

5 **MULTIPLE CATALYST AND REACTOR SYSTEM FOR OLEFIN
POLYMERIZATION AND POLYMERS PRODUCED THEREFROM**

Field of the Invention

10 This invention relates to a process to polymerize olefins using multiple catalysts and multiple reactors and polymers produced therefrom. In particular this invention relates to a process to produce polyolefin adhesives and the adhesives so produced.

15 Background of the Invention

For some applications such as adhesives individual polymers do not possess the necessary combination of properties. Individual polyolefins having certain characteristics are often blended together in the hope of combining the positive attributes of the individual components. Typically the result is a blend which displays an average of the individual properties of the individual resins. For example EP 0 527 589 discloses blends of flexible, low molecular weight amorphous polypropylene with higher molecular weight isotactic polypropylene to obtain compositions with balanced mechanical strength and flexibility. These compositions show better flexibility compared to that of the isotactic polypropylene alone, but are still lacking in other physical attributes. Physical blends also have the problems of inadequate miscibility. Unless the components are selected for their compatibility they can phase separate or smaller components can migrate to the surface. Reactor blends, also called intimate blends (a composition comprising two or more polymers made in the same reactor or in a series of reactors) are often used to address these issues, however finding catalyst systems that will operate under the same environments to produce different polymers has been a challenge.

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5 Multiple catalyst systems have been used in the past to produce reactor blends (also called intimate blends) of various polymers and other polymer compositions. Reactor blends and other one-pot polymer compositions are often regarded as superior to physical blends of similar polymers. For example US 6,248,832 discloses a polymer composition produced in the presence of one or
10 more stereospecific metallocene catalyst systems and at least one non-stereospecific metallocene catalyst system. The resultant polymer has advantageous properties over the physical blends disclosed in EP 0 527 589 and US 5,539,056.

15 Thus there has been interest in the art in developing multiple catalyst systems to produce new polymer compositions. For example, US Patent 5,516,848 discloses the use of two different cyclopentadienyl based transition metal compounds activated with alumoxane or non-coordinating anions. In particular, the examples disclose, among other things, catalyst compounds in
20 combination, such as $(\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-c-C}_{12}\text{H}_{23})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(\text{H}_4\text{Ind})\text{ZrCl}_2$, or $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-c-C}_{12}\text{H}_{23})\text{TiCl}_2$ and $\text{Me}_2\text{Si}(\text{Ind})_2\text{HfMe}_2$, (Ind = indenyl) activated with activators such as methylalumoxane or N,N-dimethyl anilinium tetrakis(pentafluorophenyl) borate to produce polypropylenes having bimodal molecular weight distributions (M_w/M_n), varying amounts of
25 isotacticity (from 12 to 52 weight % isotactic PP in the product in Ex 2, 3 and 4), and having weight average molecular weights over 100,000, and some even as high as 1,200,000 for use as thermoplastics. Likewise, US 6,184,327 discloses a thermoplastic elastomer comprising a branched olefin polymer having crystalline sidechains and an amorphous backbone wherein at least 90 mole percent of the
30 sidechains are isotactic or syndiotactic polypropylene and at least 80 mole percent of the backbone is atactic polypropylene produced by a process comprising: a) contacting, in solution, at a temperature from about 90°C to about 120°C, propylene monomers with a catalyst composition comprising a chiral, stereorigid transition metal catalyst compound capable of producing isotactic or syndiotactic

5 polypropylene; b) copolymerizing the product of a) with propylene and, optionally, one or more copolymerizable monomers, in a polymerization reactor using an achiral transition metal catalyst capable of producing atactic polypropylene; and c) recovering a branched olefin polymer. Similarly US 6,147,180 discloses the synthesis of a thermoplastic polymer composition, which
10 is produced by first polymerizing monomers to produce at least 40% vinyl terminated macromonomers and then copolymerizing the macromonomers with ethylene. In addition US 6,323,284 discloses a method to produce thermoplastic compositions (mixtures of crystalline and amorphous polyolefin copolymers) by copolymerizing alpha-olefins and alpha, omega dienes using two separate catalyst
15 systems.

Likewise others have experimented with multiple stage processes to produce new polymer compositions. For example EP 0 366 411 discloses a graft polymer having an EPDM backbone with polypropylene grafted thereto at one or more of the diene monomer sites through the use of a two-step process using a
20 different Ziegler-Natta catalyst system in each step. This graft polymer is stated to be useful for improving the impact properties in blended polypropylene compositions.

Although each of the polymers described in the above references has interesting combinations of properties, there remains a need for new composition
25 that offer other new and different property balances tailored for a variety of end uses. In particular, it would be desirable to find a composition that is strong yet has adhesive characteristics and the ability to be applied using adhesive technology and equipment.

For general information in this area, one may refer to:

- 30 1. DeSouza and Casagrande, in 2001 addressed the issue of binary catalyst systems in "Recent Advances in Olefin Polymerization Using Binary Catalyst Systems", Macromol. Rapid Commun. 2001, 22, No. 16 (pages 1293 to 1301). At page 1299 they report propylene systems that produce a "gooey" product.

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2. Studies with respect to the production of stereoblock polypropylene by using in-situ mixtures of metallocene catalysts with different stereoselectivity were recently performed by Lieber and Brintzinger in "Propene Polymerization with Catalyst Mixtures Containing Different Ansa-Zirconocenes: Chain Transfer to Alkylaluminum Cocatalysts and Formation of Stereoblock Polymers",
10 Macromolecules 2000, 33, No.25 (pages 9192-9199). Propylene polymerization reactions were performed using metallocene catalysts $H_4C_2(Flu)_2ZrCl_2$, $rac-Me_2Si(2-Me-4-tBu-C_5H_2)_2ZrCl_2$ and $rac-Me_2Si(2-MeInd)_2ZrCl_2$ in the presence of either MAO (methylalumoxane) or triisobutylaluminium (Al^iBu_3)
15 /triphenylcarbenium tetrakis(perfluorophenylborate) (trityl borate) as the cocatalyst. Propylene polymerization using the mixed catalysts, $H_4C_2(Flu)_2ZrCl_2$ and $rac-Me_2Si(2-MeInd)_2ZrCl_2$ in the presence of either MAO or Al^iBu_3 /trityl borate produced waxy solids, which are completely separable into an atactic (diethyl ether-soluble) and an isotactic (insoluble) fraction. Neither fraction
20 contained any combination of isotactic and atactic pentad patterns indicating that these catalyst mixtures did not form stereoblock polymers.

3. Aggarwal addressed the various polymers produced in "Structures and Properties of Block Polymers and Multiphase Polymer Systems: An Overview
25 of Present Status and Future Potential", S. L. Aggarwal, Sixth Biennial Manchester Polymer Symposium (UMIST Manchester, March 1976)

4. "Selectivity in Propene Polymerization with Metallocene Catalysts" Resconi, et al, Chem Rev. 2000, 100, 1253-1345.

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None of the references above has directly addressed the need for polyolefin based adhesives containing both amorphous and crystalline components. Such adhesives are desired in the industry as a replacement for blends requiring significant amount of hydrocarbon resin tackifiers.

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Additional references that are of interest include:

- 1) EP Patents: EP 0 619 325 B1, and EP 719 802 B1;
- 2) US Patents/Publications: 6,207,606, 6,258,903, 6,271,323, 6,340,703, 6,297,301, US 2001/0007896 A1, 6,184,327, 6,225,432, 6,342,574, 10 6,147,180, 6,114,457, 6,143,846, 5,998,547, 5,696,045; 5,350,817, and 6,659,965.
- 3) PCT Publications: WO 00/37514, WO 01/81493, WO 98/49229, WO 98/32784; and WO 01/09200
- 4) "Metallocene-Based Branch-Block thermoplastic Elastomers," 15 Markel, et al. Macromolecules 2000, Volume 33, No. 23. pgs 8541-8548.

Summary of the Invention

This invention relates to a continuous process to produce a branched olefin
20 polymer comprising:

- 1) selecting a first catalyst component capable of producing a polymer having an Mw of 100,000 or less and a crystallinity of 20% or less under selected polymerization conditions;
- 2) selecting a second catalyst component capable of producing
25 polymer having an Mw of 100,000 or less and a crystallinity of 20% or more, (preferably 40% or more) at the selected polymerization conditions;
- 3) contacting a catalyst component, one or more activators and one or more C2 to C40 olefins in a first reaction zone, at a temperature of greater than 70°C, and at a residence time of 120 minutes or less; and
- 30 4) transferring the contents of the first reaction zone to a second reaction zone and further contacting the contents with a catalyst component, an activator and or one or more C2 to C40 olefins, at a temperature of greater than 70°C, and at a residence time of 120 minutes or less; and

- 5 5) optionally, transferring the contents of the second reaction zone to a third reaction zone and further contacting the contents with a catalyst compound, an activator and or one or more C2 to C40 olefins, at a temperature of greater than 70°C, and at a residence time of 120 minutes or less; and
- 6) recovering a branched olefin polymer comprising at least 50
- 10 mole% of one or more C3 to C40 olefins,
- where the first catalyst component is present in at least one reaction zone and the second catalyst component is present in a second reaction zone and where in at least one reaction zone the C2 to C40 olefin is a C3 to C40 alpha-olefin.

15 Detailed Description

 For the purposes of this invention and the claims thereto and for ease of reference when a polymer is referred to as comprising an olefin, the olefin present in the polymer is the polymerized form of the olefin.

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 In another embodiment this invention relates to a polymer comprising one or more C3 to C40 olefins, preferably propylene, and less than 50 mole % of ethylene, having:

- a) a Dot T-Peel between 1 Newton and the 10,000 Newtons; and
- 25 b) a Mz/Mn of 2 to 200; and/or
- c) an Mw of X and a g' of Y (measured at the Mz of the polymer)

 according to the following Table C:

Table C

X (Mw)	Y (g')
100,000 or less, preferably 80,000 or less, preferably 70,000 or less, more preferably 60,000 or less, more preferably 50,000 or less, more preferably 40,000 or less, more preferably 30,000 or less, more preferably 20,000 or less, more preferably 10,000 or less. In some embodiments X is also at least 1000, preferably at least 2000, more preferably at least 3000, more preferably at least	0.9 or less, preferably 0.7 or less Preferably between

4000, more preferably at least 5000, more preferably at least 7000, more preferably 10,000, more preferably at least 15,000.	0.5-0.9
75,000 or less, preferably 70,000 or less, more preferably 60,000 or less, more preferably 50,000 or less, more preferably 40,000 or less, more preferably 30,000 or less, more preferably 20,000 or less, more preferably 10,000 or less. In some embodiments A is also at least 1000, preferably at least 2000, more preferably at least 3000, more preferably at least 4000, more preferably at least 5000, more preferably at least 7000, more preferably 10,000, more preferably at least 15,000.	0.92 or less, preferably, 0.6 or less preferably between 0.4-0.6
50,000 or less, more preferably 40,000 or less, more preferably 30,000 or less, more preferably 20,000 or less, more preferably 10,000 or less. In some embodiments A is also at least 1000, preferably at least 2000, more preferably at least 3000, more preferably at least 4000, more preferably at least 5000, more preferably at least 7000, more preferably 10,000, more preferably at least 15,000.	0.95 or less, preferably 0.7 or less preferably between 0.5-0.7
30,000 or less, preferably 25,000 or less, more preferably 20,000 or less, more preferably 15,000 or less, more preferably 10,000 or less. In some embodiments A is also at least 1000, preferably at least 2000, more preferably at least 3000, more preferably at least 4000, more preferably at least 5000, more preferably at least 7000, more preferably 10,000, more preferably at least 15,000.	0.98 or less preferably between 0.7-0.98

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In a some embodiments the g' is 0.9 or less, 0.8 or less, 0.7 or less, 0.6 or less, 0.5 or less, 0.4 or less, 0.3 or less measured at the M_z of the polymer.

In another embodiment the polymer described above also has a peak
 10 melting point (T_m) between 40 and 250°C, or between 60 and 190°C, or between about 60 and 150°C, or between 80 and 130°C. In some embodiments the peak melting point is between 60 and 160°C. In other embodiments the peak melting

- 5 point is between 124-140°C. In other embodiments the peak melting temperature is between 40-130°C.

In another embodiment the polymer described above also has a viscosity of 90,000 mPa·sec or less at 190°C (as measured by ASTM D 3236 at 190°C); or
10 80,000 or less, or 70,000 or less, or 60,000 or less, or 50,000 or less, or 40,000 or less, or 30,000 or less, or 20,000 or less, or 10,000 or less, or 8,000 or less, or 5000 or less, or 4000 or less, or 3000 or less, or 1500 or less, or between 250 and 6000 mPa·sec, or between 500 and 5500 mPa·sec, or between 500 and 3000 mPa·sec, or between 500 and 1500 mPa·sec, and/or a viscosity of 8000 mPa·sec
15 or less at 160°C (as measured by ASTM D 3236 at 160°C); or 7000 or less, or 6000 or less, or 5000 or less, or 4000 or less, or 3000 or less, or 1500 or less, or between 250 and 6000 mPa·sec, or between 500 and 5500 mPa·sec, or between 500 and 3000 mPa·sec, or between 500 and 1500 mPa·sec. In other embodiments the viscosity is 200,000 mPa·sec or less at 190 ° C, depending on the application.
20 In other embodiments the viscosity is 50,000 mPa·sec or less depending on the applications.

In another embodiment the polymer described above also has a heat of fusion of 70 J/g or less, or 60 J/g or less, or 50 J/g or less; or 40 J/g or less, or 30
25 J/g or less, or 20 J/g or less and greater than zero, or greater than 1 J/g, or greater than 10 J/g, or between 20 and 50 J/g.

In another embodiment the polymer described above also has a Shore A Hardness (as measured by ASTM 2240) of 95 or less, 70 or less, or 60 or less, or
30 50 or less, or 40 or less or 30 or less, or 20 or less. In other embodiments the Shore A Hardness is 5 or more, 10 or more, or 15 or more. In certain applications, such as packaging, the Shore A Hardness is preferably 60-70.

5 In another embodiment the polymer of this invention has an M_z/M_n of 2 to 200, preferably 2 to 150, preferably 10 to 100.

 In another embodiment the polymer described above also has a Shear Adhesion Fail Temperature (SAFT - as measured by ASTM 4498) of 200°C or
10 less, or of 40 to 150°C, or 60 to 130°C, or 65 to 110°C, or 70-80°C. In certain embodiments SAFT's of 130-140°C are preferred.

 In another embodiment the polymer described above also has a Dot T-Peel
of between 1 Newton and 10,000 Newtons, or 3 and 4000 Newtons, or between 5
15 and 3000 Newtons, or between 10 and 2000 Newtons, or between 15 and 1000
Newtons. Dot T-Peel is determined according to ASTM D 1876, except that the
specimen is produced by combining two 1 inch by 3 inch (2.54 cm x 7.62 cm)
Kraft paper substrate cut outs with a dot of adhesive with a volume that, when
compressed under a 500 gram weight occupies about 1 square inch of area (1 inch
20 = 2.54 cm). Once made all the specimens are pulled apart in side by side testing
(at a rate of 2 inches per minute) by a machine that records the destructive force of
the insult being applied. The maximum force achieved for each sample tested was
recorded and averaged, thus producing the Average Maximum Force which is
reported as the Dot T-Peel.

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 In another embodiment the polymer described above also has a set time of
several days to 1 second, or 60 seconds or less, or 30 seconds or less, or 20
seconds or less, or 15 seconds or less, or 10 seconds or less, or 5 seconds or less,
or 4 seconds or less, or 3 seconds or less, more or 2 seconds or less, or 1 second or
30 less.

 In another embodiment the polymer described above also has an M_w/M_n
of 2 to 75, or 4 to 60, or 5 to 50, or 6 to 20.

5 In another embodiment the polymer described above also has an M_z of 500,000 or less, preferably 15,000 to 500,000, or 20,000 to 400,000, or 25,000 to 350,000.

 In another embodiment the polymer described above may also have a
10 strain at break (as measured by ASTM D-1708 at 25 °C) of 50 to 1000%, preferably 80 to 200%. In some other embodiments the strain at break is 100 to 500%.

 In another embodiment, the polymer described herein has a tensile strength
15 at break (as measured by ASTM D-1708 at 25 °C) of 0.5 MPa or more, alternatively 0.75 MPa or more, alternatively 1.0 MPa or more, alternatively 1.5 MPa or more, alternatively 2.0 MPa or more, alternatively 2.5 MPa or more, alternatively 3.0 MPa or more, alternatively 3.5 MPa or more.

20 In another embodiment the polymer described above also has a crystallization point (T_c) between 20 and 110°C. In some embodiments the T_c is between 70 to 100°C. In other embodiments the T_c is between 30 to 80°C. In other embodiments the T_c is between 20 to 50°C.

25 In some embodiment the polymers described above has a slope of -0.1 or less, preferably -0.15 or less, more preferably -0.25 or less in the trace of complex viscosity versus temperature (as measured by ARES dynamic mechanical spectrometer operating at a frequency of 10 rad/s, with a strain of 20 % under a nitrogen atmosphere, and a cooling rate of 10°C/min) over the range of
30 temperatures from $T_c + 10$ °C to $T_c + 40$ °C. See U.S. Patent Application Publication No. US 2004-0138392, published July 15, 2004. The slope is defined as a derivative of log (complex viscosity) with respect to temperature.

5 In another embodiment the polymer described above has a T_c that is at least 10 °C below the T_m , preferably at least 20 °C below the T_m , preferably at least 30 °C below the T_m , more preferably at least 35 °C below the T_m .

10 In another embodiment some polymers described above have a melt index ratio (I_{10}/I_2) of 6.5 or less, preferably 6.0 or less, preferably 5.5 or less, preferably 5.0 or less, preferably 4.5 or less, preferably between 1 and 6.0. (I_{10} and I_2 are measured according to ASTM 1238 D, 2.16kg, 190°C).

15 In another embodiment some polymers described above have a melt index (as determined by ASTM 1238 D, 2.16 kg, 190 deg. C) of 25 dg/min or more, preferably 50 dg/min or more, preferably 100 dg/min or more, more preferably 200dg/min or more, more preferably 500 dg/min or more, more preferably 2000 dg/min or more.

20 In another embodiment the polymer has a melt index of 900 dg/min or more.

25 In another embodiment the polymer described above has a range of crystallization of 10 to 60°C wide, preferably 20 to 50 °C, preferably 30 to 45 °C in the DSC traces. In DSC traces where there are two or more non-overlapping peaks, then each peak has a range of crystallization of 10 to 60 °C wide, preferably 20 to 50 °C, preferably 30 to 45 °C in the DSC traces.

30 In another embodiment the polymer produced by this invention has a molecular weight distribution (M_w/M_n) of at least 2, preferably at least 5, preferably at least 10, even more preferably at least 20.

 In another embodiment the polymer produced may have a unimodal, bimodal, or multimodal molecular weight distribution of polymer species as

5 determined by Size Exclusion Chromatography (SEC). By bimodal or multimodal is meant that the SEC trace has more than one peak or inflection points. An inflection point is that point where the second derivative of the curve changes in sign (e.g., from negative to positive or vice versus).

10 In another embodiment the polymer described above has an Energy of activation of 8 to 15 cal/mol. Energy of activation was calculated using the relationships of complex viscosity and temperature over the region where thermal effects are responsible for viscosity increase (assuming an Arrhenius- like relationship).

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In another embodiment the polymers of this invention may have a crystallinity of at least 5%.

In another embodiment the polymer described above may also have one or
20 more of the following:

- a) a peak melting point between 60 and 190°C, or between about 60 and 150°C, or between 80 and 130°C; and/or
- b) a viscosity of 8000 mPa·sec or less at 190°C (as measured by ASTM D 3236 at 190°C); or 5000 or less, or 4000 or less, or 3000 or less, or 1500
25 or less, or between 250 and 6000 mPa·sec, or between 500 and 5500 mPa·sec, or between 500 and 3000 mPa·sec, or between 500 and 1500 mPa·sec, or a viscosity of 8000 mPa·sec or less at 160°C (as measured by ASTM D 3236 at 160°C); or 7000 or less, or 6000 or less, or 5000 or less, or 4000 or less, or 3000 or less, or 1500 or less, or between 250 and 6000 mPa·sec, or between 500 and 5500
30 mPa·sec, or between 500 and 3000 mPa·sec, or between 500 and 1500 mPa·sec; and/or
- c) an Hf (Heat of fusion) of 70 J/g or less, or 60 J/g or less, or 50 J/g or less; or 40 J/g or less, or 30 J/g or less, or 20 J/g or less and greater than zero, or greater than 1 J/g, or greater than 10 J/g, or between 20 and 50 J/g; and or

- 5 d) a Shore A Hardness (as measured by ASTM 2240) of 70 or less, or
60 or less, or 50 or less, or 40 or less or 30 or less, or 20 or less; and or
- e) a Shear Adhesion Fail Temperature (SAFT - as measured by
ASTM 4498) of 40 to 150°C, or 60 to 130°C, or 65 to 110°C, or 70-80°C; and or;
- f) a Dot T-Peel of between 1 Newton and 10,000 Newtons , or 3 and
10 4000 Newtons, or between 5 and 3000 Newtons, or between 10 and 2000
Newtons, or between 15 and 1000 Newtons; and/or
- g) a set time of several days to 0.1 second, or 60 seconds or less, or 30
seconds or less, or 20 seconds or less, or 15 seconds or less, or 10 seconds or less,
or 5 seconds or less, or 4 seconds or less, or 3 seconds or less, more or 2 seconds
15 or less, or 1 second or less; and or
- h) an Mw/Mn of 1 to 75, or 2 to 60, or 2 to 50, or 3 to 20; and/or
- i) an Mz of 500,000 or less, preferably 15,000 to 500,000, or 20,000
to 400,000, or 25,000 to 350,000.

20 Useful combinations of features include polymers as described above
having a Dot T-Peel of between 1 Newton and 10,000 Newtons , or 3 and 4000
Newtons, or between 5 and 3000 Newtons, or between 10 and 2000 Newtons, or
between 15 and 1000 Newtons and:

1. an Mw of 30,000 or less, a peak melting point between 60 and
25 190°C, a Heat of fusion of 1 to 70 J/g, a branching index (g') of 0.90 or less
measured at the Mz of the polymer; and a melt viscosity of 8000 mPa·sec or less
at 190°C; or
2. an Mz of 20,000 to 500,000 and a SAFT of 60 to 150°C; or
3. an Mz/Mn of 2-200 and a set time of 2 seconds or less; or
- 30 4. an Hf (heat of fusion) of 20 to 50 J/g, an Mz or 20,000-500,000 and
a shore hardness of 50 or less; or
5. an Mw/Mn of greater than 1 to 50, a viscosity of 5000 or less
mPa·sec at 190°C; or

5 6. an Mw of 50,000 or less, a peak melting point between 60 and 190°C, a heat of fusion of 2 to 70 J/g, a branching index (g') of 0.70 or less measured at the Mz of the polymer, and a melt viscosity of 8000 mPa·sec or less at 190°C.

10 In a preferred embodiment, the polymer of this invention comprises amorphous, crystalline and branch-block molecular structures.

 In a preferred embodiment the polymer comprises at least 50 weight % propylene, preferably at least 60% propylene, alternatively at least 70% propylene, alternatively at least 80% propylene. In another embodiment the polymer
15 comprises propylene and 50 mole % ethylene or less, preferably 45 mole % ethylene or less, more preferably 40 mole % ethylene or less, more preferably 35 mole % ethylene or less, more preferably 30 mole % ethylene or less, more preferably 25 mole % ethylene or less, more preferably 20 mole % ethylene or
20 less, more preferably 15 mole % ethylene or less, more preferably 10 mole % ethylene or less, more preferably 5 mole % ethylene or less.

 In another embodiment the polymer produced has a glass transition temperature (Tg) as measured by ASTM E 1356 of 5 °C or less, preferably -5 °C
25 or less, alternatively between -5°C and -40°C, alternatively between -5 °C and -15°C.

 In another embodiment the polymer of this invention has a crystallinity of 40 % or less, alternatively 30% or less, alternatively 20% or less, even alternatively between 10% and 30%. Percent crystallinity content is determined
30 using Differential Scanning Calorimetry measurement according to ASTM D 3417-99. Percent amorphous content is determined by subtracting the % crystallinity from 100.

5 In another embodiment the polymer of this invention has an amorphous content of at least 50%, alternatively at least 60%, alternatively at least 70 %, even alternatively between 50 and 95%.

 In another embodiment the polymer produced by this invention has a
10 molecular weight distribution (Mw/Mn) of at least 1.5, preferably at least 2, preferably at least 5, preferably at least 10, even alternatively at least 20. In other embodiments the Mw/Mn is 20 or less, 10 or less, even 5 or less. Molecular weight distribution generally depends on the catalysts used and process conditions such as temperature, monomer concentration, catalyst ratio, if multiple catalysts
15 are used, and the presence or absence of hydrogen. Hydrogen may be used at amounts up to 2 weight %, but is preferably used at levels of 50 to 500 ppm.

 In another embodiment the polymer produced is found to have at least two molecular weights fractions are present at greater than 20 weight % each based
20 upon the weight of the polymer as measured by Gel Permeation Chromatography. The fractions can be identified on the GPC trace by observing two distinct populations of molecular weights. An example would be a GPC trace showing a peak at 20,000 Mw and another peak at 50,000 Mw where the area under the first peak represents more than 20 weight % of the polymer and the area under the
25 second peak represents more than 20 weight % of the polymer.

 In another embodiment the polymer of this invention has 20 weight % or more (based upon the weight of the starting polymer) of hexane room temperature soluble fraction, and 70 weight % or less, preferably 50 weight% or less of
30 Soxhlet boiling heptane insolubles, based upon the weight of the polymer. Soxhlet heptane insoluble refers to one of the fractions obtained when a sample is fractionated using successive solvent extraction technique. The fractionations are carried out in two steps: one involves room temperature solvent extraction, the other soxhlet extraction. In the room temperature solvent extraction, about one

5 gram of polymer is dissolved in 50 ml of solvent (hexane) to isolate the amorphous or very low molecular weight polymer species. The mixture is stirred at room temperature for about 12 hours. The soluble fraction is separated from the insoluble material using filtration under vacuum. The insoluble material is then subjected to a Soxhlet extraction procedure. This involves the separation of
10 polymer fractions based on their solubility in various solvents having boiling points from just above room temperature to 110°C. The insoluble material from the room temperature solvent extraction is first extracted overnight with hexane (Soxhlet); the extracted material is recovered by evaporating the solvent and weighing the residue. The insoluble sample is then extracted with heptane and
15 after solvent evaporation, it is weighed. The insolubles and the thimble from the final stage are air-dried in a hood to evaporate most of the solvent, then dried in a nitrogen-purged vacuum oven. The amount of insoluble left in the thimble is then calculated, provided the tare weight of the thimble is known.

20 In another embodiment, the polymers produced in this invention have a heptane insoluble fraction between 20% and 70 weight%, based upon the weight of the starting polymer, and the heptane insoluble fraction has branching index g' of 0.9 (preferably 0.7) or less as measured at the M_z of the polymer. In another embodiment, the polymers produced in this invention have a heptane insoluble
25 fraction between 20% and 70 weight%, based upon the weight of the starting polymer and a M_z between 20,000 and 500,000 of the heptane insoluble portion. In a preferred embodiment the composition also has at least 20 weight% hexane soluble fraction, based upon the weight of the starting polymer. In another embodiment the polymers produced have a hexane soluble portion of at least 20
30 weight %, based upon the weight of the starting polymer and that hexane soluble portion has a T_g but not a T_m .

In another embodiment the polymer of this invention comprises less than 4.5 mole % of ethylene, preferably less than 4.0 mole % ethylene, alternatively

5 less than 3.5 mole % ethylene, alternatively less than 3.0 mole % ethylene, alternatively less than 2.5 mole % ethylene, alternatively less than 2.0 mole % ethylene, alternatively less than 1.5 mole % ethylene, alternatively less than 1.0 mole % ethylene, alternatively less than 0.5 mole % ethylene, alternatively less than 0.25 mole % ethylene, alternatively 0 mole % ethylene.

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For ease of reference the polymer produced by the second catalyst having at least 40% crystallinity may also be referred to as the "semi-crystalline polymer" and the polymer produced by the first catalyst component having a crystallinity of less than 20% may be referred to as the "amorphous polymer."

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In another embodiment of this invention the polymer produced has a characteristic three-zone complex viscosity-temperature pattern. The temperature dependence of complex viscosity was measured using ARES dynamic mechanical spectrometer operating at a frequency of 10 rad/s, with a strain of 20 % under a nitrogen atmosphere, and a cooling rate of 10°C/min. The sample was first molten then gradually cooled down to room temperature while monitoring the build-up in complex viscosity. Above the melting point, which is typical of polymer processing temperature, the complex viscosity is relatively low (Zone I) and increases gradually with decreasing temperature. In zone II, a sharp increase in complex viscosity appears as temperature is dropped. The third zone (Zone III) is the high complex viscosity zone, which appears at lower temperatures corresponding to application temperatures. In Zone III the complex viscosity is high and varies slightly with further decrease in temperature. Such a complex viscosity profile provides, in hot melt adhesive applications, a desirable combination of long opening time at processing temperatures and fast set time at lower temperatures.

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In a preferred embodiment, the polymers produced herein having less than 1 mol % ethylene, have at least 2 mol% (CH₂)₂ units, preferably 4 mol%,

5 preferably 6 mol%, more preferably 8 mol%, more preferably 10 mol%, more preferably 12 mol%, more preferably 15 mol%, more preferably 18 mol%, more preferably 20 mol% as measured by Carbon 13 NMR as described below.

In an another embodiment, the polymers produced herein having between
10 1 and 5 mol % ethylene, have at least 2 +X mol% (CH₂)₂ units, preferably 4 +X mol%, preferably 6 +X mol%, more preferably 8 +X mol%, more preferably 10+X mol%, more preferably 12 +X mol%, more preferably 15 +X mol%, more preferably 18 +X mol%, more preferably 20 +X mol%, where X is the mole % of ethylene as measured by Carbon 13 NMR as described below.

15

In a preferred embodiment, the polymers produced herein, having less than 1 Mol% ethylene, have an amorphous component (which is defined to be that portion of the polymer composition that has a crystallinity of less than 20%) which contains at least 3 mol% (CH₂)₂ units, preferably 4 mol%, preferably 6
20 mol%, more preferably 8 mol%, more preferably 10 mol%, more preferably 12 mol%, more preferably 15 mol%, more preferably 18 mol%, more preferably 20 mol% as measured by Carbon 13 NMR as described below.

In an another embodiment, the polymers produced herein having between
25 1 and 5 mol % ethylene, have an amorphous component (which is defined to be that portion of the polymer composition that has a crystallinity of less than 20%) which contains at least 3 +X mol% (CH₂)₂ units, preferably 4 +X mol%, preferably 6 +X mol%, more preferably 8 +X mol%, more preferably 10+X mol%, more preferably 12 +X mol%, more preferably 15 +X mol%, more
30 preferably 18 +X mol%, more preferably 20 +X mol%, where X is the mole % of ethylene as measured by Carbon 13 NMR as described below.

Monomers

5 In a preferred embodiment the polymer comprises an olefin homopolymer or copolymer, comprising one or more C3 to C40 alpha olefins. In another preferred embodiment the olefin polymer further comprises one or more diolefin comonomers, preferably one or more C4 to C40 diolefins.

10 In a preferred embodiment the polymer comprises an olefin homopolymer or copolymer, having less than 5 mol% ethylene, and comprising one or more C3 to C40 alpha olefins. In another preferred embodiment the olefin polymer, having less than 5 mol% ethylene, further comprises one or more diolefin comonomers, preferably one or more C4 to C40 diolefins.

15

 In a preferred embodiment the polymer produced herein is a propylene homopolymer or copolymer. The comonomer is preferably a C4 to C20 linear, branched or cyclic monomer, and in one embodiment is a C4 to C12 linear or branched alpha-olefin, preferably butene, pentene, hexene, heptene, octene,
20 nonene, decene, dodecene, 4-methyl-pentene-1, 3-methyl pentene-1, 3,5,5-trimethyl-hexene-1, and the like. Ethylene may be present at 5 mol% or less.

 In another embodiment the polymer produced herein is a copolymer of one or more linear or branched C3 to C30 prochiral alpha-olefins or C5 to C30 ring
25 containing olefins or combinations thereof capable of being polymerized by either stereospecific and non-stereospecific catalysts. Prochiral, as used herein, refers to monomers that favor the formation of isotactic or syndiotactic polymer when polymerized using stereospecific catalyst(s).

30 The polymerizable olefinic moiety can be linear, branched, cyclic-containing, or a mixture of these structures. Preferred linear alpha-olefins include C3 to C8 alpha-olefins, more preferably propylene, 1-butene, 1-hexene, and 1-octene, even more preferably propylene or 1-butene. Preferred branched alpha-olefins include 4-methyl-1-pentene, 3-methyl-1-pentene, and 3,5,5-trimethyl-1-

5 hexene, 5-ethyl-1-nonene. Preferred aromatic-group-containing monomers contain up to 30 carbon atoms. Suitable aromatic-group-containing monomers comprise at least one aromatic structure, preferably from one to three, more preferably a phenyl, indenyl, fluorenyl, or naphthyl moiety. The aromatic-group-containing monomer further comprises at least one polymerizable double bond
10 such that after polymerization, the aromatic structure will be pendant from the polymer backbone. The aromatic-group containing monomer may further be substituted with one or more hydrocarbyl groups including but not limited to C1 to C10 alkyl groups. Additionally two adjacent substitutions may be joined to form a ring structure. Preferred aromatic-group-containing monomers contain at
15 least one aromatic structure appended to a polymerizable olefinic moiety. Particularly preferred aromatic monomers include styrene, alpha-methylstyrene, para-alkylstyrenes, vinyltoluenes, vinylnaphthalene, allyl benzene, and indene, especially styrene, paramethyl styrene, 4-phenyl-1-butene and allyl benzene.

