PROCESS FOR PRODUCING A POLYOLEFIN FUNCTIONAL AT ONE END

VERFAHREN ZUR HERSTELLUNG EINES AN EINEM ENDE FUNKTIONELLEN POLYOLEFINS

PROCÉDÉ DE PRÉPARATION D'UNE POLYOLÉFINE FONCTIONNELLE À UNE EXTRÉMITÉ
[0001] The present invention relates to a process of producing a single-chain-end functionalized polyolefin.

Background Art

[0002] Polyolefins such as polyethylene (PE) and polypropylene (PP) are light and inexpensive and further have characteristics of having excellent physical properties and workability. On the other hand, high chemical stability of polyolefins is an obstacle for giving, thereto, high functionalities, typical examples of which include printability, paintability, heat resistance and impact resistance, and a function for improving compatibility thereof with other polar polymers. There are known methods for making up for such drawbacks and causing polyolefins to have functionalities. Examples thereof include a method of polymerizing an olefin with a polar monomer such as vinyl acetate or a methacrylic acid ester by radical polymerization; and a method of grafting a polar monomer such as maleic anhydride to a polyolefin in the presence of a peroxide. However, according to these methods, it is generally difficult to control minutely the structure of olefin chain moieties in the resultant polymers. As a result, excellent, original physical properties of polyolefin may be damaged.

[0003] In general, it is well known that a process using living polymerization is useful as a process for producing such a polymer. In the case of highly-controlled living polymerization, a growing terminal of the polymer quantitatively keeps reactivity. It is therefore known that the reactivity is used to cause the terminal to react directly with a polar-group-containing monomer, whereby a polymer having a functional group at its terminal position can be effectively produced.

[0004] However, in the case of polymerizing any olefin by living polymerization, chain transfer reaction of the growing polymer chain is frequently caused under ordinary conditions; therefore, it is very difficult to produce an olefin polymer by living polymerization. Some examples wherein an \( \alpha \)-olefin is subjected to living polymerization have been reported so far. However, in any one of the report examples, the polymerization is conducted at a very low temperature in order to control chain transfer reaction. The polymerization activity thereof is also a low value. The molecular weight thereof is also at most several tens of thousands. Furthermore, monomers that can be polymerized are restricted in many cases. It is particularly difficult to produce industrially important ethylene-based (co) polymers or block copolymers. Concerning stereoregular polymerization of \( \alpha \)-olefins, living polymerizations exhibiting a high regularity are hardly known (see, for example, "Kobunshi", 1988, 47(2), 74-77).

[0005] Under such situations, the Applicant already discloses a transition metal compound having a salicylaldimine ligand as a novel catalyst for olefin polymerization (see Japanese Patent Application Laid-Open No. 11-315109), and further suggests a process of using the transition metal compound to produce a novel single-terminal vinyl-group-containing copolymer or a novel polar-group-containing block copolymer (see Japanese Patent Application Laid-Open Nos. 2003-73412 and 2003-40953). However, the two published documents neither disclose any polymer having a polar functional group only at its single terminal (single-chain-end functionalized polymer) nor any process for the production thereof. The present Applicant has eagerly searched a process of producing a single-chain-end functionalized polymer which can be used for various purposes and has overcome the above-mentioned problems, and has then made the present invention.

Patent documents 1-5 disclose terminal modified polyolefins comprising polypropylene or a polypropylene/polyethylene copolymer wherein the terminal group contains, e.g. oxygen or nitrogen and \( M_w/M_n \) is less than 1.5. Patent document 6 discloses terminal modified polyolefins comprising \( \text{C}_3\text{C}_1\text{C}_2 \) polyolefines wherein the terminal group is a functional group such as a halogen, an alkoxy or an ester group. Patent document 7 discloses a terminal carbonyl group-containing polymer comprising a \( \text{C}_2\text{C}_2\text{C}_20 \) polyolefin having a \( M_w/M_n \) of 1-1.3. Patent document 8 discloses terminal modified polyolefins comprising \( \text{C}_3\text{C}_6 \) polyolefines wherein the terminal group contains, e.g. oxygen or nitrogen and \( M_w/M_n \) is less than 1.2.

Disclosure of the Invention

[0006] Accordingly, the present invention relates to a process of producing a polyolefin which has a polar functional group at its single-terminal position and is useful for various purposes.

[0007] The single-chain-end functionalized polyolefin (F) obtained by the present invention is represented by the following general formula (I):

\[ P - X \]  

wherein X is a group containing at least one element selected from oxygen, sulfur, nitrogen, phosphorus and halogens, P represents a polymer chain, and X is bonded to a terminal of P, wherein the molecular weight distribution (Mw/Mn) obtained by gel permeation chromatography (GPC) is from 1.0 to 1.5.

[0008] The polymer chain (P) is a polymer chain made of units of at least one olefin selected from ethylene and olefins having 3 to 20 carbon atoms.

