminum compounds of the general formula $E^1_a AlZ_{3-a}$,

5 (B2) cyclic aluminoxanes having a structure of the general formula $\{-Al(E^2)-O-\}_b$,

(B3) linear aluminoxanes having a structure of the general formula $E^3\{-Al(E^3)-O-\}_c AlE^3_2$

(wherein, each of E^1 , E^2 and E^3 represents a hydrocarbon group;
10 all E^1 's, all E^2 's or all E^3 's may be the same or different; Z represents a hydrogen atom or halogen atom; all Zs may be the same or different; a represents a number satisfying $0 < a \leq 3$; b represents an integer of 2 or more; and c represent an integer of 1 or more.), and

15 (C) one or more boron compounds selected from the following (C1) to (C3):

(C1) boron compounds represented by the general formula $BQ^1Q^2Q^3$,

(C2) boron compounds represented by the general formula
20 $G^+(BQ^1Q^2Q^3Q^4)^-$, and

(C3) boron compounds represented by the general formula $(L-H)^+(BQ^1Q^2Q^3Q^4)^-$

(wherein, B represents boron in trivalent state; Q^1 to Q^4 represent a halogen atom, hydrocarbon group, halogenated
25 hydrocarbon group, substituted silyl group, alkoxy group or di-substituted amino group; they may be the same or different; G^+ represents an inorganic or organic cation; L represents a neutral Lewis base; and $(L-H)^+$ represents a Brønsted acid.),

and a process for producing an addition polymer using this catalyst.

BRIEF DESCRIPTION OF THE DRAWING

5 Fig. 1 is a flow chart for helping understanding of the invention. This flow chart shows a typical example of embodiments of the invention, and the scope of the present invention is not restricted to this example at all.

The present invention will be illustrated in detail below.

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DETAILED DESCRIPTION OF THE INVENTION

(A) Transition metal compound

M and M' in a transition metal compound of the above-mentioned general formula [1] each independently represent a transition
15 metal atom of III to X group in the Periodic Table of Elements (IUPAC Inorganic Chemistry Nomenclature revised edition, 1989).

M is preferably a transition metal atom of III to V Group, more preferably a titanium atom, zirconium atom, hafnium atom,
20 vanadium atom, niobium atom or tantalum atom, further preferably a titanium atom or zirconium atom.

M' is preferably a transition metal atom of VI to X group, more preferably a chromium atom, molybdenum atom, tungsten atom, ruthenium atom, rhodium atom or palladium atom, further
25 preferably a chromium atom, molybdenum atom or tungsten atom.

X in the above-mentioned general formula [1] represents a hydrogen atom, halogen atom, alkyl group, aralkyl group, aryl group, substituted silyl group, alkoxy group, aralkyloxy group,

aryloxy group, di-substituted amino group, azido group, cyano group, isothiocyanate group or group having a cyclopentadiene type anion skeleton, and all Xs may be the same or different.

Among these, a hydrogen atom, halogen atom, alkyl group, aralkyl group, aryl group, substituted silyl group, alkoxy group, aralkyloxy group, aryloxy group, di-substituted amino group and group having a cyclopentadiene type anion skeleton are preferred.

As the halogen atom of the substituent X, a fluorine atom, chlorine atom, bromine atom, iodine atom and the like are listed.

The alkyl group of the substituent X is preferably an alkyl group having 1 to 20 carbon atoms, and examples thereof include a methyl group, ethyl group, n-propyl group, isopropyl group, n-butyl group, sec-butyl group, tert-butyl group, n-pentyl group, neopentyl group, isoamyl group, n-hexyl group, n-octyl group, n-decyl group, n-dodecyl group, n-pentadecyl group, n-eicosyl group and the like, and more preferable examples are a methyl group, ethyl group, isopropyl group, tert-butyl group and isoamyl group.

Any of these alkyl groups may be substituted with a halogen atom such as a fluorine atom, chlorine atom, bromine atom, iodine atom or the like. Examples of the alkyl group having 1 to 20 carbon atoms substituted with a halogen atom include a fluoromethyl group, difluoromethyl group, trifluoromethyl group, chloromethyl group, dichloromethyl group, trichloromethyl group, bromomethyl group, dibromomethyl group, tribromomethyl group, iodomethyl group, diiodomethyl group, triiodomethyl group, fluoroethyl group, difluoroethyl group,

trifluoroethyl group, tetrafluoroethyl group,
 pentafluoroethyl group, chloroethyl group, dichloroethyl group,
 trichloroethyl group, tetrachloroethyl group,
 pentachloroethyl group, bromoethyl group, dibromoethyl group,
 5 tribromoethyl group, tetrabromoethyl group, pentabromoethyl
 group, perfluoropropyl group, perfluorobutyl group,
 perfluoropentyl group, perfluorohexyl group, perfluorooctyl
 group, perfluorododecyl group, perfluoropentadecyl group,
 perfluoroeicosyl group, perchloropropyl group, perchlorobutyl
 10 group, perchloropentyl group, perchlorohexyl group,
 perchlorooctyl group, perchlorododecyl group,
 perchloropentadecyl group, perchloroeicosyl group,
 perbromopropyl group, perbromobutyl group, perbromopentyl
 group, perbromohexyl group, perbromooctyl group,
 15 perbromododecyl group, perbromopentadecyl group,
 perbromoeicosyl group and the like.

Further, any of these alkyl groups may also be partially
 substituted with an alkoxy group such as a methoxy group, ethoxy
 group and or like, an aryloxy group such as a phenoxy group
 20 or the like, an aralkyloxy group such as a benzyloxy group or
 the like, etc.

The aralkyl group of the substituent X is preferably an
 aralkyl group having 7 to 20 carbon atoms, and examples thereof
 include a benzyl group, (2-methylphenyl)methyl group,
 25 (3-methylphenyl)methyl group, (4-methylphenyl)methyl group,
 (2,3-dimethylphenyl)methyl group,
 (2,4-dimethylphenyl)methyl group,
 (2,5-dimethylphenyl)methyl group,

- (2,6-dimethylphenyl)methyl group,
 (3,4-dimethylphenyl)methyl group,
 (3,5-dimethylphenyl)methyl group,
 (2,3,4-trimethylphenyl)methyl group,
 5 (2,3,5-trimethylphenyl)methyl group,
 (2,3,6-trimethylphenyl)methyl group,
 (3,4,5-trimethylphenyl)methyl group,
 (2,4,6-trimethylphenyl)methyl group,
 (2,3,4,5-tetramethylphenyl)methyl group,
 10 (2,3,4,6-tetramethylphenyl)methyl group,
 (2,3,5,6-tetramethylphenyl)methyl group,
 (pentamethylphenyl)methyl group, (ethylphenyl)methyl group,
 (n-propylphenyl)methyl group, (isopropylphenyl)methyl group,
 (n-butylphenyl)methyl group, (sec-butylphenyl)methyl group,
 15 (tert-butylphenyl)methyl group, (n-pentylphenyl)methyl group,
 (neopentylphenyl)methyl group, (n-hexylphenyl)methyl group,
 (n-octylphenyl)methyl group, (n-decylphenyl)methyl group,
 (n-tetradecylphenyl)methyl group, naphthylmethyl group,
 anthracenylmethyl group and the like, and a benzyl group is
 20 preferable.

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

The aryl group of the substituent X is preferably an aryl group having 6 to 20 carbon atoms, and examples thereof include

a phenyl group, 2-tolyl group, 3-tolyl group, 4-tolyl group, 2,3-xylyl group, 2,4-xylyl group, 2,5-xylyl group, 2,6-xylyl group, 3,4-xylyl group, 3,5-xylyl group, 2,3,4-trimethylphenyl group, 2,3,5-trimethylphenyl group, 2,3,6-trimethylphenyl group, 2,4,6-trimethylphenyl group, 3,4,5-trimethylphenyl group, 2,3,4,5-tetramethylphenyl group, 2,3,4,6-tetramethylphenyl group, 2,3,5,6-tetramethylphenyl group, pentamethylphenyl group, ethylphenyl group, n-propylphenyl group, isopropylphenyl group, n-butylphenyl group, sec-butylphenyl group, tert-butylphenyl group, n-pentylphenyl group, neopentylphenyl group, n-hexylphenyl group, n-octylphenyl group, n-decylphenyl group, n-dodecylphenyl group, n-tetradecylphenyl group, naphthyl group, anthracenyl group and the like, and a phenyl group is more preferable.

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

The substituted silyl group of the substituent X is a silyl group substituted with a hydrocarbon group, and examples of the hydrocarbon group include alkyl groups having 1 to 10 carbon atoms such as a methyl group, ethyl group, n-propyl group, isopropyl group, n-butyl group, sec-butyl group, tert-butyl group, isobutyl group, n-pentyl group, n-hexyl group, cyclohexyl group and the like, and aryl groups such as a phenyl

group and the like, etc. As such substituted silyl group having 1 to 20 carbon atoms, for example, mono-substituted silyl groups having 1 to 20 carbon atoms such as a methylsilyl group, ethylsilyl group, phenylsilyl group and the like,

5 di-substituted silyl groups having 2 to 20 carbon atoms such as a dimethylsilyl group, diethylsilyl group, diphenylsilyl group and the like, tri-substituted silyl groups such as a trimethylsilyl group, triethylsilyl group, tri-n-propylsilyl group, triisopropylsilyl group, tri-n-butylsilyl group,

10 tri-sec-butylsilyl group, tri-tert-butylsilyl group, triisobutylsilyl group, tert-butyldimethylsilyl group, tri-n-pentylsilyl group, tri-n-hexylsilyl group, tricyclohexylsilyl group, triphenylsilyl group and the like, etc. are listed, and preferable are a trimethylsilyl group,

15 tert-butyldimethylsilyl group and triphenylsilyl group.

Any of these substituted silyl groups may also be partially substituted with a halogen atom such as a fluorine atom, chlorine atom, bromine atom, iodine atom and the like, an alkoxy group such as a methoxy group, ethoxy group and the like, an aryloxy group

20 group such as a phenoxy group and the like, an aralkyloxy group such as a benzyloxy group and the like, etc.

The alkoxy group of the substituent X is preferably an alkoxy group having 1 to 20 carbon atoms, and examples thereof include a methoxy group, ethoxy group, n-propoxy group, isopropoxy group,

25 n-butoxy group, sec-butoxy group, t-butoxy group, n-pentoxy group, neopentoxy group, n-hexoxy group, n-octoxy group, n-dodexy group, n-pentadecy group, n-icooxy group and the like, and more preferable are a methoxy group, ethoxy group,

and t-butoxy group.

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

The aralkyloxy group of the substituent X is preferably an aralkyloxy group having 7 to 20 carbon atoms, and examples
 10 thereof include a benzyloxy group, (2-methylphenyl)methoxy group, (3-methylphenyl)methoxy group, (4-methylphenyl)methoxy group, (2,3-dimethylphenyl)methoxy group, (2,4-dimethylphenyl)methoxy group, (2,5-dimethylphenyl)methoxy group,
 15 (2,6-dimethylphenyl)methoxy group, (3,4-dimethylphenyl)methoxy group, (3,5-dimethylphenyl)methoxy group, (2,3,4-trimethylphenyl)methoxy group, (2,3,5-trimethylphenyl)methoxy group,
 20 (2,3,6-trimethylphenyl)methoxy group, (2,4,5-trimethylphenyl)methoxy group, (2,4,6-trimethylphenyl)methoxy group, (3,4,5-trimethylphenyl)methoxy group, (2,3,4,5-tetramethylphenyl)methoxy group,
 25 (2,3,4,6-tetramethylphenyl)methoxy group, (2,3,5,6-tetramethylphenyl)methoxy group, (pentamethylphenyl)methoxy group, (ethylphenyl)methoxy group, (n-propylphenyl)methoxy group, (isopropylphenyl)methoxy

group, (n-butylphenyl)methoxy group,

(sec-butylphenyl)methoxy group, (tert-butylphenyl)methoxy group, (n-hexylphenyl)methoxy group, (n-octylphenyl)methoxy group, (n-decylphenyl)methoxy group,

- 5 (n-tetradecylphenyl)methoxy group, naphthylmethoxy group, anthracenylmethoxy group and the like, and more preferable is a benzyloxy group.