20 Non aromatic cyclic group containing monomers are also preferred. These monomers can contain up to 30 carbon atoms. Suitable non-aromatic cyclic group containing monomers preferably have at least one polymerizable olefinic group that is either pendant on the cyclic structure or is part of the cyclic structure. The cyclic structure may also be further substituted by one or more hydrocarbyl groups
25 such as, but not limited to, C1 to C10 alkyl groups. Preferred non-aromatic cyclic group containing monomers include vinylcyclohexane, vinylcyclohexene, vinylnorbornene, ethylidene norbornene, cyclopentadiene, cyclopentene, cyclohexene, cyclobutene, vinyladamantane and the like.

30 Preferred diolefin monomers useful in this invention include any hydrocarbon structure, preferably C4 to C30, having at least two unsaturated bonds, wherein at least two of the unsaturated bonds are readily incorporated into a polymer by either a stereospecific or a non-stereospecific catalyst(s). It is further preferred that the diolefin monomers be selected from alpha, omega-diene

5 monomers (i.e. di-vinyl monomers). More preferably, the diolefin monomers are linear di-vinyl monomers, most preferably those containing from 4 to 30 carbon atoms. Examples of preferred dienes include butadiene, pentadiene, hexadiene, heptadiene, octadiene, nonadiene, decadiene, undecadiene, dodecadiene, tridecadiene, tetradecadiene, pentadecadiene, hexadecadiene, heptadecadiene, 10 octadecadiene, nonadecadiene, icosadiene, heneicosadiene, docosadiene, tricosadiene, tetracosadiene, pentacosadiene, hexacosadiene, heptacosadiene, octacosadiene, nonacosadiene, triacontadiene, particularly preferred dienes include 1,6-heptadiene, 1,7-octadiene, 1,8-nonadiene, 1,9-decadiene, 1,10-undecadiene, 1,11-dodecadiene, 1,12-tridecadiene, 1,13-tetradecadiene, and low molecular 15 weight polybutadienes (Mw less than 1000 g/mol). Preferred cyclic dienes include cyclopentadiene, vinylnorbornene, norbornadiene, ethylidene norbornene, divinylbenzene, dicyclopentadiene or higher ring containing diolefins with or without substituents at various ring positions.

20 In a preferred embodiment one or more dienes are present in the polymer produced herein at up to 10 weight %, preferably at 0.00001 to 1.0 weight %, preferably 0.002 to 0.5 weight %, even more preferably 0.003 to 0.2 weight %, based upon the total weight of the composition. In some embodiments 500 ppm or less of diene is added to the polymerization, preferably 400 ppm or less, 25 preferably or 300 ppm or less. In other embodiments at least 50 ppm of diene is added to the polymerization, or 100 ppm or more, or 150 ppm or more.

In a preferred embodiment the olefin polymer is homo-polypropylene. In another preferred embodiment the olefin polymer comprises propylene, ethylene, 30 preferably less than 5 mol% ethylene, and at least one divinyl comonomer. In another preferred embodiment the olefin polymer comprises propylene and at least one divinyl comonomer.

In another embodiment, the olefin polymer comprises:

- 5 a first monomer present at from 40 to 95 mole %, preferably 50 to 90 mole
%, preferably 60 to 80 mole %, and
 a comonomer present at from 5 to 40 mole %, preferably 10 to 60 mole %,
more preferably 20 to 40 mole %, and
 a termonomer present at from 0 to 10 mole %, more preferably from 0.5 to
10 5 mole %, more preferably 1 to 3 mole %.

 In a preferred embodiment the first monomer comprises one or more of
any C3 to C8 linear, branched or cyclic alpha-olefins, including propylene, butene
(and all isomers thereof), pentene (and all isomers thereof), hexene (and all
15 isomers thereof), heptene (and all isomers thereof), and octene (and all isomers
thereof). Preferred monomers include propylene, 1-butene, 1-hexene, 1-octene,
and the like.

 In a preferred embodiment the comonomer comprises one or more of any
20 C2 to C40 linear, branched or cyclic alpha-olefins (provided ethylene, if present,
is present at 5 mole % or less), including ethylene, propylene, butene, pentene,
hexene, heptene, and octene, nonene, decene, undecene, dodecene, hexadecene,
styrene, 3,5,5-trimethylhexene-1, 3-methylpentene-1, 4-methylpentene-1,
norbornene and cyclopentene.

25

 In a preferred embodiment the termonomer comprises one or more of any
C2 to C40 linear, branched or cyclic alpha-olefins, (preferably ethylene, if present,
is present at 5 mole % or less), including, but not limited to, ethylene, propylene,
butene, pentene, hexene, heptene, and octene, nonene, decene, undecene,
30 dodecene, hexadecene, butadiene, 1,5-hexadiene, 1,6-heptadiene, 1,4-pentadiene,
1,7-octadiene, 1,8-nonadiene, 1,9-decadiene, 1,11-dodecadiene, styrene, 3,5,5-
trimethylhexene-1, 3-methylpentene-1, 4-methylpentene-1, and cyclopentadiene.

5 In a preferred embodiment the polymer comprises propylene and from 0 to 50 mole % ethylene, preferably from 0 to 30 mole % ethylene, more preferably from 0 to 15 mole % ethylene, more preferably from 0 to 10 mole % ethylene, more preferably from 0 to 5 mole % ethylene.

10 In a preferred embodiment the polymer comprises propylene and from 0 to 50 mole % butene, preferably from 0 to 30 mole % butene, more preferably from 0 to 15 mole % butene, more preferably from 0 to 10 mole % butene, more preferably from 0 to 5 mole % butene.

15 In a preferred embodiment the polymer comprises propylene and from 0 to 50 mole % hexene, preferably from 0 to 30 mole % hexene, more preferably from 0 to 15 mole % hexene, more preferably from 0 to 10 mole % hexene, more preferably from 0 to 5 mole % hexene.

20 Process

 This invention relates to a continuous process to produce a branched olefin polymer comprising:

 1) selecting a first catalyst component capable of producing a polymer
25 having an Mw of 100,000 or less and a crystallinity of 20% or less (preferably 5 % or less) under selected polymerization conditions;

 2) selecting a second catalyst component capable of producing
polymer having an Mw of 100,000 or less and a crystallinity of 40% (preferably 20% or more) or more at the selected polymerization conditions;

30 3) contacting a catalyst component, one or more activators and one or more C2 to C40 olefins in a first reaction zone, at a temperature of greater than 70°C (preferably 100°C or more), and at a residence time of 120 minutes or less (preferably 60 minutes or less); and

- 5 4) transferring the contents of the first reaction zone to a second reaction zone and further contacting the contents with a catalyst component, an activator and or one or more C2 to C40 olefins, at a temperature of greater than 70°C (preferably 100°C or more), and at a residence time of 120 minutes or less (preferably 60 minutes or less); and
- 10 5) optionally, transferring the contents of the second reaction zone to a third reaction zone and further contacting the contents with a catalyst compound, an activator and or one or more C2 to C40 olefins, at a temperature of greater than 70 °C (preferably greater than 100°C), and at a residence time of 120 minutes or less, (preferably 60 minutes or less); and
- 15 6) recovering a branched olefin polymer comprising at least 50 mole% of one or more C3 to C40 olefins,
- where the first catalyst component is present in at least one reaction zone and the second catalyst component is present in a second reaction zone and where in at least one reaction zone the C2 to C40 olefin is a C3 to C40 alpha-olefin.

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This invention further relates to a continuous process to produce a branched olefin polymer comprising:

- 1) selecting a first catalyst component capable of producing a polymer having an Mw of 100,000 or less and a heat of fusion of 70 J/g or less (preferably 10 J/g or less) under selected polymerization conditions;
- 25 2) selecting a second catalyst component capable of producing polymer having an Mw of 100,000 or less and a crystallinity of 20% or more (preferably 30% or more) at the selected polymerization conditions;
- 3) contacting a catalyst component, one or more activators and one or 30 more C2 to C40 olefins in a first reaction zone, at a temperature of greater than 70 °C (preferably greater than 100°C), and at a residence time of 120 minutes or less, (preferably 60 minutes or less); and
- 4) transferring the contents of the first reaction zone to a second reaction zone and further contacting the contents with a catalyst component, an

5 activator and or one or more C2 to C40 olefins, at a temperature of greater than 70 °C, (preferably greater than 100°C), and at a residence time of 120 minutes or less (preferably 60 minutes or less); and

5) optionally, transferring the contents of the second reaction zone to a third reaction zone and further contacting the contents with a catalyst compound,
10 an activator and or one or more C2 to C40 olefins, at a temperature of greater than 70°C (preferably greater than 100°C), and at a residence time of 120 minutes or less (preferably 60 minutes or less); and

6) recovering a branched olefin polymer comprising at least 50 mole% of one or more C3 to C40 olefins,
15 where the first catalyst component is present in at least one reaction zone and the second catalyst component is present in a second reaction zone and where in at least one reaction zone the C2 to C40 olefin is a C3 to C40 alpha-olefin.

This invention further relates to a continuous process to produce a
20 branched olefin polymer comprising:

1) selecting a first catalyst component capable of producing a polymer having an Mw of 100,000 or less and a heat of fusion of 70 J/g or less (preferably 10 J/g or less), capable of polymerizing macromonomers having reactive termini at the selected polymerization conditions;

25 2) selecting a second catalyst component capable of producing macromonomers having reactive termini, an Mw of 100,000 or less and a crystallinity of 20% or more (preferably 30% or more) at the selected polymerization conditions;

3) contacting a catalyst component, one or more activators and one or
30 more C2 to C40 olefins in a first reaction zone, at a temperature of greater than 70°C, (preferably greater than 100°C), and at a residence time of 120 minutes or less, (preferably 60 minutes or less); and

4) transferring the contents of the first reaction zone to a second reaction zone and further contacting the contents with a catalyst component, an

- 5 activator and or one or more C2 to C40 olefins, at a temperature of greater than 70°C, (preferably greater than 100°C), and at a residence time of 120 minutes or less, (preferably 60 minutes or less); and
- 5) optionally, transferring the contents of the second reaction zone to a third reaction zone and further contacting the contents with a catalyst compound,
- 10 an activator and or one or more C2 to C40 olefins, at a temperature of greater than 70°C, (preferably greater than 100°C), and at a residence time of 120 minutes or less, (preferably 60 minutes or less); and
- 6) recovering a branched olefin polymer comprising at least 50 mole% of one or more C3 to C40 olefins,
- 15 where the first catalyst component is present in at least one reaction zone and the second catalyst component is present in a second reaction zone and where in at least one reaction zone the C2 to C40 olefin is a C3 to C40 alpha-olefin.

 This invention further relates to a process to produce the olefin polymers described above comprising:

20

- 1) selecting a first catalyst component capable of producing a polymer having an Mw of 30,000 or less and a heat of fusion of 70 J/g or less (preferably 10 J/g or less), capable of polymerizing macromonomers having reactive termini;
- 2) selecting a second catalyst component capable of producing
- 25 macromonomers having reactive termini, an Mw of 30,000 or less and a crystallinity of 20% or more (preferably 30% or more);
- 3) contacting a catalyst component, one or more activators and one or more C2 to C40 olefins in a first reaction zone, at a temperature of greater than 70°C, (preferably greater than 100°C), and at a residence time of 120 minutes or
- 30 less, (preferably 60 minutes or less); and
- 4) transferring the contents of the first reaction zone to a second reaction zone and further contacting the contents with a catalyst component, an activator and or one or more C2 to C40 olefins, at a temperature of greater than

5 70°C, (preferably greater than 100°C), and at a residence time of 120 minutes or less, (preferably 60 minutes or less); and

5) optionally, transferring the contents of the second reaction zone to a third reaction zone and further contacting the contents with a catalyst compound, an activator and or one or more C2 to C40 olefins, at a temperature of greater than
10 70°C, (preferably greater than 100°C), and at a residence time of 120 minutes or less, (preferably 60 minutes or less); and

6) recovering a branched olefin polymer comprising at least 50 mole% of one or more C3 to C40 olefins, where the first catalyst component is present in at least one reaction zone and the
15 second catalyst component is present in a second reaction zone and where in at least one reaction zone the C2 to C40 olefin is a C3 to C40 alpha-olefin.

In another preferred embodiment this invention relates to a continuous process to produce a branched olefin polymer comprising:

20 1) selecting a first catalyst component capable of producing a polymer having an Mw of 100,000 or less, preferably 80,000 or less, preferably 60,000 or less and a crystallinity of 20% or less, preferably 15% or less, more preferably 10% or less, under selected polymerization conditions;

2) selecting a second catalyst component capable of producing
25 polymer having an Mw of 100,000 or less, preferably 80,000 or less, preferably 60,000 or less and a crystallinity of 20% or more, preferably 40% or more, preferably 50% or more, more preferably 60% or more at the selected polymerization conditions;

3) contacting a catalyst component, one or more activators and one or
30 more C2 to C40 olefins (preferably one or more C3 to C12 olefins, preferably C3 and one or more of ethylene and/or C4 to C20 comonomers, and, optionally one or more diolefins, preferably a C4 to C20 diene) in a first reaction zone, at a temperature of greater than 70°C, (preferably greater than 100°C, more preferably greater than 105°C, more preferably greater than 110°C, more preferably greater

5 than 115°C), and at a residence time of 120 minutes or less, (preferably 60 minutes or less, more preferably 50 minutes or less, preferably 40 minutes, preferably 30 minutes or less, preferably 25 minutes or less, more preferably 20 minutes or less, more preferably 15 minutes or less, more preferably at 10 minutes or less, more preferably at 5 minutes or less); and

10 4) transferring the contents of the first reaction zone to a second reaction zone and further contacting the contents with a catalyst component, an activator and or one or more C2 to C40 olefins (preferably one or more C3 to C12 olefins, preferably C3 and one or more of ethylene and/or C4 to C20 comonomers, and, optionally one or more diolefins, preferably a C4 to C20 diene), at a
15 temperature of greater than 70 °C, preferably greater than 100°C, more preferably greater than 105°C, more preferably greater than 110°C, more preferably greater than 115°C), and at a residence time of 120 minutes or less, preferably 60 minutes or less, more preferably 50 minutes or less, preferably 40 minutes, preferably 30 minutes or less, preferably 25 minutes or less, more preferably 20 minutes or less,
20 more preferably 15 minutes or less, more preferably at 10 minutes or less, more preferably at 5 minutes or less, more preferably at 3 minutes or less); and

5) optionally, transferring the contents of the second reaction zone to a third reaction zone and further contacting the contents with a catalyst compound, an activator and or one or more C2 to C40 olefins (preferably one or more C3 to
25 C12 olefins, preferably C3 and one or more of ethylene and/or C4 to C20 comonomers, and, optionally one or more diolefins, preferably a C4 to C20 diene), at a temperature of greater than 70°C, (preferably greater than 100°C, more preferably greater than 105°C, more preferably greater than 110°C, more preferably greater than 115°C), and at a residence time of 120 minutes or less,
30 (preferably 60 minutes or less, more preferably 50 minutes or less, preferably 40 minutes, preferably 30 minutes or less, preferably 25 minutes or less, more preferably 20 minutes or less, more preferably 15 minutes or less, more preferably at 10 minutes or less, more preferably at 5 minutes or less, more preferably at 3 minutes or less); and

- 5 6) recovering a branched olefin polymer comprising at least 50 mole% of one or more C3 to C40 olefins, where the first catalyst component is present in at least one reaction zone and the second catalyst component is present in a second (preferably different) reaction zone and where in at least one reaction zone the C2 to C40 olefin is a C3 to C40 alpha-olefin; and
- 10 a) wherein the ratio of the first catalyst to the second catalyst is from 1:1 to 50:1, preferably 1:1 to 30:1, preferably 1:1 to 20:1, more preferably 1:1 to 1:10;
- b) wherein the activity of the catalyst components is at least 3 kilograms, preferably at least 50 kilograms, more preferably at least 100
- 15 kilograms, more preferably at least 200 kilograms, more preferably, 300 kilograms, more preferably 400 kilograms, more preferably 50 kilograms of polymer per gram of the catalyst mixture; and wherein at least 80%, preferably at least 85%, more preferably at least 90%, more preferably at least 95 % of the olefins are converted to polymer.

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In another embodiment at least 20 % or more of the olefins are converted to polymer, preferably 20% or more, more preferably 60% or more, more preferably 75% or more, more preferably 85% or more, more preferably 95% or more.

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In a preferred embodiment the process described above takes place in a solution phase, slurry or bulk phase polymerization process.

By continuous is meant a system that operates (or is intended to operate)

30 without interruption or cessation. For example a continuous process to produce a polymer would be one where the reactants are continually introduced into one or more reactors and polymer product is continually withdrawn.

5 In another preferred embodiment, in the process described above the concentrations of the reactants vary by 20% or less in the reaction zone during the residence time, preferably by 15% or less, more preferably by 10% or less. In a preferred embodiment the concentration of the monomer(s) remains constant in the reaction zone during the residence time. Preferably the concentration of the monomer(s) varies by 20% or less, preferably by 15% or less, more preferably by 10% or less, more preferably by 5% or less.

 In a preferred embodiment the concentration of the catalyst components remains constant in the reaction zone during the residence time. Preferably the concentration of the monomer(s) varies by 20% or less, preferably by 15% or less, more preferably by 10% or less, more preferably by 5% or less.

 In a preferred embodiment the concentration of the activator(s) remains constant in the reaction zone during the residence time. Preferably the concentration of the monomer(s) varies by 20% or less, preferably by 15% or less, more preferably by 10% or less, more preferably by 5% or less.

 In another preferred embodiment a third catalyst (or more) may be present in the processes described above. The third catalyst may be any of the catalyst components listed herein. Preferred third catalysts include catalysts that are capable of producing waxes. Particularly preferred third catalysts include those capable of producing polymer having an Mw of 20,000 or less and a crystallinity of 10% or less at the selected polymerization conditions. In another embodiment, particularly preferred third catalysts include those capable of producing polymer having an Mw of 20,000 or less and a crystallinity of 10% or more, preferably 20% or more, at the selected polymerization conditions.

 Other preferred third catalysts may include any catalyst described herein. One may select two or more catalysts to produce various macromonomers having

5 reactive termini, used in combination with a catalyst that can polymerize such macromonomers. One may select two or more catalysts that can polymerize macromonomers and one catalyst that can produce macromonomers with reactive termini. Likewise one could also select three catalysts that produce different polymers under the same reaction conditions. For example one could select a
10 catalyst that produces a somewhat crystalline polymer, one that produces a very crystalline polymer and one that produces an amorphous polymer, any of which may produce macromonomers with reactive termini or polymerize polymers having reactive termini. Similarly one could select two catalysts, one that produces crystalline polymers and one that produces an amorphous polymer, any
15 of which may make macromonomers with reactive termini or polymerize polymers having reactive termini. Likewise one could select a catalyst that produces a somewhat crystalline polymer, one that produces a wax and one that produces an amorphous polymer, any of which may make macromonomers with reactive termini or polymerize polymers having reactive termini.

20

By reaction zone is meant an area where the activated catalyst and monomers can react.

By macromonomers having reactive termini is meant a polymer having
25 twelve or more carbon atoms (preferably 20 or more, more preferably 30 or more, more preferably between 12 and 8000 carbon atoms) and having a vinyl, vinylidene, vinylene or other terminal group that can be polymerized into a growing polymer chain. By capable of polymerizing macromonomer having reactive termini is meant a catalyst component that can incorporate a
30 macromonomer (which tend to be molecules larger than a typical single monomer such as ethylene or propylene), having reactive termini into a growing polymer chain. Vinyl terminated chains are generally more reactive than vinylene or vinylidene terminated chains.

5 In a preferred embodiment propylene is present in the first, second and or third reaction zone, preferably at an amount of 20 to 100 weight %, based upon the weight of the monomers present in the reaction zone, preferably 40 to 99 weight %, more preferably 60 to 95 weight %.

10 In a preferred embodiment ethylene is present in the first, second and or third reaction zone, preferably at an amount of up to 50 weight %, preferably at 1 to 40 weight %, preferably 5-20 weight %, preferably at 5-10 weight%, based upon the weight of the monomers in the reaction zone. In another embodiment, ethylene is not present in the reaction zone, or if present is present at 10 weight %
15 or less, preferably 5 weight % or less, preferably 3 weight % or less, preferably 2 weight % or less, preferably 1 weight % or less, preferably 0.5 weight % or less, based on the weight of the monomers in the reaction zone.

 In a preferred embodiment ethylene and propylene are present in the first,
20 second and or third reaction zone.

 In another embodiment propylene is present in the first reaction zone at 100 weight %, (based upon the weight of the monomers present in the first reaction zone) and ethylene is present in the second reaction zone at up to 50
25 weight%, (based upon the weight of the monomers present in the second reaction zone).

 In another embodiment ethylene is present in the first reaction zone at 100 weight %, (based upon the weight of the monomers present in the first reaction
30 zone).

 In another embodiment propylene is present at 100 weight %, (based upon the weight of the monomers present in the first reaction zone) in the first reaction zone and the second reaction zone.

5

In another embodiment propylene and ethylene are present in the first reaction zone and no ethylene, other than residual ethylene monomer present in the contents of the first reaction zone, is introduced into the second reaction zone.

10

In another embodiment ethylene is intermittently introduced into one or more reaction zones.

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In another embodiment propylene is present in the first reaction zone, ethylene is present in the second reaction zone, the second catalyst component is present in the first reaction zone, and the first catalyst component is present in the second reaction zone.

20

In another embodiment propylene is present in the first reaction zone, propylene and ethylene or other monomers are present in the second reaction zone, the second catalyst component is present in the first reaction zone, and the first catalyst component is present in the second reaction zone.

25

In another embodiment propylene is present in the first reaction zone, propylene and ethylene are present in the second reaction zone, the second catalyst component is present in the first reaction zone, and the first catalyst component is present in the second reaction zone.

30

In another embodiment propylene and ethylene are present in the first reaction zone, propylene is present in the second reaction zone, the first catalyst component is present in the first reaction zone, and the second catalyst component is present in the second reaction zone.

In another embodiment propylene is present in the first reaction zone, propylene and ethylene are present in the second reaction zone, the second catalyst

5 component is present in the first reaction zone, and the second catalyst component is present in the second reaction zone.

In another embodiment ethylene is present in the first reaction zone, propylene and ethylene or other monomers are present in the second reaction zone, 10 the first catalyst component is present in the first reaction zone, and the first and the second catalyst components are present in the second reaction zone, and the catalyst compound present in the first reaction zone is capable of producing polymer having an Mw of 20,000 or less and a crystallinity of 50 % or more at the selected polymerization conditions.

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In another embodiment ethylene is present in the first reaction zone, propylene is present in the second reaction zone, propylene is present in the third reaction zone, the first catalyst component is present in the second reaction zone, and the second catalyst component is present in the third reaction zone, and the 20 catalyst compound present in the first reaction zone is capable of producing polymer having an Mw of 20,000 or less and a crystallinity of 10% or less at the selected polymerization conditions.

In another embodiment ethylene is present in the first reaction zone, 25 propylene is present in the second reaction zone, propylene is present in the third reaction zone, the first catalyst component is present in the second reaction zone, and the second catalyst component is present in the third reaction zone, and the catalyst compound present in the first reaction zone is capable of producing polymer having an Mw of 20,000 or less and a crystallinity of 10% or more, 30 preferably 20% or more, preferably 30% or more, preferably 40% or more, preferably 50% or more at the selected polymerization conditions.

In another embodiment all catalyst components are only introduced into the first reaction zone and no catalyst components, other than residual catalyst

5 components present in the contents of the first reaction zone, is introduced into the second reaction zone.

In another embodiment only one catalyst component is present in all the reaction zones. The catalyst component can be introduced into the first reaction
10 zone only, or introduced into multiple reaction zones.

In another embodiment more than two catalyst components are present in one or all reaction zones, at least one of the catalyst components is capable of producing a polymer having a crystallinity of 5 % or less, and at least one other
15 of the catalyst components is capable of producing a polymer having a crystallinity of 20 % or more at the selected polymerization conditions.

In another embodiment a diolefin monomer is present in one or all of the reaction zones.

20

In another embodiment hydrogen is present in one or all of the reaction zones.

In a particular embodiment the present invention is directed to a polyolefin
25 polymer produced by copolymerizing one or more C_3 or higher alpha-olefins and/or one or more di-vinyl monomers, and optionally up to 5 mol% ethylene, in the presence of at least one stereospecific catalyst system and at least one other catalyst system. The polymer so produced may contain amorphous polymer segments and crystalline polymer segments in which at least some of the segments
30 are linked. Typically the amorphous and the crystalline polymer segments are copolymers of one or more alpha-olefins (optionally including up to 5 mol% ethylene) and/or one or more monomers having at least two olefinically unsaturated bonds. Both of these unsaturated bonds are suitable for and readily incorporated into a growing polymer chain by coordination polymerization using

5 either the first or second catalyst systems independently such that the di-olefin is incorporated into polymer segments produced by both catalysts in the mixed catalyst system according to this invention. In a preferred embodiment these monomers having at least two olefinically unsaturated bonds are di-olefins, preferably di-vinyl monomers. Crosslinking of at least a portion of the mixture of
10 polymer segments is believed to be accomplished during the polymerization of the composition by incorporation of a portion of di-vinyl comonomers into two polymer segments, thus producing a crosslink between those segments.

In another embodiment, polyolefin branch-block compositions containing
15 amorphous and semi-crystalline components may be prepared in two or more reactors to yield desired property balance. In particular, aPP-g-scPP branch structures may be produced in-situ in two or more continuous solution reactors using mixed catalysts and propylene as the preferred feed. In one embodiment stereospecific bridged bis-indenyl group 4 catalysts can be selected to produce
20 semicrystalline PP macromonomers. (All references to the Periodic Table of the Elements are to the Table published in Chemical and Engineering News, 63(5), 27, 1985.) A bridged mono-cyclopentadienyl heteroatom group 4 catalyst can be used to build amorphous PP (aPP) backbone while simultaneously incorporating some of the semi-crystalline macromonomers (scPP). This is believed to produce
25 a aPP-g-scPP structure where the "-g-" indicates that the polymer types are at least partially grafted. By selecting the catalysts, the polymerization reaction conditions, and/or by introducing a diene modifier, the amorphous and crystalline components can be linked together to produce various branch-block structures. To effectively incorporate into a growing chain, a macromonomer with vinyl end
30 group is preferred. Other types of chain end unsaturations (vinylene and vinylidene) can also be used. While not wishing to be bound by theory, branch-block copolymer is believed to comprise an amorphous backbone having crystalline side chains originating from the scPP macromonomers and the sidechains are believed to be polypropylene macromonomers, which can be

- 5 prepared under solution polymerization conditions with catalysts suitable for preparing either of isotactic or syndiotactic polypropylene.

A preferred reaction process to produce polypropylene macromonomers having high levels of terminal vinyl unsaturation is described in U.S. patent
10 6,117,962. Typically used catalysts are stereorigid, chiral or asymmetric, bridged metallocenes. See, for example, U.S. Pat. No. 4,892,851, U.S. Pat. No. 5,017,714, U.S. Pat. No. 5,132,281, U.S. Pat. No. 5,296,434, U.S. Pat. No. 5,278,264, U.S. Pat. No. 5,304,614, U.S. Pat. No. 5,510,502, WO-A-(PCT/US92/10066) WO-A-93/19103, EP-A2-0 577 581, EP-A1-0 578 838, and academic literature "The
15 Influence of Aromatic Substituents on the Polymerization Behavior of Bridged Zirconocene Catalysts", Spaleck, W., et al, Organometallics 1994, 13, 954-963, and "ansa-Zirconocene Polymerization Catalysts with Annelated Ring Ligands- Effects on Catalytic Activity and Polymer Chain Lengths", Brinzinger, H., et al, Organometallics 1994, 13, 964-970, and documents referred to therein.

20

In some embodiments, the first catalyst which comprises a stereorigid transition metal pre-catalyst compound used to produce the semi-crystalline polypropylene macromonomers of the present invention is selected from the group consisting of racemic bridged bis(indenyl) zirconocenes or hafnocenes. In a
25 another embodiment, the transition metal pre-catalyst compound is a rac-dimethylsilyl-bridged bis(indenyl) zirconocene or hafnocene. In another embodiment, the transition metal pre-catalyst compound is rac-dimethylsilyl bis(2-methyl-4-phenylindenyl) zirconium or hafnium dichloride or dimethyl. In another preferred embodiment, the transition metal catalyst is a rac-dimethylsilyl-
30 bridged bis(indenyl) hafnocene such as rac-dimethylsilyl bis(indenyl)hafnium dimethyl or dichloride.

It is believed that the fraction of branch-block and the level of branching depend on the availability of macromonomers with unsaturated chain end and

5 macromonomer incorporation capability of the specific catalyst. To increase the population of aPP-g-scPP branch-block composition, one typically operates within a process window that favors macromonomer production and insertion. Such conditions have been described in U.S. patent 6, 117, 962 and the journal article by W. Weng et. al, Macromol. Rapid Commun., 2000, 21, 1103-1107 and are further
10 illustrated by the examples therein.

It is also believed that the higher the population of vinyl terminated scPP macromonomers the higher the probability of getting them incorporated into aPP backbone and therefore the higher the branch-block population.

15

To further increase the population of macromonomers having vinyl chain ends diolefin monomers can be introduced into the reaction medium. The resultant product is typically a blend comprised of isotactic polypropylene segments, atactic polypropylene segments, and increased population of branch-
20 block species resulting from the additional couplings brought about by the diolefin crosslinking agent.

Crosslinking typically refers to the connection of two polymer segments by incorporation of each double bond of a diolefin monomer into two different
25 polymer segments. The polymer segments so connected can be the same or different, with respect to their crystallinity. Three or more polymer segments may also be connected via incorporation of two or more diolefins in on polymer segment into two other polymer segments.

30 A consideration for selection of the monomer, or combinations of monomers, is that, both crystalline and amorphous polymer segments can be formed with the selection of two or more different catalyst systems. In some embodiments it is further desired that the level of incorporation of the diolefin monomer, if present, into the crystalline segments be limited to an amount that

- 5 will not substantially alter its crystallinity. The diolefin coupling agent is typically kept minimum to insure the overall composition has a viscosity of 80000 mPa•s or less for some adhesive applications.

As mentioned above, to increase the population of aPP-g-scPP branch-
10 block composition, one typically operates within a process window that favors macromonomer production and insertion. Favorable conditions include:

1. High concentration of catalyst producing the semi-crystalline vinyl terminated macromonomers, and or
2. Adjusting the Al/metal ratio; and or
- 15 3. High operating temperature; and or
4. Catalyst structure that has a high affinity for macromonomer incorporation; and or
5. Relatively long residence time; and or
6. High monomer conversion (monomer starvation condition enhances
20 the insertion of macromonomer); and or
7. Addition of modifier (diene) to enhance the population of vinyl terminated macromonomers.

Another method of enhancing aPP-g-scPP branch block compositions is to
25 add in a chain transfer agent that transfers a vinyl group to the end of the polymer chain while deactivating the catalyst. Such chain transfer agents include, but are not limited to, vinyl chloride, vinyl fluoride, vinyl bromide. In the process, the catalyst is reactivated by the presence of an aluminum alkyl activator such as an alumoxane (typically methylalumoxane).

5 Similarly, melting and crystallization characteristics can be controlled through catalyst selection, comonomer addition and changes in process conditions such as temperature and catalyst ratio if more than one catalyst is used.

Catalyst Compounds

10

Any catalyst compound that can produce the desired polymer species (i.e. a polymer having an Mw of 100,000 or less and a heat of fusion of 70 J/g or less, or a polymer having an Mw of 100,000 or less and a crystallinity of 70% or less) may be used in the practice of this invention. In the description herein the
15 transition metal compound may be described as a catalyst precursor, a pre-catalyst compound or a catalyst compound, and these terms are used interchangeably. A catalyst system is combination of a catalyst precursor and an activator.

Catalyst Compounds and Selection

20

Any pre-catalyst compound (catalyst precursor compound) that can produce the desired polymer species (i.e. a polymer having an Mw of 100,000 or less and crystallinity of 5% or less, or a polymer having an Mw of 100,000 or less and a crystallinity of 20% or more) may be used in the practice of this invention.
25 Pre-catalyst compounds which may be utilized in the process of the invention include metallocene transition metal compounds (containing one, two, or three cyclopentadienyl ligands per metal atom), non-metallocene early transition metal compounds (including those with amide and/or phenoxide type ligands), non-metallocene late transition metal compounds (including those with diimine or
30 diiminepyridyl ligands), and other transition metal compounds.

Generally, bulky ligand metallocene compounds (pre-catalysts) useful in this invention include half and full sandwich compounds having one or more bulky ligands bonded to at least one metal atom. Typical bulky ligand
35 metallocene compounds are generally described as containing one or more bulky

5 ligand(s) and one or more leaving group(s) bonded to at least one metal atom. The bulky ligands are generally represented by one or more open, acyclic, or fused ring(s) or ring system(s) or a combination thereof. These bulky ligands, preferably the ring(s) or ring system(s) are typically composed of atoms selected from Groups 13 to 16 atoms of the Periodic Table of Elements, preferably the
10 atoms are selected from the group consisting of carbon, nitrogen, oxygen, silicon, sulfur, phosphorous, germanium, boron and aluminum or a combination thereof. Most preferably, the ring(s) or ring system(s) are composed of carbon atoms such as but not limited to those cyclopentadienyl ligands or cyclopentadienyl-type ligand structures or other similar functioning ligand structure such as a
15 pentadienyl, a cyclooctatetraendiyl, a cyclobutadienyl, or a substituted allyl ligand. Other ligands that can function similarly to a cyclopentadienyl-type ligand include amides, phosphides, imines, phosphinimines, amidinates, and ortho-substituted phenoxides. The metal atom is preferably selected from Groups 3 through 15 and or lanthanide or actinide series of the Periodic Table of Elements.
20 Preferably the metal is a transition metal from Groups 3 through 12, more preferably Groups 4, 5 and 6, and most preferably the transition metal is from Group 4.