[0009] The single-chain-end functionalized polyolefin of the invention is obtained by: performing the following steps 1 and 2 and subsequently performing the following step 3 if necessary:

[steps 1] the step of contacting an olefin polymerizing catalyst containing a compound (A) which contains a transition metal in the groups IV to V with a polar-group-containing olefin (C) represented by the following general formula (II):

\[ \text{CHA}=\text{C(R)}-\text{Q}-\text{Y} \]  

wherein \( Y' \) is a group containing at least one element from oxygen, sulfur, nitrogen, phosphorus and halogens, Q is an alkylene group which may have a substituent, a carbonyl group, or bivalent oxygen, A and R each represent a hydrogen atom or a hydrocarbon group which may have a substituent, and A or R may be bonded together to Q to form a ring,

[step 2] the step of contacting the resultant with at least one olefin (D) selected from ethylene and olefins having 3 to 20 carbon atoms n times wherein n is an integer of 1 or more, so as to mix them, provided that when n is an integer of 2 or more, the olefins (D) used in the respective contact operations are different in kind or composition, and

[step 3] the step of chemical conversion of group \( Y' \) to another group.

**Best Modes for Carrying Out the Invention**

[0010] The following will describe the single-chain-end functionalized polyolefin produced by the production process of the present invention in detail.

Single-chain-end functionalized polyolefin

[0011] The single-chain-end functionalized polyolefin (F) produced by the process of the invention is represented by the following general formula (I):

\[ P - X \]  

[0012] In the formula (I), X is a group containing at least one element selected from oxygen, sulfur, nitrogen, phosphorus and halogens, that is, a polar functional group. Specific examples thereof include an oxy group; a peroxy group; a hydroxy group; a hydroperoxy group; alkoxy groups such as methoxy, ethoxy, propoxy and butoxy; aryloxy groups such as phenoxy, methylenephenoxy, dimethylenephenoxy, and naphthoxy; arylalkoxy groups such as phenylmethoxy, and phenylethoxy; an acetoxy group; a carbonyl group; groups wherein an element in the group XIII or XIV is bonded to an oxygen, such as silyloxy, boryloxy, and alumiinox; an amino group; N-mono-substituted amino groups such as methylamino, N-benzylamino, and N-cyclohexylamino; N,N-di-substituted alkylamino groups such as dimethylamino, diethylamino, dipropylamino, dibutylamino, dicyclohexylamino, dibenzylamino, piperidino, and morpholino; arylamino or alkylarylamino groups such as phenylamino, diphenylamino, ditolylamino, dinaphthylamino, and methylphenylamino; N,N-disilyl-substituted amino groups such as N,N-bis(trimethylsilyl)amino, N,N-bis(triethyldimethylsilyl)amino; other nitrogen-containing groups such as imine, amid, imide, ammonium, nitrile and sulfonamide; sulfonate groups such as methylsulfonate, trifluoromethanesulfonate, phenylsulfonate, benzylsulfonate, p-toluenesulfonate, trimethylbenzenesulfonate, triisobutylberizenesulfonate, p-chlorobenzenesulfonate, and pentafluorobenzenesulfonate; sulfinate groups such as methylsulfinate, phenylsulfinate, benzylsulfinate, p-toluenesulfinate, trimethylbenzenesulfinate, and pentafluorobenzenesulfinate; alkythio groups; arylthio groups; a sulfate group; a sulfide group; a polysu-
fide group; and a thiolate group. Examples of the phosphorus-containing group include phosphines such as phenylphosphino, methylphosphino, ethylphosphino, diphenylphosphino, dimethylphosphino, diethylphosphino, methylphenylphosphino, and dibenzylphosphino; phosphate oxides; phosphine sulfides; and phosphinous acids. Examples of the halogens include fluorine, chlorine, bromine, and iodine. P represents a polymer chain made of structural units derived from at least one selected from ethylene and olefins having 3 to 10 carbon atoms. In the formula (I), X is bonded to a terminal of P.

[0013] The molecular weight distribution (Mw/Mn) of the single-chain-end functionalized polyolefin produced by the process of the invention represented by the general formula (I), the distribution being obtained by gel permeation chromatography (GPC), is from 1.0 to 1.5. However, when the single-chain-end functionalized polyolefin produced by the process of the invention is produced by adopting a production process (n = 1 in the step 2) which will be described later, the molecular weight distribution (Mw/Mn) is usually 1.2 or less.

[0014] When the polymer chain (P) is a polyethylene chain, that is, a chain wherein the concentration of a skeleton originating from ethylene is 80% or more by mol, the weight-average molecular weight (Mw) of the single-chain-end functionalized polyolefin produced by the process of the invention is 5,000 or more, preferably 7,000 or more.