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

The aryloxy group of the substituent X is preferably an
15 aryloxy group having 1 to 20 carbon atoms, and examples thereof include aryloxy groups having 6 to 20 carbon atoms such as a phenoxy group, 2-methylphenoxy group, 3-methylphenoxy group, 4-methylphenoxy group, 2,3-dimethylphenoxy group, 2,4-dimethylphenoxy group, 2,5-dimethylphenoxy group,
20 2,6-dimethylphenoxy group, 3,4-dimethylphenoxy group, 3,5-dimethylphenoxy group, 2,3,4-trimethylphenoxy group, 2,3,5-trimethylphenoxy group, 2,3,6-trimethylphenoxy group, 2,4,5-trimethylphenoxy group, 2,4,6-trimethylphenoxy group, 3,4,5-trimethylphenoxy group, 2,3,4,5-tetramethylphenoxy
25 group, 2,3,4,6-tetramethylphenoxy group, 2,3,5,6-tetramethylphenoxy group, pentamethylphenoxy group, ethylphenoxy group, n-propylphenoxy group, isopropylphenoxy group, n-butylphenoxy group, sec-butylphenyl group,

tert-butylphenoxy group, n-hexylphenoxy group, n-octylphenoxy group, n-decylphenoxy group, n-tetradecylphenoxy group, naphthoxy group, anthracenoxy group and the like.

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

The di-substituted amino group of the substituent X is an amino group substituted with two hydrocarbon groups, and examples of the hydrocarbon group include alkyl groups having 1 to 10 carbon atoms such as a methyl group, ethyl group, n-propyl group, isopropyl group, n-butyl group, sec-butyl group, tert-butyl group, isobutyl group, n-pentyl group, n-hexyl group, cyclohexyl group and the like, and aryl groups such as a phenyl group, and the like. Examples of such di-substituted amino groups include a dimethylamino group, diethylamino group, di-n-propylamino group, diisopropylamino group, di-n-butylamino group, di-sec-butylamino group, di-tert-butylamino group, diisobutylamino group, tert-butylisopropylamino group, di-n-hexylamino group, di-n-octylamino group, di-n-decylamino group, diphenylamino group, bistrimethylsilylamino group, bis-tert-butyl dimethylsilylamino group and the like, and preferable are a dimethylamino group, diethylamino group.

As the group having a cyclopentadiene-type anion skeleton of the substituent X, there are listed 7)

- ⁵-(substituted)cyclopentadienyl group, η
⁵-(substituted)indenyl group, η^5 -(substituted)fluorenyl
 group and the like. Specific examples thereof are a η
⁵-cyclopentadienyl group, η^5 -methylcyclopentadienyl group, η
 5 ⁵-dimethylcyclopentadienyl group, η
⁵-trimethylcyclopentadienyl group, η
⁵-tetramethylcyclopentadienyl group, η
⁵-pentamethylcyclopentadienyl group, η
⁵-ethylcyclopentadienyl group, η^5 -n-propylcyclopentadienyl
 10 group, η^5 -isopropylcyclopentadienyl group, η
⁵-n-butylcyclopentadienyl group, η
⁵-sec-butylcyclopentadienyl group, η
⁵-tert-butylcyclopentadienyl group, η
⁵-n-pentylcyclopentadienyl group, η
 15 ⁵-neopentylcyclopentadienyl group, η
⁵-n-hexylcyclopentadienyl group, η^5 -n-octylcyclopentadienyl
 group, η^5 -phenylcyclopentadienyl group, η
⁵-naphthylcyclopentadienyl group, η
⁵-trimethylsilylcyclopentadienyl group, η
 20 ⁵-triethylsilylcyclopentadienyl group, η
⁵-tert-butyldimethylsilylcyclopentadienyl group, η^5 -indenyl
 group, η^5 -methylindenyl group, η^5 -dimethylindenyl group, η
⁵-ethylindenyl group, η^5 -n-propylindenyl group, η
⁵-isopropylindenyl group, η^5 -n-butylindenyl group, η
 25 ⁵-sec-butylindenyl group, η^5 -tert-butylindenyl group, η
⁵-n-pentylindenyl group, η^5 -neopentylindenyl group, η
⁵-n-hexylindenyl group, η^5 -n-octylindenyl group, η
⁵-n-decylindenyl group, η^5 -phenylindenyl group, η

- η^5 -methylphenylindenyl group, η^5 -naphthylindenyl group, η
 η^5 -trimethylsilylindenyl group, η^5 -triethylsilylindenyl group,
 η^5 -tert-butyl dimethylsilylindenyl group, η
 η^5 -tetrahydroindenyl group, η^5 -fluorenyl group, η
5 η^5 -methylfluorenyl group, η^5 -dimethylfluorenyl group, η
 η^5 -ethylfluorenyl group, η^5 -diethylfluorenyl group, η
 η^5 -n-propylfluorenyl group, η^5 -di-n-propylfluorenyl group, η
 η^5 -isopropylfluorenyl group, η^5 -diisopropylfluorenyl group, η
 η^5 -n-butylfluorenyl group, η^5 -sec-butylfluorenyl group, η
10 η^5 -tert-butylfluorenyl group, η^5 -di-n-butylfluorenyl group, η
 η^5 -di-sec-butylfluorenyl group, η^5 -di-tert-butylfluorenyl
group, η^5 -n-pentylfluorenyl group, η^5 -neopentylfluorenyl
group, η^5 -n-hexylfluorenyl group, η^5 -n-octylfluorenyl group,
 η^5 -n-decylfluorenyl group, η^5 -n-dodecylfluorenyl group, η
15 η^5 -phenylfluorenyl group, η^5 -di-phenylfluorenyl group, η
 η^5 -methylphenylfluorenyl group, η^5 -naphthylfluorenyl group, η
 η^5 -trimethylsilylfluorenyl group, η
 η^5 -bis-trimethylsilylfluorenyl group, η
 η^5 -triethylsilylfluorenyl group, η
20 η^5 -tert-butyl dimethylsilylfluorenyl group and the like, and
preferable are a η^5 -cyclopentadienyl group, η
 η^5 -methylcyclopentadienyl group, η
 η^5 -tert-butylcyclopentadienyl group, η
 η^5 -tetramethylcyclopentadienyl group, η
25 η^5 -pentamethylcyclopentadienyl group, η^5 -indenyl group and η
 η^5 -fluorenyl group.

" η^5 -" is sometimes omitted.

X' in the above-mentioned general formula [1] represents

a counter anion, and is an anionic group which is not covalent-bonded to but free ionically from M, and for example, conjugated bases of Brønsted acids are listed. X' is preferably F^- , Cl^- , Br^- , I^- , BF_4^- , $B(phenyl)_4^-$ or PF_6^- .

5 L in the above-mentioned general formula [1] represents a group which bonds to M or M' by lone pair or π electron.

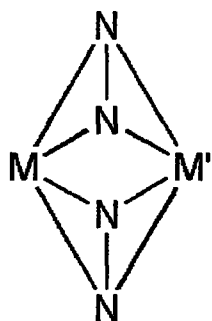
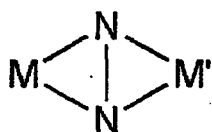
The group which bonds to M or M' by a lone electron-pair is a neutral ligand which bonds to M or M' by a coordinate bond, and examples thereof include ethers such as diethyl ether, 10 tetrahydrofuran, diethoxyethane; amines such as triethylamine, N,N,N',N'-tetramethylethylenediamine; pyridines such as pyridine, 2,6-dimethylpyridine, quinoline; phosphines such as trimethylphosphine, triethylphosphine, triphenylphosphine, dimethylphenylphosphine, methyldiphenylphosphine, 15 1,2-bis(dimethylphosphino)ethane, 1,2-bis(diethylphosphino)ethane, 1,2-bis(diphenylphosphino)ethane, 1,3-bis(diphenylphosphino)propane; nitriles such as acetonitrile, benzonitrile and the like, an end-on type nitrogen 20 molecule and carbon mono-oxide, and the like, and preferable are tetrahydrofuran, N,N,N',N'-tetramethylethylenediamine, pyridine, dimethylphenylphosphine, 1,2-bis(dimethylphosphino)ethane, 1,2-bis(diethylphosphino)ethane, 25 1,2-bis(diphenylphosphino)ethane, 1,3-bis(diphenylphosphino)propane, acetonitrile and carbon mono-oxide.

The group which bonds to M or M' by a π electron is a neutral

ligand which bonds to M or M' by a multi bonding orbital, and examples thereof include olefins such as ethylene, propylene and the like, dienes such as butadiene, 2,4-hexadiene, 1,4-diphenylbutadiene; ketones such as acetone, benzophenone; a side-on type nitrogen molecule, and the like, and preferable are olefins and dienes, and more preferable are ethylene, butadiene, 2,4-hexadiene and 1,4-diphenylbutadiene.

k, l, m, o and p each independently represent an integer of 0 to 5, and n represents an integer of 1 to 3 in the above-mentioned general formula [1]. l, m, o and p are preferably selected so as to satisfy $q \geq o+n$; $r \geq m+n$; $n+o+p \geq 6$ and $n+l+m \geq 6$ when group numbers of M and M' in the Periodic Table are represented by q and r, respectively. k is preferably selected so as to satisfy $k=s+t-o-m-2$ when the formal oxidizing numbers of M and M' are represented by s and t, respectively.

As the partial structure $M(N_2)_nM'$ in a transition metal compound of the above-mentioned general formula [1], the following structures are exemplified.



A transition metal compound of the general formula [1] used in the present invention is produced by ,for example, a method described in Organometallics, Vol. 13, p.3764-3766 (1994).

- 5 Specific examples of a compound of the general formula [1] include transition metal compounds such as
- [chlorotetrakis(trimethylphosphine)tungsten]
- (μ -dinitrogen)[trichlorotitanium],
- [chlorotris(trimethylphosphine)(pyridine)tungsten](μ
- 10 -dinitrogen)[trichlorotitanium],
- [chlorotetrakis(triethylphosphine)tungsten] (μ
- dinitrogen)[trichlorotitanium],
- [chlorotris(triethylphosphine)(pyridine)tungsten] (μ
- dinitrogen) [trichlorotitanium],

- [chlorotetrakis(triphenylphosphine)tungsten] (μ -dinitrogen)[trichlorotitanium],
- [chlorotris(triphenylphosphine)(pyridine)tungsten] (μ -dinitrogen) [trichlorotitanium],
- 5 [chlorotetrakis(dimethylphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
- [chlorotris(dimethylphenylphosphine)(pyridine)tungsten] (μ -dinitrogen) [trichlorotitanium],
- [chlorotetrakis(methyldiphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
- 10 [chlorotris(methyldiphenylphosphine)(pyridine)tungsten] (μ -dinitrogen) [trichlorotitanium],
- [chlorobis{1,2-bis(dimethylphosphino)ethane}tungsten] (μ -dinitrogen) [trichlorotitanium],
- 15 [chlorobis{1,2-bis(diethylphosphino)ethane}tungsten] (μ -dinitrogen) [trichlorotitanium],
- [chlorobis{1,2-bis(diphenylphosphino)ethane}tungsten] (μ -dinitrogen)[trichlorotitanium],
- [chlorobis{1,3-bis(diphenylphosphino)propane}tungsten] (μ -dinitrogen)[trichlorotitanium],
- 20 [chlorotris(trimethylphosphine)(2,6-dimethylpyridine)tungsten](μ -dinitrogen)[trichlorotitanium],
- [chlorotris(triethylphosphine)(2,6-dimethylpyridine)tungsten](μ -dinitrogen)[trichlorotitanium],
- 25 [chlorotris(triphenylphosphine)(2,6-dimethylpyridine)tungsten] (μ -dinitrogen) [trichlorotitanium],
- [chlorotris(dimethylphenylphosphine)(2,6-dimethylpyridine)tungsten] (μ -dinitrogen) [trichlorotitanium],

- [chlorotris(trimethylphosphine)(quinoline)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chlorotris(triethylphosphine)(quinoline)tungsten] (μ -dinitrogen) [trichlorotitanium],
 5 [chlorotris(triphenylphosphine)(quinoline)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chlorotris(dimethylphenylphosphine)(quinoline)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(tetrahydrofuran)tris(trimethylphosphine)tungsten]
 10 (μ -dinitrogen) [trichlorotitanium],
 [chloro(tetrahydrofuran)tris(triethylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(tetrahydrofuran)tris(triphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 15 [chloro(tetrahydrofuran)tris(dimethylphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(diethyl ether)tris(dimethylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium], [chloro(diethyl ether)tris(triethylphosphine)tungsten] (μ -dinitrogen)
 20 [trichlorotitanium], [chloro(diethyl ether)tris(triphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium], [chloro(diethyl ether)tris(dimethylphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 25 [chloro(acetonitrile)tris(trimethylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(acetonitrile)tris(triethylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],

- [chloro(acetonitrile)tris(triphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(acetonitrile)tris(dimethylphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
- 5 [chloro(benzonitrile)tris(trimethylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(benzonitrile)tris(triethylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(benzonitrile)tris(triphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
- 10 [chloro(benzonitrile)tris(dimethylphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(carbonyl)tris(trimethylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
- 15 [chloro(carbonyl)tris(triethylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(carbonyl)tris(triphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(carbonyl)tris(dimethylphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
- 20 [chloro(dimethoxyethane)bis(trimethylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(dimethoxyethane)bis(triethylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
- 25 [chloro(dimethoxyethane)bis(triphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],
 [chloro(dimethoxyethane)bis(dimethylphenylphosphine)tungsten] (μ -dinitrogen) [trichlorotitanium],