In one embodiment, the catalyst composition useful in the invention
25 includes one or more bulky ligand metallocene catalyst compounds represented by the formula:



30 where M is a metal atom from the Periodic Table of the Elements and may be a Group 3 to 12 metal or from the lanthanide or actinide series of the Periodic Table of Elements, preferably M is a Group 4, 5 or 6 transition metal, more preferably M is a Group 4 transition metal, even more preferably M is zirconium, hafnium or titanium. The bulky ligands, L^A and L^B , are open, acyclic or fused ring(s) or ring

5 system(s) and are any ancillary ligand system, including unsubstituted or substituted, cyclopentadienyl ligands or cyclopentadienyl-type ligands, heteroatom substituted and/or heteroatom containing cyclopentadienyl-type ligands. Non-limiting examples of bulky ligands include cyclopentadienyl ligands, cyclopentaphenanthrenyl ligands, indenyl ligands, benzindenyl ligands, fluorenyl ligands, dibenzo[b,h]fluorenyl ligands, benzo[b]fluorenyl ligands, 10 cyclooctatetraendiyl ligands, cyclopentacyclododecene ligands, azenyl ligands, azulene ligands, pentalene ligands, phosphoyl ligands, phosphinimine (WO 99/40125), pyrrolyl ligands, pyrozolyl ligands, carbazolyl ligands, boratobenzene ligands and the like, including hydrogenated versions thereof, for example 15 tetrahydroindenyl ligands. In one embodiment, L^A and L^B may be any other ligand structure capable of π -bonding to M. In yet another embodiment, the atomic molecular weight (MW) of L^A or L^B exceeds 60 a.m.u., preferably greater than 65 a.m.u. In another embodiment, L^A and L^B may comprise one or more heteroatoms, for example, nitrogen, silicon, boron, germanium, sulfur and 20 phosphorous, in combination with carbon atoms to form an open, acyclic, or preferably a fused, ring or ring system, for example, a hetero-cyclopentadienyl ancillary ligand. Other L^A and L^B bulky ligands include but are not limited to bulky amides, phosphides, alkoxides, aryloxides, imides, carbolides, borollides, porphyrins, phthalocyanines, corrins and other polyazomacrocycles.

25 Independently, each L^A and L^B may be the same or different type of bulky ligand that is bonded to M. In one embodiment of Formula 1 only one of either L^A or L^B is present.

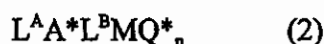
Independently, each L^A and L^B may be unsubstituted or substituted with a combination of substituent groups R^* . Non-limiting examples of substituent 30 groups R^* include one or more from the group selected from hydrogen, or linear or branched alkyl radicals, alkenyl radicals, alkynyl radicals, cycloalkyl radicals, aryl radicals, acyl radicals, aroyl radicals, alkoxy radicals, aryloxy radicals, alkylthio radicals, dialkylamino radicals, alkoxycarbonyl radicals, aryloxycarbonyl radicals, carbomoyl radicals, alkyl- or dialkyl- carbamoyl

5 radicals, acyloxy radicals, acylamino radicals, aroylamino radicals or combination thereof. In a preferred embodiment, substituent groups R^* have up to 50 non-hydrogen atoms, preferably from 1 to 30 carbon, that can also be substituted with halogens or heteroatoms or the like. Non-limiting examples of alkyl substituents R^* include methyl, ethyl, propyl, butyl, pentyl, hexyl, cyclopentyl, cyclohexyl, 10 benzyl or phenyl groups and the like, including all their isomers, for example tertiary butyl, isopropyl, and the like. Other hydrocarbyl radicals include fluoromethyl, fluoroethyl, difluoroethyl, iodopropyl, bromohexyl, chlorobenzyl and hydrocarbyl substituted organometalloid radicals including trimethylsilyl, trimethylgermyl, methyldiethylsilyl and the like; and halocarbyl-substituted 15 organometalloid radicals including tris(trifluoromethyl)silyl, methyl-bis(difluoromethyl)silyl, bromomethyldimethylgermyl and the like; and disubstituted boron radicals including dimethylboron for example; and disubstituted pnictogen radicals including dimethylamine, dimethylphosphine, diphenylamine, methylphenylphosphine, chalcogen radicals including methoxy, 20 ethoxy, propoxy, phenoxy, methylsulfide and ethylsulfide. Non-hydrogen substituents R^* include the atoms carbon, silicon, boron, aluminum, nitrogen, phosphorous, oxygen, tin, sulfur, germanium and the like, including olefins such as but not limited to olefinically unsaturated substituents including vinyl-terminated ligands, for example but-3-enyl, prop-2-enyl, hex-5-enyl and the like.

25 Also, at least two R^* groups, preferably two adjacent R groups, are joined to form a ring structure having from 3 to 30 atoms selected from carbon, nitrogen, oxygen, phosphorous, silicon, germanium, aluminum, boron or a combination thereof. Also, a substituent group, R^* , may also be a diradical bonded to L at one end and forming a carbon sigma bond to the metal M . Other ligands may be bonded to the 30 metal M , such as at least one leaving group Q^* . In one embodiment, Q^* is a monoanionic labile ligand having a sigma-bond to M . Depending on the oxidation state of the metal, the value for n is 0, 1 or 2 such that Formula 1 above represents a neutral bulky ligand metallocene catalyst compound. Non-limiting examples of Q^* ligands include weak bases such as amines, phosphines, ethers, carboxylates,

5 dienes, hydrocarbyl radicals having from 1 to 20 carbon atoms, hydrides or halogens and the like or a combination thereof. In another embodiment, two or more Q*'s form a part of a fused ring or ring system. Other examples of Q* ligands include those substituents for R* as described above and including cyclobutyl, cyclohexyl, heptyl, tolyl, trifluoromethyl, tetramethylene (both Q*),
 10 pentamethylene (both Q*), methyldiene (both Q*), methoxy, ethoxy, propoxy, phenoxy, bis(N-methylanilide), dimethylamide, dimethylphosphide radicals and the like.

In another embodiment, the catalyst composition useful in the invention
 15 may include one or more bulky ligand metallocene catalyst compounds where L^A and L^B of Formula 1 are bridged to each other by at least one bridging group, A*, as represented by Formula 2.



20

The compounds of Formula 2 are known as bridged, bulky ligand metallocene catalyst compounds. L^A, L^B, M, Q* and n are as defined above. Non-limiting examples of bridging group A* include bridging groups containing at least one Group 13 to 16 atom, often referred to as a divalent moiety such as but not limited
 25 to at least one of a carbon, oxygen, nitrogen, silicon, aluminum, boron, germanium and tin atom or a combination thereof. Preferably bridging group A* contains a carbon, silicon or germanium atom, most preferably A* contains at least one silicon atom or at least one carbon atom. The bridging group A* may also contain substituent groups R* as defined above including halogens and iron.
 30 Non-limiting examples of bridging group A* may be represented by R'₂C, R'₂CCR'₂, R'₂Si, R'₂SiCR'₂, R'₂SiSiR'₂, R'₂Ge, R'P, R'N, R'B where R' is independently, a radical group which is hydride, hydrocarbyl, substituted hydrocarbyl, halocarbyl, substituted halocarbyl, hydrocarbyl-substituted organometalloid, halocarbyl-substituted organometalloid, disubstituted boron,

5 disubstituted pnictogen, substituted chalcogen, or halogen or two or more R' may be joined to form a ring or ring system. In one embodiment, the bridged, bulky ligand metallocene catalyst compounds of Formula 2 have two or more bridging groups A* (EP 664 301 B1).

10 In another embodiment, the bulky ligand metallocene catalyst compounds are those where the R* substituents on the bulky ligands L^A and L^B of Formulas 1 and 2 are substituted with the same or different number of substituents on each of the bulky ligands. In another embodiment, the bulky ligands L^A and L^B of Formulas 1 and 2 are different from each other.

15

 Other bulky ligand metallocene catalyst compounds and catalyst systems useful in the invention may include those described in U.S. Patent Nos. 5,064,802, 5,145,819, 5,149,819, 5,243,001, 5,239,022, 5,276,208, 5,296,434, 5,321,106, 5,329,031, 5,304,614, 5,677,401, 5,723,398, 5,753,578, 5,854,363, 5,856,547
20 5,858,903, 5,859,158, 5,900,517 and 5,939,503 and PCT publications WO 93/08221, WO 93/08199, WO 95/07140, WO 98/11144, WO 98/41530, WO 98/41529, WO 98/46650, WO 99/02540 and WO 99/14221 and European publications EP-A-0 578 838, EP-A-0 638 595, EP-B-0 513 380, EP-A1-0 816 372, EP-A2-0 839 834, EP-B1-0 632 819, EP-B1-0 748 821 and EP-B1-0 757
25 996, all of which are herein fully incorporated by reference.

 In another embodiment, the catalyst compositions useful in the invention may include bridged heteroatom, mono-bulky ligand metallocene compounds. These types of catalysts and catalyst systems are described in, for example, PCT
30 publication WO 92/00333, WO 94/07928, WO 91/ 04257, WO 94/03506, WO96/00244, WO 97/15602 and WO 99/20637 and U.S. Patent Nos. 5,057,475, 5,096,867, 5,055,438, 5,198,401, 5,227,440 and 5,264,405 and European publication EP-A-0 420 436, all of which are herein fully incorporated by reference.

5

In another embodiment, the catalyst composition useful in the invention includes one or more bulky ligand metallocene catalyst compounds represented by Formula 3:



where M is a Group 3 to 16 metal atom or a metal selected from the Group of actinides and lanthanides of the Periodic Table of Elements, preferably M is a Group 3 to 12 transition metal, and more preferably M is a Group 4, 5 or 6 transition metal, and most preferably M is a Group 4 transition metal in any oxidation state, and is especially titanium; L^C is a substituted or unsubstituted bulky ligand bonded to M; J^* is bonded to M; A^* is bonded to J^* and L^C ; J^* is a heteroatom ancillary ligand; and A^* is a bridging group; Q^* is a univalent anionic ligand; and n is the integer 0, 1 or 2. In Formula 3 above, L^C , A^* and J^* form a fused ring system. In an embodiment, L^C of Formula 3 is as defined above for L^A . A^* , M and Q^* of Formula 3 are as defined above in Formula 1. In Formula 3, J^* is a heteroatom containing ligand in which J^* is an element with a coordination number of three from Group 15 or an element with a coordination number of two from Group 16 of the Periodic Table of Elements. Preferably J^* contains a nitrogen, phosphorus, oxygen or sulfur atom with nitrogen being most preferred. In an embodiment of the invention, the bulky ligand metallocene catalyst compounds are heterocyclic ligand complexes where the bulky ligands, the ring(s) or ring system(s), include one or more heteroatoms or a combination thereof. Non-limiting examples of heteroatoms include a Group 13 to 16 element, preferably nitrogen, boron, sulfur, oxygen, aluminum, silicon, phosphorous and tin. Examples of these bulky ligand metallocene catalyst compounds are described in WO 96/33202, WO 96/34021, WO 97/17379 and WO 98/22486 and EP-A1-0 874 005 and U.S. Patent No. 5,637,660, 5,539,124, 5,554,775,

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5 5,756,611, 5,233,049, 5,744,417, and 5,856,258 all of which are herein incorporated by reference.

In one embodiment, the bulky ligand metallocene compounds (pre-catalysts) are those complexes based on bidentate ligands containing pyridine or quinoline moieties, such as those described in U.S. Application Serial No. 10 09/103,620 filed June 23, 1998, which is herein incorporated by reference. In another embodiment, the bulky ligand metallocene catalyst compounds are those described in PCT publications WO 99/01481 and WO 98/42664, which are fully incorporated herein by reference.

15

In another embodiment, the bulky ligand metallocene catalyst compound is a complex of a metal, preferably a transition metal, a bulky ligand, preferably a substituted or unsubstituted pi-bonded ligand, and one or more heteroallyl moieties, such as those described in U.S. Patent Nos. 5,527,752 and 5,747,406 and 20 EP-B1-0 735 057, all of which are herein fully incorporated by reference.

In another embodiment, the bulky ligand metallocene catalyst compounds are those described in PCT publications WO 99/01481 and WO 98/42664, which are fully incorporated herein by reference.

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Useful Group 6 bulky ligand metallocene catalyst systems are described in U.S. Patent No. 5,942,462, which is incorporated herein by reference.

Still other useful catalysts include those multinuclear metallocene catalysts 30 as described in WO 99/20665 and 6,010,794 which are herein incorporated herein by reference. Other metallocene catalysts include those described in EP 0 950 667 A1, double cross-linked metallocene catalysts (EP 0 970 074 A1), tethered

- 5 metallocenes (EP 970 963 A2) and those sulfonyl catalysts described in U.S. Patent No. 6,008,394, which are incorporated herein by reference.

It is also contemplated that in one embodiment the bulky ligand metallocene catalysts, described above, include their structural or optical or
10 enantiomeric isomers (meso and racemic isomers, for example see U.S. Patent No. 5,852,143, incorporated herein by reference) and mixtures thereof.

It is further contemplated that any one of the bulky ligand metallocene catalyst compounds, described above, have at least one fluoride or fluorine
15 containing leaving group as described in U.S. Application Serial No. 09/191,916 filed November 13, 1998.

The Group 15 containing metal compounds utilized in the catalyst composition of the invention are prepared by methods known in the art, such as
20 those disclosed in EP 0 893 454 A1, U.S. Patent No. 5,889,128 and the references cited in U.S. Patent No. 5,889,128 which are all herein incorporated by reference. U.S. Application Serial Number 09/312,878, filed May 17, 1999, discloses a gas or slurry phase polymerization process using a supported bisamide catalyst, which is also incorporated herein by reference. For additional information of Group 15
25 containing metal compounds, please see Mitsui Chemicals, Inc. in EP 0 893 454 A1 which discloses transition metal amides combined with activators to polymerize olefins.

In one embodiment the Group 15 containing metal compound is allowed to
30 age prior to use as a polymerization. It has been noted on at least one occasion that one such catalyst compound (aged at least 48 hours) performed better than a newly prepared catalyst compound.

- 5 It is further contemplated that bis-amide based pre-catalysts may be used. Exemplary compounds include those described in the patent literature. International patent publications WO 96/23010, WO 97/48735 and Gibson, et al., Chem. Comm., pp. 849-850 (1998), which disclose diimine-based ligands for Group-8-10 compounds that undergo ionic activation and polymerize olefins.
- 10 Polymerization catalyst systems from Group-5-10 metals, in which the active center is highly oxidized and stabilized by low-coordination-number, polyanionic, ligand systems, are described in U.S. Patent 5,502,124 and its divisional U.S. Patent 5,504,049. See also the Group-5 organometallic catalyst compounds of U.S. Patent 5,851,945 and the tridentate-ligand-containing, Group-5-10,
- 15 organometallic catalysts of U.S. Patent No. 6, 294, 495. Group-11 catalyst precursor compounds, activatable with ionizing cocatalysts, useful for olefin and vinylic polar molecules are described in WO 99/30822.

- Other useful catalyst compounds are those Group 5 and 6 metal imido
- 20 complexes described in EP-A2-0 816 384 and U.S. Patent No. 5,851,945, which is incorporated herein by reference. In addition, metallocene catalysts include bridged bis(arylamido) Group 4 compounds described by D.H. McConville, et al., in *Organometallics* 1995, 14, 5478-5480, which is herein incorporated by reference. In addition, bridged bis(amido) catalyst
- 25 compounds are described in WO 96/27439, which is herein incorporated by reference. Other useful catalysts are described as bis(hydroxy aromatic nitrogen ligands) in U.S. Patent No. 5,852,146, which is incorporated herein by reference. Other useful catalysts containing one or more Group 15 atoms include those described in WO 98/46651, which is herein incorporated herein
- 30 by reference.

 U.S. Patent 5,318,935 describes bridged and unbridged, bisamido catalyst compounds of Group-4 metals capable of α -olefins polymerization. Bridged bi(arylamido)-Group-4 compounds for olefin polymerization are described by D.

5 H. McConville, et al., in *Organometallics* 1995, 14, 5478-5480. This reference presents synthetic methods and compound characterizations. Further work appearing in D. H. McConville, et al, *Macromolecules* 1996, 29, 5241-5243, describes bridged bis(arylamido)-Group-4 compounds that are polymerization catalysts for 1-hexene. Additional invention-suitable transition metal compounds
10 include those described in WO 96/40805. Cationic Group-3- or Lanthanide-metal olefin polymerization complexes are disclosed in copending U.S. Application Ser. No. 09/408,050, filed 29 September 1999. A monoanionic bidentate ligand and two monoanionic ligands stabilize those catalyst precursors, which can be activated with this invention's ionic cocatalysts.

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The literature describes many additional suitable catalyst-precursor compounds. Compounds that contain abstractable ligands or that can be alkylated to contain abstractable ligands suit this invention. See, for instance, V. C. Gibson, et al; "The Search for New-Generation Olefin Polymerization Catalysts: Life
20 Beyond Metallocenes", *Angew. Chem. Int. Ed.*, 38, 428-447 (1999).

This invention may also be practiced with the catalysts containing phenoxide ligands such as those disclosed in EP 0 874 005 A1, which in incorporated by reference herein.

25

In another embodiment, conventional-type transition metal catalysts may be used in the practice of this invention. Conventional-type transition metal catalysts are those traditional Ziegler-Natta, vanadium and Phillips-type catalysts well known in the art. Such as, for example Ziegler-Natta catalysts as described in
30 Ziegler-Natta Catalysts and Polymerizations, John Boor, Academic Press, New York, 1979. Examples of conventional-type transition metal catalysts are also discussed in U.S. Patent Nos. 4,115,639, 4,077,904, 4,482,687, 4,564,605, 4,721,763, 4,879,359 and 4,960,741, all of which are herein fully incorporated by reference. The conventional-type transition metal catalyst compounds that may be

- 5 used in the present invention include transition metal compounds from Groups 3 to 17, preferably 4 to 12, more preferably 4 to 6 of the Periodic Table of Elements.

Preferred conventional-type transition metal catalysts may be represented by the formula: MR_x , where M is a metal from Groups 3 to 17, preferably Group 4 to 6, more preferably Group 4, most preferably titanium; R is a halogen or a hydrocarbyloxy group; and x is the oxidation state of the metal M. Non-limiting examples of R include alkoxy, phenoxy, bromide, chloride and fluoride. Non-limiting examples of conventional-type transition metal catalysts where M is titanium include $TiCl_4$, $TiBr_4$, $Ti(OC_2H_5)_3Cl$, $Ti(OC_2H_5)_3Cl_3$, $Ti(OC_4H_9)_3Cl$, $Ti(OC_3H_7)_2Cl_2$, $Ti(OC_2H_5)_2Br_2$, $TiCl_3 \cdot 1/3 AlCl_3$ and $Ti(OC_{12}H_{25})Cl_3$.

Conventional-type transition metal catalyst compounds based on magnesium/titanium electron-donor complexes that are useful in the invention are described in, for example, U.S. Patent Nos. 4,302,565 and 4,302,566, which are herein fully incorporate by reference. The $MgTiCl_6$ (ethyl acetate)₄ derivative is particularly preferred.

British Patent Application 2,105,355 and U.S. Patent No. 5,317,036, herein incorporated by reference, describes various conventional-type vanadium catalyst compounds. Non-limiting examples of conventional-type vanadium catalyst compounds include vanadyl trihalide, alkoxy halides and alkoxides such as $VOCl_3$, $VOCl_2(OBu)$ where Bu =butyl and $VO(OC_2H_5)_3$; vanadium tetra-halide and vanadium alkoxy halides such as VCl_4 and $VCl_3(OBu)$; vanadium and vanadyl acetyl acetonates and chloroacetyl acetonates such as $V(AcAc)_3$ and $VOCl_2(AcAc)$ where (AcAc) is an acetyl acetonate. The preferred conventional-type vanadium catalyst compounds are $VOCl_3$, VCl_4 and $VOCl_2-OR$ where R is a hydrocarbon radical, preferably a C_1 to C_{10} aliphatic or aromatic hydrocarbon radical such as ethyl, phenyl, isopropyl, butyl, propyl, n-butyl, iso-butyl, tertiary-butyl, hexyl, cyclohexyl, naphthyl, etc., and vanadium acetyl acetonates.

5 Conventional-type chromium catalyst compounds, often referred to as Phillips-type catalysts, suitable for use in the present invention include CrO_3 , chromocene, silyl chromate, chromyl chloride (CrO_2Cl_2), chromium-2-ethylhexanoate, chromium acetylacetonate ($\text{Cr}(\text{AcAc})_3$), and the like. Non-limiting examples are disclosed in U.S. Patent Nos. 3,709,853, 3,709,954, 3,231,550,
10 3,242,099 and 4,077,904, which are herein fully incorporated by reference.

Still other conventional-type transition metal catalyst compounds and catalyst systems suitable for use in the present invention are disclosed in U.S. Patent Nos. 4,124,532, 4,302,565, 4,302,566, 4,376,062, 4,379,758, 5,066,737,
15 5,763,723, 5,849,655, 5,852,144, 5,854,164 and 5,869,585 and published EP-A2 0 416 815 A2 and EP-A1 0 420 436, which are all herein incorporated by reference.

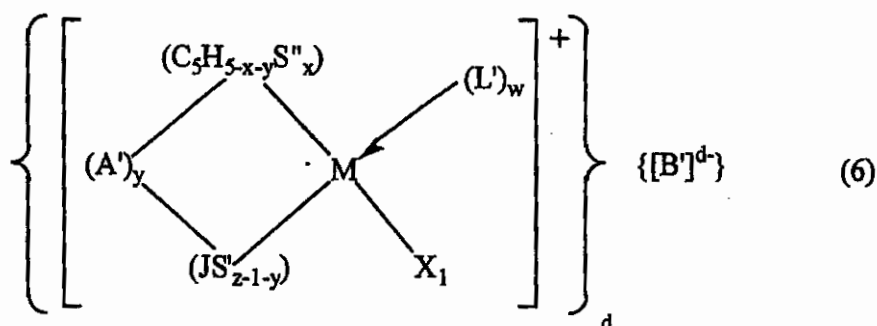
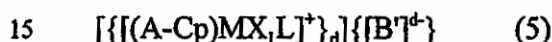
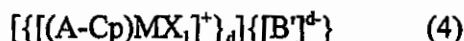
Other catalysts may include cationic catalysts such as AlCl_3 , and other cobalt, iron, nickel and palladium catalysts well known in the art. See for example
20 U.S. Patent Nos. 3,487,112, 4,472,559, 4,182,814 and 4,689,437, all of which are incorporated herein by reference.

It is also contemplated that other catalysts can be combined with the catalyst compounds in the catalyst composition useful in the invention. For
25 example, see U.S. Patent Nos. 4,937,299, 4,935,474, 5,281,679, 5,359,015, 5,470,811, and 5,719,241 all of which are herein fully incorporated herein reference.

It is further contemplated that one or more of the catalyst compounds
30 described above or catalyst systems may be used in combination with one or more conventional catalyst compounds or catalyst systems. Non-limiting examples of mixed catalysts and catalyst systems are described in U.S. Patent Nos. 4,159,965, 4,325,837, 4,701,432, 5,124,418, 5,077,255, 5,183,867, 5,391,660, 5,395,810,

5 5,691,264, 5,723,399 and 5,767,031 and PCT Publication WO 96/23010 published August 1, 1996, all of which are herein fully incorporated by reference.

Preferred metallocene catalysts used in this invention can more specifically be represented by one of the following general formulae (all references to Groups being the new Group notation of the Period Table of the Elements as described by Chemical and Engineering News, 63(5), 27, 1985):



wherein:

(A-Cp) is either (Cp), (Cp*) or Cp-A'-Cp*; Cp and Cp* are the same or different cyclopentadienyl rings substituted with from zero to five substituent groups S'', each substituent group S'' being, independently, a radical group which is a hydrocarbyl, substituted-hydrocarbyl, halocarbyl, substituted-halocarbyl, hydrocarbyl-substituted organometalloid, halocarbyl-substituted organometalloid, disubstituted boron, disubstituted pnictogen, substituted chalcogen or halogen radicals, or Cp and Cp* are cyclopentadienyl rings in which any two adjacent S'' groups are joined forming a C₄ to C₂₀ ring to give a saturated or unsaturated polycyclic cyclopentadienyl ligand; Cp and Cp* may also have one or two carbon atoms within the ring replaced by a Group 15 or 16 element especially, S, O, N or P;

5 A' is a bridging group;

(C₅H_{5-y-x}S^{''})_x is a cyclopentadienyl ring substituted with from zero to five S^{''} radicals as defined above;

10 x is from 0 to 5 denoting the degree of substitution;

M is titanium, zirconium or hafnium;

15 X₁ is a hydride radical, hydrocarbyl radical, substituted- hydrocarbyl radical, hydrocarbyl-substituted organometalloid radical or halocarbyl-substituted organometalloid radical which radical may optionally be covalently bonded to both or either M and L or L' or all or any M, S^{''} or S', and provided that X₁ is not a substituted or unsubstituted cyclopentadienyl ring;

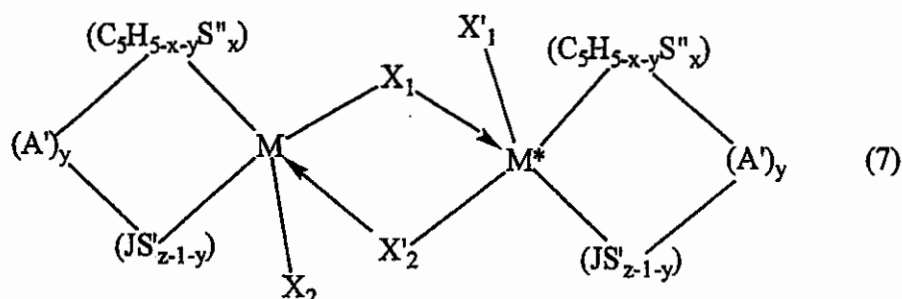
20 (JS'_{z-1-y}) is a heteroatom ligand in which J is an element from Group 15 of the Periodic Table of Elements with a coordination number of 3 or an element from Group 16 with a coordination number of 2; S' is a radical group which is a hydrocarbyl, substituted hydrocarbyl, halocarbyl, substituted halocarbyl, hydrocarbyl-substituted organometalloid, or halocarbyl-substituted organometalloid; and z is the coordination number of the element J;

25

y is 0 or 1;

30 L is an olefin, diolefin or aryne ligand. L' is the same as L, and can additionally be an amine, phosphine, ether, or sulfide ligand, or any other neutral Lewis base; L' can also be a second transition metal compound of the same type such that the two metal center M and M* are bridged by X₁ and X'₁, wherein M* has the same meaning as M, X'₁, X₂ and X'₂ have the same meaning as X₁, where such dimeric

- 5 compounds which are precursors to the cationic portion of the catalyst are represented by the formula:



wherein

- 10 w is an integer from 0 to 3;

- B' is a chemically stable, non-nucleophilic anionic complex having a molecular diameter about or greater than 4 Angstroms or an anionic Lewis-acid activator resulting from the reaction of a Lewis-acid activator with the precursor to the
- 15 cationic portion of the catalyst system described in formulae 1-4. When B' is a Lewis-acid activator, X₁ can also be an alkyl group donated by the Lewis-acid activator; and

d is an integer representing the charge of B'.

20

- The catalysts are preferably prepared by combining at least two components. In one preferred method, the first component is a cyclopentadienyl derivative of a Group 4 metal compound containing at least one ligand which will combine with the second component or at least a portion thereof such as a cation
- 25 portion thereof. The second component is an ion-exchange compound comprising a cation which will irreversibly react with at least one ligand contained in said Group 4 metal compound (first component) and a non-coordinating anion which is either a single coordination complex comprising a plurality of lipophilic radicals covalently coordinated to and shielding a central formally charge-bearing metal or

5 metalloid atom or an anion comprising a plurality of boron atoms such as polyhedral boranes, carboranes and metallocarboranes.

 In general, suitable anions for the second component may be any stable and bulky anionic complex having the following molecular attributes: 1) the anion
10 should have a molecular diameter greater than 4 Angstroms; 2) the anion should form stable ammonium salts; 3) the negative charge on the anion should be delocalized over the framework of the anion or be localized within the core of the anion; 4) the anion should be a relatively poor nucleophile; and 5) the anion should not be a powerful reducing or oxidizing agent. Anions meeting these
15 criteria-- such as polynuclear boranes, carboranes, metallocarboranes, polyoxoanions and anionic coordination complexes are well described in the chemical literature.

 The cation portion of the second component may comprise Bronsted acids
20 such as protons or protonated Lewis bases or may comprise Lewis acids such as ferricinium, tropylium, triphenylcarbenium or silver cations.

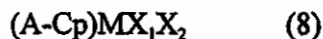
 In another preferred method, the second component is a Lewis-acid complex which will react with at least one ligand of the first component, thereby
25 forming an ionic species described in formulae 4-6 with the ligand abstracted from the first component now bound to the second component. Alumoxanes and especially methylalumoxane, the product formed from the reaction of trimethylaluminum in an aliphatic or aromatic hydrocarbon with stoichiometric quantities of water, are particularly preferred Lewis-acid second components.
30 Modified alumoxanes are also preferred. Alumoxanes are well known in the art and methods for their preparation are illustrated by U.S. Pat. Nos. 4,542,199; 4,544,762; 5,015,749; and 5,041,585. A technique for preparing modified alumoxanes has been disclosed in U.S. Pat. No. 5,041,584, in EPA 0 516 476, and in EPA 0 561 476, which are incorporated by reference herein.

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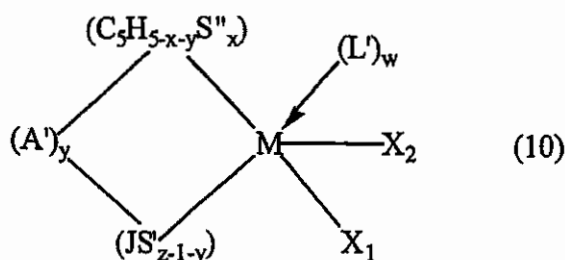
Upon combination of the first and second components, the second component reacts with one of the ligands of the first component, thereby generating an anion pair consisting of a Group 4 metal cation and the aforementioned anion, which anion is compatible with and non-coordinating towards the Group 4 metal cation formed from the first component. The anion of the second compound must be capable of stabilizing the Group 4 metal cation's ability to function as a catalyst and must be sufficiently labile to permit displacement by an olefin, diolefin or an acetylenically unsaturated monomer during polymerization. The catalysts of this invention may be supported. U.S. Pat. Nos. 4,808, 561, issued Feb. 28, 1989; 4,897,455 issued Jan. 3, 1990; 5,057,475 issued Oct. 15, 1991; U.S. patent application Ser. No. 459,921 (published as PCT International publication WO 91/09882), Canadian Patent 1,268,753, U.S. Pat. No. 5,240,894 and WO 94 03506 disclose such supported catalysts and the methods to produce such and are herein incorporated by reference.

20

The Group 4 metal compounds; i.e., titanium, zirconium and hafnium metallocene compounds, useful as first compounds (pre-catalysts) in the preparation of the preferred metallocene catalysts of this invention are cyclopentadienyl derivatives of titanium, zirconium and hafnium. In general, useful titanocenes, zirconocenes and hafnocenes may be represented by the following general formulae:



5



wherein:

- (A-Cp) is either (Cp)(Cp*) or Cp-A'-Cp*; Cp and Cp* are the same or different
 10 cyclopentadienyl rings substituted with from zero to five substituent groups S'',
 each substituent group S'' being, independently, a radical group which is a
 hydrocarbyl, substituted-hydrocarbyl, halocarbyl, substituted-halocarbyl,
 hydrocarbyl-substituted organometalloid, halocarbyl-substituted organometalloid,
 disubstituted boron, disubstituted pnictogen, substituted chalcogen or halogen
 15 radicals, or Cp and Cp* are cyclopentadienyl rings in which any two adjacent S''
 groups are joined forming a C₄ to C₂₀ ring to give a saturated or unsaturated
 polycyclic cyclopentadienyl ligand;

A' is a bridging group;

20

y is 0 or 1;

(C₅H_{5-y-x}S''_x) is a cyclopentadienyl ring substituted with from zero to five S''
 radicals as defined above;

25

x is from 0 to 5 denoting the degree of substitution;

(JS'_{z-1-y}) is a heteroatom ligand in which J is an element from Group 15 of the
 Periodic Table of Elements with a coordination number of 3 or an element from
 30 Group 16 with a coordination number of 2, S' is a radical group which is a

5 hydrocarbyl, substituted hydrocarbyl, halocarbyl, substituted halocarbyl, hydrocarbyl-substituted organometalloid, or halocarbyl-substituted organometalloid; and z is the coordination number of the element J ;

L is an olefin, diolefin or aryne ligand. L' is the same as L and can additionally be
10 an amine, phosphine, ether, or sulfide ligand, or any other neutral Lewis base; L' can also be a second transition metal compound of the same type such that the two metal centers M and M^* are bridged by X_1 and X'_1 , wherein M^* has the same meaning as M , X'_1 has the same meaning as X_1 and X'_2 has the same meaning as X_2 where such dimeric compounds which are precursors to the cationic portion of
15 the catalyst are represented by formula 7 above;

w is an integer from 0 to 3; and

X_1 and X_2 are, independently, hydride radicals, hydrocarbyl radicals, substituted
20 hydrocarbyl radicals, halocarbyl radicals, substituted halocarbyl radicals, and hydrocarbyl- and halocarbyl-substituted organometalloid radicals, substituted pnictogen radicals, or substituted chalcogen radicals; or X_1 and X_2 are joined and bound to the metal atom to form a metallacycle ring containing from about 3 to about 20 carbon atoms; or X_1 and X_2 together can be an olefin, diolefin or aryne
25 ligand; or when Lewis-acid activators, such as methylalumoxane, which are capable of donating an X_1 ligand as described above to the transition metal component are used, X_1 and X_2 may independently be a halogen, alkoxide, aryloxide, amide, phosphide or other univalent anionic ligand or both X_1 and X_2 can also be joined to form an anionic chelating ligand and with the proviso that X_1
30 and X_2 are not a substituted or unsubstituted cyclopentadienyl ring.