[0015] In the case that the polymer chain P in the single-chain-end functionalized polyolefin (F) produced by the process of the invention contains an α-olefin chain having 3 to 20 carbon atoms, the α-olefin chain has a feature of exhibiting syndiotacticity. The fact that the α-olefin chain is syndiotactic can be identified by various spectral analyses. The following will describe the fact that the polymer chain (P) of the single-chain-end functionalized polyolefin produced by the process of the invention is syndiotactic on the basis of analytic findings, giving a case in which the α-olefin is propylene as an example.

[0016] The 13C NMR spectrum of polypropylene is measured, and attention is paid to a range of methyl groups of side chains (19.5-21.7 ppm). The syndiotacticity [rr] of a triad can be obtained by substituting an integrated value of plural peaks (19.5-20.2 ppm) corresponding to an rr triad in this range and an integrated value of peaks (20.2-21.7 ppm) corresponding to a different mm or mr triad for I (rr) /{I (rr) + I (mr) + I (mm) } wherein I represents the integrated intensity of each chain in the 13C NMR. In the case that polypropylene has no regularity, a statistically-random distribution is generated; therefore, values close to the following are obtained: I (rr) :I (mr) :I (mm) = 1:2:1 and [rr] = 0.25. In the single-chain-end functionalized polyolefin produced by the process of the invention, the [rr] can be controlled into any value from 0.25 to 1.0 by catalytic structure or other polymerizing conditions. In the case that the regularity is particularly high ([rr] > 0.80), a sharp peak (20.0-20.1 ppm) corresponding to an rrrr pentad makes its appearance at a higher intensity than peaks resulting from other chains. Accordingly, the syndiotacticity can be more precisely evaluated by the [rrrr] . In a copolymer made from ethylene and propylene also, syndiotacticity is kept when chains of propylene are present therein. In this case, the value of the [rr] can be obtained from a value obtained by amending overlap of methyl groups which originates from chains of EPE and EPP wherein E and P represent an ethylene unit and a propylene unit, respectively, in each polymer.

[0017] Out of single-chain-end functionalized polyolefins (F), preferred are polyolefins wherein X is an oxygen-containing group or nitrogen-containing group or is the two groups from the viewpoint of exhibiting high reactivity with various chemical species.

Single-chain-end functionalized polyolefin produced by a specific production process

[0018] The single-chain-end functionalized polyolefin produced by the process of the invention can be effectively obtained by carrying out steps which will be detailed below successively in the presence of an olefin polymerizing catalyst containing a compound (A) which contains a transition metal in the groups IV to V in the periodic table.

[0019] As the group IV to V transition metal containing compound (A), transitionmetal compounds described in the above-mentioned Japanese PatentApplication Laid-Open No. 2003-40953, which was filed by the Applicant, can be used without any limitation. Of these transition metal compounds, preferred transition metal compounds are illustrated below.
In the production process according to the invention, an organic aluminum oxy compound (B) can be used together with the group IV to V transition metal containing compound (A). The organic aluminum oxy compound (B) may be an aluminoxane known in the prior art, or an organic aluminum oxy compound insoluble in benzene, as exemplified in Japanese Patent Application Laid-Open No. 2-78687.

The known aluminoxane can be produced by, for example, a process as described below, and is usually obtained as a solution containing a solvent of a hydrocarbon.

1. A process of adding an organic aluminum compound, such as trialkylaluminum, to a suspension of a compound containing absorbed water or a salt containing crystal water, such as magnesium chloride hydrate, copper sulfate hydrate, aluminum sulfate hydrate, nickel sulfate hydrate or cerium (I) chloride hydrate, in a hydrocarbon medium, so as to cause the absorbed water or crystal water to react with the organic aluminum compound.
2. A process of causing water, ice or water vapor to act directly on an organic aluminum compound, such as trialkylaluminum, in a solvent such as benzene, toluene, ethyl ether or tetrahydrofuran.
3. A process of causing an organic tin oxide such as dimethyltin oxide or dibutyltin oxide to react with an organic aluminum compound, such as trialkylaluminum, in a solvent such as decane, benzene or toluene.
In the general formula (II), Q is an alkylene group which may have a substituent, a carbonyl group, or bivalent oxygen, A and R each represent a hydrogen atom or a hydrocarbon group which may have a substituent, and A or R may be bonded together to Q to form a ring.

[step 1] the step of bringing it into contact with a polar-group-containing olefin (C) represented by the following general formula (II):

\[ \text{CHA} = \text{C(R)} - \text{Q} - \text{Y'} \]  

wherein Y' is a group containing at least one element from oxygen, sulfur, nitrogen, phosphorus and halogens, Q is an alkylene group which may have a substituent, a carbonyl group, or bivalent oxygen, and A or R may be bonded together to Q to form a ring.

[step 2] the step of bringing the resultant into contact with at least one olefin (D) selected from ethylene and olefins having 3 to 20 carbon atoms n times wherein n is an integer of 1 or more, so as to mix them (provided that when n is an integer of 2 or more, the olefins (D) used in the respective contact operations are different in kind or composition), and

[step 3] the step of chemical conversion of group Y' to another group.