- [chloro(dimethoxyethane)bis(methyldiphenylphosphine)tungsten](μ -dinitrogen)[trichlorotitanium],
 [chloro(diphenylphosphinoethane)(dimethoxyethane)tungsten](μ -dinitrogen)[trichlorotitanium],
 5 [chloro(diphenylphosphinopropane)(dimethoxyethane)tungsten](μ -dinitrogen)[trichlorotitanium],
 [chloro(N,N,N',N'-tetramethylethylenediamine)bis(trimethylphosphine)tungsten](μ -dinitrogen)[trichlorotitanium],
 [chloro(N,N,N',N'-tetramethylethylenediamine)bis(triethylphosphine)tungsten](μ -dinitrogen)[trichlorotitanium],
 10 [chloro(N,N,N',N'-tetramethylethylenediamine)bis(triphenylphosphine)tungsten](μ -dinitrogen)[trichlorotitanium],
 [chloro(N,N,N',N'-tetramethylethylenediamine)bis(dimethylphenylphosphine)tungsten](μ -dinitrogen)[trichlorotitanium],
 15 [chloro(N,N,N',N'-tetramethylethylenediamine)bis(methyldiphenylphosphine)tungsten](μ -dinitrogen)[trichlorotitanium],
 [chloro(N,N,N',N'-tetramethylethylenediamine)(diphenylphosphinoethane)tungsten](μ -dinitrogen)[trichlorotitanium],
 [chloro(N,N,N',N'-tetramethylethylenediamine)(diphenylphosphinopropane)tungsten](μ -dinitrogen)[trichlorotitanium]
 20 and the like, compounds obtained by substituting tungsten in these compounds by molybdenum, chromium, ruthenium, rhodium or palladium, compounds obtained by substituting titanium in these compounds by zirconium, hafnium, vanadium, niobium,
 25 tantalum or scandium, and compounds obtained by substituting chloro in these compounds by fluoro, bromo, iodo, methyl, benzyl, methoxy or phenoxy.

The transition metal compound of the general formula [1]

is preferably a transition metal compound in which o represents an integer of 1 to 5 and at least one X is a group having a cyclopentadiene-type anion skeleton. Specific examples of such a preferable transition metal compound include transition

- 5 metal compounds such as
- [chlorotetrakis(trimethylphosphine)tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- [chlorotris(trimethylphosphine)tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- 10 [chlorotetrakis(triethylphosphine)(pyridine)tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- [chlorotris(triethylphosphine)(pyridine)tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- [chlorotetrakis(triphenylphosphine)tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- 15 [chlorotris(triphenylphosphine)(pyridine)tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- [chlorotetrakis(dimethylphenylphosphine)tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- 20 [chlorotris(dimethylphenylphosphine)(pyridine)tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- [chlorotetrakis(methyldiphenylphosphine)tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- [chlorotris(methyldiphenylphosphine)(pyridine)tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- 25 [chlorobis{1,2-bis(dimethylphosphino)ethane}tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- [chlorobis{1,2-bis(diethylphosphino)ethane}tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],

- dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chlorobis{1,2-bis(diphenylphosphino)ethane}tungsten] (μ -
 -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chlorobis{1,3-bis(diphenylphosphino)propane}tungsten] (μ -
 5 -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chlorotris(trimethylphosphine)(2,6-dimethylpyridine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chlorotris(triethylphosphine)(2,6-dimethylpyridine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 10 [chlorotris(triethylphosphine)(2,6-dimethylpyridine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chlorotris(dimethylphenylphosphine)(2,6-dimethylpyridine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium], [chlorotris
 15 (trimethylphosphine)(quinoline)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chlorotris(triethylphosphine)(quinoline)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chlorotris(triphenylphosphine)(quinoline)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 20 -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chlorotris(dimethylphenylphosphine)(quinoline)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(tetrahydrofuran)tris(trimethylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 25 [chloro(tetrahydrofuran)tris(triethylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(tetrahydrofuran)tris(triphenylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],

- [chloro(tetrahydrofuran)tris(dimethylphenylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(diethyl ether)tris(trimethylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- 5 [chloro(diethyl ether)tris(triethylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(diethyl ether)tris(triphenylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(diethyl
- 10 ether)tris(dimethylphenylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(acetonitrile)tris(trimethylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(acetonitrile)tris(triethylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- 15 [chloro(acetonitrile)tris(triphenylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(acetonitrile)tris(dimethylphenylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- 20 [chloro(benzonitrile)tris(trimethylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(benzonitrile)tris(triethylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(benzonitrile)tris(triphenylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
- 25 [chloro(benzonitrile)tris(dimethylphenylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(carbonyl)tris(trimethylphosphine)tungsten(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],

- dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(carbonyl)tris(triethylphosphine)tungsten(μ -
 -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(carbonyl)tris(triphenylphosphine)tungsten(μ -
 5 -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(carbonyl)tris(dimethylphenylphosphine)tungsten(μ -
 -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(dimethoxyethane)bis(trimethylphosphine)tungsten(μ -
 -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 10 [chloro(dimethoxyethane)bis(triethylphosphine)tungsten(μ -
 -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(dimethoxyethane)bis(triphenylphosphine)tungsten(μ -
 -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(dimethoxyethane)bis(dimethylphenylphosphine)tungst
 15 en(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(dimethoxyethane)bis(dimethylphenylphosphine)tungst
 en(μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(diphenylphosphinoethane)(dimethoxyethane)tungsten(
 μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 20 [chloro(diphenylphosphinopropane)(dimethoxyethane)tungsten
 (μ -dinitrogen) [dichloro(cyclopentadienyl)titanium],
 [chloro(N,N,N',N'-tetramethylethylenediamine)bis(trimethyl
 phosphine)tungsten] (μ -dinitrogen)
 [dichloro(cyclopentadienyl)titanium],
 25 [chloro(N,N,N',N'-tetramethylethylenediamine)bis(triethylp
 hosphine)tungsten] (μ -dinitrogen)
 [dichloro(cyclopentadienyl)titanium],
 [chloro(N,N,N',N'-tetramethylethylenediamine)bis(triphenyl

- phosphine)tungsten] (μ -dinitrogen)
 [dichloro(cyclopentadienyl)titanium],
 [chloro(N,N,N',N'-tetramethylethylenediamine)bis(dimethylp
 henylphosphine)tungsten] (μ -dinitrogen)
- 5 [dichloro(cyclopentadienyl)titanium],
 [chloro(N,N,N',N'-tetramethylethylenediamine)bis(methyldip
 henylphosphine)tungsten] (μ -dinitrogen)
 [dichloro(cyclopentadienyl)titanium],
 [chloro(N,N,N',N'-tetramethylethylenediamine)(diphenylphos
 10 phinoethane)tungsten] (μ -dinitrogen)
 [dichloro(cyclopentadienyl)titanium],
 [chloro(N,N,N',N'-tetramethylethylenediamine)(diphenylphos
 phinopropane)tungsten] (μ -dinitrogen)
 [dichloro(cyclopentadienyl)titanium], compounds obtained by
 15 substituting tungsten in these compounds by molybdenum,
 chromium, ruthenium, rhodium or palladium, compounds obtained
 by substituting titanium in these compounds by zirconium,
 hafnium, vanadium, niobium, tantalum or scandium, compounds
 obtained by substituting dichloro in these compounds by difluoro,
 20 dibromo, diiodo, dimethyl, dibenzyl, dimethoxy, diphenoxy,
 1,3-butadiene, 2,4-hexadiene, diphenyl-1,3-butadiene,
 (chloro)(methyl), (benzyl)(chloro), (chloro)(methoxy) or
 (chloro)(phenoxy), transition metal compounds obtained by
 substituting cyclopentadienyl in these compounds by
 25 methylcyclopentadienyl, dimethylcyclopentadienyl,
 trimethylcyclopentadienyl, tetramethylcyclopentadienyl,
 n-butylcyclopentadienyl,
 tert-butyldimethylsilylcyclopentadienyl, indenyl,

tetrahydroindenyl, fluorenyl or octahydrofluorenyl.

The catalyst component for addition polymerization of the present invention is a catalyst component for addition polymerization composed of a transition metal compound of the general formula [1], and catalysts for addition polymerization having a high activity are obtained by contacting this component with a co-catalyst component for activation.

The co-catalyst component for activation is preferably the following (B) and/or (C), and the catalyst for addition polymerization of the present invention is preferably a catalyst for addition polymerization obtained by bringing a transition metal compound (A) of the general formula [1] into contact with an organoaluminum compound selected from the group consisting of the following (B1), and an aluminoxane selected from the group consisting of the following (B2) and (B3) and/or a boron compound selected from the group consisting of the following (C), or with an aluminoxane selected from the group consisting of the following (B2) and (B3) and/or the following (C):

(B1) organic aluminum compounds of the general formula $E^1_a AlZ_{3-a}$,

(B2) cyclic aluminoxanes having a structure of the general formula $\{-Al(E^2)-O-\}_b$,

(B3) linear aluminoxanes having a structure of the general formula $E^3\{-Al(E^3)-O-\}_c AlE^3_2$

(wherein, each of E^1 , E^2 and E^3 represents a hydrocarbon group, and all E^1 's, all E^2 's or all E^3 's may be the same or different. Z represents a hydrogen atom or halogen atom, and all Zs may be the same or different. a represents a number satisfying

$0 < a \leq 3$, b represents an integer of 2 or more, and c represent an integer of 1 or more.),

(C) one or more boron compounds selected from the following (C1) to (C3):

5 (C1) boron compounds represented by the general formula $BQ^1Q^2Q^3$,

(C2) boron compounds represented by the general formula $G^+(BQ^1Q^2Q^3Q^4)^-$,

10 (C3) boron compounds represented by the general formula $(L-H)^+(BQ^1Q^2Q^3Q^4)^-$

(wherein, B represents a trivalent boron atom in valance state, and Q^1 to Q^4 represent a halogen atom, hydrocarbon group, halogenated hydrocarbon group, substituted silyl group, alkoxy group or di-substituted amino group, and they may be the same or different. G^+ represents an inorganic or organic cation, and L represents a neutral Lewis base and $(L-H)^+$ represents a Brønsted acid.).

The catalyst for addition polymerization will be further illustrated in detail below.

20 (B) Aluminum compound

The aluminum compound (B) includes organoaluminum compounds of the following (B1) and aluminoxanes of the following (B2) and (B3).

(B1) organic aluminum compounds represented by the general
25 formula $E^1_aAlZ_{3-a}$,

(B2) cyclic aluminoxanes having a structure represented by the general formula $\{-Al(E^2)-O-\}_b$,

(B3) linear aluminoxanes having a structure represented

by the general formula $E^3\{-Al(E^3)-O-\}_cAlE^3_2$
 (wherein, each of E^1 , E^2 and E^3 represents a hydrocarbon group;
 all E^1 's, all E^2 's or all E^3 's may be the same or different; Z
 represents a hydrogen atom or halogen atom; all Zs may be the
 5 same or different; a represents a number satisfying $0 < a \leq 3$;
 b represents an integer of 2 or more; and c represent an integer
 of 1 or more.).

As the hydrocarbon group represented by E^1 , E^2 and E^3 , a
 hydrocarbon group having 1 to 8 carbon atoms is preferable,
 10 and an alkyl group is more preferable.

Specific examples of the organic aluminum compound (B1)
 represented by the general formula $E^1_aAlZ_{3-a}$ include
 trialkylaluminums such as trimethylaluminum, triethylaluminum,
 tripropylaluminum, triisobutylaluminum, trihexylaluminum and
 15 the like; dialkylaluminum chlorides such as dimethylaluminum
 chloride, diethylaluminum chloride, dipropylaluminum chloride,
 diisobutylaluminum chloride, dihexylaluminum chloride and the
 like; alkylaluminum dichlorides such as methylaluminum
 dichloride, ethylaluminum dichloride, propylaluminum
 20 dichloride, isobutylaluminum dichloride, hexylaluminum
 dichloride and the like; dialkylaluminum hydrides such as
 dimethylaluminum hydride, diethylaluminum hydride,
 dipropylaluminum hydride, diisobutylaluminum hydride,
 dihexylaluminum hydride and the like, etc.