Table A depicts representative constituent moieties for the metallocene components of formulae 7-10. The list is for illustrative purposes only and should not be construed to be limiting in any way. A number of final components may be

5 formed by permuting all possible combinations of the constituent moieties with each other. When hydrocarbyl radicals including alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, cycloalkynyl and aromatic radicals are disclosed in this application the term includes all isomers. For example, butyl includes n-butyl, 2-methylpropyl, 1-methylpropyl, tert-butyl, and cyclobutyl; pentyl includes n-
10 pentyl, 1-methylbutyl, 2-methylbutyl, 3-methylbutyl, 1-ethylpropyl, neopentyl, cyclopentyl and methylcyclobutyl; butenyl includes E and Z forms of 1-butenyl, 2-butenyl, 3-butenyl, 1-methyl-1-propenyl, 1-methyl-2-propenyl, 2-methyl-1-propenyl and 2-methyl-2-propenyl. This includes when a radical is bonded to another group, for example, propylcyclopentadienyl include n-
15 propylcyclopentadienyl, isopropylcyclopentadienyl and cyclopropylcyclopentadienyl. In general, the ligands or groups illustrated in Table A include all isomeric forms. For example, dimethylcyclopentadienyl includes 1,2-dimethylcyclopentadienyl and 1,3-dimethylcyclopentadienyl; methylindenyl includes 1-methylindenyl, 2-methylindenyl, 3-methylindenyl, 4-methylindenyl, 5-
20 methylindenyl, 6-methylindenyl and 7-methylindenyl; methylethylphenyl includes ortho-methylethylphenyl, meta-methylethylphenyl and para-methylethylphenyl. Examples of specific invention catalyst precursors take the following formula where some components are listed in Table A. To illustrate members of the transition metal component, select any combination of the species listed in Tables
25 A. For nomenclature purposes, for the bridging group, A', the words "silyl" and "silylene" are used interchangeably, and represent a diradical species. For the bridging group A', "ethylene" refers to a 1,2-ethylene linkage and is distinguished from ethene-1,1-diyl. Thus, for the bridging group A', "ethylene" and "1,2-ethylene" are used interchangeably. For compounds processing a bridging group,
30 A', the bridge position on the cyclopentadienyl-type ring is always considered the 1-position. Thus, for example, the use of "1-fluorenyl" is interchangeable with the use of "fluorenyl"

- 5 Illustrative compounds of the formula 8 type are:
 bis(cyclopentadienyl)hafnium dimethyl, ethylenebis(tetrahydroindenyl)zirconium
 dihydride, bis(pentamethyl)zirconium diethyl, dimethylsilyl(1-
 fluorenyl)(cyclopentadienyl)titanium dichloride and the like. Illustrative
 compounds of the formula 9 type are: bis(cyclopentadienyl)(1,3-
 10 butadiene)zirconium, bis(cyclopentadienyl)(2,3-dimethyl-1,3-
 butadiene)zirconium, bis(pentamethylcyclopentadienyl)(benzene)zirconium,
 bis(pentamethylcyclopentadienyl)titanium ethylene and the like. Illustrative
 compounds of the formula 10 type are:
 dimethylsilyl(tetramethylcyclopentadienyl)(t-butylamido)zirconium dichloride,
 15 ethylene(methylcyclopentadienyl)(phenylamido)titanium dimethyl,
 methylphenylsilyl(indenyl)(phenylphosphido)hafnium dihydride and
 (pentamethylcyclopentadienyl)(di-t-butylamido)hafnium dimethoxide.

- The conditions under which complexes containing neutral Lewis base
 20 ligands such as ether or those which form dimeric compounds is determined by
 the steric bulk of the ligands about the metal center. For example, the t-butyl
 group in $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-Bu})\text{ZrCl}_2$ has greater steric requirements than the
 phenyl in $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{NPh})\text{ZrCl}_2 \cdot \text{Et}_2\text{O}$ thereby not permitting ether
 coordination in the former compound in its solid state. Similarly, due to the
 25 decreased steric bulk of the trimethylsilylcyclopentadienyl group in
 $[\text{Me}_2\text{Si}(\text{Me}_3\text{SiC}_5\text{H}_3)(\text{N-t-Bu})\text{ZrCl}_2]_2$ versus that of the tetramethylcyclopentadienyl
 group in $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-Bu})\text{ZrCl}_2$, the former compound is dimeric and the
 latter is not.

30 TABLE A

A'	Cp, Cp*, CpR or (C ₅ H _{5-x} S ^x)
Dimethylsilylene	Cyclopentadienyl
Diethylsilylene	methylcyclopentadienyl
Dipropylsilylene	Dimethylcyclopentadienyl
Dibutylsilylene	Trimethylcyclopentadienyl

dipentylsilylene	Tetramethylcyclopentadienyl
dihexylsilylene	Pentamethylcyclopentadienyl (no A')
diheptylsilylene	Ethylcyclopentadienyl
dioctylsilylene	Diethylcyclopentadienyl
dinonylsilylene	Propylcyclopentadienyl
didecylsilylene	Dipropylcyclopentadienyl
diundecylsilylene	Butylcyclopentadienyl
didodecylsilylene	Dibutylcyclopentadienyl
ditridecylsilylene	Pentylcyclopentadienyl
ditetradecylsilylene	Dipentylcyclopentadienyl
dipentadecylsilylene	Hexylcyclopentadienyl
dihexadecylsilylene	Dihexylcyclopentadienyl
diheptadecylsilylene	Heptylcyclopentadienyl
dioctadecylsilylene	Diheptylcyclopentadienyl
dinonadecylsilylene	Octylcyclopentadienyl
dieicosylsilylene	Dioctylcyclopentadienyl
diheneicosylsilylene	Nonylcyclopentadienyl
didocosylsilylene	Dinonylcyclopentadienyl
ditricosylsilylene	Decylcyclopentadienyl
ditetracosylsilylene	Didecylcyclopentadienyl
dipentacosylsilylene	Undecylcyclopentadienyl
dihexacosylsilylene	Dodecylcyclopentadienyl
diheptacosylsilylene	Tridecylcyclopentadienyl
dioctacosylsilylene	Tetradecylcyclopentadienyl
dinonacosylsilylene	Pentadecylcyclopentadienyl (no A')
ditriacontylsilylene	Hexadecylcyclopentadienyl
dicyclohexylsilylene	Heptadecylcyclopentadienyl
dicyclopentylsilylene	Octadecylcyclopentadienyl
dicycloheptylsilylene	Nonadecylcyclopentadienyl
dicyclooctylsilylene	Eicosylcyclopentadienyl
dicyclodecylsilylene	Heneicosylcyclopentadienyl
dicyclododecylsilylene	Docosylcyclopentadienyl
dinapthylsilylene	Tricosylcyclopentadienyl
diphenylsilylene	Tetracosylcyclopentadienyl
ditolylsilylene	Pentacosylcyclopentadienyl
dibenzylsilylene	Hexacosylcyclopentadienyl
diphenethylsilylene	Heptacosylcyclopentadienyl
di(butylphenethyl)silylene	Octacosylcyclopentadienyl
methylethylsilylene	Nonacosylcyclopentadienyl
methylpropylsilylene	Triacontylcyclopentadienyl
methylbutylsilylene	Cyclohexylcyclopentadienyl

methylhexylsilylene	Phenylcyclopentadienyl
methylphenylsilylene	Diphenylcyclopentadienyl
ethylphenylsilylene	triphenylcyclopentadienyl
ethylpropylsilylene	tetraphenylcyclopentadienyl
ethylbutylsilylene	pentaphenylcyclopentadienyl
propylphenylsilylene	tolylcyclopentadienyl
dimethylgermylene	benzylcyclopentadienyl
diethylgermylene	phenethylcyclopentadienyl
diphenylgermylene	cyclohexylmethylcyclopentadienyl
methylphenylgermylene	naphylcyclopentadienyl
cyclotetramethylenesilylene	methylphenylcyclopentadienyl
cyclopentamethylenesilylene	methyltolylcyclopentadienyl
cyclotrimethylenesilylene	methylethylcyclopentadienyl
cyclohexylazanediy1	methylpropylcyclopentadienyl
butylazanediy1	methylbutylcyclopentadienyl
methylazanediy1	methylpentylcyclopentadienyl
phenylazanediy1	methylhexylcyclopentadienyl
perfluorophenylazanediy1	methylheptylcyclopentadienyl
methylphosphanediy1	methyloctylcyclopentadienyl
ethylphosphanediy1	methylnonylcyclopentadienyl
propylphosphanediy1	methyldecylcyclopentadienyl
butylphosphanediy1	vinylcyclopentadienyl
cyclohexylphosphanediy1	propenylcyclopentadienyl
phenylphosphanediy1	butenylcyclopentadienyl
methylboranediy1	indenyl
phenylboranediy1	methyindenyl
methylene	dimethylindenyl
dimethylmethylene	trimethylindenyl
diethylmethylene	methylpropylindenyl
dibutylmethylene	dimethylpropylindenyl
dipropylmethylene	methyldipropylindenyl
diphenylmethylene	methylethylindenyl
ditolylmethylene	methylbutylindenyl
di(butylphenyl)methylene	ethylindenyl
di(trimethylsilylphenyl)methylene	propylindenyl
di(triethylsilylphenyl)methylene	butylindenyl
dibenzylmethylene	pentylindenyl
cyclotetramethylenemethylene	hexylindenyl
cyclopentamethylenemethylene	heptylindenyl
ethylene	octylindenyl
methylethylene	nonylindenyl

dimethylethylene	decylindenyl
trimethylethylene	phenylindenyl
tetramethylethylene	(fluorophenyl)indenyl
cyclopentylene	(methylphenyl)indenyl
cyclohexylene	biphenylindenyl
cycloheptylene	(bis(trifluoromethyl)phenyl)indenyl
cyclooctylene	napthylindenyl
propanediyl	phenanthrylindenyl
methylpropanediyl	benzylindenyl
dimethylpropanediyl	benzindenyl
trimethylpropanediyl	cyclohexylindenyl
tetramethylpropanediyl	methylphenylindenyl
pentamethylpropanediyl	ethylphenylindenyl
hexamethylpropanediyl	propylphenylindenyl
tetramethyldisiloxylene	methylnapthylindenyl
vinylene	ethylnapthylindenyl
ethene-1,1-diyl	propylnapthylindenyl
divinylsilylene	(methylphenyl)indenyl
dipropenylsilylene	(dimethylphenyl)indenyl
dibutenylsilylene	(ethylphenyl)indenyl
methylvinylsilylene	(diethylphenyl)indenyl
methylpropenylsilylene	(propylphenyl)indenyl
methylbutenylsilylene	(dipropylphenyl)indenyl
dimethylsilylmethylene	methyltetrahydroindenyl
diphenylsilylmethylene	dimethyltetrahydroindenyl
dimethylsilylethylene	dimethyldihydroindenyl
diphenylsilylethylene	dimethyltrihydroindenyl
dimethylsilylpropylene	methylphenyltetrahydroindenyl
diphenylsilylpropylene	methylphenyldihydroindenyl
dimethylstannylene	methylphenyltrihydroindenyl
diphenylstannylene	ethyltetrahydroindenyl
	propyltetrahydroindenyl
	butyltetrahydroindenyl
	phenyltetrahydroindenyl
	fluorenyl
	methylfluorenyl
	dimethylfluorenyl
	trimethylfluorenyl
	ethylfluorenyl
	propylfluorenyl
	butylfluorenyl

	dibutylfluorenyl
	pentylfluorenyl
	hexylfluorenyl
	heptylfluorenyl
	octylfluorenyl
	nonylfluorenyl
	decylfluorenyl
	phenylfluorenyl
	napthylfluorenyl
	benzylfluorenyl
	methylphenylfluorenyl
	ethylphenylfluorenyl
	propylphenylfluorenyl
	methylnapthylfluorenyl
	ethylnapthylfluorenyl
	propylnapthylfluorenyl
	octahydrofluorenyl
	tetrahydrofluorenyl
	octamethyloctahydrodibenzo[b,h]fluorenyl
	tetramethyltetrahydrobenzo[b]fluorenyl
	diphenylmethylcyclopentadienyl
	trimethylsilylcyclopentadienyl
	triethylsilylcyclopentadienyl
	trimethylgermylcyclopentadienyl
	trimethylstannylcyclopentadienyl
	triethylplumbylcyclopentadienyl
	trifluoromethylcyclopentadienyl
	N,N-dimethylamidocyclopentadienyl
	P,P-dimethylphosphidocyclopentadienyl
	N,N-diethylamidocyclopentadienyl
	methoxycyclopentadienyl
	ethoxycyclopentadienyl
	trimethylsiloxy-cyclopentadienyl
	(N,N-dimethylamidomethyl)cyclopentadienyl
	methoxyindenyl
	dimethoxyindenyl
	N,N-dimethylaminoindenyl
	trimethylsiloxyindenyl
	butyldimethylsiloxyindenyl
	bis(N,N-dimethylamino)indenyl
	di(trimethylsiloxy)indenyl

	di(butyldimethylsiloxy)indenyl
	methoxyfluorenyl
	dimethoxyfluorenyl
	N,N-dimethylamino fluorenyl
	trimethylsiloxyfluorenyl
	butyldimethylsiloxyfluorenyl
	dimethoxyfluorenyl
	bis(N,N-dimethylamino)fluorenyl
	di(trimethylsiloxy)fluorenyl
	di(butyldimethylsiloxy)fluorenyl

5

Table A (continued)

(JS' _{z-1-y}) (y = 1)	X ₁ or X ₂	<u>M</u>
Methylamido	chloride	titanium
Ethylamido	bromide	zirconium
Propylamido	iodide	hafnium
Butylamido	fluoride	
Pentylamido	hydride	<u>L or L' (optional)</u>
Hexylamido	methyl	ethylene
Heptylamido	ethyl	propylene
Octylamido	propyl	butene
Nonylamido	butyl	hexene
Decylamido	pentyl	styrene
Eicosylamido	hexyl	hexadiene
Heneicosylamido	heptyl	butadiene
Docosylamido	octyl	dimethylbutadiene
Tricosylamido	nonyl	pentadiene
Tetracosylamido	decyl	methylhexadiene
Pentacosylamido	undecyl	dimethylhexadiene
Hexacosylamido	dodecyl	acetylene
Heptacosylamido	tridecyl	methylacetylene
Octacosylamido	tetradecyl	ethylacetylene
Nonacosylamido	pentadecyl	benzyne
Triacontylamido	hexadecyl	cyclopentene
Phenylamido	heptadecyl	cyclohexene
Tolylamido	octadecyl	
Phenethylamido	nonadecyl	<u>L' (optional)</u>
Benzylamido	eicosyl	diethylether
Cyclobutylamido	heneicosyl	dimethylether
Cyclopentylamido	docosyl	trimethylamine

Cyclohexylamido	tricosyl	triphenylamine
Cycloheptylamido	tetracosyl	triethylamine
Cyclooctylamido	pentacosyl	tricyclohexylphosphine
Cyclononylamido	hexacosyl	triphenylphosphine
Cyclodecylamido	heptacosyl	trimethylphosphine
Cyclododecylamido	octacosyl	tetrahydrofuran
Adamantylamido	nonacosyl	furan
Norbornylamido	triacontyl	thiophene
Perfluorophenylamido	phenyl	dimethylsulfide
Fluorophenylamido	benzyl	diphenylsulfide
Difluorophenylamido	phenethyl	
Oxo	tolyl	
Sulfido	methoxy	
	ethoxy	
(JS' _{x-1-y}) (y = 0)	propoxy	
Methoxide	butoxy	
Ethoxide	dimethylamido	
Phenoxide	diethylamido	
Dimethylphenoxide	methylethylamido	
Dipropylphenoxide	phenoxy	
Methylthio	benzoxo	
Ethylthio	allyl	
Phenylthio		
Dimethylphenylthio		
Dipropylphenylthio		
	X₁ and X₂ together	
	methylidene	
	ethylidene	
	propylidene	
	tetramethylene	
	pentamethylene	
	hexamethylene	
	ethylenedihydroxy	
	butadiene	
	methylbutadiene	
	dimethylbutadiene	
	pentadiene	
	methylpentadiene	
	dimethylpentadiene	
	hexadiene	
	methylhexadiene	

	dimethylhexadiene	

5

Additional preferred catalysts include those described in WO 01/48034, which is incorporated herein by reference. Particularly preferred catalyst compounds include those disclosed at page 9, line 38 to page 25, line 42, page 28, lines 5 to 17, and page 30, line 37 to page 35, line 28.

10

Activators and Activation Methods for Catalyst Compounds

The polymerization pre-catalyst compounds, described above, are typically activated in various ways to yield compounds having a vacant coordination site that will coordinate, insert, and polymerize olefin(s). For the purposes of this patent specification and appended claims, the terms "cocatalyst" and "activator" are used herein interchangeably and are defined to be any compound which can activate any one of the catalyst compounds described above by converting the neutral catalyst compound to a catalytically active catalyst compound cation.

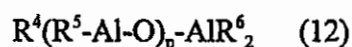
Non-limiting activators, for example, include alumoxanes, aluminum alkyls, ionizing activators, which may be neutral or ionic, and conventional-type cocatalysts. Preferred activators typically include alumoxane compounds, modified alumoxane compounds, and ionizing anion precursor compounds that abstract one reactive, σ -bond, metal ligand making the metal complex cationic and providing a charge-balancing noncoordinating or weakly coordinating anion.

Aluminoxane and Aluminum Alkyl Activators

In one embodiment, alumoxane activators are utilized as an activator in the catalyst composition useful in the invention. Alumoxanes are generally oligomeric compounds containing $-Al(R^1)-O-$ sub-units, where R^1 is an alkyl group. Examples of alumoxanes include methylalumoxane (MAO), modified methylalumoxane (MMAO), ethylalumoxane and isobutylalumoxane.

- 5 Alkylalumoxanes and modified alkylalumoxanes are suitable as catalyst activators, particularly when the abstractable ligand is a halide, alkoxide or amide. Mixtures of different alumoxanes and modified alumoxanes may also be used.

The activator compounds comprising Lewis-acid activators and in
10 particular alumoxanes are represented by the following general formulae:



15



An alumoxane is generally a mixture of both the linear and cyclic compounds. In the general alumoxane formula, R^3 , R^4 , R^5 and R^6 are,
20 independently a C_1-C_{30} alkyl radical, for example, methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, nonyl, decyl, and "p" is an integer from 1 to about 50. Most preferably, R^3 , R^4 , R^5 and R^6 are each methyl and "p" is a least 4. When an alkyl aluminum halide or alkoxide is employed in the preparation of the alumoxane, one or more R^{3-6} groups may be halide or alkoxide. M' is a metal or
25 metalloid, and Q' is a partially or fully fluorinated hydrocarbyl.

It is recognized that alumoxane is not a discrete material. A typical alumoxane will contain free trisubstituted or trialkyl aluminum, bound trisubstituted or trialkyl aluminum, and alumoxane molecules of varying degree of
30 oligomerization. Those methylalumoxanes most preferred contain lower levels of trimethylaluminum. Lower levels of trimethylaluminum can be achieved by reaction of the trimethylaluminum with a Lewis base or by vacuum distillation of the trimethylaluminum or by any other means known in the art. It is also recognized that after reaction with the transition metal compound, some

5 alumoxane molecules are in the anionic form as represented by the anion in equations 4-6, thus for our purposes are considered "non-coordinating" anions.

For further descriptions, see U.S. Patents 4,665,208, 4,952,540, 5,041,584, 5,091,352, 5,206,199, 5,204,419, 4,874,734, 4,924,018, 4,908,463, 4,968,827,
10 5,329,032, 5,248,801, 5,235,081, 5,157,137, 5,103,031 and EP 0 561 476 A1, EP 0 279 586 B1, EP 0 516 476 A, EP 0 594 218 A1 and WO 94/10180.

When the activator is an alumoxane (modified or unmodified), some embodiments select the maximum amount of activator at a 5000-fold molar excess Al/M over the catalyst precursor (per metal catalytic site). The minimum
15 activator-to-catalyst-precursor is a 1:1 molar ratio.

Alumoxanes may be produced by the hydrolysis of the respective trialkylaluminum compound. MMAO may be produced by the hydrolysis of trimethylaluminum and a higher trialkylaluminum such as triisobutylaluminum. MMAO's are generally more soluble in aliphatic solvents and more stable during
20 storage. There are a variety of methods for preparing alumoxane and modified alumoxanes, non-limiting examples of which are described in U.S. Patent No. 4,665,208, 4,952,540, 5,091,352, 5,206,199, 5,204,419, 4,874,734, 4,924,018, 4,908,463, 4,968,827, 5,308,815, 5,329,032, 5,248,801, 5,235,081, 5,157,137, 5,103,031, 5,391,793, 5,391,529, 5,693,838, 5,731,253, 5,731,451, 5,744,656,
25 5,847,177, 5,854,166, 5,856,256 and 5,939,346 and European publications EP-A-0 561 476, EP-B1-0 279 586, EP-A-0 594-218 and EP-B1-0 586 665, and PCT publications WO 94/10180 and WO 99/15534, all of which are herein fully incorporated by reference. It may be preferable to use a visually clear methylalumoxane. A cloudy or gelled alumoxane can be filtered to produce a clear
30 solution or clear alumoxane can be decanted from the cloudy solution. Another alumoxane is a modified methyl alumoxane (MMAO) cocatalyst type 3A (commercially available from Akzo Chemicals, Inc. under the trade name

- 5 Modified Methylalumoxane type 3A, covered under patent number US 5,041,584).

Aluminum alkyl or organoaluminum compounds which may be utilized as activators (or scavengers) include trimethylaluminum, triethylaluminum,
10 triisobutylaluminum, tri-n-hexylaluminum, tri-n-octylaluminum and the like.

Ionizing Activators

It is within the scope of this invention to use an ionizing or stoichiometric
15 activator, neutral or ionic, such as tri (n-butyl) ammonium tetrakis (pentafluorophenyl) boron, a trisperfluorophenyl boron metalloid precursor or a trisperfluoronaphthyl boron metalloid precursor, polyhalogenated heteroborane anions (WO 98/43983), boric acid (U.S. Patent No. 5,942,459) or combination thereof. It is also within the scope of this invention to use neutral or ionic
20 activators alone or in combination with alumoxane or modified alumoxane activators.

Examples of neutral stoichiometric activators include tri-substituted boron, tellurium, aluminum, gallium and indium or mixtures thereof. The three
25 substituent groups are each independently selected from alkyls, alkenyls, halogen, substituted alkyls, aryls, arylhalides, alkoxy and halides. Preferably, the three groups are independently selected from halogen, mono or multicyclic (including halosubstituted) aryls, alkyls, and alkenyl compounds and mixtures thereof, preferred are alkenyl groups having 1 to 20 carbon atoms, alkyl groups having 1 to
30 20 carbon atoms, alkoxy groups having 1 to 20 carbon atoms and aryl groups having 3 to 20 carbon atoms (including substituted aryls). More preferably, the three groups are alkyls having 1 to 4 carbon groups, phenyl, naphthyl or mixtures thereof. Even more preferably, the three groups are halogenated, preferably

5 fluorinated, aryl groups. Most preferably, the neutral stoichiometric activator is tris(perfluorophenyl) boron or tris(perfluoronaphthyl) boron.

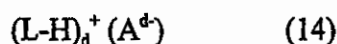
Ionic stoichiometric activator compounds may contain an active proton, or some other cation associated with, but not coordinated to, or only loosely
10 coordinated to, the remaining ion of the ionizing compound. Such compounds and the like are described in European publications EP-A-0 570 982, EP-A-0 520 732, EP-A-0 495 375, EP-B1-0 500 944, EP-A-0 277 003 and EP-A-0 277 004, and U.S. Patent Nos. 5,153,157, 5,198,401, 5,066,741, 5,206,197, 5,241,025, 5,384,299 and 5,502,124 and U.S. Patent Application Serial No. 08/285,380, filed
15 August 3, 1994, all of which are herein fully incorporated by reference.

Ionic catalysts can be prepared by reacting a transition metal compound with some neutral Lewis acids, such as $B(C_6F_5)_3$, which upon reaction with the hydrolyzable ligand (X) of the transition metal compound forms an anion, such as
20 $[B(C_6F_5)_3(X)]^-$, which stabilizes the cationic transition metal species generated by the reaction. The catalysts can be, and preferably are, prepared with activator components which are ionic compounds or compositions. However preparation of activators utilizing neutral compounds is also contemplated by this invention.

25 Compounds useful as an activator component in the preparation of the ionic catalyst systems used in the process of this invention comprise a cation, which is preferably a Bronsted acid capable of donating a proton, and a compatible non-coordinating anion which anion is relatively large (bulky), capable of stabilizing the active catalyst species (the Group 4 cation) which is formed
30 when the two compounds are combined and said anion will be sufficiently labile to be displaced by olefinic diolefinic and acetylenically unsaturated substrates or other neutral Lewis bases such as ethers, nitriles and the like. Two classes of compatible non-coordinating anions have been disclosed in EPA 277,003 and EPA 277,004 published 1988: 1) anionic coordination complexes comprising a

- 5 plurality of lipophilic radicals covalently coordinated to and shielding a central charge-bearing metal or metalloid core, and 2) anions comprising a plurality of boron atoms such as carboranes, metallocarboranes and boranes.

In a preferred embodiment, the stoichiometric activators include a cation
10 and an anion component, and may be represented by the following formula:



wherein L is a neutral Lewis base;

15 H is hydrogen;

(L-H)⁺ is a Bronsted acid

A^{d-} is a non-coordinating anion having the charge d-

d is an integer from 1 to 3.

20 The cation component, (L-H)_d⁺ may include Bronsted acids such as protons or protonated Lewis bases or reducible Lewis acids capable of protonating or abstracting a moiety, such as an alkyl or aryl, from the bulky ligand metallocene containing transition metal catalyst precursor, resulting in a cationic transition metal species.

25

The activating cation (L-H)_d⁺ may be a Bronsted acid, capable of donating a proton to the transition metal catalytic precursor resulting in a transition metal cation, including ammoniums, oxoniums, phosphoniums, silyliums, and mixtures thereof, preferably ammoniums of methylamine, aniline, dimethylamine,
30 diethylamine, N-methylaniline, diphenylamine, trimethylamine, triethylamine, N,N-dimethylaniline, methyldiphenylamine, pyridine, p-bromo N,N-dimethylaniline, p-nitro-N,N-dimethylaniline, phosphoniums from triethylphosphine, triphenylphosphine, and diphenylphosphine, oxoniums from ethers such as dimethyl ether diethyl ether, tetrahydrofuran and dioxane,
35 sulfoniums from thioethers, such as diethyl thioethers and tetrahydrothiophene,

5 and mixtures thereof. The activating cation $(L-H)_d^+$ may also be a moiety such as silver, tropylium, carbeniums, ferroceniums and mixtures, preferably carboniums and ferroceniums. Most preferably $(L-H)_d^+$ is triphenyl carbonium.

The anion component A^d include those having the formula $[M^{k+}Q_n]^{d-}$
 10 wherein k is an integer from 1 to 3; n is an integer from 2-6; $n - k = d$; M is an element selected from Group 13 of the Periodic Table of the Elements, preferably boron or aluminum, and Q is independently a hydride, bridged or unbridged dialkylamido, halide, alkoxide, aryloxy, hydrocarbyl, substituted hydrocarbyl, halocarbyl, substituted halocarbyl, and halosubstituted-hydrocarbyl radicals, said
 15 Q having up to 20 carbon atoms with the proviso that in not more than 1 occurrence is Q a halide. Preferably, each Q is a fluorinated hydrocarbyl group having 1 to 20 carbon atoms, more preferably each Q is a fluorinated aryl group, and most preferably each Q is a pentafluorophenyl group. Examples of suitable A^d also include diboron compounds as disclosed in U.S. Pat. No. 5,447,895, which is
 20 fully incorporated herein by reference.

Illustrative, but not limiting examples of boron compounds which may be used as an activating cocatalyst in the preparation of the improved catalysts of this invention are tri-substituted ammonium salts such as:

25 trimethylammonium tetraphenylborate, triethylammonium tetraphenylborate, tripropylammonium tetraphenylborate, tri(n-butyl)ammonium tetraphenylborate, tri(t-butyl)ammonium tetraphenylborate, N,N-dimethylanilinium tetraphenylborate, N,N-diethylanilinium tetraphenylborate, N,N-dimethyl-(2,4,6-trimethylanilinium) tetraphenylborate, tropylium tetraphenylborate,
 30 triphenylcarbenium tetraphenylborate, triphenylphosphonium tetraphenylborate, triethylsilylium tetraphenylborate, benzene(diazonium)tetraphenylborate, trimethylammonium tetrakis(pentafluorophenyl)borate, triethylammonium tetrakis(pentafluorophenyl)borate, tripropylammonium tetrakis(pentafluorophenyl)borate, tri(n-butyl)ammonium

- 5 tetrakis(pentafluorophenyl)borate, tri(sec-butyl)ammonium
tetrakis(pentafluorophenyl)borate, N,N-dimethylanilinium
tetrakis(pentafluorophenyl)borate, N,N-diethylanilinium
tetrakis(pentafluorophenyl)borate, N,N-dimethyl-(2,4,6-trimethylanilinium)
tetrakis(pentafluorophenyl)borate, tropillium tetrakis(pentafluorophenyl)borate,
- 10 triphenylcarbenium tetrakis(pentafluorophenyl)borate, triphenylphosphonium
tetrakis(pentafluorophenyl)borate, triethylsilylium
tetrakis(pentafluorophenyl)borate, benzene(diazonium)
tetrakis(pentafluorophenyl)borate, trimethylammonium tetrakis-(2,3,4,6-
tetrafluorophenyl) borate, triethylammonium tetrakis-(2,3,4,6-
- 15 tetrafluorophenyl)borate, tripropylammonium tetrakis-(2,3,4,6-
tetrafluorophenyl)borate, tri(n-butyl)ammonium tetrakis-(2,3,4,6-tetrafluoro-
phenyl)borate, dimethyl(t-butyl)ammonium tetrakis-(2,3,4,6-
tetrafluorophenyl)borate, N,N-dimethylanilinium tetrakis-(2,3,4,6-
tetrafluorophenyl)borate, N,N-diethylanilinium tetrakis-(2,3,4,6-
- 20 tetrafluorophenyl)borate, N,N-dimethyl-(2,4,6-trimethylanilinium) tetrakis-
(2,3,4,6-tetrafluorophenyl)borate, tropillium tetrakis-(2,3,4,6-
tetrafluorophenyl)borate, triphenylcarbenium tetrakis-(2,3,4,6-
tetrafluorophenyl)borate, triphenylphosphonium tetrakis-(2,3,4,6-
tetrafluorophenyl)borate, triethylsilylium tetrakis-(2,3,4,6-
- 25 tetrafluorophenyl)borate, benzene(diazonium) tetrakis-(2,3,4,6-
tetrafluorophenyl)borate, trimethylammonium tetrakis(perfluoronaphthyl)borate,
triethylammonium tetrakis(perfluoronaphthyl)borate, tripropylammonium
tetrakis(perfluoronaphthyl)borate, tri(n-butyl)ammonium
tetrakis(perfluoronaphthyl)borate, tri(t-butyl)ammonium
- 30 tetrakis(perfluoronaphthyl)borate, N,N-dimethylanilinium
tetrakis(perfluoronaphthyl)borate, N,N-diethylanilinium
tetrakis(perfluoronaphthyl)borate, N,N-dimethyl-(2,4,6-trimethylanilinium)
tetrakis(perfluoronaphthyl)borate, tropillium tetrakis(perfluoronaphthyl)borate,
triphenylcarbenium tetrakis(perfluoronaphthyl)borate, triphenylphosphonium

- 5 tetrakis(perfluoronaphthyl)borate, triethylsilylium tetrakis(perfluoronaphthyl)borate, benzene(diazonium) tetrakis(perfluoronaphthyl)borate, trimethylammonium tetrakis(perfluorobiphenyl)borate, triethylammonium tetrakis(perfluorobiphenyl)borate, tripropylammonium tetrakis(perfluorobiphenyl)borate, tri(n-butyl)ammonium
- 10 tetrakis(perfluorobiphenyl)borate, tri(t-butyl)ammonium tetrakis(perfluorobiphenyl)borate, N,N-dimethylanilinium tetrakis(perfluorobiphenyl)borate, N,N-diethylanilinium tetrakis(perfluorobiphenyl)borate, N,N-dimethyl-(2,4,6-trimethylanilinium) tetrakis(perfluorobiphenyl)borate, tropillium tetrakis(perfluorobiphenyl)borate,
- 15 triphenylcarbenium tetrakis(perfluorobiphenyl)borate, triphenylphosphonium tetrakis(perfluorobiphenyl)borate, triethylsilylium tetrakis(perfluorobiphenyl)borate, benzene(diazonium) tetrakis(perfluorobiphenyl)borate, trimethylammonium tetrakis(3,5-
- 20 bis(trifluoromethyl)phenyl)borate, triethylammonium tetrakis(3,5-bis(trifluoromethyl)phenyl)borate, tripropylammonium tetrakis(3,5-bis(trifluoromethyl)phenyl)borate, tri(n-butyl)ammonium tetrakis(3,5-bis(trifluoromethyl)phenyl)borate, tri(t-butyl)ammonium tetrakis(3,5-bis(trifluoromethyl)phenyl)borate, N,N-dimethylanilinium tetrakis(3,5-bis(trifluoromethyl)phenyl)borate, N,N-diethylanilinium tetrakis(3,5-
- 25 bis(trifluoromethyl)phenyl)borate, N,N-dimethyl-(2,4,6-trimethylanilinium) tetrakis(3,5-bis(trifluoromethyl)phenyl)borate, tropillium tetrakis(3,5-bis(trifluoromethyl)phenyl)borate, triphenylcarbenium tetrakis(3,5-bis(trifluoromethyl)phenyl)borate, triphenylphosphonium tetrakis(3,5-bis(trifluoromethyl)phenyl)borate, triethylsilylium tetrakis(3,5-
- 30 bis(trifluoromethyl)phenyl)borate, benzene(diazonium) tetrakis(3,5-bis(trifluoromethyl)phenyl)borate, and dialkyl ammonium salts such as: di-(i-propyl)ammonium tetrakis(pentafluorophenyl)borate, and dicyclohexylammonium tetrakis(pentafluorophenyl)borate; and additional tri-substituted phosphonium

- 5 salts such as tri(o-tolyl)phosphonium tetrakis(pentafluorophenyl)borate, and tri(2,6-dimethylphenyl)phosphonium tetrakis(pentafluorophenyl)borate.