In the production process according to the invention, at least one selected from the following can be caused to be present together with the group IV to V transition metal containing compound (A) and the organic aluminum compound (B) : an organic metal compound, a compound which can react with the transition metal compound (A) to form an ion pair, a carrier, and an organic compound. About the four components used if necessary, ones described in the Japanese Patent Application Laid-Open No. 2003-40953 can be used without any limitation.

In the general formula (II) used in the step 1 is a group containing at least one element from oxygen, sulfur, nitrogen, phosphorus and halogens. Examples of such a group include an oxy group; a peroxy group; a hydroxy group; a hydroperoxy group; alkoxo groups such as methoxy, ethoxy, propoxy and butoxy; arylxyloxy groups such as phenoxo, methylphenoxo, dimethylphenoxo, and naphthoxy; aryalkoxo groups such as phenylmethoxy, and phenylethoxy; an acetoxy group; a carbonyl group; nitrogen-containing groups such as imine, amide, imide, ammonium, nitrile and sulfonamide; sulfonate groups such as methylsulfonate, trifluoromethanesulfonate, phenylsulfonate, benzylsulfonate, p-toluenesulfonate, trimethylbenzenesulfonate, trifluorobenzenesulfonate, p-chlorobenzenesulfonate, and pentfluorobenzenesulfonate; sulfinate groups such as methylsulfinate, p-nitrobenzenesulfinate, benzylsulfinate, p-toluenesulfinate, trimethylbenzenesulfinate, and pentfluorobenzenesulfinate; alkylthio groups; arylthio groups; a sulfate group; a sulfide group; a polysulfide group; and a thiolate group. Examples of the phosphorus-containing group include phosphines such as phenylphosphino, methylphosphino, ethylphosphino, diphenylphosphino, diethylphosphino, methylphenylphosphino, and dibenzylphosphino; phosphine oxides; phosphine sulfides; and phosphinous acids. Examples of the halogens include fluoride, chlorine, bromine, and iodine. Of these, preferred are silyloxy, aluminoxy, boryloxy, and N,N-disilyl-substituted amino groups, which do not poison the catalyst easily and which generate active hydrogen in the case that hydrolysis is performed after the end of the step 2.

In the general formula (II), Q is an alkylene group which may have a substituent, a carbonyl group, or bivalent oxygen. Q is usually an alkylene group which may have a substituent wherein the total number of carbon atoms is from 1 to 20. Of alkylene groups having such a requirement, an unsubstituted linear alkylene group represented by the

Specific examples of the organic aluminum compound used when the aluminoxane is prepared include tri-n-alkylaluminums such as trimethylaluminum, triethylaluminum, tri-n-butylaluminum, tripropylaluminum, triarylaluminum, and trietylaluminum; branched-trialkyl aluminums such as trisopropylaluminum, trisobutylaluminum, tri-sec-butylaluminum, and tri-2-ethylhexylaluminum; tricycloalkylaluminums such as tricyclohexylaluminum, and tricyclo-cloctylaluminum; triarylamines such as triphenylaluminum, and tritolylaluminum; and trialkenylaluminums such as triisoprenylaluminum represented by (i-C₃H₇)₂Al[(C₅H₉)₂], wherein x, y and z are each a positive number and z ≥ 2x. Of these, trialkylaluminums and tricycloaluminums are preferred and trimethylaluminum is particularly preferred. The above-mentioned organic aluminum compounds may be used alone or in combination of two or more thereof.

In the production process according to the invention, at least one selected from the following can be caused to be present together with the group IV to V transition metal containing compound (A) and the organic aluminum compound (B) : an organic metal compound, a compound which can react with the transition metal compound (A) to form an ion pair, a carrier, and an organic compound. About the four components used if necessary, ones described in the Japanese Patent Application Laid-Open No. 2003-40953 can be used without any limitation.

The single-chain-end functionalized polyolefin (F) produced by the process of the invention is obtained by performing the following steps 1 and 2 in the presence of an olefin polymerizing catalyst containing a compound (A) which contains a transition metal in the groups IV to V, and subsequently performing the following step 3 if necessary; that is, the single-chain-endfunctionalized polyolefin (F) produced by the process of the invention is obtained by carrying out the steps 1 and 2 in this order, and carrying out the step 3 if necessary.

[step 1] the step of bringing it into contact with a polar-group-containing olefin (C) represented by the following general formula (II):

\[ \text{CHA} = \text{C(R)} - \text{Q} - \text{Y'} \]  

wherein Y' is a group containing at least one element from oxygen, sulfur, nitrogen, phosphorus and halogens, Q is an alkylene group which may have a substituent, a carbonyl group, or bivalent oxygen, and A or R may be bonded together to Q to form a ring.