25 Trialkylaluminums are preferable, and triethylaluminum or
 triisobutylaluminum is more preferable.

As specific examples of E^2 and E^3 in cyclic aluminoxanes
 (B2) having a structure of the general formula $\{-Al(E^2)-O-\}_b$,

and linear aluminoxanes (B3) having a structure of the general formula $E^3\{-Al(E^3)-O-\}_cAlE^3_2$, alkyl groups such as a methyl group, ethyl group, n-propyl group, isopropyl group, n-butyl group, isobutyl group, n-pentyl group, neopentyl group and the like
 5 can be exemplified. b is an integer of 2 or more, and c is an integer of 1 or more. Preferably, E^2 and E^3 are a methyl group or isobutyl group, n is 2 to 40, and c is 1 to 40.

The above-mentioned aluminoxane is produced by various methods. The method is not particularly restricted, and it
 10 may be advantageously carried out according to a known method. For example, a trialkylaluminum (for example, trimethylaluminum and the like) is dissolved in a suitable organic solvent (benzene, aliphatic hydrocarbon and the like) to prepare a solution which is allowed to contact with water.
 15 Alternatively, there is exemplified a method in which a trialkylaluminum (for example, trimethylaluminum and the like) is allowed to contact with a metal salt containing crystal water (for example, copper sulfate hydrate and the like).

The aluminoxane produced by such methods is usually supposed
 20 to be a mixture of a cyclic aluminoxane and a linear aluminoxane.

(C) Boron compound

As the boron compound (C), one or more boron compounds selected from (C1) boron compounds represented by the general formula $BQ^1Q^2Q^3$, (C2) boron compounds represented by the general
 25 formula $G^+(BQ^1Q^2Q^3Q^4)^-$, and (C3) boron compounds represented by the general formula $(L-H)^+(BQ^1Q^2Q^3Q^4)^-$ are used, in the present invention.

In the boron compound (C1) of the general formula $BQ^1Q^2Q^3$,

B represents a trivalent boron atom, and Q^1 to Q^3 represent a halogen atom, hydrocarbon group, halogenated hydrocarbon group, substituted silyl group, alkoxy group or di-substituted amino group, and they may be the same or different. Q^1 to Q^3 preferably represent a halogen atom, a hydrocarbon group having 1 to 20 carbon atoms, a halogenated hydrocarbon group having 1 to 20 carbon atoms, a substituted silyl group having 1 to 20 carbon atoms, an alkoxy group having 1 to 20 carbon atoms, or an amino group having 1 to 20 carbon atoms, and Q^1 to Q^3 more preferably represent a halogen atom, a hydrocarbon group having 1 to 20 carbon atoms or a halogenated hydrocarbon group having 1 to 20 carbon atoms. Further preferably, Q^1 to Q^4 represent a fluorinated hydrocarbon group having 1 to 20 carbon atoms containing at least one fluorine atom, and particularly preferably, Q^1 to Q^4 represent a fluorinated aryl group having 6 to 20 carbon atoms containing at least fluorine atom.

Specific examples of the compound (C1) include tris(pentafluorophenyl)borane, tris(2,3,5,6-tetrafluorophenyl)borane, tris(2,3,4,5-tetrafluorophenyl)borane, tris(3,4,5-trifluorophenyl)borane, tris(2,3,4-trifluorophenyl)borane, phenylbis(pentafluorophenyl)borane and the like, and tris(pentafluorophenyl)borane is most preferable.

In the boron compound (C2) of the general formula $G^+(BQ^1Q^2Q^3Q^4)^-$, G^+ represents an inorganic or organic cation, B represents a trivalent boron atom, and Q^1 to Q^4 are as defined for Q^1 to Q^3 in the above-mentioned (C1).

- As specific examples of an inorganic cation G^+ in the compound of the general formula $G^+(BQ^1Q^2Q^3Q^4)^-$, a ferrocenium cation, alkyl-substituted ferrocenium cation, silver cation and the like are listed, and as specific examples of an organic cation G^+ in the compound, a triphenylmethyl cation and the like are listed. G^+ represents preferably a carbenium cation, and particularly preferably a triphenylmethyl cation. As $(BQ^1Q^2Q^3Q^4)^-$, there are listed
- tetrakis(pentafluorophenyl)borate,
 - 10 tetrakis(2,3,5,6-tetrafluorophenyl)borate,
 - tetrakis(2,3,4,5-tetrafluorophenyl)borate,
 - tetrakis(3,4,5-trifluorophenyl)borate,
 - tetrakis(2,3,4-trifluorophenyl)borate,
 - phenyltris(pentafluorophenyl)borate,
 - 15 tetrakis(3,5-bistrifluoromethylphenyl)borate and the like.

- As specific combinations thereof, ferrocenium tetrakis(pentafluorophenyl)borate, 1,1'-dimethylferrocenium tetrakis(pentafluorophenyl)borate, silver tetrakis(pentafluorophenyl)borate,
- 20 triphenylmethyltetrakis(pentafluorophenyl)borate,
 - triphenylmethyltetrakis(3,5-bistrifluoromethylphenyl)borate, and the like are listed, and
 - triphenylmethyltetrakis(pentafluorophenyl)borate is most preferable.

- 25 In the boron compound (C3) of the general formula $(L-H)^+(BQ^1Q^2Q^3Q^4)^-$, L represents a neutral Lewis base and $(L-H)^+$ represents a Brønsted acid, B represents a trivalent boron atom, and Q^1 to Q^4 are as defined for Q^1 to Q^3 in the above-mentioned

Lewis acid (Cl).

As specific examples of the Brønsted acid $(L-H)^+$ in the compound of the general formula $(L-H)^+(BQ^1Q^2Q^3Q^4)^-$,

trialkyl-substituted ammoniums, N,N-dialkylaniliniums,

- 5 dialkylammoniums, triarylphosponiums and the like are listed, and as $(BQ^1Q^2Q^3Q^4)^-$, the same moieties as described above are listed.

As specific combinations thereof,

- triethylammoniumtetrakis(pentafluorophenyl)borate,
 10 tripropylammonium tetrakis(pentafluorophenyl)borate,
 tri(n-butyl)ammoniumtetrakis(pentafluorophenyl)borate,
 tri(n-butyl)ammonium
 tetrakis(3,5-bistrifluoromethylphenyl)borate,
 N,N-dimethylaniliniumtetrakis(pentafluorophenyl)borate,
 15 N,N-diethylanilinium tetrakis(pentafluorophenyl)borate,
 N,N-2,4,6-pentamethylaniliniumtetrakis
 (pentafluorophenyl)borate, N,N-dimethylaniliniumtetrakis
 (3,5-bistrifluoromethylphenyl)borate,
 diisopropylammoniumtetrakis(pentafluorophenyl)borate,
 20 dicyclohexylammoniumtetrakis(pentafluorophenyl)borate,
 triphenylphosponiumtetrakis(pentafluorophenyl)borate,
 tri(methylphenyl)phosponiumtetrakis(pentafluorophenyl)
 borate, tri(dimethylphenyl)phosponiumtetrakis
 (pentafluorophenyl)borate and the like are listed, and most
 25 preferable is tri(n-butyl)ammonium
 tetrakis(pentafluorophenyl)borate, or
 N,N-dimethylaniliniumtetrakis(pentafluorophenyl)borate.

[Production of addition polymer]

The process for producing an addition polymer of the present invention is a process for producing an addition polymer comprising polymerizing an addition polymerizable monomer with the above-mentioned catalyst for addition polymerization.

5 Contact in producing a catalyst for addition polymerization by bringing the above-mentioned transition metal compound into contact with a co-catalyst component for activation (components (B1)-(B3), and (C)) may be conducted by any means providing the transition metal compound and the co-catalyst for activation
10 are in contact to form a catalyst, and there are adopted a method in which a transition metal compound and a co-catalyst for activation are previously diluted by a solvent or not diluted, before they are mixed for mutual contact, and a method in which a transition metal compound and a co-catalyst for activation
15 are separately fed into a polymerization reactor for mutual contact thereof. As the co-catalyst for activation, a plurality of compounds may be combined and used, however, it is needless to say that a part of which may be previously mixed and used, or they may be separately fed into a polymerization reactor
20 and used.

It is desirable to use components so that the molar ratio of (B)/transition metal compound (A) is from 0.1 to 10000, preferably from 5 to 2000, and the molar ratio of (C)/transition metal compound (A) is from 0.01 to 100, preferably from 0.5
25 to 10.

The concentrations of components when they are used in the form of a solution, or suspension or slurry in a solvent are appropriately selected depending on the ability of an apparatus

for feeding components into a polymerization reactor, and it is generally desirable that the concentration of the transition metal compound (A) is usually from 0.001 to 200 mmol/L, more preferably from 0.01 to 100 mmol/L, further preferably from 0.05 to 50 mmol/L, the concentration of (B) is, in terms of an Al atom, usually from 0.01 to 5000 mmol/L, more preferably from 0.1 to 2500 mmol/L, further preferably 0.1 to 2000 mmol/L, and the concentration of (C) is usually from 0.001 to 500 mmol/L, more preferably from 0.01 to 250 mmol/L, further preferably from 0.05 to 100 mmol/L.

The process for producing an addition polymer of the present invention can be applied to various polymerization methods of addition polymerizable monomers, and of them, is suitable as a process for producing a polymer of an olefin and/or alkenyl aromatic hydrocarbon.

As the olefins herein referred to, olefins having 2 to 20 carbon atoms, particularly, ethylene, α -olefins having 3 to 20 carbon atoms, diolefins having 4 to 20 carbon atoms, and the like can be used, and two or more olefins can be used simultaneously. Specific examples of the olefin include, for example, linear olefins such as ethylene, propylene, 1-butene, 1-pentene, 1-hexene, 1-heptene, 1-octene, 1-nonene, 1-decene and the like, branched olefins such as 3-methyl-1-butene, 3-methyl-1-pentene, 4-methyl-1-pentene, 5-methyl-1-hexene and the like, vinylcyclohexane, and the like. Specific examples of combinations of monomers in conducting copolymerization, include, for example, ethylene and propylene, ethylene and 1-butene, ethylene and 1-hexene, ethylene and 1-octane,

propylene and 1-butene, and the like.

As the above-mentioned alkenyl aromatic hydrocarbon, alkenyl aromatic hydrocarbons having an aromatic hydrocarbon group of 6 to 25 carbon atoms are preferable, and specific
5 examples thereof include, for example, a phenyl group, tolyl group, xylyl group, tert-butylphenyl group, vinylphenyl group, naphthyl group, phenanthryl group, anthracenyl group and the like. The aromatic hydrocarbon group is preferably a phenyl group, tolyl group, xylyl group, tert-butylphenyl group,
10 vinylphenyl group or naphthyl group. Examples of the alkenyl aromatic hydrocarbon include alkylstyrenes such as p-methylstyrene, m-methylstyrene, o-methylstyrene, p-ethylstyrene, m-ethylstyrene, o-ethylstyrene, 2,4-dimethylstyrene, 2,5-dimethylstyrene,
15 3,4-dimethylstyrene, 3,5-dimethylstyrene, 3-methyl-5-ethylstyrene, p-tert-butylstyrene, p-sec-butylstyrene and the like; alkenylbenzenes such as styrene, 2-phenylpropylene, 2-phenylbutene and the like; and vinylnaphthalenes such as 1-vinylnaphthalene and the like.
20 Among them, styrene, p-methylstyrene, m-methylstyrene, o-methylstyrene, p-tert-butylstyrene, 2-phenylpropylene and 1-vinylnaphthalene are preferred, and styrene is particularly preferred.

The process for producing an addition polymer of the present
25 invention is suitable for producing a copolymer of the above-mentioned olefin with the above-mentioned alkenyl aromatic hydrocarbon, and as the combination of monomers in this process, combinations of ethylene and styrene, propylene

and styrene, ethylene, propylene and styrene are listed, and particularly, this process is suitable for producing a copolymer of ethylene with styrene.

The process for producing an addition polymer of the present invention is particularly suitable for producing copolymers of ethylene and α -olefins having 3 to 8 carbon atoms, particularly, linear low density polyethylenes.

The polymerization method should also not be limited particularly, and for example, solvent polymerization using, as a solvent, an aliphatic hydrocarbon such as butane, pentane, hexane, heptane, octane or the like, an aromatic hydrocarbon such as benzene, toluene or the like, or a halogenated hydrocarbon such as methylene dichloride or the like, or slurry polymerization, gas phase polymerization in gaseous monomer(s), and the like can be applied, and any of continuous polymerization and batch-wise polymerization is possible.

The polymerization temperature can be usually from -50°C to 200°C , and is particularly, preferably from -20°C to 100°C , and the polymerization pressure is usually preferably from normal pressure to 60 kg/cm^2 . The polymerization time is generally determined appropriately depending on the kind of the intended polymer, and the reaction apparatus, and usually, it can be from 1 minute to 20 hours. Further, in the present invention, a chain transfer agent such as hydrogen or the like can also be added for controlling the molecular weight of a copolymer.

EXAMPLE

The present invention is illustrated using the following Examples and Comparative Examples in more detail below, but not limited thereto.