Most preferably, the ionic stoichiometric activator $(L-H)_d^+ (A^d^-)$ is N,N-dimethylanilinium tetra(perfluorophenyl)borate, N,N-dimethylanilinium
 10 tetrakis(perfluoronaphthyl)borate, N,N-dimethylanilinium tetrakis(perfluorobiphenyl)borate, N,N-dimethylanilinium tetrakis(3,5-bis(trifluoromethyl)phenyl)borate, triphenylcarbenium
 tetrakis(perfluoronaphthyl)borate, triphenylcarbenium tetrakis(perfluorobiphenyl)borate, triphenylcarbenium tetrakis(3,5-
 15 bis(trifluoromethyl)phenyl)borate, or triphenylcarbenium tetra(perfluorophenyl)borate.

In one embodiment, an activation method using ionizing ionic compounds not containing an active proton but capable of producing a bulky ligand
 20 metallocene catalyst cation and their non-coordinating anion are also contemplated, and are described in EP-A- 0 426 637, EP-A- 0 573 403 and U.S. Patent No. 5,387,568, which are all herein incorporated by reference.

The term "non-coordinating anion" (NCA) means an anion which either
 25 does not coordinate to said cation or which is only weakly coordinated to said cation thereby remaining sufficiently labile to be displaced by a neutral Lewis base. "Compatible" non-coordinating anions are those which are not degraded to neutrality when the initially formed complex decomposes. Further, the anion will not transfer an anionic substituent or fragment to the cation so as to cause it to
 30 form a neutral four coordinate metallocene compound and a neutral by-product from the anion. Non-coordinating anions useful in accordance with this invention are those that are compatible, stabilize the metallocene cation in the sense of balancing its ionic charge at +1, yet retain sufficient lability to permit displacement by an ethylenically or acetylenically unsaturated monomer during

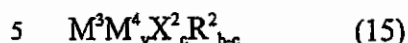
- 5 polymerization. These types of cocatalysts sometimes use tri-isobutyl aluminum or tri-octyl aluminum as a scavenger.

Invention process also can employ cocatalyst compounds or activator compounds that are initially neutral Lewis acids but form a cationic metal
10 complex and a noncoordinating anion, or a zwitterionic complex upon reaction with the invention compounds. For example, tris(pentafluorophenyl) boron or aluminum act to abstract a hydrocarbyl or hydride ligand to yield an invention cationic metal complex and stabilizing noncoordinating anion, see EP-A-0 427 697 and EP-A-0 520 732 for illustrations of analogous Group-4 metallocene
15 compounds. Also, see the methods and compounds of EP-A-0 495 375. For formation of zwitterionic complexes using analogous Group 4 compounds, see U.S. Patents 5,624,878; 5,486,632; and 5,527,929.

When the cations of noncoordinating anion precursors are Bronsted acids
20 such as protons or protonated Lewis bases (excluding water), or reducible Lewis acids such as ferrocenium or silver cations, or alkali or alkaline earth metal cations such as those of sodium, magnesium or lithium, the catalyst-precursor-to-activator molar ratio may be any ratio. Combinations of the described activator compounds may also be used for activation. For example, tris(perfluorophenyl) boron can be
25 used with methylalumoxane.

Conventional-Type Cocatalysts (Activators)

Typically, conventional transition metal catalyst compounds excluding
30 some conventional-type chromium catalyst compounds are activated with one or more of the conventional cocatalysts which may be represented by the formula:



wherein M^3 is a metal from Group 1 to 3 and 12 to 13 of the Periodic Table of Elements; M^4 is a metal of Group 1 of the Periodic Table of Elements; v is a number from 0 to 1; each X^2 is any halogen; c is a number from 0 to 3; each R^2 is
 10 a monovalent hydrocarbon radical or hydrogen; b is a number from 1 to 4; and wherein b minus c is at least 1. Other conventional-type organometallic cocatalyst compounds for the above conventional-type transition metal catalysts have the formula $M^3R^2_k$, where M^3 is a Group IA, IIA, IIB or IIIA metal, such as lithium, sodium, beryllium, barium, boron, aluminum, zinc, cadmium, and gallium; k
 15 equals 1, 2 or 3 depending upon the valency of M^3 which valency in turn normally depends upon the particular Group to which M^3 belongs; and each R^2 may be any monovalent hydrocarbon radical.

Non-limiting examples of conventional-type organometallic cocatalyst
 20 compounds useful with the conventional-type catalyst compounds described above include methyllithium, butyllithium, dihexylmercury, butylmagnesium, diethylcadmium, benzylpotassium, diethylzinc, tri-*n*-butylaluminum, diisobutyl ethylboron, diethylcadmium, di-*n*-butylzinc and tri-*n*-amylboron, and, in particular, the aluminum alkyls, such as tri-hexyl-aluminum, triethylaluminum,
 25 trimethylaluminum, and tri-isobutylaluminum. Other conventional-type cocatalyst compounds include mono-organohalides and hydrides of Group 2 metals, and mono- or di-organohalides and hydrides of Group 3 and 13 metals. Non-limiting examples of such conventional-type cocatalyst compounds include di-isobutylaluminum bromide, isobutylboron dichloride, methyl magnesium
 30 chloride, ethylberyllium chloride, ethylcalcium bromide, di-isobutylaluminum hydride, methylcadmium hydride, diethylboron hydride, hexylberyllium hydride, dipropylboron hydride, octylmagnesium hydride, butylzinc hydride, dichloroboron hydride, di-bromo-aluminum hydride and bromocadmium hydride. Conventional-type organometallic cocatalyst compounds are known to those in the art and a

- 5 more complete discussion of these compounds may be found in U.S. Patent Nos., 3,221,002 and 5,093,415, which are herein fully incorporated by reference.

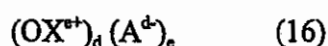
Additional Activators

- 10 Other activators include those described in PCT publication WO 98/07515 such as tris (2, 2', 2'' - nonafluorobiphenyl) fluoroaluminate, which publication is fully incorporated herein by reference. Combinations of activators are also contemplated by the invention, for example, alumoxanes and ionizing activators in combinations, see for example, EP-B1 0 573 120, PCT publications WO 94/07928
15 and WO 95/14044 and U.S. Patent Nos. 5,153,157 and 5,453,410 all of which are herein fully incorporated by reference.

- Other suitable activators are disclosed in WO 98/09996, incorporated herein by reference, which describes activating bulky ligand metallocene catalyst
20 compounds with perchlorates, periodates and iodates including their hydrates. WO 98/30602 and WO 98/30603, incorporated by reference, describe the use of lithium (2,2'-bisphenyl-ditrimethylsilicate)•4THF as an activator for a bulky ligand metallocene catalyst compound. WO 99/18135, incorporated herein by reference, describes the use of organo-boron-aluminum activators. EP-B1-0 781
25 299 describes using a silylium salt in combination with a non-coordinating compatible anion. Also, methods of activation such as using radiation (see EP-B1-0 615 981 herein incorporated by reference), electro-chemical oxidation, and the like are also contemplated as activating methods for the purposes of rendering the neutral bulky ligand metallocene catalyst compound or precursor to a bulky
30 ligand metallocene cation capable of polymerizing olefins. Other activators or methods for activating a bulky ligand metallocene catalyst compound are described in for example, U.S. Patent Nos. 5,849,852, 5,859,653 and 5,869,723 and WO 98/32775, WO 99/42467 (dioctadecylmethylammonium-

5 bis(tris(pentafluorophenyl)borane) benzimidazolidine), which are herein incorporated by reference.

Another suitable ion forming, activating cocatalyst comprises a salt of a cationic oxidizing agent and a noncoordinating, compatible anion represented by
10 the formula:



wherein OX^{e+} is a cationic oxidizing agent having a charge of $e+$; e is an integer
15 from 1 to 3; and A^{d-} , and d are as previously defined. Examples of cationic oxidizing agents include: ferrocenium, hydrocarbyl-substituted ferrocenium, Ag^+ , or Pb^{+2} . Preferred embodiments of A^{d-} are those anions previously defined with respect to the Bronsted acid containing activators, especially tetrakis(pentafluorophenyl)borate.

20

It is within the scope of this invention that catalyst compounds can be combined one or more activators or activation methods described above. For example, a combination of activators have been described in U.S. Patent Nos. 5,153,157 and 5,453,410, European publication EP-B1 0 573 120, and PCT
25 publications WO 94/07928 and WO 95/14044. These documents all discuss the use of an alumoxane and an ionizing activator with a bulky ligand metallocene catalyst compound.

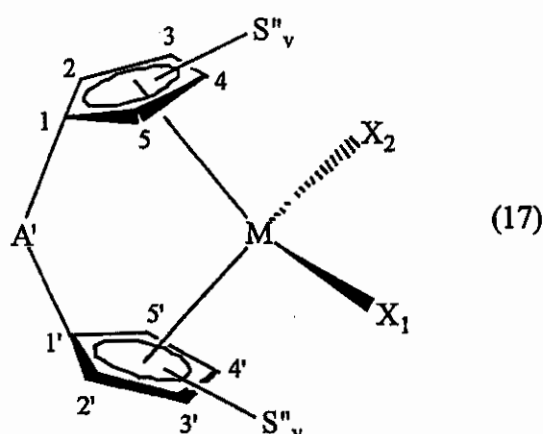
The Choice of Transition Metal Catalyst Components

30

The catalyst system of this invention comprises two or more transition metal compounds as described above. At least one of the compounds must be capable of producing a crystalline poly- α -olefin, preferably isotactic polypropylene or syndiotactic polypropylene, having a crystallinity of 40% or

- 5 more. The other compound must be capable of producing an amorphous poly- α -olefin, preferably atactic polypropylene, having a crystallinity of 20% or less.

The choice of transition metal component for the crystalline polymer fraction is a subset of the transition metal component of equations 8-9. This
 10 preferred component is illustrated in equation 17:



wherein A', M, X₁ and X₂ are as previously defined. Substituents S''_v are
 independently defined as S'' in equations 8-9 where the subscript "v" denotes the
 15 carbon atom on the Cp-ring to which the substituent is bonded.

Preferably metallocene precursors for producing poly- α -olefins having
 enhanced isotactic character are those of Equation 17 where S''_v are independently
 chosen such that the metallocene framework 1) has no plane of symmetry
 20 containing the metal center, and 2) has a C₂-axis of symmetry through the metal
 center. Such complexes, such as rac-Me₂Si(indenyl)₂ZrMe₂ and rac-
 Me₂Si(indenyl)₂HfMe₂ are well known in the art and generally produce isotactic
 polymers with higher degrees of stereoregularity than the less symmetric chiral
 systems. Likewise another preferred class of transition metal compounds that can
 25 produce isotactic polymers useful in this invention are those
 monocyclopentadienyl catalysts disclosed in U.S. Pat. No. 5,026,798, which is
 incorporated by reference herein.

5

Preferred chiral racemic metallocene compounds which, according to the present invention, provide catalyst systems which are specific to the production of isotactic poly- α -olefins include the racemic versions of:

- dimethylsilylbis(indenyl) zirconium dichloride, dimethylsilylbis(indenyl)
10 zirconium dimethyl, diphenylsilylbis(indenyl) zirconium dichloride,
diphenylsilylbis(indenyl) zirconium dimethyl, methylphenylsilylbis(indenyl)
zirconium dichloride,
methylphenylsilylbis(indenyl) zirconium dimethyl, ethylenebis(indenyl)
zirconium dichloride, ethylenebis(indenyl) zirconium dimethyl,
15 methylenebis(indenyl) zirconium dichloride, methylenebis(indenyl) zirconium
dimethyl, dimethylsilylbis(indenyl) hafnium dichloride, dimethylsilylbis(indenyl)
hafnium dimethyl, diphenylsilylbis(indenyl) hafnium dichloride,
diphenylsilylbis(indenyl) hafnium dimethyl, methylphenylsilylbis(indenyl)
hafnium dichloride, methylphenylsilylbis(indenyl) hafnium dimethyl,
20 ethylenebis(indenyl) hafnium dichloride, ethylenebis(indenyl) hafnium dimethyl,
methylenebis(indenyl) hafnium dichloride, methylenebis(indenyl) hafnium
dimethyl, dimethylsilylbis(tetrahydroindenyl) zirconium dichloride,
dimethylsilylbis(tetrahydroindenyl) zirconium dimethyl,
diphenylsilylbis(tetrahydroindenyl) zirconium dichloride,
25 diphenylsilylbis(tetrahydroindenyl) zirconium dimethyl,
methylphenylsilylbis(tetrahydroindenyl) zirconium dichloride,
methylphenylsilylbis(tetrahydroindenyl) zirconium dimethyl,
ethylenebis(tetrahydroindenyl) zirconium dichloride,
ethylenebis(tetrahydroindenyl) zirconium dimethyl,
30 methylenebis(tetrahydroindenyl) zirconium dichloride,
methylenebis(tetrahydroindenyl) zirconium dimethyl,
dimethylsilylbis(tetrahydroindenyl) hafnium dichloride,
dimethylsilylbis(tetrahydroindenyl) hafnium dimethyl,
diphenylsilylbis(tetrahydroindenyl) hafnium dichloride,

- 5 diphenylsilylbis(tetrahydroindenyl) hafnium dimethyl,
methylphenylsilylbis(tetrahydroindenyl) hafnium dichloride,
methylphenylsilylbis(tetrahydroindenyl) hafnium dimethyl,
ethylenebis(tetrahydroindenyl) hafnium dichloride, ethylenebis(tetrahydroindenyl)
hafnium dimethyl, methylenebis(tetrahydroindenyl) hafnium dichloride,
10 methylenebis(tetrahydroindenyl) hafnium dimethyl, dimethylsilylbis(2-
methylindenyl) zirconium dichloride, dimethylsilylbis(2-methylindenyl)
zirconium dimethyl, diphenylsilylbis(2-methylindenyl) zirconium dichloride,
diphenylsilylbis(2-methylindenyl) zirconium dimethyl, methylphenylsilylbis(2-
methylindenyl) zirconium dichloride, methylphenylsilylbis(2-methylindenyl)
15 zirconium dimethyl, ethylenebis(2-methylindenyl) zirconium dichloride,
ethylenebis(2-methylindenyl) zirconium dimethyl, methylenebis(2-methylindenyl)
zirconium dichloride, methylenebis(2-methylindenyl) zirconium dimethyl,
dimethylsilylbis(2-methylindenyl) hafnium dichloride, dimethylsilylbis(2-
methylindenyl) hafnium dimethyl, diphenylsilylbis(2-methylindenyl) hafnium
20 dichloride, diphenylsilylbis(2-methylindenyl) hafnium dimethyl,
methylphenylsilylbis(2-methylindenyl) hafnium dichloride,
methylphenylsilylbis(2-methylindenyl) hafnium dimethyl, ethylenebis(2-
methylindenyl) hafnium dichloride, ethylenebis(2-methylindenyl) hafnium
dimethyl, methylenebis(2-methylindenyl) hafnium dichloride, methylenebis(2-
25 methylindenyl) hafnium dimethyl, rac-dimethylsilylbis(2-methyl-4-
phenylindenyl) zirconium dichloride, diphenylsilylbis(2-methyl-4-phenylindenyl)
zirconium dimethyl, diphenylsilylbis(2-methyl-4-phenylindenyl) zirconium
dichloride, diphenylsilylbis(2-methyl-4-phenylindenyl) zirconium dimethyl,
methylphenylsilylbis(2-methyl-4-phenylindenyl) zirconium dichloride,
30 methylphenylsilylbis(2-methyl-4-phenylindenyl) zirconium dimethyl,
ethylenebis(2-methyl-4-phenylindenyl) zirconium dichloride, ethylenebis(2-
methyl-4-phenylindenyl) zirconium dimethyl, methylenebis(2-methyl-4-
phenylindenyl) zirconium dichloride, methylenebis(2-methyl-4-phenylindenyl)
zirconium dimethyl, dimethylsilylbis(2-methyl-4-phenylindenyl) hafnium

- 5 dichloride, dimethylsilylbis(2-methyl-4-phenylindenyl) hafnium dimethyl,
diphenylsilylbis(2-methyl-4-phenylindenyl) hafnium dichloride,
diphenylsilylbis(2-methyl-4-phenylindenyl) hafnium dimethyl,
methylphenylsilylbis(2-methyl-4-phenylindenyl) hafnium dichloride,
methylphenylsilylbis(2-methyl-4-phenylindenyl) hafnium dimethyl,
- 10 ethylenebis(2-methyl-4-phenylindenyl) hafnium dichloride, ethylenebis(2-methyl-
4-phenylindenyl) hafnium dimethyl, methylenebis(2-methyl-4-phenylindenyl)
hafnium dichloride, methylenebis(2-methyl-4-phenylindenyl) hafnium dimethyl,
dimethylsilylbis(4,7-dimethylindenyl) zirconium dichloride, dimethylsilylbis(4,7-
dimethylindenyl) zirconium dimethyl, diphenylsilylbis(4,7-dimethylindenyl)
- 15 zirconium dichloride, diphenylsilylbis(4,7-dimethylindenyl) zirconium dimethyl,
methylphenylsilylbis(4,7-dimethylindenyl) zirconium dichloride,
methylphenylsilylbis(4,7-dimethylindenyl) zirconium dimethyl, ethylenebis(4,7-
dimethylindenyl) zirconium dichloride, ethylenebis(4,7-dimethylindenyl)
zirconium dimethyl, methylenebis(4,7-dimethylindenyl) zirconium dichloride,
- 20 methylenebis(4,7-dimethylindenyl) zirconium dimethyl, dimethylsilylbis(4,7-
dimethylindenyl) hafnium dichloride, dimethylsilylbis(4,7-dimethylindenyl)
hafnium dimethyl, diphenylsilylbis(4,7-dimethylindenyl) hafnium dichloride,
diphenylsilylbis(4,7-dimethylindenyl) hafnium dimethyl,
methylphenylsilylbis(4,7-dimethylindenyl) hafnium dichloride,
- 25 methylphenylsilylbis(4,7-dimethylindenyl) hafnium dimethyl, ethylenebis(4,7-
dimethylindenyl) hafnium dichloride, ethylenebis(4,7-dimethylindenyl) hafnium
dimethyl, methylenebis(4,7-dimethylindenyl) hafnium dichloride,
methylenebis(4,7-dimethylindenyl) hafnium dimethyl, dimethylsilylbis(2-methyl-
4-naphthylindenyl) zirconium dichloride, dimethylsilylbis(2-methyl-4-
- 30 naphthylindenyl) zirconium dimethyl, diphenylsilylbis(2-methyl-4-naphthylindenyl)
zirconium dichloride, diphenylsilylbis(2-methyl-4-naphthylindenyl) zirconium
dimethyl, methylphenylsilylbis(2-methyl-4-naphthylindenyl) zirconium dichloride,
methylphenylsilylbis(2-methyl-4-naphthylindenyl) zirconium dimethyl,
ethylenebis(2-methyl-4-naphthylindenyl) zirconium dichloride, ethylenebis(2-

- 5 methyl-4-naphthylindenyl) zirconium dimethyl, methylenebis(2-methyl-4-naphthylindenyl) zirconium dichloride, methylenebis(2-methyl-4-naphthylindenyl) zirconium dimethyl, dimethylsilylbis(2-methyl-4-naphthylindenyl) hafnium dichloride, dimethylsilylbis(2-methyl-4-naphthylindenyl) hafnium dimethyl, diphenylsilylbis(2-methyl-4-naphthylindenyl) hafnium dichloride,
- 10 diphenylsilylbis(2-methyl-4-naphthylindenyl) hafnium dimethyl, methylphenylsilylbis(2-methyl-4-naphthylindenyl) hafnium dichloride, methylphenylsilylbis(2-methyl-4-naphthylindenyl) hafnium dimethyl, ethylenebis(2-methyl-4-naphthylindenyl) hafnium dichloride, ethylenebis(2-methyl-4-naphthylindenyl) hafnium dimethyl, methylenebis(2-methyl-4-
- 15 naphthylindenyl) hafnium dichloride, methylenebis(2-methyl-4-naphthylindenyl) hafnium dimethyl, dimethylsilylbis(2,3-dimethylcyclopentadienyl) zirconium dichloride, dimethylsilylbis(2,3-dimethylcyclopentadienyl) zirconium dimethyl, diphenylsilylbis(2,3-dimethylcyclopentadienyl) zirconium dichloride, diphenylsilylbis(2,3-dimethylcyclopentadienyl) zirconium dimethyl,
- 20 methylphenylsilylbis(2,3-dimethylcyclopentadienyl) zirconium dichloride, methylphenylsilylbis(2,3-dimethylcyclopentadienyl) zirconium dimethyl, ethylenebis(2,3-dimethylcyclopentadienyl) zirconium dichloride, ethylenebis(2,3-dimethylcyclopentadienyl) zirconium dimethyl, methylenebis(2,3-dimethylcyclopentadienyl) zirconium dichloride, methylenebis(2,3-
- 25 dimethylcyclopentadienyl) zirconium dimethyl, dimethylsilylbis(2,3-dimethylcyclopentadienyl) hafnium dichloride, dimethylsilylbis(2,3-dimethylcyclopentadienyl) hafnium dimethyl, diphenylsilylbis(2,3-dimethylcyclopentadienyl) hafnium dichloride, diphenylsilylbis(2,3-dimethylcyclopentadienyl) hafnium dimethyl, methylphenylsilylbis(2,3-
- 30 dimethylcyclopentadienyl) hafnium dichloride, methylphenylsilylbis(2,3-dimethylcyclopentadienyl) hafnium dimethyl, ethylenebis(2,3-dimethylcyclopentadienyl) hafnium dichloride, ethylenebis(2,3-dimethylcyclopentadienyl) hafnium dimethyl, methylenebis(2,3-dimethylcyclopentadienyl) hafnium dichloride, methylenebis(2,3-

- 5 dimethylcyclopentadienyl) hafnium dimethyl, dimethylsilylbis(3-trimethylsilylcyclopentadienyl) zirconium dichloride, dimethylsilylbis(3-trimethylsilylcyclopentadienyl) zirconium dimethyl, diphenylsilylbis(3-trimethylsilylcyclopentadienyl) zirconium dichloride, diphenylsilylbis(3-trimethylsilylcyclopentadienyl) zirconium dimethyl, methylphenylsilylbis(3-trimethylsilylcyclopentadienyl) zirconium dichloride, methylphenylsilylbis(3-trimethylsilylcyclopentadienyl) zirconium dimethyl, ethylenebis(3-trimethylsilylcyclopentadienyl) zirconium dichloride, ethylenebis(3-trimethylsilylcyclopentadienyl) zirconium dimethyl, methylenebis(3-trimethylsilylcyclopentadienyl) zirconium dichloride, methylenebis(3-trimethylsilylcyclopentadienyl) zirconium dimethyl, dimethylsilylbis(3-trimethylsilylcyclopentadienyl) hafnium dichloride, dimethylsilylbis(3-trimethylsilylcyclopentadienyl) hafnium dimethyl, diphenylsilylbis(3-trimethylsilylcyclopentadienyl) hafnium dichloride, diphenylsilylbis(3-trimethylsilylcyclopentadienyl) hafnium dimethyl, methylphenylsilylbis(3-trimethylsilylcyclopentadienyl) hafnium dichloride, methylphenylsilylbis(3-trimethylsilylcyclopentadienyl) hafnium dimethyl, ethylenebis(3-trimethylsilylcyclopentadienyl) hafnium dichloride, ethylenebis(3-trimethylsilylcyclopentadienyl) hafnium dimethyl, methylenebis(3-trimethylsilylcyclopentadienyl) hafnium dichloride, methylenebis(3-trimethylsilylcyclopentadienyl) hafnium dimethyl,
- 10
- 15
- 20
- 25
- 30
- dimethylsiladiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;

- 5 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
- 10 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
dimethylsiladiyl(2-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
- 15 9-silafluorendiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
9-silafluorendiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
- 20 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
- 25 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
9-silafluorendiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
- 30 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;

- 5 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium
dichloride;
dimethylsiladiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
dimethylsiladiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
- 10 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
- 15 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
dimethylsiladiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
- 20 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl
dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
- 25 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
- 30 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;

- 5 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
- 10 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
- 15 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
dimethylsiladiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
- 20 dichloride;
dimethylsiladiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
- 25 dimethylsiladiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
- 30 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;

- 5 dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
- dimethylsiladiyl(2-n-propyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂hafnium dichloride;
- dimethylsiladiyl(2-iso-propyl, 4-[3',5'-bis-
- 10 trifluoromethylphenyl]indenyl)₂hafnium dichloride;
- dimethylsiladiyl(2-n-butyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂hafnium dichloride;
- dimethylsiladiyl(2-iso-butyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂hafnium dichloride;
- 15 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂hafnium dichloride;
- dimethylsiladiyl(2-tert-butyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂hafnium dichloride;
- 9-silafluorendiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
- 20 dichloride;
- dimethylsiladiyl(2-ethyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂hafnium dichloride;
- 9-silafluorendiyl(2-n-propyl, 4-[3',5'-bis-
- trifluoromethylphenyl]indenyl)₂zirconium dichloride;
- 25 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-bis-
- trifluoromethylphenyl]indenyl)₂zirconium dichloride;
- 9-silafluorendiyl(2-n-butyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂zirconium dichloride;
- 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-bis-
- 30 trifluoromethylphenyl]indenyl)₂zirconium dichloride;
- 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-bis-
- trifluoromethylphenyl]indenyl)₂zirconium dichloride;
- 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-bis-
- trifluoromethylphenyl]indenyl)₂zirconium dichloride;

- 5 9-silafluorendiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-n-propyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂
- 10 hafnium dichloride;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂ hafnium dichloride;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
- 15 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium
- 20 dichloride;
dimethylsiladiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
- 25 dimethylsiladiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
dimethylsiladiyl(2-iso-propyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
dimethylsiladiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
- 30 dimethyl;
dimethylsiladiyl(2-iso-butyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
dimethylsiladiyl(2-sec-butyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂zirconium dimethyl;

- 5 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
dimethylsiladiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium
dimethyl;
dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium
10 dimethyl;
dimethylsiladiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium
dimethyl;
dimethylsiladiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
15 dimethylsiladiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium
dimethyl;
dimethylsiladiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium
dimethyl;
dimethylsiladiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium
20 dimethyl;
dimethylsiladiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium
dimethyl;
9-silafluorendiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
dimethyl;
25 dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium
dimethyl;
9-silafluorendiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
30 9-silafluorendiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
dimethyl;
9-silafluorendiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;

- 5 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium
10 dimethyl;
9-silafluorendiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
15 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium
20 dimethyl;
9-silafluorendiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
25 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride
dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
30 dichloride;
dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;

- 5 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dichloride;
dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dichloride;
dimethylsiladiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium
10 dichloride;
dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium
dichloride;
dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di- iso-propylphenyl]indenyl)₂hafnium
dichloride;
15 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di- iso-propylphenyl]indenyl)₂hafnium
dichloride;
dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di- iso-propylphenyl]indenyl)₂hafnium
dichloride;
dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di- iso-propylphenyl]indenyl)₂hafnium
20 dichloride;
dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di- iso-propylphenyl]indenyl)₂hafnium
dichloride;
9-silafluorendiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dichloride;
25 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dichloride;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dichloride;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
30 dichloride;
9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dichloride;
9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dichloride;

- 5 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dichloride;
9-silafluorendiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium
dichloride;
9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium
10 dichloride;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium
dichloride;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium
dichloride;
15 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium
dichloride;
9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium
dichloride;
9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium
20 dichloride;
dimethylsiladiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl
25 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
dimethylsiladiyl(2-isobutyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
30 dimethyl;
dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;

- 5 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
dimethylsiladiyl(2-n-propyl, 4-[3',5'-di- iso-propylphenyl]indenyl)₂hafnium
dimethyl;
dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di- iso-propylphenyl]indenyl)₂hafnium
dimethyl;
- 10 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di- iso-propylphenyl]indenyl)₂hafnium
dimethyl;
dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
- 15 dimethyl;
dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
9-silafluorendiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
- 20 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
- 25 dimethyl;
9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
- 30 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
dimethyl;
9-silafluorendiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium
dimethyl;

- 5 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
- 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
- 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium
- 10 dimethyl;
- 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
- 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
- 15 dimethylsiladiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
- dimethylsiladiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
- dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
- 20 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
- dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
- dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
- 25 dichloride;
- dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
- dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
- 30 dimethylsiladiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
- dimethylsiladiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
- dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
- dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;

- 5 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium
dichloride;
dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium
dichloride;
- 10 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium
dichloride;
9-silafluorendiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
9-silafluorendiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
- 15 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
- 20 dichloride;
9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
- 25 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
9-silafluorendiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium
- 30 dichloride;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium
dichloride;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;

- 5 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium
dichloride;
9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium
dichloride;
9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium
10 dichloride;
dimethylsiladiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
dimethylsiladiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dimethyl;
15 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dimethyl;
dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dimethyl;
20 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dimethyl;
dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dimethyl;
dimethylsiladiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
25 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium
dimethyl;
dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
30 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;

- 5 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
- 10 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
- 15 dimethyl;
9-silafluorendiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium
- 20 dimethyl;
9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium
- 25 dimethyl;
dimethylamidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
- 30 dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
- 35 dichloride;

- 5 dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-ethyl, 4-[3',5'-bis-
- 10 trifluoromethylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
- 15 dimethylamidoborane(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-sec-butyl, 4-[3',5'-bis-
- 20 trifluoromethylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
- 25 dimethylamidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride
dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
- 30 dichloride;
dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
- 35 dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;

- 5 dimethylamidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
dimethylamidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
dimethylamidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
10 dichloride;
dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
dimethylamidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
15 dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
20 dichloride;
dimethylamidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ zirconium
dimethyl;
dimethylamidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
25 dimethylamidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
dimethylamidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
30 dimethyl;
dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
35 dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;

- 5 dimethylamidoborane(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
10 dimethylamidoborane(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
15 dimethylamidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
20 dimethyl;
dimethylamidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
25 dimethylamidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
30 dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
35 dimethylamidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;

- 5 dimethylamidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
- 10 dimethyl;
dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
- 15 dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
- 20 dichloride;
diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
- 25 diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
- 30 dichloride;
diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
- 35 diisopropylamidoborane(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;

- 5 diisopropylamidoborane(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
10 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
15 diisopropylamidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride
diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
20 diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
25 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
30 dichloride;
diisopropylamidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
35 diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

- 5 diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
10 dichloride;
diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
dichloride;
diisopropylamidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
15 diisopropylamidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
20 dimethyl;
diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
25 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
dimethyl;
diisopropylamidoborane(2-ethyl, 4-[3',5'-bis-
30 trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-n-propyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-iso-propyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
35 diisopropylamidoborane(2-n-butyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dimethyl;

- 5 diisopropylamidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
10 diisopropylamidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl
15 diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
20 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
25 diisopropylamidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium
30 dimethyl;
diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
35 diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;

- 5 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-methyl, 4-[3',5'-di-
10 tbutylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dichloride;
15 bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-
20 tbutylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dichloride;
25 bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-bis-
30 trifluoromethylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dichloride;
35 bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dichloride;

- 5 bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-iso-
10 propylphenyl]indenyl)₂zirconium dichloride
bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
15 bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-iso-
20 propylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
25 bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-
30 phenylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
35 bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

- 5 bis(trimethylsilyl)amidoborane(2-methyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-
10 tbutylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dimethyl;
15 bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-
tbutylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-
20 tbutylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
25 bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-bis-
30 trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-bis-
trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
35 bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-iso-
propylphenyl]indenyl)₂zirconium dimethyl;

- 5 bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl
 bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 10 bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 15 bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 bis(trimethylsilyl)amidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 20 bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 25 bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl; and
 30 bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl; and the like.