[step 2] the step of bringing the resultant into contact with at least one olefin (D) selected from ethylene and olefins having 3 to 20 carbon atoms n times wherein n is an integer of 1 or more, so as to mix them (provided that when n is an integer of 2 or more, the olefins (D) used in the respective contact operations are different in kind or composition), and

[step 3] the step of chemical conversion of group Y' to another group.
following formula (III) is preferably used:

\[-[\text{CH}_2]^n-\quad (\text{III})\]

wherein \(n\) is a positive integer of 1 to 15.

[0027] In the general formula (II), \(A\) and \(R\) each represent a hydrogen atom or a hydrocarbon group which may have a substituent, and \(A\) or \(R\) may be bonded together to \(Q\) to form a ring. Of structures satisfying such requirements, a cycloolefin represented by the following formula (IV) or (IV’) is preferably used:

![Diagram](image)

wherein \(p\) represents an integer of 1 to 10, and is bonded to \(Y'\) at any position, \(q\) is an integer of 0 to 10, and when \(q\) is 0, the cycloolefin is a monocycloolefin.

Examples of the olefin having 3 to 20 carbon atoms, used in the step 2, include linear or branched \(\alpha\)-olefins having 3 to 20 carbon atoms, such as propylene, 1-butene, 2-butene, 1-pentene, 3-methyl-1-butene, 1-hexene, 4-methyl-1-pentene, 3-methyl-1-pentene, 1-octene, 1-decene, 1-dodecene, 1-tetradecene, 1-hexadecene, 1-octadecene, and 1-eicocene; and cyclic olefins having 3 to 20 carbon atoms, such as cyclopentene, cycloheptene, norbornene, 5-methyl-2-norbornene, tetracyclododecene, and 2-methyl 1,4,5,8-dimethano-1,2,3,4,4a,5,8,8a-octahydronaphthalene. Other examples of the olefin having 3 to 20 carbon atoms include vinylcyclohexane, dienes and polyenes. Additional examples of the olefin include such as aromatic vinyl compounds styrene, \(\alpha\)-methylstyrene, \(\alpha\)-methylstyrene, \(\alpha\)-methylstyrene, \(\alpha\)-methylstyrene, and \(\alpha\)-methylstyrene. These olefins may each contain in the molecule thereof a heteroatom such as an oxygen, nitrogen, or silicon atom. The olefins maybe used alone or in combination of two or more thereof.

[0028] In the case that \(n\) is set to 2 in the [step 2] in the production of the single-chain-end functionalized polyolefin (F) and then olefins (D) used in the first and second contacts in this production are made different from each other in kind or composition, the polymer chain (P) in the single-chain-end functionalizedpolyolefin (F) represented by the general formula (I) can be rendered a block type chain composed of two kinds of olefin chains each having a controlled molecular weight (the wording “different in kind or composition” in the step 2 related to the invention means the following: “different in kind”; “different in composition”; and further “different in kind and composition”). For example, in the case that after the step 1 is carried out the olefin (D) used in the first olefin-contact in the step 2 is ethylene and the olefin (D) used in the second step 2 is propylene, the resultant single-chain-end functionalizedpolyolefin is a single-chain-end functionalized block polymer represented by the following general formula (V):

\[X-\text{PE}-\text{PP}\quad \text{(V)}\]

wherein \(X\) has the same meanings as in the formula (I), and PE and PP represent a polyethylene chain and a polypropylene chain, respectively.

[0029] In the case that after the step 1 is carried out the olefin (D) used in the first olefin-contact in the step 2 is ethylene and the olefin (D) used in the second step 2 is ethylene and propylene, the resultant single-chain-end functionalized polyolefin is a single-chain-end functionalized polyolefin represented by the following formula (VI):

\[\text{X-PE-PP}\quad \text{(VI)}\]
The single-chain-end functionalized polyolefin produced by the process of the invention can be developed into various applications. The polyolefin can be applied to, for example, a high molecular weight additive; a compatibility accelerator; a diblock copolymer useful as a compatibility accelerator or modifier for polymer; a precursor of a triblock copolymer useful as thermoplastic elastomer or the above-mentioned articles; or a surface modifier for improving paintability, adhesive property and other properties of resin. The polyolefin can be used, in the form of a macromonomer, as raw material of a polymer having a specific structure such as a comb-shaped or star-shaped structure, and applied to various applications. The polyolefin can be applied to, for example, a high molecular weight additive; a compatibility accelerator; a diblock copolymer useful as a compatibility accelerator or modifier for polymer; a precursor of a triblock copolymer useful as thermoplastic elastomer or the above-mentioned articles; or a surface modifier for improving paintability, adhesive property and other properties of resin. The polyolefin can be used, in the form of a macromonomer, as raw material of a polymer having a specific structure such as a comb-shaped or star-shaped structure, and applied to various applications.