The properties of polymers in Examples were measured by the following methods.

(1) Intrinsic viscosity $[\eta]$ was measured by using a Ubbelohde viscometer in a tetralin solution at 135°C.

(2) Weight-average molecular weight (M_w), number-average molecular weight (M_n) and molecular weight distribution (M_w/M_n): The molecular weights were measured by gel permeation chromatography (GPC) under the following conditions. A calibration curve was made by using standard polystyrene. The molecular weight distribution was evaluated as a ratio (M_w/M_n) of the weight-average molecular weight (M_w) to the number-average molecular weight (M_n).

Machine 150CV type, manufactured by Millipore Waters

Column Shodex M/S 80

Measurement temperature 145°C, solvent:
orthodichlorobenzene

Sample concentration 5 mg/8 ml

(3) Content of repeating units derived from α -olefin in ethylene- α -olefin copolymer (α -olefin content):

It was measured from the specific absorptions of ethylene and α -olefin using an infrared spectrophotometer (IR-810, manufactured by Nippon Bunko Kogyo K.K.), and represented in terms of short chain branch (SCB) numbers per 1000 carbons.

(4) Melting point of copolymer:

(In the case of Example 9)

It was measured by using Seiko SSC-5200 under the following conditions.

Cooling: 20°C to -50°C (20°C/min.), kept for 5 minutes

Heating: -50°C to 200°C (20°C/min.), kept for 5 minutes

5 Cooling: 200°C to -50°C (20°C/min.), kept for 5 minutes

Measurement: -50°C to 300°C (20°C/min.)

(In other case than Example 9)

It was measured by using Seiko SSC-5200 under the following conditions.

10 Heating: 40°C to 150°C (10°C/min.), kept for 5 minutes

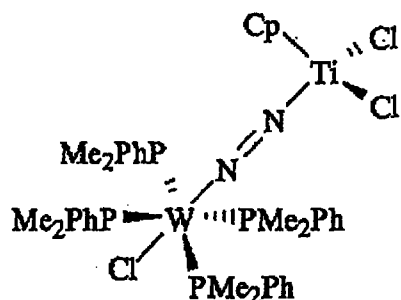
Cooling: 150°C to 10°C (5°C/min.), kept for 10 minutes

Measurement: 10°C to 160°C (5°C/min.)

Example 1(1)

15 Synthesis of [chlorotetrakis(dimethylphenylphosphine)
tungsten](μ -dinitrogen)[dichloro(cyclopentadienyl)
titanium]

225.2 mg (0.284 mmol) of cis-bis(dinitrogen)tetrakis
(dimethylphenylphosphine)tungsten and 62.2 mg (0.284 mmol) of
20 trichloro(cyclopentadienyl)titanium were dissolved in 5 ml of
benzene, to recognize generation of a gas from the solution,
and the color of the solution changed from orange to dark green.
This solution was stirred for 8 hours, then, the solvent was
distilled off to obtain dark green solid. The dark green solid
25 was subjected to extraction with methylene chloride, and the
resulted extracted substance was re-crystallized by using a
methylene chloride/ether mixed solvent, to obtain 196.0 mg of
a black crystal. Yield: 70%.



$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 1.63 (s, 24H, PMe), 6.44 (s, 5H, Cp),
7.27-7.46 (m, 20H, Ph);

$^{31}\text{P}\{^1\text{H}\}\text{-NMR}(\text{CDCl}_3)$ δ : -24.5 (s with ^{183}W satellites,

5 $J_{\text{WP}} = 277$ Hz);

IR(KBr) 1408(m) cm^{-1} .

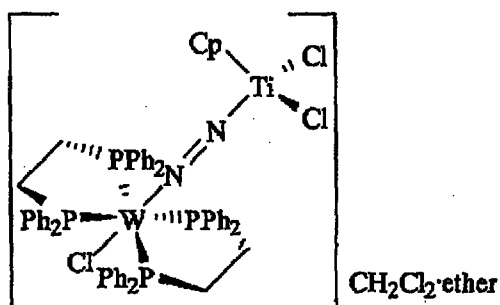
Anal. Calcd. for $\text{C}_{37}\text{H}_{49}\text{Cl}_3\text{N}_2\text{P}_4\text{TiW}$: C, 45.17; H, 5.02;
N, 2.85. Found: C, 45.50; H, 5.29; N, 2.70

Example 1(2)

10 Synthesis of [chlorobis(diphenylphosphinoethane)
tungsten](μ -dinitrogen)[dichloro(cyclopentadienyl)
titanium] (diethyl ether)(dichloromethane);

198.0 mg (0.191 mmol) of trans-bis(N_2)bis
(diphenylphosphinoethane)tungsten and 41.9 mg (0.191 mmol) of
15 trichloro(cyclopentadienyl)titanium were dissolved in 5 ml of
tetrahydrofuran under room temperature, to recognize
generation of a gas from the solution, and the color of this
solution changed from orange to dark green. This solution was
stirred for 8 hours, then, the solvent was distilled off to
20 obtain dark green solid. The dark green solid was subjected
to extraction with methylene chloride, and the resulted
extracted substance was re-crystallized by using a methylene

chloride/ether mixed solvent, to obtain 166.2 mg of a black crystal. Yield: 63%.



- 5 ¹H NMR (CDCl₃) δ : 1.21 (s, 6H, J=6.8Hz, Me of ether), 2.46, 2.89 (br, 4H, each, CH₂ of dppe), 3.48 (q, 4H, J=7.3Hz, CH₂ of ether), 5.29 (s, 2H, CH₂Cl₂), 5.70 (s, 5H, Cp), 6.66-7.67 (m, 40H, Ph);
- ³¹P{¹H} NMR (CDCl₃) δ : 36.0 (s with ¹⁸³W satellites,
- 10 J_{WP} = 284 Hz);
- IR (KBr) 1412 (m) cm⁻¹.
- Anal. Calcd for C₆₃H₆₅Cl₅N₂OP₄TiW: C, 53.69; H, 4.72; N, 2.02. Found: C, 53.49; H, 4.60; N, 2.19

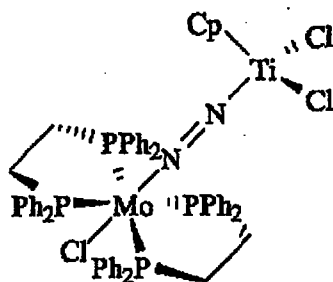
Example 1(3)

- 15 Synthesis of [chlorobis(diphenylphosphinoethane)molybdenum] (μ-dinitrogen) [dichloro(cyclopentadienyl)titanium];

194.7 mg (0.205 mmol) of trans-bis(dinitrogen)bis(diphenylphosphinoethane)molybdenum and 45.0 mg (0.205 mmol) of trichloro(cyclopentadienyl)titanium were dissolved in 5 ml of tetrahydrofuran under room temperature, to recognize generation of a gas from the solution, and the color of this gas changed from orange to dark green. This solution was stirred

20

for 8 hours, then, the solvent was distilled off to obtain dark green solid. The dark green solid was subjected to extraction with methylene chloride, and the resulted extracted substance was re-crystallized by using a methylene chloride/ether mixed
 5 solvent, to obtain 182.3 mg of a black crystal. Yield: 78%.



$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 2.31, 2.66 (br, 4H each, CH_2 of dppe), 5.73 (s, 5H, Cp), 6.78-7.77 (m, 40H, Ph);

10 $^{31}\text{P}\{^1\text{H}\}\text{NMR}(\text{CDCl}_3)$ δ : 52.9 (s);

IR(KBr) 1416(m) cm^{-1} .

Anal. Calcd for $\text{C}_{57}\text{H}_{53}\text{Cl}_3\text{MoN}_2\text{P}_4\text{Ti}$: C, 60.05; H, 4.69; N, 2.46. Found: C, 60.13; H, 5.06; N, 2.22

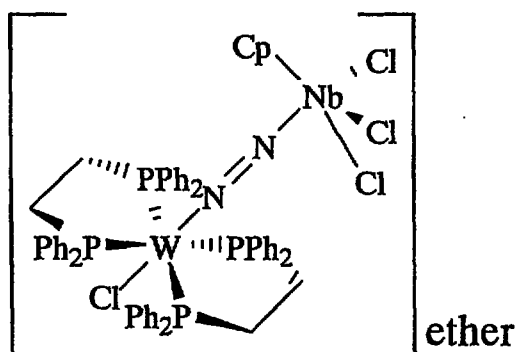
Example 1(4)

15 Synthesis of

[chlorobis(diphenylphosphinoethane)tungsten] (μ -dinitrogen)[trichloro(cyclopentadienyl)niobium];

164.6 mg (0.159 mmol) of trans-bis(dinitrogen)bis (diphenylphosphinoethane)tungsten and 47.6 mg (0.159 mmol) of
 20 tetrachloro(cyclopentadienyl)niobium were dissolved in 5 ml of tetrahydrofuran under room temperature, to recognize generation of a gas from the solution, and the color of this gas changed from orange to black. This solution was stirred

for 1 hour, then, the solvent was distilled off to obtain dark green solid. The dark green solid was subjected to extraction with methylene chloride, and the resulted extracted substance was re-crystallized by using a methylene chloride/ether mixed solvent, to obtain 164.8 mg of a black crystal. Yield: 75%.



$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 1.12 (t, 6H, $J = 7.0$ Hz, Me of ether), 2.68, 2.90 (br, 4H each, CH_2 of dppe), 3.38 (t, 4H, $J = 7.2$ Hz, CH_2 of ether), 5.86 (s, 5H, Cp), 6.67-7.66 (m, 40H, Ph);

$^{31}\text{P}\{^1\text{H}\}\text{-NMR}(\text{CDCl}_3)$ δ : 34.1 (s with ^{183}W satellites, $J_{\text{WP}} = 275$ Hz);

IR(KBr) 1383(m) cm^{-1} .

Anal. Calcd for $\text{C}_{61}\text{H}_{63}\text{Cl}_4\text{N}_2\text{NbOP}_4\text{W}$: C, 52.99; H, 4.59; N, 2.03. Found: C, 52.88; H, 4.22; N, 2.32

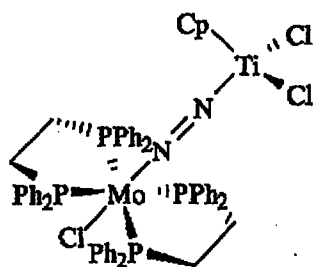
Example 1(5)

Synthesis of [chlorobis(diethylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(cyclopentadienyl)titanium];

326 mg (0.50 mmol) of trans-bis(dinitrogen)bis(diethylphosphinoethane)tungsten and 110 mg (0.50 mmol) of trichloro(cyclopentadienyl)titanium were dissolved in 7.5 ml of tetrahydrofuran under room temperature, and the mixture was

stirred for 21 hours. The solvent was distilled off from the resulted violet black solution and the resultant solution was subjected to extraction with 8 ml of dichloromethane, then, the resulted extracted solution was concentrated to 1.5 ml.

5 To this was added slowly 8.5 ml of hexane, to obtain 320 mg of a black crystal. Yield: 76%.



$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 6.17 (s, 5H, C₅H₅), 2.18-1.76 (m, 24H, PCH₂), 1.29-1.13 (m, 24H, PCH₂CH₃);

10 $^{31}\text{P}\{^1\text{H}\}\text{-NMR}(\text{CDCl}_3)$ δ : 30.99 (s with 183W satellites, $J(\text{PW}) = 270 \text{ Hz}$);

IR(KBr) 1406.3(s) cm^{-1} .

Anal. Calcd. for C₂₅H₅₃Cl₃N₂P₄TiW: C, 35.59; H, 6.33;

15 N, 3.32. Found: C, 35.50; H, 6.44; N, 3.50.

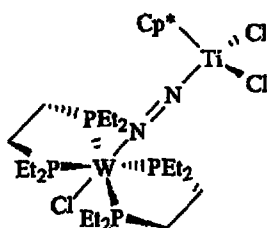
Example 1(6)

Synthesis of [chlorobis(diethylphosphinoethane) tungsten] (μ -dinitrogen) [dichloro(cyclopentadienyl) titanium];

0.50 mg (0.77 mmol) of trans-bis(dinitrogen)bis

20 (diethylphosphinoethane)tungsten and 0.22 g (0.76 mmol) of trichloro(pentamethylcyclopentadienyl)titanium were dissolved in 15 ml of tetrahydrofuran under room temperature, and the mixture was stirred for 15 hours. In this procedure,

foaming was caused by gas generation from the red solution, and it changed to dark green suspension. The solvent was distilled off from this suspension and the suspension was subjected to extraction with 10 ml of dichloromethane, then, the resulted extracted solution was concentrated, and to this was added slowly diethyl ether to cause re-crystallization giving 0.48 g of a dark green crystal. Yield: 68%.