The most preferred species are the racemic versions of:

- 35 dimethylsilylbis(indenyl) zirconium dichloride, dimethylsilylbis(indenyl) zirconium dimethyl, ethylenebis(indenyl) zirconium dichloride,

- 5 ethylenebis(indenyl) zirconium dimethyl, dimethylsilylbis(tetrahydorindenyl) zirconium dichloride, dimethylsilylbis(tetrahydorindenyl) zirconium dimethyl, ethylenebis(tetrahydorindenyl) zirconium dichloride, ethylenebis(tetrahydorindenyl) zirconium dimethyl, dimethylsilylbis(2-methylindenyl) zirconium dichloride, dimethylsilylbis(2-methylindenyl) zirconium dimethyl, ethylenebis(2-methylindenyl) zirconium dichloride, ethylenebis(2-methylindenyl) zirconium dimethyl, dimethylsilylbis(2-methyl-4-phenylindenyl) zirconium dichloride, dimethylsilylbis(2-methyl-4-phenylindenyl) zirconium dimethyl, ethylenebis(2-methyl-4-phenylindenyl) zirconium dichloride, ethylenebis(2-methyl-4-phenylindenyl) zirconium dimethyl, dimethylsilylbis(4,7-dimethylindenyl) zirconium dichloride, dimethylsilylbis(4,7-dimethylindenyl) zirconium dimethyl, ethylenebis(4,7-dimethylindenyl) zirconium dichloride, ethylenebis(4,7-dimethylindenyl) zirconium dimethyl, dimethylsilylbis(indenyl) hafnium dichloride, dimethylsilylbis(indenyl) hafnium dimethyl, ethylenebis(indenyl) hafnium dichloride, ethylenebis(indenyl) hafnium dimethyl, dimethylsilylbis(tetrahydorindenyl) hafnium dichloride, dimethylsilylbis(tetrahydorindenyl) hafnium dimethyl, ethylenebis(tetrahydorindenyl) hafnium dichloride, ethylenebis(tetrahydorindenyl) hafnium dimethyl, dimethylsilylbis(2-methylindenyl) hafnium dichloride, dimethylsilylbis(2-methylindenyl) hafnium dimethyl, ethylenebis(2-methylindenyl) hafnium dichloride, ethylenebis(2-methylindenyl) hafnium dimethyl, dimethylsilylbis(2-methyl-4-phenylindenyl) hafnium dichloride, dimethylsilylbis(2-methyl-4-phenylindenyl) hafnium dimethyl, ethylenebis(2-methyl-4-phenylindenyl) hafnium dichloride, ethylenebis(2-methyl-4-phenylindenyl) hafnium dimethyl, dimethylsilylbis(4,7-dimethylindenyl) hafnium dichloride, dimethylsilylbis(4,7-dimethylindenyl) hafnium dimethyl, ethylenebis(4,7-dimethylindenyl) hafnium dichloride, and ethylenebis(4,7-dimethylindenyl) hafnium dimethyl.

Similarly, metallocene precursors providing tacticity control exist where (A-Cp) is (Cp) (Cp*), both Cp and Cp* having substituents on the cyclopentadienyl rings of sufficient steric bulk to restrict rotation of the

- 5 cyclopentadienyl ligands such that the aforementioned symmetry conditions are satisfied. Preferable chiral racemic metallocenes of this type include bis(tricyclo[5.2.1.0^{2,6}]deca-2,5-dienyl) zirconium and -hafnium dimethyl, bis((1R)-9,9-dimethyltricyclo[6.1.1.0^{2,6}]deca-2,5-dienyl) zirconium dimethyl, bis(tricyclo[5.2.1.0^{2,6}]deca-2,5,8-trienyl) zirconium dimethyl,
- 10 bis(tricyclo[5.2.2.0^{2,6}]undeca-2,5,8-trienyl)zirconium and - hafnium dimethyl and bis((1R,8R)-7,7,9,9-tetramethyl[6.1.1.0^{2,6}]deca-2,5-dienyl) zirconium and - hafnium dimethyl.

- Preferably metallocene precursors for the production of poly- α -olefins
- 15 having enhanced syndiotactic character are also those of Equation 17 where S" are independently chosen such that the two Cp-ligands have substantially different steric bulk. In order to produce a syndiotactic polymer the pattern of the groups substituted on the Cp-rings is important. Thus, by steric difference or sterically different as used herein, it is intended to imply a difference between the steric
- 20 characteristics of the Cp and Cp* rings that renders each to be symmetrical With respect to the A bridging group but different with respect to each other that controls the approach of each successive monomer unit that is added to the polymer chain. The steric difference between the Cp and Cp* rings act to block the approaching monomer from a random approach such that the monomer is
- 25 added to the polymer chain in the syndiotactic configuration.

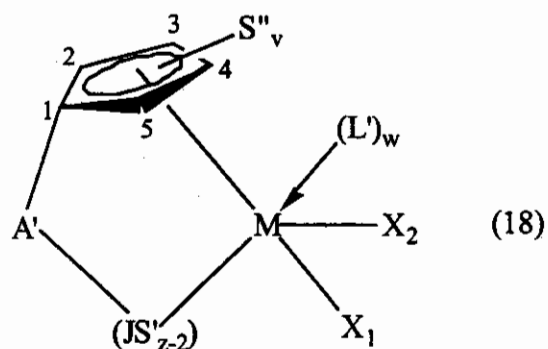
- Preferable metallocene precursors for the production of syndiotactic polymers are those of Equation 17 where S" are independently chosen such that 1) the steric difference between the two Cp-ligands is maximized and 2) there
- 30 remains a plane of symmetry through the metal center and the C₁ and C_{1'} carbon atoms of the Cp-rings in Equation 17. Thus, complexes such as Me₂C(η^5 -C₅H₄)(1-fluorenyl)MMe₂ (where M = Ti, Zr, or Hf) which possess this symmetry are preferred, and generally produce the syndiotactic polymer with higher degrees of stereoregularity than similar, but less symmetric, systems. Additionally, in the
- 35 above equation, 1-fluorenyl may be substituted with 3,8-di-t-butylfluorenyl, octahydrofluorenyl or 3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,10,10,11,11-

5 octahydrodibenzo[b,h]fluorene. Because pre-catalysts of this type often lose there ability to control the stereoregularity of the polymer under high temperature reaction conditions, to insure higher crystallinity in the material requires using these catalysts at lower reactor temperatures, preferably at temperatures below 80°C.

10

Preferred catalysts that can produce the lower molecular weight isotactic polypropylene are those described in U.S. Pat. No. 5,120,867, which is incorporated by reference herein. Any mixture of catalysts, including supported catalysts, which can be used together in a single reactor or in a series reactor
15 configuration, that can also produce the desired polypropylene can be utilized in this invention to produce the *in situ* blend. Preferred catalysts include cyclopentadienyl transition metals compounds and derivatives thereof used in conjunction with an alumoxane and/or a compatible non-coordinating anion. Additional preferred catalysts that produce crystalline polypropylene are discussed
20 in Chem. rev. 2000, 100, 1253-1345, which is incorporated by reference herein.

The preferred choice of transition metal component for the amorphous polymer fraction is the mono-cyclopentadienyl transition metal component of equation 10 where y is equal to 1. This preferred component is illustrated in
25 equation 18:



where A', J, S', X₁, X₂, L', z and w as are previously defined and M is titanium. Substituent S''_v is defined to be the same as S'' in equation 10 where the subscript

5 "v" denotes the carbon atom on the cyclopentadienyl ring to which the substituent is bonded and where there can be zero, two or four substituents, S'', on the cyclopentadienyl ring provided that the cyclopentadienyl ring is symmetrically substituted. Symmetrically substituted is defined to mean that the cyclopentadienyl ring is substituted in the 2 and 5 positions and/or 3 and 4
10 positions with S'' groups that are of approximately of the same steric bulk. Typically the size of these S'' groups are within 2 carbons of each other. Thus a cyclopentadienyl substituted at the 2 and the 5 positions with methyl and ethyl respectively, or substituted at the 3 and the 4 positions with hexyl and octyl, respectively, would be considered symmetric. Likewise, the cyclopentadienyl ring
15 may be substituted at all four sites with S'' groups and be considered symmetric as long as each of the symmetrical pairs are of similar steric bulk. Additionally, two adjacent S''-groups in the 3 and 4 position may be linked to form a ring provided that the new ring is also symmetrically substituted.

20 Catalyst systems of this type are known to impart 2,1-mistakes when incorporating C3 and higher α -olefins. The pre-catalysts where S' is bonded to the nitrogen ligand (J) via a 3° carbon (for example when S' is tert-butyl or 1-adamantyl) have fewer 2,1-mistakes than when S' is bonded to the nitrogen ligand (J) via a 1° carbon (for example when S' is n-butyl, methyl, or benzyl) or 2°
25 carbon (for example when S' is cyclododecyl, cyclohexyl, or sec-butyl). The 2,1-mistakes in the polymer backbone impart (CH₂)₂ units that can be beneficial to the polymer properties. Polymers of this type, the characterization of such polymers and the catalyst systems used to produce such polymers are described in US5,723,560 and is incorporated herein by reference. Lower Mw versions of
30 such polymers can be produced by changing process condition, for example, by increasing reactor temperature.

Preferred mono-cyclopentadienyl transition metal compounds which, according to the present invention, provide catalyst systems which are specific to
35 the production of atactic poly- α -olefins include:

- 5 dimethylsilyl(tetramethylcyclopentadienyl)(cyclododecylamido) titanium dichloride,
dimethylsilyl(tetramethylcyclopentadienyl)(cyclohexylamido) titanium dichloride,
dimethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dichloride,
dimethylsilyl(tetramethylcyclopentadienyl)(t-butylamido) titanium dichloride,
dimethylsilyl(tetramethylcyclopentadienyl)(s-butylamido) titanium dichloride,
- 10 dimethylsilyl(tetramethylcyclopentadienyl)(n-butylamido) titanium dichloride,
dimethylsilyl(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dichloride,
diethylsilyl(tetramethylcyclopentadienyl)(cyclododecylamido) titanium dichloride,
diethylsilyl(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dichloride,
diethylsilyl(tetramethylcyclopentadienyl)(cyclohexylamido) titanium dichloride,
- 15 diethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dichloride,
diethylsilyl(tetramethylcyclopentadienyl)(t-butylamido) titanium dichloride,
methylene(tetramethylcyclopentadienyl)(cyclododecylamido) titanium dichloride,
methylene(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dichloride,
methylene(tetramethylcyclopentadienyl)(cyclohexylamido) titanium dichloride,
- 20 methylene(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dichloride,
methylene(tetramethylcyclopentadienyl)(t-butylamido) titanium dichloride,
ethylene(tetramethylcyclopentadienyl)(cyclododecylamido) titanium dichloride,
ethylene(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dichloride,
ethylene(tetramethylcyclopentadienyl)(cyclohexylamido) titanium dichloride,
- 25 ethylene(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dichloride,
ethylene(tetramethylcyclopentadienyl)(t-butylamido) titanium dichloride,
dimethylsilyl(tetramethylcyclopentadienyl)(cyclododecylamido) titanium dimethyl,
dimethylsilyl(tetramethylcyclopentadienyl)(cyclohexylamido) titanium dimethyl,
dimethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,
- 30 dimethylsilyl(tetramethylcyclopentadienyl)(t-butylamido) titanium dimethyl,
dimethylsilyl(tetramethylcyclopentadienyl)(s-butylamido) titanium dimethyl,
dimethylsilyl(tetramethylcyclopentadienyl)(n-butylamido) titanium dimethyl,
dimethylsilyl(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
diethylsilyl(tetramethylcyclopentadienyl)(cyclododecylamido) titanium dimethyl,
- 35 diethylsilyl(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
diethylsilyl(tetramethylcyclopentadienyl)(cyclohexylamido) titanium dimethyl,
diethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,
diethylsilyl(tetramethylcyclopentadienyl)(t-butylamido) titanium dimethyl,
methylene(tetramethylcyclopentadienyl)(cyclododecylamido) titanium dimethyl,
- 40 methylene(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
methylene(tetramethylcyclopentadienyl)(cyclohexylamido) titanium dimethyl,
methylene(tetramethylcyclopentadienyl)(1-adamantylamido)

- 5 titanium dimethyl, methylene(tetramethylcyclopentadienyl)(t-butylamido)
 titanium dimethyl, ethylene(tetramethylcyclopentadienyl)(cyclododecylamido)
 titanium dimethyl, ethylene(tetramethylcyclopentadienyl)(exo-2-norbornylamido)
 titanium dimethyl, ethylene(tetramethylcyclopentadienyl)(cyclohexylamido)
 titanium dimethyl, ethylene(tetramethylcyclopentadienyl)(1-adamantylamido)
- 10 titanium dimethyl, ethylene(tetramethylcyclopentadienyl)(t-butylamido) titanium
 dimethyl, dimethylsilyl(2,5-dimethylcyclopentadienyl)(cyclododecylamido)
 titanium dichloride, dimethylsilyl(2,5-dimethylcyclopentadienyl)(exo-2-
 norbornylamido) titanium dichloride, dimethylsilyl(2,5-
 dimethylcyclopentadienyl)(cyclohexylamido) titanium dichloride,
- 15 dimethylsilyl(2,5-dimethylcyclopentadienyl)(1-adamantylamido) titanium
 dichloride, dimethylsilyl(2,5-dimethylcyclopentadienyl)(t-butylamido) titanium
 dichloride, dimethylsilyl(3,4-dimethylcyclopentadienyl)(cyclododecylamido)
 titanium dichloride, dimethylsilyl(3,4-dimethylcyclopentadienyl)(exo-2-
 norbornylamido) titanium dichloride, dimethylsilyl(3,4-
- 20 dimethylcyclopentadienyl)(cyclohexylamido) titanium dichloride,
 dimethylsilyl(3,4-dimethylcyclopentadienyl)(1-adamantylamido) titanium
 dichloride, dimethylsilyl(3,4-dimethylcyclopentadienyl)(t-butylamido) titanium
 dichloride, dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(cyclododecylamido)
 titanium dichloride, dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(exo-2-
- 25 norbornylamido) titanium dichloride, dimethylsilyl(2-ethyl-5-
 methylcyclopentadienyl)(cyclohexylamido) titanium dichloride, dimethylsilyl(2-
 ethyl-5-methylcyclopentadienyl)(1-adamantylamido) titanium dichloride,
 dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(t-butylamido) titanium
 dichloride, dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(cyclododecylamido)
- 30 titanium dichloride, dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(exo-2-
 norbornylamido) titanium dichloride, dimethylsilyl(3-ethyl-4-
 methylcyclopentadienyl)(cyclohexylamido) titanium dichloride, dimethylsilyl(3-
 ethyl-4-methylcyclopentadienyl)(1-adamantylamido) titanium dichloride,
 dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(t-butylamido) titanium
- 35 dichloride, dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-
 octylcyclopentadienyl)(cyclododecylamido) titanium dichloride, dimethylsilyl(2-

- 5 ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(exo-2-norbornylamido) titanium
dichloride, dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-
octylcyclopentadienyl)(cyclohexylamido) titanium dichloride, dimethylsilyl(2-
ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(1-adamantylamido) titanium
dichloride, dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(t-
10 butylamido) titanium dichloride, dimethylsilyl(2-
tetrahydroindenyl)(cyclododecylamido) titanium dichloride, dimethylsilyl(2-
tetrahydroindenyl)(exo-2-norbornylamido) titanium dichloride, dimethylsilyl(2-
tetrahydroindenyl)(cyclohexylamido) titanium dichloride, dimethylsilyl(2-
tetrahydroindenyl)(1-adamantylamido) titanium dichloride, dimethylsilyl(2-
15 tetrahydroindenyl)(t-butylamido) titanium dichloride, dimethylsilyl(2,5-
dimethylcyclopentadienyl)(cyclododecylamido) titanium dimethyl,
dimethylsilyl(2,5-dimethylcyclopentadienyl)(exo-2-norbornylamido) titanium
dimethyl, dimethylsilyl(2,5-dimethylcyclopentadienyl)(cyclohexylamido) titanium
dimethyl, dimethylsilyl(2,5-dimethylcyclopentadienyl)(1-adamantylamido)
20 titanium dimethyl, dimethylsilyl(2,5-dimethylcyclopentadienyl)(t-butylamido)
titanium dimethyl, dimethylsilyl(3,4-
dimethylcyclopentadienyl)(cyclododecylamido) titanium dimethyl,
dimethylsilyl(3,4-dimethylcyclopentadienyl)(exo-2-norbornylamido) titanium
dimethyl, dimethylsilyl(3,4-dimethylcyclopentadienyl)(cyclohexylamido) titanium
25 dimethyl, dimethylsilyl(3,4-dimethylcyclopentadienyl)(1-adamantylamido)
titanium dimethyl, dimethylsilyl(3,4-dimethylcyclopentadienyl)(t-butylamido)
titanium dimethyl, dimethylsilyl(2-ethyl-5-
methylcyclopentadienyl)(cyclododecylamido) titanium dimethyl, dimethylsilyl(2-
ethyl-5-methylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
30 dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(cyclohexylamido) titanium
dimethyl, dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(1-adamantylamido)
titanium dimethyl, dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(t-butylamido)
titanium dimethyl, dimethylsilyl(3-ethyl-4-
methylcyclopentadienyl)(cyclododecylamido) titanium dimethyl, dimethylsilyl(3-
35 ethyl-4-methylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(cyclohexylamido) titanium

5 dimethyl, dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(1-adamantylamido)
 titanium dimethyl, dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(t-butylamido)
 titanium dimethyl, dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-
 octylcyclopentadienyl)(cyclododecylamido) titanium dimethyl, dimethylsilyl(2-
 ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(exo-2-norbornylamido) titanium
 10 dimethyl, dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-
 octylcyclopentadienyl)(cyclohexylamido) titanium dimethyl, dimethylsilyl(2-
 ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(1-adamantylamido) titanium
 dimethyl, dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(t-
 butylamido) titanium dimethyl, dimethylsilyl(2-
 15 tetrahydroindenyl)(cyclododecylamido) titanium dimethyl, dimethylsilyl(2-
 tetrahydroindenyl)(exo-2-norbornylamido) titanium dimethyl, dimethylsilyl(2-
 tetrahydroindenyl)(cyclohexylamido) titanium dimethyl, dimethylsilyl(2-
 tetrahydroindenyl)(1-adamantylamido) titanium dimethyl, dimethylsilyl(2-
 tetrahydroindenyl)(t-butylamido) titanium dimethyl and the like.

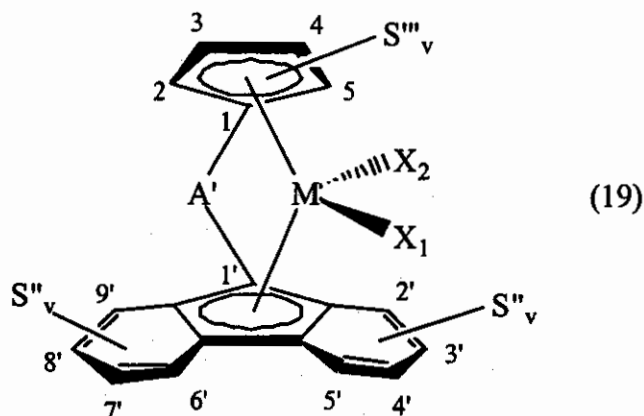
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The most preferred species are:

dimethylsilyl(tetramethylcyclopentadienyl)(cyclododecylamido) titanium
 dichloride, dimethylsilyl(tetramethylcyclopentadienyl)(t-butylamido) titanium
 dichloride, dimethylsilyl(tetramethylcyclopentadienyl)(cyclohexylamido) titanium
 25 dichloride, dimethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido)
 titanium dichloride, dimethylsilyl(tetramethylcyclopentadienyl)(exo-2-
 norbornylamido) titanium dichloride,
 dimethylsilyl(tetramethylcyclopentadienyl)(cyclododecylamido) titanium
 dimethyl, dimethylsilyl(tetramethylcyclopentadienyl)(t-butylamido) titanium
 30 dimethyl, dimethylsilyl(tetramethylcyclopentadienyl)(cyclohexylamido) titanium
 dimethyl, dimethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido)
 titanium dimethyl, and dimethylsilyl(tetramethylcyclopentadienyl)(exo-2-
 norbornylamido) titanium dimethyl.

35 Additionally, at higher reaction temperatures, some catalysts that produce
 syndiotactic poly- α -olefin at lower temperatures, will produce virtually non-

- 5 crystalline poly- α -olefins at higher temperatures. The choice of transition metal component for this amorphous polymer fraction is a subset of the transition metal component of equations 8-9. Preferred components of this type are illustrated in equation 19:



- 10 wherein A', M, X₁ and X₂ are as previously defined. Substituents S''_v and S'''_v are independently defined as S'' in equations 8-9 where the subscript "v" denotes the carbon atom on the Cp-ring or Flu-ring (fluorenyl-ring) to which the substituent is bonded.

- 15 Preferably metallocene precursors for producing poly- α -olefins having largely amorphous character (when used as catalysts under higher reactor temperature conditions) are those of Equation 19 where S'''_v are independently chosen such that the metallocene framework has a plane of symmetry containing the metal center and bisecting the Flu- and Cp-rings. The A' ligand need not be
- 20 symmetrical – for example dimethylsilyl or methylphenylsilyl will not effect the stereochemistry of the polymer produced. Substituent S'''_v is defined to be the same as S'' in equation 8-9 where the subscript "v" denotes the carbon atom on the cyclopentadienyl ring to which the substituent is bonded and where there can be zero, two or four substituents, S''', on the cyclopentadienyl ring provided that the
- 25 cyclopentadienyl ring is symmetrically substituted. Symmetrically substituted is defined to mean that the cyclopentadienyl ring is substituted in the 2 and 5 positions and/or 3 and 4 positions with S''' groups that are of approximately of the same steric bulk. Typically the size of these S''' groups are within 2 carbons of

5 each other. Thus a cyclopentadienyl substituted at the 2 and the 5 positions with methyl and ethyl respectively, or substituted at the 3 and the 4 positions with hexyl and octyl, respectively, would be considered symmetric. Likewise, the cyclopentadienyl ring may be substituted at all four sites with S''' groups and be considered symmetric as long as each of the symmetrical pairs are of similar steric
 10 bulk. Additionally, two adjacent S'''-groups in the 3 and 4 position may be linked to form a ring provided that the new ring is also symmetrically substituted. Because of the distant placement of the S'' substituents on the fluorenyl ring, these substituents need not be symmetrically placed on the fluorenyl ring. Hence, the fluorenyl ring may be substituted with from 0-7 substituents that may be the
 15 same or different. Two or more adjacent S''-groups may optionally be linked to form a ring.

Preferred metallocene transition metal compounds which, according to the present invention, provide catalyst systems which are specific to the production of
 20 amorphous or low crystallinity poly- α -olefins include:
 isopropylidene(cyclopentadienyl)(fluorenyl) zirconium dichloride,
 isopropylidene(cyclopentadienyl)(fluorenyl) zirconium dimethyl,
 methylene(cyclopentadienyl)(fluorenyl) zirconium dichloride,
 methylene(cyclopentadienyl)(fluorenyl) zirconium dimethyl,
 25 diphenylmethylene(cyclopentadienyl)(fluorenyl) zirconium dichloride,
 diphenylmethylene(cyclopentadienyl)(fluorenyl) zirconium dimethyl, di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(fluorenyl) zirconium dichloride,
 di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(fluorenyl) zirconium dimethyl, di(p-trimethylsilylphenyl)methylene(cyclopentadienyl)(fluorenyl)
 30 zirconium dichloride, di(p-trimethylsilylphenyl)methylene(cyclopentadienyl)(fluorenyl) zirconium dimethyl,
 diphenylsilyl(cyclopentadienyl)(fluorenyl) zirconium dichloride,
 diphenylsilyl(cyclopentadienyl)(fluorenyl) zirconium dimethyl,
 dimethylsilyl(cyclopentadienyl)(fluorenyl) zirconium dichloride,
 35 dimethylsilyl(cyclopentadienyl)(fluorenyl) zirconium dimethyl,
 methylphenylsilyl(cyclopentadienyl)(fluorenyl) zirconium dichloride,

- 5 methylphenylsilyl(cyclopentadienyl)(fluorenyl) zirconium dimethyl,
isopropylidene(cyclopentadienyl)(fluorenyl) hafnium dichloride,
isopropylidene(cyclopentadienyl)(fluorenyl) hafnium dimethyl,
methylene(cyclopentadienyl)(fluorenyl) hafnium dichloride,
methylene(cyclopentadienyl)(fluorenyl) hafnium dimethyl,
- 10 diphenylmethylene(cyclopentadienyl)(fluorenyl) hafnium dichloride,
diphenylmethylene(cyclopentadienyl)(fluorenyl) hafnium dimethyl, di(p-
triethylsilylphenyl)methylene(cyclopentadienyl)(fluorenyl) hafnium dichloride,
di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(fluorenyl) hafnium
dimethyl, di(p-trimethylsilylphenyl)methylene(cyclopentadienyl)(fluorenyl)
- 15 hafnium dichloride,
di(p-trimethylsilylphenyl)methylene(cyclopentadienyl)(fluorenyl) hafnium
dimethyl, diphenylsilyl(cyclopentadienyl)(fluorenyl) hafnium dichloride,
diphenylsilyl(cyclopentadienyl)(fluorenyl) hafnium dimethyl,
dimethylsilyl(cyclopentadienyl)(fluorenyl) hafnium dichloride,
- 20 dimethylsilyl(cyclopentadienyl)(fluorenyl) hafnium dimethyl,
methylphenylsilyl(cyclopentadienyl)(fluorenyl) hafnium dichloride,
methylphenylsilyl(cyclopentadienyl)(fluorenyl) hafnium dimethyl,
isopropylidene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) zirconium dichloride,
isopropylidene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) zirconium dimethyl,
- 25 methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) zirconium dichloride,
methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) zirconium dimethyl,
diphenylmethylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) zirconium
dichloride, diphenylmethylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl)
zirconium dimethyl,
- 30 di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl)
zirconium dichloride, di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,8-
di-t-butylfluorenyl) zirconium dimethyl, di(p-
trimethylsilylphenyl)methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl)
zirconium dichloride, di(p-trimethylsilylphenyl)methylene(cyclopentadienyl)(3,8-
di-t-butylfluorenyl) zirconium dimethyl, diphenylsilyl(cyclopentadienyl)(3,8-di-t-
butylfluorenyl) zirconium dichloride, diphenylsilyl(cyclopentadienyl)(3,8-di-t-
- 35

- 5 butylfluorenyl) zirconium dimethyl, dimethylsilyl(cyclopentadienyl)(3,8-di-t-butylfluorenyl) zirconium dichloride, dimethylsilyl(cyclopentadienyl)(3,8-di-t-butylfluorenyl) zirconium dimethyl, methylphenylsilyl(cyclopentadienyl)(3,8-di-t-butylfluorenyl) zirconium dichloride, methylphenylsilyl(cyclopentadienyl)(3,8-di-t-butylfluorenyl) zirconium dimethyl, isopropylidene(cyclopentadienyl)(3,8-di-t-
- 10 butylfluorenyl) hafnium dichloride, isopropylidene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dimethyl, methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dichloride, methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dimethyl, diphenylmethylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dichloride, diphenylmethylene(cyclopentadienyl)(3,8-di-
- 15 t-butylfluorenyl) hafnium dimethyl, di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dichloride, di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dimethyl, di(p-trimethylsilylphenyl)methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl)
- 20 hafnium dichloride, di(p-trimethylsilylphenyl)methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dimethyl, diphenylsilyl(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dichloride, diphenylsilyl(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dimethyl, dimethylsilyl(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dichloride, dimethylsilyl(cyclopentadienyl)(3,8-di-t-
- 25 butylfluorenyl) hafnium dimethyl, methylphenylsilyl(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dichloride, methylphenylsilyl(cyclopentadienyl)(3,8-di-t-butylfluorenyl) hafnium dimethyl,
- isopropylidene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) zirconium dichloride,
- 30 isopropylidene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) zirconium dimethyl,
- methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) zirconium dichloride,
- methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
- 35 octahydrodibenzyl[b,h]fluorenyl) zirconium dimethyl,
- diphenylmethylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-

- 5 4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) zirconium dichloride,
 diphenylmethylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-
 4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) zirconium dimethyl,
 di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-
 octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) zirconium dichloride,
- 10 di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-
 octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) zirconium dimethyl,
 di(p-trimethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-
 octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) zirconium dichloride,
 di(p-trimethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-
 15 octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) zirconium dimethyl,
 diphenylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) zirconium dichloride,
 diphenylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) zirconium dimethyl,
- 20 dimethylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) zirconium dichloride,
 dimethylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) zirconium dimethyl,
 methylphenylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 25 octahydrodibenzyl[b,h]fluorenyl) zirconium dichloride,
 methylphenylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) zirconium dimethyl,
 isopropylidene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) hafnium dichloride,
- 30 isopropylidene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) hafnium dimethyl,
 methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) hafnium dichloride,
 methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 35 octahydrodibenzyl[b,h]fluorenyl) hafnium dimethyl,
 diphenylmethylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-

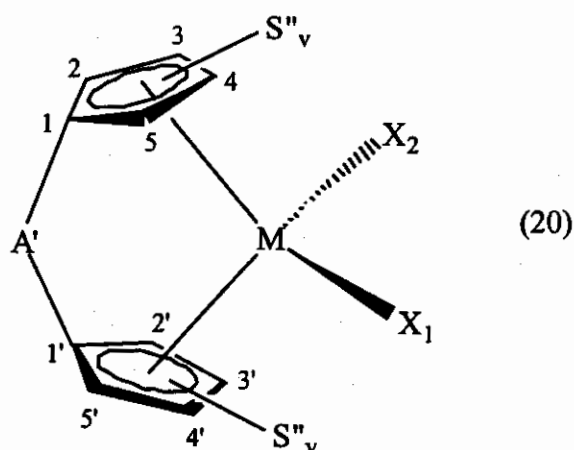
- 5 4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) hafnium dichloride,
 diphenylmethylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-
 4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) hafnium dimethyl,
 di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-
 octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) hafnium dichloride,
 10 di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-
 octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) hafnium dimethyl,
 di(p-trimethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-
 octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) hafnium dichloride,
 di(p-trimethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-
 15 octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) hafnium dimethyl,
 diphenylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) hafnium dichloride,
 diphenylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) hafnium dimethyl,
 20 dimethylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) hafnium dichloride,
 dimethylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) hafnium dimethyl,
 methylphenylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 25 octahydrodibenzyl[b,h]fluorenyl) hafnium dichloride,
 methylphenylsilyl(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-
 octahydrodibenzyl[b,h]fluorenyl) hafnium dimethyl, and the like.

- The most preferred species are: di(p-
 30 triethylsilylphenyl)methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl)
 zirconium dichloride, di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,8-
 di-t-butylfluorenyl) hafnium dichloride, di(p-
 triethylsilylphenyl)methylene(cyclopentadienyl)(3,8-di-t-butylfluorenyl)
 zirconium dimethyl, di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,8-di-
 35 t-butylfluorenyl) hafnium dimethyl, di(p-
 triethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-

- 5 4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) zirconium dichloride, di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) hafnium dichloride, di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) zirconium dimethyl, and di(p-triethylsilylphenyl)methylene(cyclopentadienyl)(3,3,6,6,9,9,12,12-octamethyl-4,4,5,5,8,8,9,9-octahydrodibenzyl[b,h]fluorenyl) hafnium dimethyl.
- 10

Additionally, compounds of formula 20 may be used to produce the amorphous polymer fraction.

15



- In this case, S''_v are independently chosen such that the metallocene framework has a plane of symmetry that bisects M and A'. Substituents S''_v are independently defined to be the same as S'' in equation 8-9 where the subscript "v" denotes the carbon atom on the cyclopentadienyl ring to which the substituent is bonded and where there can be zero to four substituents, S'' , on the cyclopentadienyl ring provided that the cyclopentadienyl ring is symmetrically substituted. Symmetrically substituted is defined to mean that the cyclopentadienyl ring is substituted in the 2 and 2' positions and/or 3 and 3' positions and/or 4 and 4' positions and/or 5 and 5' positions with S'' groups that are of approximately of the same steric bulk. Typically the size of these S'' groups are within 2 carbons of each other. Thus a cyclopentadienyl substituted at the 2 and the 2' positions with methyl and ethyl respectively, or substituted at the 3 and
- 20
- 25

- 5 the 3' positions with hexyl and octyl, respectively, would be considered symmetric. Likewise, the cyclopentadienyl ring may be substituted at all four sites with S'' groups and be considered symmetric as long as each of the symmetrical pairs are of similar steric bulk. Additionally, two adjacent S''-groups may be linked to form a ring provided that the new ring is also symmetrically substituted.
- 10 Such complexes such as meso-Me₂Si(indenyl)₂ZrMe₂ meso-CH₂CH₂(indenyl)₂ZrCl₂ are well known in the art and generally produce amorphous polymers useful in this invention.