The single-chain-end functionalized polyolefin produced by the process of the invention can be developed into various applications. The polyolefin can be used, in the form of a macromonomer, as raw material of a polymer having a specific structure such as a comb-shaped or star-shaped structure, and applied to various applications. The polyolefin can be used, in the form of a macromonomer, as raw material of a polymer having a specific structure such as a comb-shaped or star-shaped structure, and applied to various applications.
bottom of the reactor to the inside thereof so as to cause the components to react at 0°C for 5 minutes. Thereafter, the supply of ethylene was stopped and methanol was added thereto, thereby terminating the polymerization. After the termination of the polymerization, the reactant was poured into 600 mL of methanol containing a small amount of hydrochloric acid to precipitate the entire amount of a polymer. The polymer was collected by filtration. The polymer was dried at 80°C under a reduced pressure for 10 hours so as to be yielded in an amount of 0.266 g. The polymerization activity per mmol of titanium was 30.9 g, the number-average molecular weight (Mn) of the polymer was 13,000, the ratio of the weight-average molecular weight (Mw) to the number-average molecular weight (Mn), (Mw/Mn), was 1.08, and the melting peak temperature based on DSC was 133.8°C. In the 1H NMR spectrum (FT, 270 MHz, in C2D2Cl4, at 120°C) of this polymer, a triplet corresponding to a methylene group adjacent to an OH group made its appearance near 3.64 ppm, and an overlap of methyl groups of two types at terminals made its appearance near 0.95 ppm. The integration ratio therebetween was 2:6. In the 13C NMR spectrum (FT, 67.5 MHz, in C2D2Cl4, at 120°C), the methyl groups made their appearance at 13.9 ppm and 19.7 ppm, and a signal corresponding to the methylene group adjacent to the OH group made its appearance at 62.9 ppm. From the above, a structure of a polymer of the following formula was identified:

\[
\begin{align*}
&\text{[Example 2]} \\
&\text{[0042]} \\
&\text{Into a glass reactor having an internal volume of 500 mL and purged sufficiently with nitrogen were charged 250 mL of toluene and 10.0 mmol of methylaluminoxane, the amount being an amount in terms of aluminum atoms therein. The reaction solution was cooled to 0°C, and then thereto was added a solution of 10.7 mg (0.0685 mmol) of Me2AlO-(CH2)4CH=CH2 in toluene. Thereto was added a solution of 58.4 mg (containing the weight of diethyl ether, 0.0666 mmol) of a titanium complex, bis[N-(3-t-butylsalicylidene)-2,3,9,5,6-pentafluoronilinate]titanium dichloride in toluene, and then the components were caused to react at 0°C for 30 minutes. Thereafter, propylene (gas flow rate: 100 L/h), the pressure of which was a normal pressure, was blown from the bottom of the reactor to the inside thereof so as to cause the components to react at 0°C for 105 minutes. Thereafter, the supply of propylene was stopped, and methanol was added thereto, thereby terminating the polymerization. After the termination of the polymerization, the reactant was poured into 600 mL of methanol containing a small amount of hydrochloric acid to precipitate the entire amount of a polymer. The polymer was collected by filtration. The polymer was dried at 80°C under a reduced pressure for 10 hours so as to be yielded in an amount of 0.354 g. The polymerization activity per mmol of titanium was 3.04 g, the number-average molecular weight (Mn) of the polymer was 8,820, the ratio of the weight-average molecular weight (Mw) to the number-average molecular weight (Mn), (Mw/Mn), was 1.05, and the melting peak temperature based on DSC was 144.4°C. In the 1H NMR spectrum (FT, 270 MHz, in C2D2Cl4, at 120°C) of this polymer, a triplet corresponding to a methylene group adjacent an OH group made its appearance near 3.64 ppm. In the 13C NMR spectrum (FT, 67.5 MHz, in C2D2Cl4, at 120°C), a signal corresponding to the methylene group adjacent to the OH group made its appearance at 62.9 ppm. Peaks at 22.5-24.0 ppm corresponding to isopentyl and isobutyl groups, which were unreacted initiating ends, hardly made their appearance.}
\end{align*}
\]
Propylene was polymerized under the same conditions as in Example 2 except that Me$_2$AlO-(CH$_2$)$_9$CH=CH$_2$ was used instead of Me$_2$AlO-(CH$_2$)$_4$CH=CH$_2$. The polymerization activity per mmol of titanium was 3.03 g, the number-average molecular weight (Mn) of the polymer was 8,200, and the ratio of the weight-average molecular weight (Mw) to the number-average molecular weight (Mn), (Mw/Mn), was 1.09. In the $^1$H NMR spectrum (FT, 270 MHz in C$_2$D$_2$Cl$_4$, at 120°C) of this polymer, a triplet corresponding to a methylene group adjacent to an OH group made its appearance near 3.64 ppm.