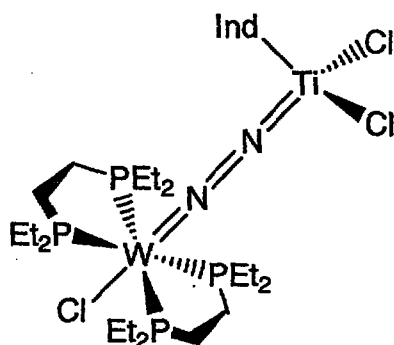


$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 2.22-1.70 (m, 24H, PCH₂), 1.97 (s, 15H, Cp-CH₃), 1.30-1.10 (m, 24H, PCH₂CH₃);
 IR(KBr) 1406.5(s)cm⁻¹.

Example 1(7)

Synthesis of [chlorobis(diethylphosphinoethane) tungsten](μ -dinitrogen)[dichloro(indenyl)titanium];
 0.252 mg (0.387 mmol) of trans-bis(dinitrogen)bis(diethylphosphinoethane)tungsten and 0.105 g (0.390 mmol) of trichloro(indenyl)titanium were dissolved in 10 ml of tetrahydrofuran under room temperature, and the mixture was stirred for 22 hours. In this procedure, foaming was caused by gas generation from the red solution, and it changed to dark green suspension. The solvent was distilled off from this suspension and the resulted solid was subjected to extraction with 9 ml of dichloromethane, then, the resulted extracted

solution was concentrated, and to this was added slowly diethyl ether to cause re-crystallization giving 0.25 g of a dark green crystal. Yield: 72%.



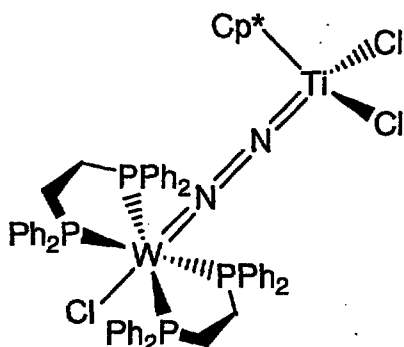
- 5 $^1\text{H-NMR}(\text{CDCl}_3)$ δ : 7.34(s, 2H, Ind), 6.98(s, 2H, Ind), 6.62(s, 1H, Ind), 6.17(s, 2H, Ind), 2.18-1.60(m, 24H, P-CH₂), 1.26-1.03(m, 24H, CH₃-CH₂).

Example 1(8)

- Synthesis of [chlorobis(diphenylphosphinoethane)
10 tungsten](μ -dinitrogen)[dichloro
(pentamethylcyclopentadienyl) titanium];

- 480 mg (0.46 mmol) of trans-bis(dinitrogen)bis
(diphenylphosphinoethane)molybdenum and 140 mg (0.48 mmol) of
trichloro(pentamethylcyclopentadienyl)titanium were
15 dissolved in 25 ml of tetrahydrofuran under room temperature,
to recognize generation of a gas from the solution, and the
color of this solution changed from red to dark green. This
solution was stirred for 18 hours, then, the solvent was
distilled off to obtain dark green solid. The dark green solid
20 was subjected to extraction with methylene chloride, and the
resulted extracted substance was re-crystallized by using a
methylene chloride/ether mixed solvent, to obtain 490 mg of

a dark green fine crystal. Yield: 82%.



$^1\text{H-NMR}(\text{CDCl}_3)$ δ : 1.72 (s, 15H, Cp-CH₃), 2.36, 2.87 (m, 4H, each, CH₂ of dppe), 6.57 (m, 8H, Ph), 6.89 (t, 8H, Ph), 7.09 (t, 4H, Ph), 7.38 (t, 4H, Ph), 7.45 (t, 8H, Ph), 7.76 (m, 8H, Ph); $^{31}\text{P}\{^1\text{H}\}$ NMR (CDCl₃) δ : 38.7 (s with ^{183}W satellites, $J_{\text{WP}} = 287$ Hz);

IR(KBr) 1409(m)cm⁻¹.

Anal. Calcd for C₆₃H₆₅Cl₅N₂OP₄TiW: C, 57.36; H, 4.89; N, 2.16. Found: C, 57.06; H, 5.16; N, 2.17

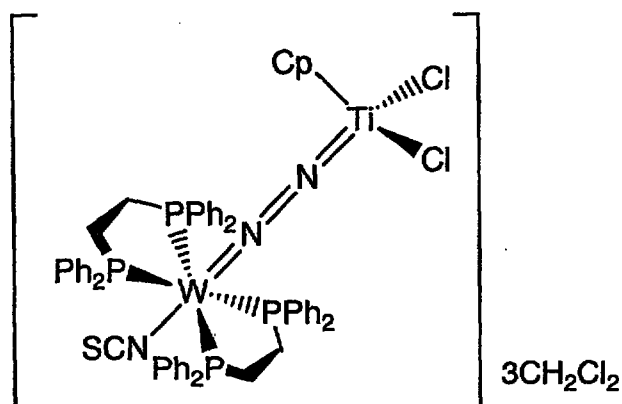
Example 1(9)

Synthesis of

[bis(diphenylphosphinoethane)(isothiocyanide)tungusten](μ -dinitrogen)[dichloro(cyclopentadienyl)titanium];

109.7 mg (0.084 mmol) of (tetra-n-butylammonium) [trans-(dinitrogen)bis (diphenylphosphinoethane)tungusten and 18.4 mg (0.084 mmol) of trichloro(cyclopentadienyl)titanium were dissolved in 3 ml of benzene under room temperature, and the solution was stirred for 17 hours, then, the solvent was distilled off to obtain a green solid. The green solid was subjected to extraction with methylene chloride, and the resulted extracted substance was re-crystallized by using a methylene chloride/ether mixed

solvent, to obtain 68.0 mg of a dark green crystal. Yield: 54%.



- 5 $^1\text{H-NMR}(\text{CDCl}_3)$ δ : 1.72 (s, 15H, Cp-CH₃), 2.49, 2.81 (m, 4H, each, CH₂ of dppe), 5.80 (s, 5H, Cp), 6.55 - 7.63 (m, 40H, Ph);
 $^{31}\text{P}\{^1\text{H}\}$ NMR (CDCl_3) δ : 38.2 (s with ^{183}W satellites, $J_{\text{WP}} = 284$ Hz);
 IR(KBr) 1441 (m) cm^{-1} .

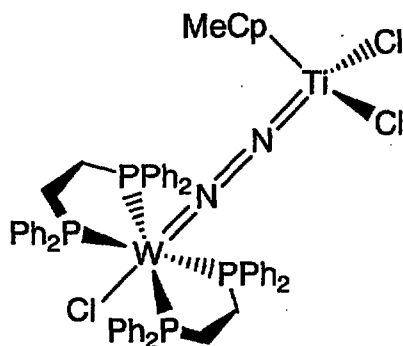
- 10 Anal. Calcd for C₆₃H₆₅Cl₅N₂OP₄TiW: C, 48.67; H, 3.95; N, 2.79.
 Found: C, 49.02; H, 3.77; N, 2.88.

Example 1(10)

Synthesis of [chlorobis(diphenylphosphinoethane) tungusten](μ -dinitrogen)[dichloro(methylcyclopentadienyl) titanium];

- 15 500 mg (0.48 mmol) of [trans-bis(dinitrogen)bis(diphenylphosphinoethane)tungusten and 110 mg (0.48 mmol) of trichloro(methylcyclopentadienyl)titanium were dissolved in 25 ml of tetrahydrofuran under room temperature, to recognize generation of a gas from the solution, and the color of this
 20 solution changed from orange to dark brown. The solution was stirred for 24 hours, then, the solvent was distilled off to obtain a black solid. The black solid was subjected to extraction with methylene chloride, and the resulted extracted

substance was re-crystallized by using a methylene chloride/ether mixed solvent, to obtain 540 mg of a bright black cubic crystal. Yield: 90%.



5

$^1\text{H-NMR}$ (toluene- d_8) δ : 2.00 (s, 3H, Cp- CH_3), 2.39, 2.81 (m, 4H, each, CH_2 of dppe), 5.27, 5.47 (t, 2H, each, Cp-H), 6.59 (s, 8H, Ph), 6.87 (m, 8H, Ph), 7.04 (m, 4H, Ph), 7.30 (m, 12H, Ph), 7.60 (s, 8H, Ph);

10 $^{31}\text{P}\{^1\text{H}\}$ -NMR(CD_2Cl_2) δ : 39.9 (s with ^{183}W satellites, $J_{\text{WP}} = 286$ Hz);

IR(KBr) 1411(m) cm^{-1} .

Anal. Calcd. for $\text{C}_{63}\text{H}_{65}\text{Cl}_5\text{N}_2\text{OP}_4\text{TiW}$: C, 56.09; H, 4.46; N, 2.26. Found: C, 56.02; H, 4.56; N, 2.21.

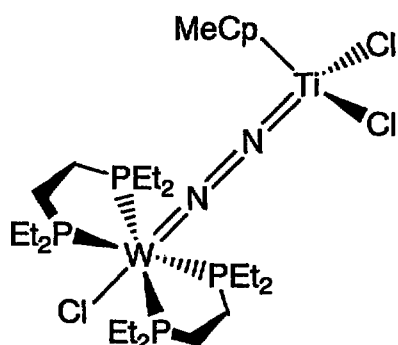
15 Example 1(11)

Synthesis of [chlorobis(diethylphosphinoethane) tungusten](μ -dinitrogen)[dichloro(methylcyclopentadienyl) titanium];

260 mg (0.40 mmol) of [trans-bis(dinitrogen)bis(diethylphosphinoethane)tungusten and 90 mg (0.39 mmol) of trichloro(methylcyclopentadienyl)titanium were dissolved in 15 ml of tetrahydrofuran under room temperature, to recognize generation of a gas from the solution, and the color of this solution changed from orange to dark brown. The solution was stirred for 14 hours, then, the solvent was distilled off to

20

obtain a black solid. The black solid was subjected to extraction with methylene chloride, and the resulted extracted substance was re-crystallized by using a methylene chloride/ether mixed solvent, to obtain 308 mg of a dark blue
 5 needle-like crystal. Yield: 92%.



$^1\text{H-NMR}$ (toluene- d_8) δ : 0.93, 1.15 (m, 4H, each, CH_2 of depe), 1.68, 1.85 (m, 4H, each, CH_2 of depe), 1.41, 1.97 (m, 8H, each, CH_2 of depe), 2.27 (s, 3H, Cp-CH_3), 5.82, 6.01 (t, 2H, each, Cp-H);

$^{31}\text{P}\{^1\text{H}\}$ -NMR(CD_2Cl_2) δ : 36.0 (s with ^{183}W satellites, $J_{\text{WP}} = 269$ Hz);

IR(KBr) 1407(m) cm^{-1} .

15 Anal. Calcd. for $\text{C}_{63}\text{H}_{65}\text{Cl}_5\text{N}_2\text{OP}_4\text{TiW}$: C, 36.41; H, 6.46; N, 3.27. Found: C, 36.82; H, 6.69; N, 3.21.

Example 2

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 200 ml of toluene as a solvent was charged into
 20 this, and the reactor was heated up to 60°C . After heating, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.50 mmol (mol number in terms of aluminum atom; hereinafter the same) of a

solution of methylisobutylaluminumoxane in toluene (MMAO3A manufactured by Tosoh Akzo Ltd.; hereinafter, abbreviated simply as MMAO") was charged, and subsequently, 0.1 μ mol of [chlorotetrakis(dimethylphenylphosphine)tungsten](μ -dinitrogen)[dichloro(cyclopentadienyl)titanium] synthesized in the above-mentioned Example 1(1) was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, an ethylene polymer having a melting point of 136.7°C was produced at a rate of 2.3×10^7 g per one hour per 1 mol of a titanium atom.

Example 3

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 200 ml of toluene as a solvent was charged into this, and the reactor was heated up to 60°C. After heating, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.25 mmol of triisobutylaluminum was charged, subsequently, 0.1 μ mol of [chlorotetrakis(dimethylphenylphosphine)tungsten](μ -dinitrogen) [dichloro(cyclopentadienyl)titanium] was charged, and subsequently, 1.0 μ mol of triphenylmethyltetrakis(pentafluorophenyl)borate (called as "[triphenylcarbenium][tetrakis(pentafluorophenyl)borate]) was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, an ethylene polymer having a melting point of 136.8°C was produced at a rate of 1.8×10^7

g per one hour per 1 mol of a titanium atom.

Example 4

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the reaction vessel was heated up to 60°C. After temperature raise, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.05 mmol of MMAO was charged, and subsequently, 0.5 μ mol of [chlorobis(diphenylphosphinoethane)molybdenum](μ -dinitrogen)[dichloro(cyclopentadienyl)titanium] synthesized in the above-mentioned Example 1(3) was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 18.3, $[\eta]$ of 4.9 dl/g, a M_w of 3.5×10^5 , a M_w/M_n of 2.14 and melting points of 92.0°C and 119.6°C was produced at a rate of 1.7×10^7 g per one hour per 1 mol of a titanium atom.