- Preferred meso-metallocene compounds which, according to the present
- 15 invention, provide catalyst systems which are specific to the production of amorphous poly- α -olefins include the meso versions of:
- dimethylsilylbis(indenyl) zirconium dichloride, dimethylsilylbis(indenyl) zirconium dimethyl, diphenylsilylbis(indenyl) zirconium dichloride, diphenylsilylbis(indenyl) zirconium dimethyl, methylphenylsilylbis(indenyl) zirconium dichloride, methylphenylsilylbis(indenyl) zirconium dimethyl,
- 20 ethylenebis(indenyl) zirconium dichloride, ethylenebis(indenyl) zirconium dimethyl, methylenebis(indenyl) zirconium dichloride, methylenebis(indenyl) zirconium dimethyl, dimethylsilylbis(indenyl) hafnium dichloride, dimethylsilylbis(indenyl) hafnium dimethyl, diphenylsilylbis(indenyl) hafnium dichloride, diphenylsilylbis(indenyl) hafnium dimethyl,
- 25 methylphenylsilylbis(indenyl) hafnium dichloride, methylphenylsilylbis(indenyl) hafnium dimethyl, ethylenebis(indenyl) hafnium dichloride, ethylenebis(indenyl) hafnium dimethyl, methylenebis(indenyl) hafnium dichloride, methylenebis(indenyl) hafnium dimethyl, dimethylsilylbis(tetrahydroindenyl) zirconium dichloride, dimethylsilylbis(tetrahydroindenyl) zirconium dimethyl,
- 30 diphenylsilylbis(tetrahydroindenyl) zirconium dichloride, diphenylsilylbis(tetrahydroindenyl) zirconium dimethyl, methylphenylsilylbis(tetrahydroindenyl) zirconium dichloride, methylphenylsilylbis(tetrahydroindenyl) zirconium dimethyl,
- 35 ethylenebis(tetrahydroindenyl) zirconium dichloride, ethylenebis(tetrahydroindenyl) zirconium dimethyl,

- 5 methylenebis(tetrahydroindenyl) zirconium dichloride,
methylenebis(tetrahydroindenyl) zirconium dimethyl,
dimethylsilylbis(tetrahydroindenyl) hafnium dichloride,
dimethylsilylbis(tetrahydroindenyl) hafnium dimethyl,
diphenylsilylbis(tetrahydroindenyl) hafnium dichloride,
10 diphenylsilylbis(tetrahydroindenyl) hafnium dimethyl,
methylphenylsilylbis(tetrahydroindenyl) hafnium dichloride,
methylphenylsilylbis(tetrahydroindenyl) hafnium dimethyl,
ethylenebis(tetrahydroindenyl) hafnium dichloride, ethylenebis(tetrahydroindenyl)
hafnium dimethyl, methylenebis(tetrahydroindenyl) hafnium dichloride,
15 methylenebis(tetrahydroindenyl) hafnium dimethyl, dimethylsilylbis(2-
methylindenyl) zirconium dichloride, dimethylsilylbis(2-methylindenyl)
zirconium dimethyl, diphenylsilylbis(2-methylindenyl) zirconium dichloride,
diphenylsilylbis(2-methylindenyl) zirconium dimethyl, methylphenylsilylbis(2-
methylindenyl) zirconium dichloride, methylphenylsilylbis(2-methylindenyl)
20 zirconium dimethyl, ethylenebis(2-methylindenyl) zirconium dichloride,
ethylenebis(2-methylindenyl) zirconium dimethyl, methylenebis(2-methylindenyl)
zirconium dichloride, methylenebis(2-methylindenyl) zirconium dimethyl,
dimethylsilylbis(2-methylindenyl) hafnium dichloride, dimethylsilylbis(2-
methylindenyl) hafnium dimethyl, diphenylsilylbis(2-methylindenyl) hafnium
25 dichloride, diphenylsilylbis(2-methylindenyl) hafnium dimethyl,
methylphenylsilylbis(2-methylindenyl) hafnium dichloride,
methylphenylsilylbis(2-methylindenyl) hafnium dimethyl, ethylenebis(2-
methylindenyl) hafnium dichloride, ethylenebis(2-methylindenyl) hafnium
dimethyl, methylenebis(2-methylindenyl) hafnium dichloride, methylenebis(2-
30 methylindenyl) hafnium dimethyl, dimethylsilylbis(2-methyl-4-phenylindenyl)
zirconium dichloride, diphenylsilylbis(2-methyl-4-phenylindenyl) zirconium
dimethyl, diphenylsilylbis(2-methyl-4-phenylindenyl) zirconium dichloride,
diphenylsilylbis(2-methyl-4-phenylindenyl) zirconium dimethyl,
methylphenylsilylbis(2-methyl-4-phenylindenyl) zirconium dichloride,
35 methylphenylsilylbis(2-methyl-4-phenylindenyl) zirconium dimethyl,
ethylenebis(2-methyl-4-phenylindenyl) zirconium dichloride, ethylenebis(2-

- 5 methyl-4-phenylindenyl) zirconium dimethyl, methylenebis(2-methyl-4-phenylindenyl) zirconium dichloride, methylenebis(2-methyl-4-phenylindenyl) zirconium dimethyl, dimethylsilylbis(2-methyl-4-phenylindenyl) hafnium dichloride, dimethylsilylbis(2-methyl-4-phenylindenyl) hafnium dimethyl, diphenylsilylbis(2-methyl-4-phenylindenyl) hafnium dichloride,
10 diphenylsilylbis(2-methyl-4-phenylindenyl) hafnium dimethyl, methylphenylsilylbis(2-methyl-4-phenylindenyl) hafnium dichloride, methylphenylsilylbis(2-methyl-4-phenylindenyl) hafnium dimethyl, ethylenebis(2-methyl-4-phenylindenyl) hafnium dichloride, ethylenebis(2-methyl-4-phenylindenyl) hafnium dimethyl, methylenebis(2-methyl-4-phenylindenyl) hafnium dichloride, methylenebis(2-methyl-4-phenylindenyl) hafnium dimethyl,
15 dimethylsilylbis(4,7-dimethylindenyl) zirconium dichloride, dimethylsilylbis(4,7-dimethylindenyl) zirconium dimethyl, diphenylsilylbis(4,7-dimethylindenyl) zirconium dichloride, diphenylsilylbis(4,7-dimethylindenyl) zirconium dimethyl, methylphenylsilylbis(4,7-dimethylindenyl) zirconium dichloride,
20 methylphenylsilylbis(4,7-dimethylindenyl) zirconium dimethyl, ethylenebis(4,7-dimethylindenyl) zirconium dichloride, ethylenebis(4,7-dimethylindenyl) zirconium dimethyl, methylenebis(4,7-dimethylindenyl) zirconium dichloride, methylenebis(4,7-dimethylindenyl) zirconium dimethyl, dimethylsilylbis(4,7-dimethylindenyl) hafnium dichloride, dimethylsilylbis(4,7-dimethylindenyl) hafnium dimethyl, diphenylsilylbis(4,7-dimethylindenyl) hafnium dichloride,
25 diphenylsilylbis(4,7-dimethylindenyl) hafnium dimethyl, methylphenylsilylbis(4,7-dimethylindenyl) hafnium dichloride, methylphenylsilylbis(4,7-dimethylindenyl) hafnium dimethyl, ethylenebis(4,7-dimethylindenyl) hafnium dichloride, ethylenebis(4,7-dimethylindenyl) hafnium dimethyl,
30 dimethyl, methylenebis(4,7-dimethylindenyl) hafnium dichloride, methylenebis(4,7-dimethylindenyl) hafnium dimethyl, and the like.

The most preferred species are the racemic versions of:

- dimethylsilylbis(indenyl) zirconium dichloride, dimethylsilylbis(indenyl) zirconium dimethyl, ethylenebis(indenyl) zirconium dichloride,
35 ethylenebis(indenyl) zirconium dimethyl, dimethylsilylbis(indenyl) hafnium

- 5 dichloride, dimethylsilylbis(indenyl) hafnium dimethyl, ethylenebis(indenyl) hafnium dichloride, ethylenebis(indenyl) hafnium dimethyl, dimethylsilylbis(tetrahydroindenyl) zirconium dichloride, dimethylsilylbis(tetrahydroindenyl) zirconium dimethyl, ethylenebis(tetrahydroindenyl) zirconium dichloride,
- 10 ethylenebis(tetrahydroindenyl) zirconium dimethyl, dimethylsilylbis(tetrahydroindenyl) hafnium dichloride, dimethylsilylbis(tetrahydroindenyl) hafnium dimethyl, ethylenebis(tetrahydroindenyl) hafnium dichloride, ethylenebis(tetrahydroindenyl) hafnium dimethyl, dimethylsilylbis(2-methylindenyl) zirconium dichloride,
- 15 dimethylsilylbis(2-methylindenyl) zirconium dimethyl, ethylenebis(2-methylindenyl) zirconium dichloride, ethylenebis(2-methylindenyl) zirconium dimethyl, dimethylsilylbis(2-methylindenyl) hafnium dichloride, dimethylsilylbis(2-methylindenyl) hafnium dimethyl, ethylenebis(2-methylindenyl) hafnium dichloride, and ethylenebis(2-methylindenyl) hafnium
- 20 dimethyl

When two transition metal compound based catalysts are used in one reactor as a mixed catalyst system, the two transition metal compounds should be chosen such that the two are compatible. A simple screening method such as by

25 ^1H or ^{13}C NMR, known to those of ordinary skill in the art, can be used to determine which transition metal compounds are compatible.

It is preferable to use the same activator for the transition metal compounds, however, two different activators, such as a non-coordinating anion

30 activator and an alumoxane, can be used in combination. If one or more transition metal compounds contain an X_1 or X_2 ligand which is not a hydride, hydrocarbyl, or substituted hydrocarbyl, then the alumoxane should be contacted with the transition metal compounds prior to addition of the non-coordinating anion activator.

35

5 Particularly preferred combinations of transition metal compounds
include:

(1) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-c-C}_{12}\text{H}_{23})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrCl}_2$
(2-Me-4-PhInd = 2-methyl-4-phenylindenyl, $\text{c-C}_{12}\text{H}_{23}$ = cyclododecyl, Me_4C_5 -
10 tetramethylcyclopentadienyl) activated with an alumoxane, such as
methylalumoxane or modified methylalumoxane;

(2) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-c-C}_{12}\text{H}_{23})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$
activated with a non-coordinating anion activator, such as N,N-
15 dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

(3) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-c-C}_{12}\text{H}_{23})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrCl}_2$ (2-
MeInd = 2-methyl-indenyl) activated with an alumoxane, such as
20 methylalumoxane or modified methylalumoxane;

(4) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-c-C}_{12}\text{H}_{23})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$
activated with a non-coordinating anion activator, such as N,N-dimethylanilinium
tetrakis(pentafluorophenyl)boron or triphenylcarbonium
25 tetrakis(pentafluorophenyl)boron;

(5) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-1-adamantyl})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrCl}_2$
activated with an alumoxane, such as methylalumoxane or modified
methylalumoxane;

30
(6) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-1-adamantyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$
activated with a non-coordinating anion activator, such as N,N-
dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

35

5 (7) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-1-adamantyl})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrCl}_2$
activated with an alumoxane, such as methylalumoxane or modified
methylalumoxane;

(8) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-1-adamantyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$
10 activated with a non-coordinating anion activator, such as N,N-dimethylanilinium
tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

(9) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-butyl})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrCl}_2$
15 activated with an alumoxane, such as methylalumoxane or modified
methylalumoxane;

(10) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-butyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$
activated with a non-coordinating anion activator, such as N,N-dimethylanilinium
20 tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

(11) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-butyl})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})$ activated
with an alumoxane, such as methylalumoxane or modified methylalumoxane;
25

(12) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-butyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$
activated with a non-coordinating anion activator, such as N,N-dimethylanilinium
tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;
30

(13) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-exo-norbornyl})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrCl}_2$
activated with an alumoxane, such as methylalumoxane or modified
methylalumoxane;

35 (14) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-exo-norbornyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$
activated with a non-coordinating anion activator, such as N,N-

- 5 dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

(15) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-exo-norbornyl})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrCl}_2$
activated with an alumoxane, such as methylalumoxane or modified
10 methylalumoxane;

(16) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-exo-norbornyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium
15 tetrakis(pentafluorophenyl)boron;

(17) $(\text{p-Et}_3\text{SiPh})_2\text{C}(\text{Cp})(3,8\text{-di-t-BuFlu})\text{HfCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrCl}_2$ (3,6-di-t-BuFlu = 3,8-di-tert-butylfluorenyl, Cp = cyclopentadienyl)
activated with an alumoxane, such as methylalumoxane or modified
20 methylalumoxane;

(18) $(\text{p-Et}_3\text{SiPh})_2\text{C}(\text{Cp})(3,8\text{-di-t-BuFlu})\text{HfMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium
25 tetrakis(pentafluorophenyl)boron;

(19) $(\text{p-Et}_3\text{SiPh})_2\text{C}(\text{Cp})(3,8\text{-di-t-BuFlu})\text{HfCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrCl}_2$ activated with an alumoxane, such as methylalumoxane or modified
methylalumoxane;

30

(20) $(\text{p-Et}_3\text{SiPh})_2\text{C}(\text{Cp})(3,8\text{-di-t-BuFlu})\text{HfMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

35

5 (21) meso-CH₂CH₂(Ind)₂ZrCl₂ and rac-Me₂Si(H₄Ind)₂ZrCl₂ (Ind = indenyl, H₄Ind = tetrahydroindenyl) activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

(22) meso-CH₂CH₂(Ind)₂ZrMe₂ and rac-Me₂Si(H₄Ind)₂ZrMe₂ activated
10 with a non-coordinating anion activator, such as N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(23) meso-CH₂CH₂(Ind)₂ZrCl₂ and rac-Me₂Si(2-MeInd)₂ZrCl₂ activated
15 with an alumoxane, such as methylalumoxane or modified methylalumoxane;

(24) meso-CH₂CH₂(Ind)₂ZrMe₂ and rac-Me₂Si(2-MeInd)₂ZrMe₂ activated
with a non-coordinating anion activator, such as N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium
20 tetrakis(pentafluorophenyl)boron;

(25) meso-Me₂Si(Ind)₂ZrCl₂ and rac-Me₂Si(H₄Ind)₂ZrCl₂ activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

25 (26) meso-Me₂Si(Ind)₂ZrMe₂ and rac-Me₂Si(H₄Ind)₂ZrMe₂ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

30 (27) meso-Me₂Si(Ind)₂ZrCl₂ and rac-Me₂Si(2-MeInd)₂ZrCl₂ activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

(28) meso-Me₂Si(Ind)₂ZrMe₂ and rac-Me₂Si(2-MeInd)₂ZrMe₂ activated
with a non-coordinating anion activator, such as N,N-dimethylanilinium
35 tetrakis(pentafluorophenyl)boron or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

5

(29) $\text{meso-Me}_2\text{Si(2-MeInd)}_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si(2-Me-4-PhInd)}_2\text{ZrCl}_2$ activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

10

(30) $\text{meso-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si(2-Me-4-PhInd)}_2\text{ZrMe}_2$ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

15

(31) $\text{meso-Me}_2\text{Si(2-MeInd)}_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si(2-MeInd)}_2\text{ZrCl}_2$ activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

20

(32) $\text{meso-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

25

(33) $\text{meso-CH}_2\text{CH}_2\text{(2-MeInd)}_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si(2-Me-4-PhInd)}_2\text{ZrCl}_2$ activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

30

(34) $\text{meso-CH}_2\text{CH}_2\text{(2-MeInd)}_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si(2-Me-4-PhInd)}_2\text{ZrMe}_2$ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

35

(35) $\text{meso-CH}_2\text{CH}_2\text{(2-MeInd)}_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si(2-MeInd)}_2\text{ZrCl}_2$ activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

5 (36) meso-CH₂CH₂(2-MeInd)₂ZrMe₂ and rac-Me₂Si(2-MeInd)₂ZrMe₂
activated with a non-coordinating anion activator, such as N,N-dimethylanilinium
tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

10 (37) meso-Me₂Si(2-Me-4-PhInd)₂ZrCl₂ and rac-Me₂Si(2-Me-4-
PhInd)₂ZrCl₂ activated with an alumoxane, such as methylalumoxane or modified
methylalumoxane;

(38) meso-Me₂Si(2-Me-4-PhInd)₂ZrMe₂ and rac-Me₂Si(2-Me-4-
15 PhInd)₂ZrMe₂ activated with a non-coordinating anion activator, such as N,N-
dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

(39) meso-CH₂CH₂(2-Me-4-PhInd)₂ZrCl₂ and rac-CH₂CH₂(2-Me-4-
20 PhInd)₂ZrCl₂ activated with an alumoxane, such as methylalumoxane or modified
methylalumoxane;

(40) meso-CH₂CH₂(2-Me-4-PhInd)₂ZrMe₂ and rac-CH₂CH₂(2-Me-4-
PhInd)₂ZrMe₂ activated with a non-coordinating anion activator, such as N,N-
25 dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

(41) meso-CH₂CH₂(2-MeInd)₂ZrCl₂ and rac-CH₂CH₂(2-MePhInd)₂ZrCl₂
activated with an alumoxane, such as methylalumoxane or modified
30 methylalumoxane;

(42) meso-CH₂CH₂(2-MeInd)₂ZrMe₂ and rac-CH₂CH₂(2-MeInd)₂ZrMe₂
activated with a non-coordinating anion activator, such as N,N-dimethylanilinium
tetrakis(pentafluorophenyl)boron or triphenylcarbonium
35 tetrakis(pentafluorophenyl)boron;

5 (43) meso-CH₂CH₂(Ind)₂ZrCl₂ and rac-CH₂CH₂(Ind)₂ZrCl₂ activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

(44) meso-CH₂CH₂(Ind)₂ZrMe₂ and rac-CH₂CH₂(Ind)₂ZrMe₂ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium
10 tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

(45) meso-Me₂Si(Ind)₂ZrCl₂ and rac-Me₂Si(Ind)₂ZrCl₂ activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

15 (46) meso-Me₂Si(Ind)₂ZrMe₂ and rac-Me₂Si(Ind)₂ZrMe₂ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium
tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

20 (47) meso-CH₂CH₂(Ind)₂ZrCl₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrCl₂ (4,7-Me₂Ind = 4,7-dimethylindenyl) activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

25 (48) meso-CH₂CH₂(Ind)₂ZrMe₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrMe₂ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium
tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

30 (49) meso-Me₂Si(Ind)₂ZrCl₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrCl₂ activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

(50) meso-Me₂Si(Ind)₂ZrMe₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrMe₂ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium
35 tetrakis(pentafluorophenyl)boron or triphenylcarbonium
tetrakis(pentafluorophenyl)boron;

5

(51) $\text{meso-CH}_2\text{CH}_2(2\text{-MeInd})_2\text{ZrCl}_2$ and $\text{rac-CH}_2\text{CH}_2(4,7\text{-Me}_2\text{Ind})_2\text{ZrCl}_2$ ($4,7\text{-Me}_2\text{Ind} = 4,7\text{-dimethylindenyl}$) activated with an alumoxane, such as methylalumoxane or modified methylalumoxane;

10

(52) $\text{meso-CH}_2\text{CH}_2(2\text{-MeInd})_2\text{ZrMe}_2$ and $\text{rac-CH}_2\text{CH}_2(4,7\text{-Me}_2\text{Ind})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

15

(53) $\text{meso-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrCl}_2$ and $\text{rac-CH}_2\text{CH}_2(4,7\text{-Me}_2\text{Ind})_2\text{ZrCl}_2$ activated with an alumoxane, such as methylalumoxane or modified methylalumoxane; and

20

(54) $\text{meso-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ and $\text{rac-CH}_2\text{CH}_2(4,7\text{-Me}_2\text{Ind})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator, such as N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron or triphenylcarbonium tetrakis(pentafluorophenyl)boron.

The two transition metal compounds (pre-catalysts) may be used in any
25 ratio. Preferred molar ratios of (A) transition metal compound to produce amorphous polymer to (B) transition metal compound to produce crystalline polymer fall within the range of (A:B) 1:1000 to 1000:1, alternatively 1:100 to 500:1, alternatively 1:10 to 200:1, alternatively 1:1 to 100:1, and alternatively 1:1 to 75:1, and alternatively 5:1 to 50:1. The particular ratio chosen will depend on
30 the exact pre-catalysts chosen, the method of activation, and the end product desired. In a particular embodiment, when using the two pre-catalysts (A-“amorphous polymer producing precatalyst” and B-“crystalline polymer producing catalyst”), where both are activated with the same activator, the preferred mole percents, based upon the molecular weight of the pre-catalysts, are
35 10 to 99.9% A to 0.1 to 90%B, alternatively 25 to 99% A to 0.5 to 50% B,

- 5 alternatively 50 to 99% A to 1 to 25% B, and alternatively 75 to 99% A to 1 to 10%B.

In general the combined pre-catalyst compounds and the activator are combined in ratios of about 1:10,000 to about 10:1. When alumoxane or
10 aluminum alkyl activators are used, the combined pre-catalyst-to-activator molar ratio is from 1:5000 to 10:1, alternatively from 1:1000 to 10:1; alternatively, 1:500 to 2:1; or 1:300 to 1:1. When ionizing activators are used, the combined pre-catalyst-to-activator molar ratio is from 10:1 to 1:10; 5:1 to 1:5; 2:1 to 1:2; or 1.2:1 to 1:1. Multiple activators may be used, including using mixes of
15 alumoxanes or aluminum alkyls with ionizing activators.

In another preferred embodiment a third catalyst (pre-catalyst plus activator) is present in the processes described above. The third catalyst may be any of the pre-catalyst components listed herein. Preferred third pre-catalysts
20 include those that are capable of producing waxes. Preferred examples include: rac-dimethylsilylbis(4,7-dimethylindenyl) hafnium dichloride, rac-dimethylsilylbis(4,7-dimethylindenyl) hafnium dimethyl, rac-dimethylsilylbis(4,7-dimethylindenyl) zirconium dichloride, rac-dimethylsilylbis(4,7-dimethylindenyl) zirconium dimethyl, rac-dimethylsilylbis(indenyl) hafnium dichloride, rac-
25 dimethylsilylbis(indenyl) hafnium dimethyl, rac-dimethylsilylbis(indenyl) zirconium dichloride, rac-dimethylsilylbis(indenyl) zirconium dimethyl, rac-dimethylsilylbis(tetrahydroindenyl) hafnium dichloride, rac-dimethylsilylbis(tetrahydroindenyl) hafnium dimethyl, rac-dimethylsilylbis(tetrahydroindenyl) zirconium dichloride, rac-
30 dimethylsilylbis(tetrahydroindenyl) zirconium dimethyl, rac-diphenylsilylbis(4,7-dimethylindenyl) hafnium dichloride, rac-diphenylsilylbis(4,7-dimethylindenyl) hafnium dimethyl, rac-diphenylsilylbis(4,7-dimethylindenyl) zirconium dichloride, rac-diphenylsilylbis(4,7-dimethylindenyl) zirconium dimethyl, rac-diphenylsilylbis(indenyl) hafnium dichloride, rac-diphenylsilylbis(indenyl)
35 hafnium dimethyl,

- 5 rac-diphenylsilylbis(indenyl) zirconium dichloride, rac-diphenylsilylbis(indenyl) zirconium dimethyl, rac-diphenylsilylbis(tetrahydroindenyl) hafnium dichloride, rac-diphenylsilylbis(tetrahydroindenyl) hafnium dimethyl, rac-diphenylsilylbis(tetrahydroindenyl) zirconium dichloride, rac-diphenylsilylbis(tetrahydroindenyl) zirconium dimethyl, rac-
- 10 methylphenylsilylbis(4,7-dimethylindenyl) hafnium dichloride, rac-methylphenylsilylbis(4,7-dimethylindenyl) hafnium dimethyl, rac-methylphenylsilylbis(4,7-dimethylindenyl) zirconium dichloride, rac-methylphenylsilylbis(4,7-dimethylindenyl) zirconium dimethyl, rac-methylphenylsilylbis(indenyl) hafnium dichloride, rac-
- 15 methylphenylsilylbis(indenyl) hafnium dimethyl, rac-methylphenylsilylbis(indenyl) zirconium dichloride, rac-methylphenylsilylbis(indenyl) zirconium dimethyl, rac-methylphenylsilylbis(tetrahydroindenyl) hafnium dichloride, rac-methylphenylsilylbis(tetrahydroindenyl) hafnium dimethyl, rac-
- 20 methylphenylsilylbis(tetrahydroindenyl) zirconium dichloride, rac-methylphenylsilylbis(tetrahydroindenyl) zirconium dimethyl, rac-ethylenebis(4,7-dimethylindenyl) hafnium dichloride, rac-ethylenebis(4,7-dimethylindenyl) hafnium dimethyl, rac-ethylenebis(4,7-dimethylindenyl) zirconium dichloride, rac-ethylenebis(4,7-dimethylindenyl) zirconium dimethyl, rac-
- 25 ethylenebis(indenyl) hafnium dichloride, rac-ethylenebis(indenyl) hafnium dimethyl, rac-ethylenebis(indenyl) zirconium dichloride, rac-ethylenebis(indenyl) zirconium dimethyl, rac-ethylenebis(tetrahydroindenyl) hafnium dichloride, rac-ethylenebis(tetrahydroindenyl) hafnium dimethyl, rac-ethylenebis(tetrahydroindenyl) zirconium dichloride, and rac-
- 30 ethylenebis(tetrahydroindenyl) zirconium dimethyl

- Three transition metal compounds (pre-catalysts) may be used in any ratio. Preferred molar ratios of (A) transition metal compound to produce amorphous polypropylene to (B) transition metal compound to produce crystalline
- 35 polypropylene to (C) transition metal compound to produce wax fall within the range of (A:B:C) 1:1000:500 to 1000:1:1, alternatively 1:100:50 to 500:1:1,

5 alternatively 1:10:10 to 200:1:1, alternatively 1:1:1 to 100:1:50, and alternatively 1:1:10 to 75:1:50, and alternatively 5:1:1 to 50:1:50. The particular ratio chosen will depend on the exact pre-catalysts chosen, the method of activation, and the end product desired.

10 Additional preferred catalysts and process are described in US Patents 6,376,410 and 6,380,122, which are incorporated by reference herein.

In another embodiment the catalyst compositions of this invention include a support material or carrier. For example, the one or more catalyst components and/or one or more activators may be deposited on, contacted with, vaporized
15 with, bonded to, or incorporated within, adsorbed or absorbed in, or on, one or more supports or carriers.

The support material is any of the conventional support materials.
20 Preferably the supported material is a porous support material, for example, talc, inorganic oxides and inorganic chlorides. Other support materials include resinous support materials such as polystyrene, functionalized or crosslinked organic supports, such as polystyrene divinyl benzene polyolefins or polymeric compounds, zeolites, clays, or any other organic or inorganic support material and
25 the like, or mixtures thereof.

The preferred support materials are inorganic oxides that include those Group 2, 3, 4, 5, 13 or 14 metal oxides. The preferred supports include silica, which may or may not be dehydrated, fumed silica, alumina (WO 99/60033),
30 silica-alumina and mixtures thereof. Other useful supports include magnesia, titania, zirconia, magnesium chloride (U.S. Patent No. 5,965,477), montmorillonite (European Patent EP-B1 0 511 665), phyllosilicate, zeolites, talc, clays (U.S. Patent No. 6,034,187) and the like. Also, combinations of these support materials may be used, for example, silica-chromium, silica-alumina,
35 silica-titania and the like. Additional support materials may include those porous acrylic polymers described in EP 0 767 184 B1, which is incorporated herein by

5 reference. Other support materials include nanocomposites as described in PCT WO 99/47598, aerogels as described in WO 99/48605, spherulites as described in U.S. Patent No. 5,972,510 and polymeric beads as described in WO 99/50311, which are all herein incorporated by reference.

10 It is preferred that the support material, most preferably an inorganic oxide, has a surface area in the range of from about 10 to about 700 m²/g, pore volume in the range of from about 0.1 to about 4.0 cc/g and average particle size in the range of from about 5 to about 500 μm. More preferably, the surface area of the support material is in the range of from about 50 to about 500 m²/g, pore
15 volume of from about 0.5 to about 3.5 cc/g and average particle size of from about 10 to about 200 μm. Most preferably the surface area of the support material is in the range is from about 100 to about 400 m²/g, pore volume from about 0.8 to about 3.0 cc/g and average particle size is from about 5 to about 100 μm. The average pore size of the carrier useful in the invention typically has pore size in
20 the range of from 10 to 1000Å, preferably 50 to about 500Å, and most preferably 75 to about 350Å.

As is well known in the art, the catalysts may also be supported together on one inert support, or the catalysts may be independently placed on two inert
25 supports and subsequently mixed. Of the two methods, the former is preferred.

In another embodiment the support may comprise one or more types of support material which may be treated differently. For example one could use two different silicas that had different pore volumes or had been calcined at different
30 temperatures. Likewise one could use a silica that had been treated with a scavenger or other additive and a silica that had not.

The stereospecific catalysts may be used to prepare macromonomer having a Mw of 100,000 or less and a crystallinity of 30% or more preferably having vinyl
35 termini. As a specific example, a method for preparing propylene-based macromonomers having a high percentage of vinyl terminal bonds involves:

- 5 a) contacting, in solution, propylene, optionally a minor amount of copolymerizable monomer, with a catalyst composition containing the stereorigid, activated transition metal catalyst compound at a temperature from about 80°C. to about 140°C.; and
- b) recovering isotactic or syndiotactic polypropylene chains having number average
10 molecular weights of about 2,000 to about 30,000 Daltons.

Preferably, the solution comprises a hydrocarbon solvent. More preferably, the hydrocarbon solvent is aliphatic or aromatic. Also, the propylene monomers are preferably contacted at a temperature from 90°C. to 120°C. More preferably, a temperature from 95°C to 115°C is used. Most preferably, the propylene monomers
15 are contacted at a temperature from 100°C to 110°C. Reactor pressure generally can vary from atmospheric to 345 MPa, preferably to 182 MPa. The reactions can be run in batch or in continuous mode. Conditions for suitable slurry-type reactions will also be suitable and are similar to solution conditions, the polymerization typically being run in liquid propylene under pressures suitable to such.

20

The catalyst pair selection criteria were discussed earlier. One catalyst typically is stereospecific with the ability to produce significant population of vinyl-terminated macromonomers, the other typically is aspecific and capable of incorporating the reactive macromonomers. In general it is believed that C2
25 symmetric bulky ligand metallocene catalysts can produce vinyl terminated isotactic polypropylene macromonomers. Catalysts that favor betamethyl-elimination also often appear to also favor isotactic polypropylene macromonomer formation. Rac-dimethylsilyl bis(indenyl)hafnium dimethyl, dimethylsilyl bis(2-methyl-4-phenylindenyl) zirconium dichloride, and rac-ethylene bis(4,7-
30 dimethylindenyl) hafnium dimethyl are catalysts capable of producing isotactic polypropylene having high vinyl chain termination for use in this invention. High temperatures, typically above 80°C, appear to positively influence vinyl termination. Likewise, $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-c-C}_{12}\text{H}_{23})\text{TiMe}_2$ and $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-c-C}_{12}\text{H}_{23})\text{TiMe}_2$ produce amorphous polypropylene useful in this invention and are
35 believed to incorporate the vinyl terminated macromonomers to also produce a grafted structure of scPP side chains on an amorphous backbone.

5

In alternate embodiments dienes such as 1,9-decadiene are introduced into the reaction zone to promote the production of vinyl-terminated aPP and scPP macromonomers that help increase the population of branch-block species.

10 Polymerization Processes

The catalysts and catalyst systems described above are suitable for use in a solution, bulk, gas or slurry polymerization process or a combination thereof, preferably solution phase or bulk phase polymerization process.

15

In one embodiment, this invention is directed toward the solution, bulk, slurry or gas phase polymerization reactions involving the polymerization of one or more of monomers having from 3 to 30 carbon atoms, preferably 3-12 carbon atoms, and more preferably 3 to 8 carbon atoms. Preferred monomers include one or more of propylene, butene-1, pentene-1, 4-methyl-pentene-1, hexene-1, octene-1, decene-1, 3-methyl-pentene-1, and cyclic olefins or a combination thereof. Other monomers can include vinyl monomers, diolefins such as dienes, polyenes, norbornene, norbornadiene, vinyl norbornene, ethylidene norbornene monomers. Preferably a homopolymer or copolymer of propylene is produced. In another embodiment, both a homopolymer of propylene and a copolymer of propylene and one or more of the monomers listed above are produced.

20

25

One or more reactors in series or in parallel may be used in the present invention. Catalyst component and activator may be delivered as a solution or slurry, either separately to the reactor, activated in-line just prior to the reactor, or preactivated and pumped as an activated solution or slurry to the reactor. A preferred operation is two solutions activated in-line. For more information on methods to introduce multiple catalysts into reactors, please see US 6,399,722, and WO0130862A1. While these references may emphasize gas phase reactors, the techniques described are equally applicable to other types of reactors, including continuous stirred tank reactors, slurry loop reactors and the like.

30

35

- 5 Polymerizations are carried out in either single reactor operation, in which monomer, comonomers, catalyst/activator, scavenger, and optional modifiers are added continuously to a single reactor or in series reactor operation, in which the above components are added to each of two or more reactors connected in series. The catalyst components can be added to the first reactor in the series. The
- 10 catalyst component may also be added to both reactors, with one component being added to first reaction and another component to other reactors.

- In one embodiment 500 ppm or less of hydrogen is added to the polymerization, or 400ppm or less, or 300 ppm or less. In other embodiments at
- 15 least 50 ppm of hydrogen is added to the polymerization, or 100 ppm or more, or 150 ppm or more.

Gas phase polymerization

- 20 Generally, in a fluidized gas bed process used for producing polymers, a gaseous stream containing one or more monomers is continuously cycled through a fluidized bed in the presence of a catalyst under reactive conditions. The gaseous stream is withdrawn from the fluidized bed and recycled back into the reactor. Simultaneously, polymer product is withdrawn from the reactor and fresh
- 25 monomer is added to replace the polymerized monomer. (See for example U.S. Patent Nos. 4,543,399, 4,588,790, 5,028,670, 5,317,036, 5,352,749, 5,405,922, 5,436,304, 5,453,471, 5,462,999, 5,616,661 and 5,668,228 all of which are fully incorporated herein by reference.)

30 Slurry phase polymerization

- A slurry polymerization process generally operates between 1 to about 50 atmosphere pressure range (15 psi to 735 psi, 103 kPa to 5068 kPa) or even greater and temperatures in the range of 0°C to about 120°C. In a slurry
- 35 polymerization, a suspension of solid, particulate polymer is formed in a liquid polymerization diluent medium to which monomer and comonomers along with

5 catalyst are added. The suspension including diluent is intermittently or continuously removed from the reactor where the volatile components are separated from the polymer and recycled, optionally after a distillation, to the reactor. The liquid diluent employed in the polymerization medium is typically an alkane having from 3 to 7 carbon atoms, preferably a branched alkane. The
10 medium employed should be liquid under the conditions of polymerization and relatively inert. When a propane medium is used the process must be operated above the reaction diluent critical temperature and pressure. Preferably, a hexane or an isobutane medium is employed.