Propylene was polymerized under the same conditions as in Example 2 except that Me$_3$SiO-(CH$_2$)$_9$CH=CH$_2$ was used instead of Me$_2$AlO-(CH$_2$)$_4$CH=CH$_2$. The polymerization activity per mmol of titanium was 2.88 g, the number-average molecular weight (Mn) of the polymer was 7,700, the ratio of the weight-average molecular weight (Mw) to the number-average molecular weight (Mn), (Mw/Mn), was 1.06, the weight-average molecular weight (Mw) of the polymer was 9,250, the ratio of the weight-average molecular weight to the number-average molecular weight (Mn) was 1.06, and the melting peak temperature based on DSC was 142.0°C. In the $^1$H NMR spectrum (FT, 270 MHz, in C$_2$D$_2$Cl$_4$, at 120°C) of this polymer, a triplet corresponding to a methylene group adjacent to an OH group made its appearance near 3.64 ppm.

Into a glass reactor having an internal volume of 500 mL and purged sufficiently with nitrogen were charged 250 mL of toluene and 4.82 mmol of methylaluminoxane, the amount being an amount in terms of aluminum atoms therein. Thereto was added a solution of 15.5 mg (0.0532 mmol) of (Me$_3$Si)$_2$N-m-C$_6$H$_4$-(CH$_2$)$_2$CH=CH$_2$ in toluene. Thereto was added a solution of 42.3 mg (containing diethyl ether, 0.0482 mmol) of a titanium complex, bis[N-(3-t-butylationcylidene)-2,3,4,5,6-pentafluoronilinate] titanium dichloride in toluene, and then the components were caused to react at 0°C for 120 minutes. Thereafter, a mixed gas of ethylene and nitrogen (gas flow rate: ethylene, 5 L/h; and nitrogen, 50 L/h), the pressure of which was a normal pressure, was blown from the bottom of the reactor to the inside thereof so as to cause the components to react at 0°C for 3.5 minutes. Thereafter, the supply of ethylene was stopped and methanol was added thereto, thereby terminating the polymerization. After the termination of the polymerization, the reactant was poured into 600 mL of methanol containing a small amount of hydrochloric acid to precipitate the entire amount of a polymer. The polymer was collected by filtration. The polymer was dried at 80°C under a reduced pressure for 10 hours so as to be yielded in an amount of 0.148 g. The polymerization activity per mmol of titanium was 52.5 g, the number-average molecular weight (Mn) of the polymer was 13,700, and the ratio of the weight-average molecular weight (Mw) to the number-average molecular weight (Mn), (Mw/Mn), was 1.15. In the $^1$H NMR spectrum (FT, 270 MHz, in C$_2$D$_2$Cl$_4$, at 120°C) of this polymer, a triplet corresponding to a methylene group adjacent to a phenyl group, aromatic protons, and an overlap of methyl groups of two types at terminals made their appearance near 2.53 ppm, 6.45-6.65 ppm and 7-7.13 ppm, at an integration ratio of 2:4:6. From the above, a structure of a polymer of the following formula was identified:
Into a glass reactor having an internal volume of 500 mL and purged sufficiently with nitrogen were charged 250 mL of toluene and 6.48 mmol of methylaluminoxane, the amount being an amount in terms of aluminum atoms therein. Thereto was added a solution of 21.0 mg (0.0669 mmol) of (Me3Si)2N-(CH2)9CH=CH2 in toluene. Thereto was added a solution of 56.8 mg (containing diethyl ether, 0.0647 mmol) of a titanium complex, bis[N-(3-t-butylsalicylidene)-2,3,4,5,6-pentafluoroanilinato] titanium dichloride in toluene, and then the components were caused to react at 0°C for 150 minutes. Thereafter, a mixed gas of ethylene and nitrogen (gas flow rate: ethylene, 5 L/h; and nitrogen, 50 L/h), the pressure of which was a normal pressure, was blown from the bottom of the reactor to the inside thereof so as to cause the components to react at 0°C for 3 minutes. Thereafter, the supply of ethylene was stopped and methanol was added thereto, thereby terminating the polymerization. After the termination of the polymerization, the reactant was poured into 600 mL of methanol containing a small amount of hydrochloric acid to precipitate the entire amount of a polymer. The polymer was collected by filtration. The polymer was dried at 80°C under a reduced pressure for 10 hours so as to be yielded in an amount of 0.143 g. The polymerization activity per mmol of titanium was 44.1 g, the number-average molecular weight (Mn) of the polymer was 15,500, and the ratio of the weight-average molecular weight (Mw) to the number-average molecular weight (Mn), (Mw/Mn), was 1.10. In the 1H NMR spectrum (FT, 270 MHz, in C2D2Cl4, at 120°C) of this polymer, a triplet corresponding to a methylene group adjacent to a NH3Cl group, a multiplet corresponding to a methylene group adjacent thereto, and an overlap of methyl groups of two types at terminals made their appearance near 3 ppm, 1.80 ppm and 0.95 ppm, at an integration ratio of 2:2:6. From the above, a structure of a polymer of the following formula was identified:

[Comparative Example 1]

Propylene was polymerized under the same conditions as in Example 2 except that Me2AlO-(CH2)4CH=CH2 was not added. The polymerization activity per mmol of titanium was 3.16 g, and the melting peak temperature based on DSC was 146.0°C. In the 1H NMR spectrum (FT, 270 MHz, in C2D2Cl4, at 120°C) of this polymer, no peak made its appearance near 3.64 ppm. In the 13CNMR spectrum (FT, 67.5 MHz, in C2D2Cl4, at 120°C), peaks at 22.5-24.0 ppm corresponding to isopentyl and isobutyl groups at the terminals made their appearance.