Comparative Example 1

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the reaction vessel was heated up to 60°C. After temperature raise, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.05 mmol of MMAO

was charged, and subsequently, 0.5 μmol of (cyclopentadienyl)titanium trichloride was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

- 5 As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 18.6, a M_w of 2.3×10^5 , a M_w/M_n of 2.43 and melting points of 99.0°C and 115.6°C was produced at a rate of 1.6×10^7 g per one hour per 1 mol of a titanium atom.

Example 5

- 10 An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the reaction vessel was heated up to 60°C. After heating, ethylene
15 was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.25 mmol of triisobutylaluminum was charged, subsequently, 0.1 μmol of [chlorobis(diphenylphosphinoethane)molybdenum](μ
2-N₂)[dichloro(cyclopentadienyl)titanium] was charged, and
20 subsequently, 1.0 μmol of triphenylmethyltetrakis(pentafluorophenyl)borate was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

- 25 As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 16.4, $[\eta]$ of 7.5 dl/g, M_w of 4.4×10^5 , M_w/M_n of 2.48 and melting point of 121.0°C was produced at a rate of 3.8×10^7 g per one hour per 1 mol of a titanium atom.

Example 6

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 185 ml of toluene as a solvent was charged into this, 15 ml of 1-hexene as an α -olefin was charged, and the reaction vessel was heated up to 180°C. After the heating, ethylene was fed while controlling the ethylene pressure at 2.5 MPa, and after the system was stabilized, 0.90 mmol of MMAO was charged, subsequently, a mixture of 0.5 μ mol of [chlorobis(diphenylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(cyclopentadienyl)titanium] and 0.10 mmol of triisobutylaluminum was charged. Polymerization was conducted for 2 minutes while controlling the temperature at 180°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 12.6, $[\eta]$ of 2.7 dl/g, and melting point of 124.6°C was produced at a rate of 1.3×10^7 g per one hour per 1 mol of a titanium atom.

Example 7

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the reactor was heated up to 60°C. After the heating, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.15 mmol of triisobutylaluminum was charged, subsequently, a mixture of 0.05 μ mol of [chlorobis(diethylphosphinoethane)tungsten](μ

-dinitrogen)[dichloro(cyclopentadienyl)titanium]
synthesized in the above-mentioned Example 1(5) and 0.10 mmol
of triisobutylaluminum was charged, and subsequently, 1.0 μ
mol of triphenylmethylnetrakis(pentafluorophenyl)borate was
5 charged. Polymerization was conducted for 60 minutes while
controlling the temperature at 60°C.

As a result of polymerization, a copolymer of ethylene and
1-hexene having a SCB of 19.0 and $[\eta]$ of 3.7 dl/g was produced
at a rate of 2.3×10^8 g per one hour per 1 mol of a titanium
10 atom.

Comparative Example 2

An autoclave reactor having an inner volume of 0.4 liter
equipped with a stirrer was dried under vacuum and purged with
argon, then, 185 ml of toluene as a solvent was charged into
15 this, 15 ml of 1-hexene as an α -olefin was charged, and the
reactor was heated up to 180°C. After the heating, ethylene
was fed while controlling the ethylene pressure at 2.5 MPa,
and after the system was stabilized, 1.0 mmol of MMAO was charged,
and subsequently, 2.5 μ mol of (cyclopentadienyl)titanium
20 trichloride was charged. Polymerization was conducted for 2
minutes while controlling the temperature at 180°C.

As a result of polymerization, a copolymer of ethylene and
1-hexene having a Mw of 2.3×10^5 , Mw/Mn of 18.8 and a melting
point of 124.3°C was produced at a rate of 2.8×10^4 g per one
25 hour per 1 mol of a titanium atom.

Example 8

A glass vessel having an inner volume of 0.1 liter was dried
under vacuum and purged with argon, then, 13.8 ml of toluene

as a solvent was charged into this, 18.3 ml of styrene was charged, and the reaction vessel was heated up to 50°C. After the heating, 1.0 mmol of triisobutylaluminum was charged, subsequently, 10 μ mol of [chlorobis(diphenylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(cyclopentadienyl)niobium] synthesized in the above-mentioned Example 1(4) was charged, and subsequently, 30 μ m of N,N-dimethylaniliniumtetrakis(pentafluorophenyl) borate was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 50°C.

As a result of polymerization, a styrene polymer having a Mw of 2.1×10^4 and Mw/Mn of 2.21 was produced at a rate of 1.1×10^6 g per one hour per 1 mol of a niobium atom.

Example 9

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 165 ml of toluene as a solvent was charged into this, 35 ml of styrene was charged, and the reaction vessel was heated up to 50°C. After the heating, ethylene was fed while controlling the ethylene pressure at 0.8 MPa, and after the system was stabilized, 2.0 mmol of MMAO was charged, subsequently, 10 μ mol of [chlorotetrakis(dimethylphenylphosphine)tungsten](μ -dinitrogen)[dichloro(cyclopentadienyl)titanium] was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 50°C.

As a result of polymerization, a copolymer of ethylene and styrene having a Mw of 4.3×10^5 , Mw/Mn of 1.96 (converted to polystyrene) and melting point of 108.8°C was produced at a

rate of 4.8×10^5 g per one hour per 1 mol of a titanium atom.

Example 10

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 40 ml of toluene as a solvent was charged into this, and the reactor was cooled down to -30°C . After the cooling, 1.0 mmol of triisobutylaluminum was charged, subsequently, 1.0 μmol of [chlorotetrakis(dimethylphenylphosphine)molybdenum](μ -dinitrogen)[dichloro(cyclopentadienyl)titanium] was charged, and subsequently, 6.0 μm of triphenylmethylnetrakis(pentafluorophenyl)borate was charged, and lastly, 80 g of propylene was introduced. After introduction of propylene, the autoclave was heated by polymerization heat, to raise the temperature up to 25°C . After introduction of propylene, polymerization was conducted for 8 minutes.

As a result of polymerization, a propylene polymer having a Mw of 7.4×10^4 , Mw/Mn of 1.98, [mm] of 8.3%, [mr] of 46.3% and [rr] of 45.4% (triad showing stereo regularity) was produced at a rate of 2.0×10^8 g per one hour per 1 mol of a titanium atom.

Example 11

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 200 ml of n-hexane as a solvent was charged into this, and the reactor was heated up to 60°C . After temperature raise, ethylene was fed while controlling the ethylene pressure

at 0.6 MPa, and after the system was stabilized, 0.50 mmol of tributylaluminum was charged, subsequently, a mixture of 0.01 μ mol of [chlorobis(diethylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(pentamethylcyclopentadienyl)titanium]
 5] synthesized in the above-mentioned Example 1(6) and 0.01 mmol of triisobutylaluminum was charged, subsequently, 1.0 μ mol triphenylmethylnetrakis(pentafluorophenyl)borate was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

10 As a result of polymerization, an ethylene polymer having a Mw of 1.1×10^6 and Mw/Mn of 2.21 was produced at a rate of 2.3×10^8 g per one hour per 1 mol of a titanium atom.

Example 12

An autoclave reactor having a content volume of 0.4 liter
 15 equipped with a stirrer was dried under vacuum and purged with argon, then, 190 ml of n-hexane as a solvent was charged into this, 10 ml of 1-hexene as an α -olefin was charged, and the reaction vessel was heated up to 60°C. After the heating, ethylene was fed while controlling the ethylene pressure at
 20 0.6 MPa, and after the system was stabilized, 0.50 mmol of tributylaluminum was charged, subsequently, a mixture of 0.01 μ mol of [chlorobis(diethylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(pentamethylcyclopentadienyl)titanium] synthesized in the above-mentioned Example 1(6) and 0.01 mmol
 25 of triisobutylaluminum was charged, subsequently, 1.0 μ mol triphenylmethylnetrakis(pentafluorophenyl)borate was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 29.2, Mw of 4.2×10^5 and Mw/Mn of 2.68 was produced at a rate of 8.0×10^7 g per one hour per 1 mol of a titanium atom.

5 Example 13

An autoclave reactor having a content volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 190 ml of toluene as a solvent was charged into this, 10 ml of 1-hexene as an α -olefin was charged, and the
 10 reaction vessel was heated up to 60°C. After the heating, ethylene was fed while controlling the ethylene pressure at 2.5 MPa, and after the system was stabilized, 0.25 mmol of tributylaluminum was charged, subsequently, 0.025 μ mol of [chlorobis(diethylphosphinoethane)tungsten](μ
 15 -dinitrogen)[dichloro(pentamethylcyclopentadienyl)titanium] synthesized in the above-mentioned Example 1(6) was charged, and subsequently, 1.2 μ mol triphenylmethylnetrakis (pentafluorophenyl)borate was charged. Polymerization was conducted for 60 minutes while controlling the temperature at
 20 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 16.5, Mw of 1.9×10^6 and Mw/Mn of 2.85 was produced at a rate of 1.9×10^9 g per one hour per 1 mol of a titanium atom.

25 Example 14

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into

this, 2 ml of 1-hexene as an α -olefin was charged, and the reaction vessel was heated up to 60°C. After temperature raise, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 1.0 mmol of MMAO
 5 was charged, and subsequently, 0.02 μ mol of [chlorobis(diethylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(indenyl)titanium] synthesized in the above-mentioned Example 1(7) was charged. Polymerization was conducted for 60 minutes while controlling the temperature at
 10 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 15.6, Mw of 6.4×10^5 , and Mw/Mn of 4.26 was produced at a rate of 8.0×10^7 g per one hour per 1 mol of a titanium atom.

15 Example 15

An autoclave reactor having a content volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the
 20 reaction vessel was heated up to 60°C. After the heating, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.25 mmol of tributylaluminum was charged, subsequently, 0.025 μ mol of [chlorobis(diphenylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(pentamethylcyclopentadienyl)titanium]
 25] synthesized in the above-mentioned Example 1(8) was charged, and subsequently, 1.0 μ mol triphenylmethylnetrakis(pentafluorophenyl)borate was charged. Polymerization was

conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 16.1, Mw of 9.2×10^5 and Mw/Mn of 5.99 was produced at a rate of 1.4×10^8 g per one hour per 1 mol of a titanium atom.

Example 16

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the reactor was heated up to 60°C. After temperature raise, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 1.0 mmol of MMAO was charged, and subsequently, 0.025 μ mol of [chlorobis(diphenylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(pentamethylcyclopentadienyl)titanium] synthesized in the above-mentioned Example 1(8) was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 16.1, Mw of 8.0×10^5 , and Mw/Mn of 3.86 was produced at a rate of 1.1×10^8 g per one hour per 1 mol of a titanium atom.

Example 17

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 40 ml of toluene as a solvent was charged into

this, 80g of propylene was charged, and the reactor was heated up to 60°C. After the heating and after the system was stabilized, 1.0 mmol of MMAO was charged, and subsequently, 0.10 μ mol of [chlorobis(diphenylphosphinoethane)tungsten](μ

5 -dinitrogen)[dichloro(pentamethylcyclopentadienyl)titanium] synthesized in the above-mentioned Example 1(8) was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, a propylene polymer having
10 a Mw of 5.6×10^5 , Mw/Mn of 4.00, and [mm] of 13.2%, [mr] of 46.4% and [rr] of 40.4% as triad showing stereo regularity was produced at a rate of 1.6×10^7 g per one hour per 1 mol of a titanium atom.

Example 18

15 An autoclave reactor having a content volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 190 ml of toluene as a solvent was charged into this, 10 ml of 1-hexene as an α -olefin was charged, and the reactor was heated up to 60°C. After the heating, ethylene was
20 fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.25 mmol of trimethylaluminum was charged, subsequently, 0.025 μ mol of [bis(diphenylphosphinoethane)(isothiocyanide)tungsten](μ -dinitrogen)[dichloro(cyclopentadienyl)titanium](3 ·
25 dichloromethane) synthesized in the above-mentioned Example 1(9) was charged, and subsequently, 1.0 μ mol triphenylmethyltetrakis (pentafluorophenyl)borate was charged. Polymerization was conducted for 60 minutes while

controlling the temperature at 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 26.4, Mw of 3.4×10^5 and Mw/Mn of 2.23 was produced at a rate of 2.9×10^8 g per one hour per 1 mol of
5 a titanium atom.