Industrial Applicability

The polyolefin having a polar functional group at its single terminal position itself, or the polyolefin subjected to a further modifying treatment is useful for various purposes.

Claims

1. Process of producing a single-chain-end functionalized polyolefine represented by the following formula (I):

   \[ P-X \]  \hspace{2cm} (I)  

   wherein X is a group containing at least one element selected from oxygen, sulfur, nitrogen, phosphorus and halogens, P represents a polymer chain made of units of at least one olefin selected from ethylene and olefins having 3 to 20 carbon atoms and X is bonded to a terminal of P, and wherein the molecular weight distribution (Mw/Mn) obtained by gel permeation chromatography (GPC) is from 1.0 to 1.5 comprising [step 1] contacting an olefin polymerizing catalyst containing a compound (A) which contains a transition metal in the groups IV to V with a polar-group-containing olefin (C) represented by the following general formula (II):

   \[ CHA=C(R)-Q-Y' \]  \hspace{2cm} (II)
wherein \( Y' \) is a group containing at least one element from oxygen, sulfur, nitrogen, phosphorus and halogens, \( Q \) is an alkylene group which may have a substituent, a carbonyl group, or bivalent oxygen, A and R each represent a hydrogen atom or a hydrocarbon group which may have a substituent, and A or R may be bonded together to Q to form a ring.

[step 2] contacting the resultant with at least one olefin (D) selected from ethylene and olefins having 3 to 20 carbon atoms \( n \) times wherein \( n \) is an integer of 1 or more, so as to mix them, provided that when \( n \) is an integer of 2 or more, the olefins (D) used in the respective contact operations are different in kind or composition.

2. The process of producing a single-chain-end functionalized polyolefine according to claim 1, wherein the process further comprises

[step 3] a chemical conversion of group \( Y' \) to another group.

3. The process of producing a single-chain-end functionalized polyolefine according to claim 2, wherein step 3 is carried out by reaction of hydrolysis, oxidation, reduction or nucleophilic substitution.

### Revendications

1. Procédé de production d’une polyoléfine fonctionnalisée sur une seule extrémité de chaîne représentée par la formule (I) suivante:
P-X  (I)

dans laquelle X est un groupe contenant au moins un élément sélectionné parmi l’oxygène, le soufre, l’azote, le phosphore et des halogènes, P représente une chaîne polymère constituée d’unités d’au moins une oléfine sélectionnée parmi l’éthylène et des oléfines ayant 3 à 20 atomes de carbone, et X est lié à une extrémité de P, et la distribution des masses moléculaires (Mw/Mn) obtenue par chromatographie par perméation de gel (GPC) est de 1,0 à 1,5, comprenant :

[étape 1] la mise en contact d’un catalyseur de polymérisation d’oléfine contenant un composé (A) qui contient un métal de transition dans les groupes IV à V avec une oléfine (C) contenant un groupe polaire représentée par la formule générale (II) suivante :

\[
\text{CHA=C(R) -Q-Y'} \quad (\text{II})
\]

dans laquelle Y’ est un groupe contenant au moins un élément parmi l’oxygène, le soufre, l’azote, le phosphore et des halogènes, Q est un groupe alkylène qui peut avoir un substituant, un groupe carboxyle, ou un oxygène divalent, A et R représentent chacun un atome d’hydrogène ou un groupe hydrocarboné qui peut avoir un substituant, et A ou R peuvent être conjointement liés à Q pour former un cycle,

[étape 2] la mise en contact du produit résultant avec au moins une oléfine (D) sélectionnée parmi l’éthylène et des oléfines ayant \( n \) fois 3 à 20 atomes de carbone, \( n \) étant un nombre entier valant 1 ou davantage, de façon à les mélanger, à la condition que lorsque \( n \) est un nombre entier valant 2 ou davantage, les oléfines (D) utilisées dans les opérations de mise en contact respectives soient différentes par leur type ou leur composition.

2. Procédé de production d’une polyoléfine fonctionnalisée sur une seule extrémité de chaîne selon la revendication 1, ledit procédé comprenant en outre :

[étape 3] une conversion chimique du groupe Y’ en un autre groupe.

3. Procédé de production d’une polyoléfine fonctionnalisée sur une seule extrémité de chaîne selon la revendication 2, dans lequel l’étape 3 est effectuée par réaction d’hydrolyse, oxydation, réduction ou substitution nucléophile.
REFERENCES CITED IN THE DESCRIPTION

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