Example 19

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into
10 this, 2 ml of 1-hexene as an α -olefin was charged, and the reactor was heated up to 60°C. After temperature raise, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 1.0 mmol of MMAO was charged, and subsequently, 0.0125 μ mol of
15 [bis(diphenylphosphinoethane)(isothiocyanide)tungsten] (μ -dinitrogen)[dichloro(cyclopentadienyl)titanium](3 · dichloromethane) synthesized in the above-mentioned Example 1(9) was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

20 As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 16.5, Mw of 7.9×10^5 , and Mw/Mn of 2.32 was produced at a rate of 3.6×10^8 g per one hour per 1 mol of a titanium atom.

Example 20

25 An autoclave reactor having a content volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the

reactor was heated up to 60°C. After the heating, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.25 mmol of tri-n-octylaluminum was charged, subsequently, 0.025 μ mol of [chlorobis(diphenylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(methylcyclopentadienyl)titanium] synthesized in the above-mentioned Example 1(10) was charged, and subsequently, 1.0 μ mol triphenylmethylnetrakis(pentafluorophenyl)borate was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 18.1, Mw of 1.3×10^6 and Mw/Mn of 3.11 was produced at a rate of 4.3×10^8 g per one hour per 1 mol of a titanium atom.

Example 21

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the reactor was heated up to 60°C. After temperature raise, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 1.0 mmol of MMAO was charged, and subsequently, 0.025 μ mol of [chlorobis(diphenylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(methylcyclopentadienyl)titanium] synthesized in the above-mentioned Example 1(10) was charged. Polymerization was conducted for 60 minutes while controlling

the temperature at 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 16.9, Mw of 9.0×10^5 , and Mw/Mn of 3.40 was produced at a rate of 2.2×10^8 g per one hour per 1 mol of a titanium atom.

Example 22

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 40 ml of toluene as a solvent was charged into this, 80g of propylene was charged, and the reactor was heated up to 60°C. After the heating and after the system was stabilized, 1.0 mmol of MMAO was charged, and subsequently, 0.10 μ mol of [chlorobis(diphenylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(methylcyclopentadienyl)titanium] synthesized in the above-mentioned Example 1(10) was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, a propylene polymer having a Mw of 1.3×10^5 , Mw/Mn of 1.80, and [mm] of 10.4%, [mr] of 44.2% and [rr] of 45.4% as triad showing stereo regularity was produced at a rate of 4.4×10^7 g per one hour per 1 mol of a titanium atom.

Example 23

An autoclave reactor having a content volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the reactor was heated up to 60°C. After the heating, ethylene was

fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.25 mmol of triisobutylaluminum was charged, subsequently, 0.025 μ mol of [chlorobis(diethylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(methylcyclopentadienyl)titanium] synthesized in the above-mentioned Example 1(11) was charged, and subsequently, 1.0 μ mol triphenylmethyltetrakis(pentafluorophenyl)borate was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 16.3, Mw of 2.8×10^5 and Mw/Mn of 1.95 was produced at a rate of 2.5×10^8 g per one hour per 1 mol of a titanium atom.

Example 24

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the reactor was heated up to 60°C. After temperature raise, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 1.0 mmol of MMAO was charged, and subsequently, 0.025 μ mol of [chlorobis(diethylphosphinoethane)tungsten](μ -dinitrogen)[dichloro(methylcyclopentadienyl)titanium] synthesized in the above-mentioned Example 1(11) was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 16.7, Mw of 4.8×10^5 , and Mw/Mn of 2.94 was produced at a rate of 1.5×10^8 g per one hour per 1 mol of a titanium atom.

5 Example 25

An autoclave reactor having an inner volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 170 ml of toluene as a solvent was charged into this, 30 ml of 1-hexene as an α -olefin was charged, and the
 10 reactor was heated up to 180°C. After temperature raise, ethylene was fed while controlling the ethylene pressure at 2.5 MPa, and after the system was stabilized, 1.0 mmol of MMAO was charged, and subsequently, 0.25 μ mol of
 [chlorobis(diethylphosphinoethane)tungsten](μ
 15 -dinitrogen)[dichloro(methylcyclopentadienyl)titanium]
 synthesized in the above-mentioned Example 1(11) was charged. Polymerization was conducted for 2 minutes while controlling the temperature at 180°C.

As a result of polymerization, a copolymer of ethylene and
 20 1-hexene having a SCB of 13.1, Mw of 6.8×10^4 , and Mw/Mn of 2.07 was produced at a rate of 6.4×10^8 g per one hour per 1 mol of a titanium atom.

Example 26

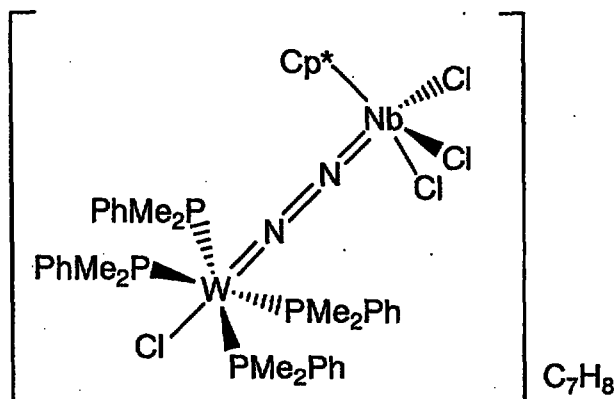
An autoclave reactor having a content volume of 0.4 liter
 25 equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the reactor was heated up to 60°C. After the heating, ethylene was

fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.35 mmol of triisobutylaluminum was charged, subsequently, 1.0 μ mol of [chlorobis(diphenylphosphinoethane)tungsten](μ -dinitrogen)[trichloro(cyclopentadienyl)niobium](diethyl ether) synthesized in the above-mentioned Example 1(4) was charged, and subsequently, 6.0 μ mol triphenylmethylnetrakis(pentafluorophenyl)borate was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C.

As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 10.1, M_w of 6.3×10^4 and M_w/M_n of 1.78 was produced at a rate of 4.9×10^6 g per one hour per 1 mol of a niobium atom.

Example 27

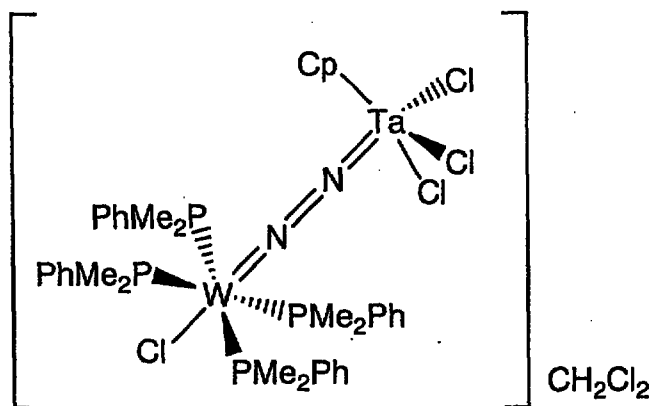
[chlorotetrakis(dimethylphenylphosphine)tungsten](μ -dinitrogen)[trichloro(pentamethylcyclopentadienyl)niobium](toluene) shown below as a transition metal compound was used.



An autoclave reactor having a content volume of 0.4 liter

equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the reactor was heated up to 60°C. After the heating, ethylene was
 5 fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.45 mmol of triisobutylaluminum was charged, subsequently, 2.0 μ mol of [chlorotetrakis(dimethylphenylphosphine)tungsten](μ -dinitrogen)[trichloro(pentamethylcyclopentadienyl)niobium
 10](toluene) synthesized in the above-mentioned Example 1(12) was charged, and subsequently, 3.0 μ mol triphenylmethyltetrakis (pentafluorophenyl)borate was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C. As a result of
 15 polymerization, a copolymer of ethylene and 1-hexene having a SCB of 8.3, Mw of 8.9×10^4 and Mw/Mn of 2.21 was produced at a rate of 1.0×10^6 g per one hour per 1 mol of a niobium atom.
 Example 28

[chlorotetrakis(dimethylphenylphosphine) tungsten](μ -dinitrogen)[trichloro(cyclopentadienyl)
 20 tantalum](dichloromethane) shown below as a transition metal compound was used.



An autoclave reactor having a content volume of 0.4 liter equipped with a stirrer was dried under vacuum and purged with argon, then, 198 ml of toluene as a solvent was charged into this, 2 ml of 1-hexene as an α -olefin was charged, and the reactor was heated up to 60°C. After the heating, ethylene was fed while controlling the ethylene pressure at 0.6 MPa, and after the system was stabilized, 0.45 mmol of triisobutylaluminum was charged, subsequently, 2.0 μ mol of [chlorotetrakis(dimethylphenylphosphine)tungsten](μ -dinitrogen)[trichloro(cyclopentadienyl)tantalum](dichloro methane) synthesized in the above-mentioned Example 1(13) was charged, and subsequently, 3.0 μ mol triphenylmethyltetrakis(pentafluorophenyl)borate was charged. Polymerization was conducted for 60 minutes while controlling the temperature at 60°C. As a result of polymerization, a copolymer of ethylene and 1-hexene having a SCB of 7.5, Mw of 6.4×10^4 and Mw/Mn of 1.74 was produced at a rate of 6.0×10^5 g per one hour per 1 mol of a tantalum atom.

20

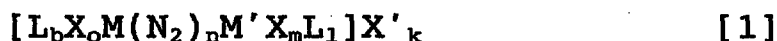
As described in detail above, the present invention provides a catalyst component for addition polymerization composed of

a transition metal compound which can manifest high activity,
a catalyst for addition polymerization prepared by using this
catalyst component for addition polymerization, and an
efficient process for producing an addition polymer, using this
5 catalyst for addition polymerization.



CLAIMS

1. A catalyst component for addition polymerization composed of a transition metal compound (A) of the following
5 general formula [1]:



wherein, M and M' each independently represent a transition metal atom of III to X group in the Periodic Table of Elements; X each independently represents a hydrogen atom, halogen atom,
10 alkyl group, aralkyl group, aryl group, substituted silyl group, alkoxy group, aralkyloxy group, aryloxy group, di-substituted amino group, azido group, cyano group, isothiocyanate group or group having a cyclopentadiene type anion skeleton; L represents a group which bonds to M or M' by a lone-paired electron
15 or a π electron; X' represents a counter anion; k, l, m, o and p each independently represent an integer of 0 to 5; and n represents an integer of 1 to 3.

2. The catalyst component according to Claim 1, wherein M represents a transition metal atom of III to V group.

20 3. The catalyst component according to Claim 1 or 2, wherein M' represents a transition metal atom of VI to X group.

4. The catalyst component according to Claim 1 or 3, wherein X is a hydrogen atom, halogen atom, alkyl group, aralkyl group, aryl group, substituted silyl group, alkoxy group, aralkyloxy
25 group, aryloxy group, di-substituted amino group or group having a cyclopentadiene type anion skeleton.

5. The catalyst component for addition polymerization according to any one of Claims 1 to 4, wherein o represents an integer



of 1 to 5 and at least one X represents a group having a cyclopentadiene type anion skeleton.

6. A catalyst for addition polymerization obtained by a process comprising bringing the transition metal compound (A) according to any one of Claims 1 to 5 into contact with an organoaluminum compound selected from the group consisting of the following (B1), and an aluminoxane selected from the group consisting of the following (B2) and (B3) and/or a boron compound selected from the group consisting of the following (C), or with an aluminoxane selected from the group consisting of the following (B2) and (B3) and/or the following (C):

(B1) organoaluminum compounds of the general formula $E^1_a AlZ_{3-a}$,

(B2) cyclic aluminoxanes having a structure of the general formula $\{-Al(E^2)-O-\}_b$,

(B3) linear aluminoxanes having a structure of the general formula $E^3\{-Al(E^3)-O-\}_c AlE^3_2$,

(wherein, each of E^1 , E^2 and E^3 represents a hydrocarbon group; all E^1 's, all E^2 's or all E^3 's may be the same or different; Z represents a hydrogen atom or halogen atom; all Zs may be the same or different; a represents a number satisfying $0 < a \leq 3$; b represents an integer of 2 or more; and c represent an integer of 1 or more.), and

(C) one or more boron compounds selected from the following (C1) to (C3):

(C1) boron compounds represented by the general formula $BQ^1Q^2Q^3$,

(C2) boron compounds represented by the general formula



$G^+(BQ^1Q^2Q^3Q^4)^-$, and

(C3) boron compounds represented by the general formula
 $(L-H)^+(BQ^1Q^2Q^3Q^4)^-$

(wherein, B represents boron in trivalent state; Q^1 to Q^4

5 represent a halogen atom, hydrocarbon group, halogenated
hydrocarbon group, substituted silyl group, alkoxy group or
di-substituted amino group; they may be the same or different;
 G^+ represents an inorganic or organic cation; L represents a
neutral Lewis base; and $(L-H)^+$ represents a Brønsted acid.).

10 7. A process for producing an addition polymer which
comprises polymerizing an addition polymerizable monomer with
the catalyst of claim 6.