15 In one embodiment, a preferred polymerization technique useful in the invention is referred to as a particle form polymerization, or a slurry process where the temperature is kept below the temperature at which the polymer goes into solution. Such technique is well known in the art, and described in for instance U.S. Patent No. 3,248,179 which is fully incorporated herein by
20 reference. The preferred temperature in the particle form process is within the range of about 85°C to about 110°C. Two preferred polymerization methods for the slurry process are those employing a loop reactor and those utilizing a plurality of stirred reactors in series, parallel, or combinations thereof. Non-limiting examples of slurry processes include continuous loop or stirred tank
25 processes. Also, other examples of slurry processes are described in U.S. Patent No. 4,613,484, which is herein fully incorporated by reference.

In another embodiment, the slurry process is carried out continuously in a loop reactor. The catalyst, as a slurry in isobutane or as a dry free flowing
30 powder, is injected regularly to the reactor loop, which is itself filled with circulating slurry of growing polymer particles in a diluent of isobutane containing monomer and comonomer. Hydrogen, optionally, may be added as a molecular weight control. (In one embodiment 500 ppm or less of hydrogen is added, or 400 ppm or less or 300 ppm or less. In other embodiments at least 50
35 ppm of hydrogen is added, or 100 ppm or more, or 150 ppm or more.)

5 The reactor is maintained at a pressure of 3620 kPa to 4309 kPa and at a temperature in the range of about 60 °C to about 104 °C depending on the desired polymer melting characteristics. Reaction heat is removed through the loop wall since much of the reactor is in the form of a double-jacketed pipe. The slurry is allowed to exit the reactor at regular intervals or continuously to a heated low
10 pressure flash vessel, rotary dryer and a nitrogen purge column in sequence for removal of the isobutane diluent and all unreacted monomer and comonomers. The resulting hydrocarbon free powder is then compounded for use in various applications.

15 In another embodiment, the reactor used in the slurry process useful in the invention is capable of and the process useful in the invention is producing greater than 2000 lbs of polymer per hour (907 Kg/hr), more preferably greater than 5000 lbs/hr (2268 Kg/hr), and most preferably greater than 10,000 lbs/hr (4540 Kg/hr). In another embodiment the slurry reactor used in the process useful in the
20 invention is producing greater than 15,000 lbs of polymer per hour (6804 Kg/hr), preferably greater than 25,000 lbs/hr (11,340 Kg/hr) to about 100,000 lbs/hr (45,500 Kg/hr).

 In another embodiment in the slurry process useful in the invention the
25 total reactor pressure is in the range of from 400 psig (2758 kPa) to 800 psig (5516 kPa), preferably 450 psig (3103 kPa) to about 700 psig (4827 kPa), more preferably 500 psig (3448 kPa) to about 650 psig (4482 kPa), most preferably from about 525 psig (3620 kPa) to 625 psig (4309 kPa).

30 In yet another embodiment in the slurry process useful in the invention the concentration of predominant monomer in the reactor liquid medium is in the range of from about 1 to 10 weight percent, preferably from about 2 to about 7 weight percent, more preferably from about 2.5 to about 6 weight percent, most preferably from about 3 to about 6 weight percent.

35

5 Another process useful in the invention is where the process, preferably a slurry process is operated in the absence of or essentially free of any scavengers, such as triethylaluminum, trimethylaluminum, tri-isobutylaluminum and tri-n-hexylaluminum and diethyl aluminum chloride, dibutyl zinc and the like. This process is described in PCT publication WO 96/08520 and U.S. Patent No.
10 5,712,352, which are herein fully incorporated by reference.

In another embodiment the process is run with scavengers. Typical scavengers include trimethyl aluminum, tri-isobutyl aluminum, tri-n-octylaluminum and an excess of alumoxane or modified alumoxane.

15

Homogeneous, bulk, or solution phase polymerization

The catalysts described herein can be used advantageously in homogeneous solution processes. Generally this involves polymerization in a
20 continuous reactor in which the polymer formed and the starting monomer and catalyst materials supplied, are agitated to reduce or avoid concentration gradients. Suitable processes operate above the melting point of the polymers at high pressures, from 1 to 3000 bar (10-30,000 MPa), in which the monomer acts as diluent or in solution polymerization using a solvent.

25

Temperature control in the reactor is obtained by balancing the heat of polymerization with reactor cooling by reactor jackets or cooling coils to cool the contents of the reactor, auto refrigeration, pre-chilled feeds, vaporization of liquid medium (diluent, monomers or solvent) or combinations of all three. Adiabatic
30 reactors with pre-chilled feeds may also be used. The reactor temperature depends on the catalyst used. In general, the reactor temperature preferably can vary between about 30°C and about 160°C, more preferably from about 90°C to about 150°C, and most preferably from about 100°C to about 140°C. Polymerization temperature may vary depending on catalyst choice. For example
35 a diimine Ni catalyst may be used at 40°C, while a metallocene Ti catalyst can be used at 100°C or more. In series operation, the second reactor temperature is

5 preferably higher than the first reactor temperature. In parallel reactor operation, the temperatures of the two reactors are independent. The pressure can vary from about 1 mm Hg to 2500 bar (25,000 MPa), preferably from 0.1 bar to 1600 bar (1-16,000 MPa), most preferably from 1.0 to 500 bar (10-5000MPa).

10 In one embodiment 500 ppm or less of hydrogen is added to the polymerization, or 400 ppm or less or 300 ppm or less. In other embodiments at least 50 ppm of hydrogen is added to the polymerization, or 100 ppm or more, or 150 ppm or more.

15 Each of these processes may also be employed in single reactor, parallel or series reactor configurations. The liquid processes comprise contacting olefin monomers with the above described catalyst system in a suitable diluent or solvent and allowing said monomers to react for a sufficient time to produce the desired polymers. Hydrocarbon solvents are suitable, both aliphatic and aromatic.
20 Alkanes, such as hexane, pentane, isopentane, and octane, are preferred.

The process can be carried out in a continuous stirred tank reactor, batch reactor or plug flow reactor, or more than one reactor operated in series or parallel. These reactors may have or may not have internal cooling or heating and
25 the monomer feed may or may not be refrigerated. See the general disclosure of U.S. patent 5,001,205 for general process conditions. See also, international application WO 96/33227 and WO 97/22639. All documents are incorporated by reference for US purposes for description of polymerization processes, metallocene selection and useful scavenging compounds.

30

This invention further relates to a continuous process to prepare an adhesive comprising:

- 1) combining monomer, optional solvent, catalyst and activator in a reactor system,
- 35 2) withdrawing polymer solution from the reactor system,
- 3) removing at least 10% solvent, if present, from the polymer

5 solution,

- 4) quenching the reaction,
 - 5) devolatilizing the polymer solution to form molten polymer,
 - 6) combining the molten polymer and one or more additives (such as those described below) in a static mixer (in a preferred embodiment tackifier is not
 - 10 added or is added in amounts of less than 30 weight%, preferably less than 20 weight%, more preferably in amounts of less than 10 weight%),
 - 7) removing the polymer combination from the static mixer, and
 - 8) pelletizing or drumming the polymer combination;
- where step 1) comprises any of the processes described above.

15

In another embodiment this invention relates to a continuous process to prepare an adhesive comprising:

- 1) combining monomer, optional solvent, catalyst and activator in a reactor system,
- 20 2) withdrawing polymer solution from the reactor system,
- 3) removing at least 10% solvent, if present, from the polymer solution,
- 4) quenching the reaction,
- 5) devolatilizing the polymer solution to form molten polymer,
- 25 6) combining the molten polymer and one or more additives in a static mixer,
- 7) removing the polymer combination from the static mixer, and
- 8) pelletizing or drumming the polymer combination.

30 In a particularly preferred embodiment, this invention relates to a continuous process to make an adhesive comprising

- 1) selecting a first catalyst component capable of producing a polymer having an Mw of 100,000 or less and a crystallinity of 20% (preferably 5 % or less) or less under selected polymerization conditions;

- 5 2) selecting a second catalyst component capable of producing polymer having an Mw of 100,000 or less and a crystallinity of 20% or more, (preferably 40% or more) at the selected polymerization conditions;
- 3) contacting, in a solvent and in a reaction zone under the selected polymerization conditions, the catalyst components in the presence of one or more
10 activators with one or more C3 to C40 olefins, and, optionally one or more diolefins;
- 4) at a temperature of greater than 70°C (preferably greater than 100 °C);
- 5) at a residence time of 120 minutes or less, (preferably 60 minutes
15 or less, more preferably 30 minutes or less);
- 6) wherein the ratio of the first catalyst to the second catalyst is from 1:1 to 50:1 (preferably 30:1);
- 7) wherein the activity of the catalyst components is at least 50 kilograms of polymer per gram of the catalyst components; and wherein at least
20 20% of the olefins are converted to polymer;
- 8) withdrawing polymer solution from the reaction zone;
- 9) removing at least 10% solvent from the polymer solution;
- 10) quenching the reaction;
- 11) devolatilizing the polymer solution to form molten polymer;
- 25 12) combining the molten polymer and one or more additives in a static mixer;
- 13) removing the polymer combination from the static mixer; and
- 14) pelletizing or drumming the polymer combination.

30 In a particularly preferred embodiment, this invention relates to a continuous process to make an adhesive comprising

- 1) selecting a first catalyst component capable of producing a polymer having an Mw of 100,000 or less and a crystallinity of 20% or less (preferably 5% or less) under selected polymerization conditions;

- 5 2) selecting a second catalyst component capable of producing polymer having an Mw of 100,000 or less and a crystallinity of 20% or more (preferably 40% or more) at the selected polymerization conditions;
- 3) contacting, in a solvent and in a reaction zone under the selected polymerization conditions, the catalyst components in the presence of one or more
10 activators with one or more C3 to C40 olefins, and, optionally one or more diolefins;
- 4) at a temperature of greater than 70°C, (preferably greater than 100°C);
- 5) at a residence time of 30 minutes or less;
- 15 6) wherein the ratio of the first catalyst to the second catalyst is from 1:1 to 50:1 (Preferably 30:1);
- 7) wherein the activity of the catalyst components is at least 50 kilograms of polymer per gram of the catalyst components; and wherein at least 50% of the olefins are converted to polymer;
- 20 8) withdrawing polymer solution from the reaction zone;
- 9) removing at least 10% solvent from the polymer solution;
- 10) quenching the reaction;
- 11) forming molten polymer
 where the polymer comprises one or more C3 to C40 olefins, and
25 less than 50 mole % of ethylene, and where the polymer has:
- a) a Dot T-Peel of 1 Newton or more; and
- b) a branching index (g') of 0.95 or less measured at the Mz of the polymer; and
- c) an Mw of 100,000 or less; and
- 30 12) combining the molten polymer and one or more additives in a static mixer;
- 13) removing the polymer combination from the static mixer; and
- 14) pelletizing or drumming the polymer combination.

5 In a particularly preferred embodiment, this invention relates to a continuous process to make an adhesive comprising

- 1) selecting a first catalyst component capable of producing a polymer having an Mw of 100,000 or less and a crystallinity of 20% or less under selected polymerization conditions;
- 10 2) selecting a second catalyst component capable of producing polymer having an Mw of 100,000 or less and a crystallinity of 40% or more at the selected polymerization conditions;
- 3) contacting, in a solvent and in a reaction zone under the selected polymerization conditions, the catalyst components in the presence of one or more
15 activators with one or more C3 to C40 olefins, and, optionally one or more diolefins;
- 4) at a temperature of greater than 100°C;
- 5) at a residence time of 30 minutes or less;
- 6) wherein the ratio of the first catalyst to the second catalyst is from
20 1:1 to 30:1;
- 7) wherein the activity of the catalyst components is at least 50 kilograms of polymer per gram of the catalyst components; and wherein at least 50% of the olefins are converted to polymer;
- 8) withdrawing polymer solution from the reaction zone;
- 25 9) removing at least 10% solvent from the polymer solution;
- 10) quenching the reaction;
- 11) forming molten polymer
 where the polymer comprises one or more C3 to C40 olefins (preferably propylene), and less than 50 mole % of
30 ethylene, and where the polymer has:
 - a) a Dot T-Peel of 3 Newton or more; and
 - b) a branching index (g') of 0.90 or less measured at the Mw of the polymer; and
 - c) an Mw of 30,000 or less;
 - 35 d) a peak melting point between 60 and 190°C,
 - e) a Heat of fusion of 1 to 70 J/g,

- 5 f) a melt viscosity of 8000 mPa·sec or less at 190°C;
and
- 12) combining the molten polymer and one or more additives in a static mixer;
- 13) removing the polymer combination from the static mixer; and
- 10 14) pelletizing or drumming the polymer combination.

In another embodiment this invention relates to a continuous process to prepare an adhesive comprising:

- 1) combining monomer, catalyst and activator in a reactor system,
- 15 2) withdrawing polymer from the reactor system,
- 3) quenching the reaction,
- 4) forming molten polymer,
- 5) combining the molten polymer and one or more additives, and
- 6) pelletizing or drumming the polymer combination.

20

Formulations of the polymers

The polymers produced herein then can be used directly as an adhesive or blended with other components to form an adhesive.

25

Tackifiers are typically not needed with the polymers of this invention. However if tackifier is desired, the tackifiers that may be blended with the polymers described above are those typically used in the art. Examples include, but are not limited to, aliphatic hydrocarbon resins, aromatic modified aliphatic hydrocarbon resins, hydrogenated polycyclopentadiene resins,

30 polycyclopentadiene resins, gum rosins, gum rosin esters, wood rosins, wood rosin esters, tall oil rosins, tall oil rosin esters, polyterpenes, aromatic modified polyterpenes, terpene phenolics, aromatic modified hydrogenated polycyclopentadiene resins, hydrogenated aliphatic resin, hydrogenated aliphatic

35 aromatic resins, hydrogenated terpenes and modified terpenes, and hydrogenated

5 rosin esters. In some embodiments the tackifier is hydrogenated. In other
embodiments the tackifier is non-polar. (Non-polar meaning that the tackifier is
substantially free of monomers having polar groups. Preferably the polar groups
are not present, however if they are preferably they are not present at more than 5
weight %, preferably not more than 2 weight %, even more preferably no more
10 than 0.5 weight %.) In some embodiments the tackifier has a softening point
(Ring and Ball, as measured by ASTM E-28) of 80 °C to 150 °C, preferably 100
°C to 130 °C.

The tackifier, if present, is typically present at about 1 weight % to about
15 80 weight %, based upon the weight of the blend, more preferably 2 weight % to
40 weight %, even more preferably 3 weight % to 30 weight %.

Preferred hydrocarbon resins for use as tackifiers or modifiers include:

1. Resins such as C5/C6 terpene resins, styrene terpenes, alpha-
20 methyl styrene terpene resins, C9 terpene resins, aromatic modified C5/C6,
aromatic modified cyclic resins, aromatic modified dicyclopentadiene based resins
or mixtures thereof Additional preferred resins include those described in WO
91/07472, US 5,571,867, US 5,171,793 and US 4,078,132. Typically these resins
are obtained from the cationic polymerization of compositions containing one or
25 more of the following monomers: C5 diolefins (such as 1-3 pentadiene, isoprene,
etc.); C5 olefins (such as 2- methylbutenes, cyclopentene, etc.); C6 olefins (such as
hexene), C9 vinylaromatics (such as styrene, alpha methyl styrene, vinyltoluene,
indene, methyl indene, etc.); cyclics (such as dicyclopentadiene,
methyldicyclopentadiene, etc.); and or terpenes (such as limonene, carene, etc).
- 30 2. Resins obtained by the thermal polymerization of
dicyclopentadiene, and/or the thermal polymerization of dimers or oligomers of
cyclopentadiene and /or methylcyclopentadiene, optionally with vinylaromatics
(such as styrene, alpha-methyl styrene, vinyl toluene, indene, methyl indene).
The resins obtained after polymerization and separation of unreacted materials,
35 can be hydrogenated if desired. Examples of preferred resins include those

- 5 described in US 4,078,132; WO 91/07472; US 4,994,516; EP 0 046 344 A; EP 0 082 726 A; and US 5,171,793.

In another embodiment an adhesive composition comprising polymer product of this invention further comprises a crosslinking agent. Preferred
10 crosslinking agents include those having functional groups that can react with the acid or anhydride group. Preferred crosslinking agents include alcohols, multiols, amines, diamines and/or triamines. Examples of crosslinking agents useful in this invention include polyamines such as ethylenediamine, diethylenetriamine, hexamethylenediamine, diethylaniinopropylamine, and/or menthanediamine.

15

In another embodiment an adhesive composition comprising the polymer product of this invention further comprises typical additives known in the art such as fillers, antioxidants, adjuvants, adhesion promoters, oils, and/or plasticizers. Preferred fillers include titanium dioxide, calcium carbonate, barium sulfate,
20 silica, silicon dioxide, carbon black, sand, glass beads, mineral aggregates, talc, clay and the like. Preferred antioxidants include phenolic antioxidants, such as Irganox 1010, Irganox1076 both available from Ciba-Geigy. Preferred oils include paraffinic or naphthenic oils such as Primol 3 52, or Primol 876 available from ExxonMobil Chemical France, S.A. in Paris, France. Preferred plasticizers include
25 polybutenes, such as Parapol 950 and Parapol 1300 available from ExxonMobil Chemical Company in Houston Texas. Other preferred additives include block, antiblock, pigments, processing aids, UV stabilizers, neutralizers, lubricants, surfactants and/or nucleating agents may also be present in one or more than one layer in the films. Preferred additives include silicon dioxide, titanium dioxide,
30 polydimethylsiloxane, talc, dyes, wax, calcium stearate, carbon black, low molecular weight resins and glass beads. Preferred adhesion promoters include polar acids, polyaminoamides (such as Versamid 115, 125, 140, available from Henkel), urethanes (such as isocyanate/hydroxy terminated polyester systems, e.g. bonding agent TN/Mondur Cb-75(Miles, Inc.), coupling agents, (such as silane esters (Z-6020 from Dow Corning)), titanate esters (such as Kr-44 available from
35 Kenrich), reactive acrylate monomers (such as sarbox SB-600 from Sartomer),

- 5 metal acid salts (such as Saret 633 from Sartomer), polyphenylene oxide, oxidized polyolefins, acid modified polyolefins, and anhydride modified polyolefins.

In another embodiment the polymers of this invention are combined with less than 3 wt % anti-oxidant, less than 3 wt % flow improver, less than 10 wt %
10 wax, and or less than 3 wt % crystallization aid.

Other optional components that may be combined with the polymer product of this invention are plasticizers or other additives such as oils, surfactants, fillers, color masterbatches, and the like. Preferred plasticizers
15 include mineral oils, polybutenes, phthalates and the like. Particularly preferred plasticizers include phthalates such as diisoundecyl phthalate (DIUP), diisononylphthalate (DINP), dioctylphthalates (DOP) and the like. Particularly preferred oils include aliphatic naphthenic oils.

20 Other optional components that may be combined with the polymer product of this invention are low molecular weight products such as wax, oil or low Mn polymer, (low meaning below Mn of 5000, preferably below 4000, more preferably below 3000, even more preferably below 2500). Preferred waxes include polar or non-polar waxes, functionalized waxes, polypropylene waxes,
25 polyethylene waxes, and wax modifiers. Preferred waxes include ESCOMER™ 101. Preferred functionalized waxes include those modified with an alcohol, an acid, a ketone, an anhydride and the like. Preferred examples include waxes modified by methyl ketone, maleic anhydride or maleic acid. Preferred oils include aliphatic naphthenic oils, white oils or the like. Preferred low Mn polymers
30 include polymers of lower alpha olefins such as propylene, butene, pentene, hexene and the like. A particularly preferred polymer includes polybutene having an Mn of less than 1000. An example of such a polymer is available under the trade name PARAPOL™ 950 from ExxonMobil Chemical Company.
PARAPOL™ 950 is a liquid polybutene polymer having an Mn of 950 and a
35 kinematic viscosity of 220cSt at 100 °C, as measured by ASTM D 445. In some

- 5 embodiments the polar and non-polar waxes are used together in the same composition.

 In some embodiments, however, wax may not be desired and is present at less than 5 weight % , preferably less than 3 weight % , more preferably less than 1
10 weight % , more preferably less than 0.5 weight % , based upon the weight of the composition.

 In another embodiment the polymers of this invention have less than 30 weight% total of any combination of additives described above, preferably less
15 than 25 weight%, preferably less than 20 weight % , preferably less than 15 weight %, preferably less than 10 weight%, preferably less than 5 weight%, based upon the weight of the polymer and the additives.

 In another embodiment the polymer produced by this invention may be
20 blended with elastomers (preferred elastomers include all natural and synthetic rubbers, including those defined in ASTM D1566). In a preferred embodiment elastomers are blended with the polymer produced by this invention to form rubber toughened compositions. In a particularly preferred embodiment the rubber toughened composition is a two (or more) phase system where the rubber
25 is a discontinuous phase and the polymer is a continuous phase. Examples of preferred elastomers include one or more of the following: ethylene propylene rubber, ethylene propylene diene monomer rubber, neoprene rubber, styrenic block copolymer rubbers (including SI, SIS, SB, SBS, SIBS and the like), butyl rubber, halobutyl rubber, copolymers of isobutylene and para-alkylstyrene,
30 halogenated copolymers of isobutylene and para-alkylstyrene. This blend may be combined with the tackifiers and/or other additives as described above.

 In another embodiment the polymer produced by this invention may be blended with impact copolymers. Impact copolymers are defined to be a blend of
35 isotactic PP and an elastomer such as an ethylene-propylene rubber. In a

- 5 preferred embodiment the blend is a two (or more) phase system where the impact copolymer is a discontinuous phase and the polymer is a continuous phase.

In another embodiment the polymer produced by this invention may be blended with ester polymers. In a preferred embodiment the blend is a two (or
10 more) phase system where the polyester is a discontinuous phase and the polymer is a continuous phase.

In a preferred embodiment the polymers of the invention described above are combined with metallocene polyethylenes (mPE's) or metallocene
15 polypropylenes (mPP's). The mPE and mPP homopolymers or copolymers are typically produced using mono- or bis-cyclopentadienyl transition metal catalysts in combination with an activator of alumoxane and/or a non-coordinating anion in solution, slurry, high pressure or gas phase. The catalyst and activator may be supported or unsupported and the cyclopentadienyl rings by may substituted or
20 unsubstituted. Several commercial products produced with such catalyst/activator combinations are commercially available from ExxonMobil Chemical Company in Baytown, Texas. under the tradenames EXCEED™, ACHIEVE™ and EXACT™. For more information on the methods and catalysts/activators to produce such mPE homopolymers and copolymers see WO 94/26816; WO
25 94/03506; EPA 277,003; EPA 277,004; U.S. Pat. No. 5,153,157; U.S. Pat. No. 5,198,401; U.S. Pat. No. 5,240,894; U.S. Pat. No. 5,017,714; CA 1,268,753; U.S. Pat. No. 5,324,800; EPA 129,368; U.S. Pat. No. 5,264,405; EPA 520,732; WO 92 00333; U.S. Pat. No. 5,096,867; U.S. Pat. No. 5,507,475; EPA 426 637; EPA 573 403; EPA 520 732; EPA 495 375; EPA 500 944; EPA 570 982; WO91/09882;
30 WO94/03506 and U.S. Pat. No. 5,055,438.

In another embodiment the olefin polymer of this invention, preferably the polypropylene homopolymer or copolymer of this invention, can be blended with another homopolymer and/or copolymer, including but not limited to,
35 homopolypropylene, propylene copolymerized with up to 50 weight % of ethylene or a C4 to C20 alpha.-olefin, isotactic polypropylene, highly isotactic

5 polypropylene, syndiotactic polypropylene, random copolymer of propylene and ethylene and/or butene and/or hexene, polybutene, ethylene vinyl acetate, low density polyethylene (density 0.915 to less than 0.935 g/cm³) linear low density polyethylene, ultra low density polyethylene (density 0.86 to less than 0.90 g/cm³), very low density polyethylene (density 0.90 to less than 0.915 g/cm³),
10 medium density polyethylene (density 0.935 to less than 0.945 g/cm³), high density polyethylene (density 0.945 to 0.98 g/cm³), ethylene vinyl acetate, ethylene methyl acrylate, copolymers of acrylic acid, polymethylmethacrylate or any other polymers polymerizable by a high-pressure free radical process, polyvinylchloride, polybutene-1, isotactic polybutene, ABS resins, elastomers
15 such as ethylene-propylene rubber (EPR), vulcanized EPR, EPDM, block copolymer elastomers such as SBS, nylons (polyamides), polycarbonates, PET resins, crosslinked polyethylene, copolymers of ethylene and vinyl alcohol (EVOH), polymers of aromatic monomers such as polystyrene, poly-1 esters, high molecular weight polyethylene having a density of 0.94 to 0.98 g/cm³ low
20 molecular weight polyethylene having a density of 0.94 to 0.98 g/cm³, graft copolymers generally, polyacrylonitrile homopolymer or copolymers, thermoplastic polyamides, polyacetal, polyvinylidene fluoride and other fluorinated elastomers, polyethylene glycols and polyisobutylene.

25 In a preferred embodiment the olefin polymer of this invention, preferably the polypropylene polymer of this invention, is present in the blend at from 10 to 99 weight %, based upon the weight of the polymers in the blend, preferably 20 to 95 weight %, even more preferably at least 30 to 90 weight %, even more preferably at least 40 to 90 weight %, even more preferably at least 50 to 90
30 weight %, even more preferably at least 60 to 90 weight %, even more preferably at least 70 to 90 weight %.

The blends described above may be produced by mixing the two or more polymers together, by connecting reactors together in series to make reactor
35 blends or by using more than one catalyst in the same reactor to produce multiple

- 5 species of polymer. The polymers can be mixed together prior to being put into the extruder or may be mixed in an extruder.

Any of the above polymers, including the polymers produced by this invention, may be functionalized. Preferred functional groups include maleic acid
10 and maleic anhydride. By functionalized is meant that the polymer has been contacted with an unsaturated acid or anhydride. Preferred unsaturated acids or anhydrides include any unsaturated organic compound containing at least one double bond and at least one carbonyl group. Representative acids include carboxylic acids, anhydrides, esters and their salts, both metallic and non-metallic.
15 Preferably the organic compound contains an ethylenic unsaturation conjugated with a carbonyl group ($-C=O$). Examples include maleic, fumaric, acrylic, methacrylic, itaconic, crotonic, alpha.methyl crotonic, and cinnamic acids as well as their anhydrides, esters and salt derivatives. Maleic anhydride is particularly preferred. The unsaturated acid or anhydride is preferably present at about 0.1
20 weight % to about 10 weight %, preferably at about 0.5 weight % to about 7 weight %, even more preferably at about 1 to about 4 weight %, based upon the weight of the hydrocarbon resin and the unsaturated acid or anhydride.

In a preferred embodiment the unsaturated acid or anhydried comprises a
25 carboxylic acid or a derivative thereof selected from the group consisting of unsaturated carboxylic acids, unsaturated carboxylic acid derivatives selected from esters, imides, amides, anhydrides and cyclic acid anhydrides or mixtures thereof.

30 Applications

The polymer product of this invention or formulations thereof may then be applied directly to a substrate or may be sprayed thereon, typically the polymer is molten. Spraying is defined to include atomizing, such as producing an even dot
35 pattern, spiral spraying such as Nordson Controlled Fiberization or oscillating a stretched filament like is done in the ITW Dynafiber/Omega heads or Summit

5 technology from Nordson, as well as melt blown techniques. Melt blown techniques are defined to include the methods described in U.S. patent 5,145,689 or any process where air streams are used to break up filaments of the extrudate and then used to deposit the broken filaments on a substrate. In general, melt blown techniques are processes that use air to spin hot melt adhesive fibers and
10 convey them onto a substrate for bonding. Fibers sizes can easily be controlled from 20-200 microns by changing the melt to air ratio. Few, preferably no, stray fibers are generated due to the inherent stability of adhesive melt blown applicators. Under UV light the bonding appears as a regular, smooth, stretched dot pattern. Atomization is a process that uses air to atomize hot melt adhesive
15 into very small dots and convey them onto a substrate for bonding.

Lamination melt coating

The adhesives of this invention can be used in any adhesive application,
20 including but not limited to, disposables, packaging, laminates, pressure sensitive adhesives, tapes labels, wood binding, paper binding, non-wovens, road marking, reflective coatings, and the like.

In a preferred embodiment the adhesives of this invention can be used for
25 disposable diaper and napkin chassis construction, elastic attachment in disposable goods converting, packaging, labeling, bookbinding, woodworking, and other assembly applications. Particularly preferred applications include: baby diaper leg elastic, diaper frontal tape, diaper standing leg cuff, diaper chassis construction, diaper core stabilization, diaper liquid transfer layer, diaper outer cover
30 lamination, diaper elastic cuff lamination, feminine napkin core stabilization, feminine napkin adhesive strip, industrial filtration bonding, industrial filter material lamination, filter mask lamination, surgical gown lamination, surgical drape lamination, and perishable products packaging.

35 The adhesives described above may be applied to any substrate. Preferred substrates include wood, paper, cardboard, plastic, thermoplastic, rubber, metal,

- 5 metal foil (such as aluminum foil and tin foil), metallized surfaces, cloth, non-wovens (particularly polypropylene spun bonded fibers or non-wovens), spunbonded fibers, cardboard, stone, plaster, glass (including silicon oxide (SiO_x) coatings applied by evaporating silicon oxide onto a film surface), foam, rock, ceramics, films, polymer foams (such as polyurethane foam), substrates
10 coated with inks, dyes, pigments, PVDC and the like or combinations thereof.

Additional preferred substrates include polyethylene, polypropylene, polyacrylates, acrylics, polyethylene terephthalate, or any of the polymers listed above as suitable for blends.

15

Any of the above substrates, and/or the polymers of this invention, may be corona discharge treated, flame treated, electron beam irradiated, gamma irradiated, microwaved, or silanized.

20

The adhesives produced herein, when coated in some fashion between two adherends, preferably perform such that the materials are held together in a sufficient fashion compared to a standard specification or a standard adhesive similarly constructed.

25

The polymer product of this invention may be used in any adhesive application described in WO 97/33921 in combination with the polymers described therein or in place of the polymers described therein.

30

The polymer product of this invention, alone or in combination with other polymers and or additives, may also be used to form hook and loop fasteners as described in WO 02/35956.

Characterization and Tests

5 Molecular weights (number average molecular weight (M_n), weight
average molecular weight (M_w), and z-average molecular weight (M_z)) were
determined using a Waters 150 Size Exclusion Chromatograph (*SEC*) equipped
with a differential refractive index detector (*DRI*), an online low angle light
scattering (*LALLS*) detector and a viscometer (*VIS*). The details of the detector
10 calibrations have been described elsewhere [Reference: T. Sun, P. Brant, R. R.
Chance, and W. W. Graessley, *Macromolecules*, Volume 34, Number 19, 6812-
6820, (2001)]; attached below are brief descriptions of the components.

The *SEC* with three Polymer Laboratories PLgel 10mm Mixed-B columns,
a nominal flow rate $0.5 \text{ cm}^3/\text{min}$, and a nominal injection volume 300 *microliters*
15 was common to both detector configurations. The various transfer lines, columns
and differential refractometer (the *DRI* detector, used mainly to determine eluting
solution concentrations) were contained in an oven maintained at 135°C .

The *LALLS* detector was the model 2040 dual-angle light scattering
photometer (Precision Detector Inc.). Its flow cell, located in the *SEC* oven, uses
20 a 690 *nm* diode laser light source and collects scattered light at two angles, 15°
and 90° . Only the 15° output was used in these experiments. Its signal was sent
to a data acquisition board (National Instruments) that accumulates readings at a
rate of 16 per second. The lowest four readings were averaged, and then a
proportional signal was sent to the *SEC-LALLS-VIS* computer. The *LALLS*
25 detector was placed after the *SEC* columns, but before the viscometer.

The viscometer was a high temperature Model 150R (Viscotek
Corporation). It consisted of four capillaries arranged in a Wheatstone bridge
configuration with two pressure transducers. One transducer measured the total
pressure drop across the detector, and the other, positioned between the two sides
30 of the bridge, measured a differential pressure. The specific viscosity for the
solution flowing through the viscometer was calculated from their outputs. The
viscometer was inside the *SEC* oven, positioned after the *LALLS* detector but
before the *DRI* detector.

5 Solvent for the *SEC* experiment was prepared by adding 6 grams of butylated hydroxy toluene (BHT) as an antioxidant to a 4 liter bottle of 1,2,4 Trichlorobenzene (TCB) (Aldrich Reagent grade) and waiting for the BHT to solubilize. The TCB mixture was then filtered through a 0.7 micron glass pre-filter and subsequently through a 0.1 micron Teflon filter. There was an
10 additional online 0.7 micron glass pre-filter/0.22 micron Teflon filter assembly between the high pressure pump and *SEC* columns. The TCB was then degassed with an online degasser (Phenomenex, Model DG-4000) before entering the *SEC*.

Polymer solutions were prepared by placing dry polymer in a glass container, adding the desired amount of TCB, then heating the mixture at 160 °C
15 with continuous agitation for about 2 hours. All quantities were measured gravimetrically. The TCB densities used to express the polymer concentration in mass/volume units were 1.463 g/ml at room temperature and 1.324 g/ml at 135 °C. The injection concentration ranged from 1.0 to 2.0 mg/ml, with lower concentrations being used for higher molecular weight samples.

20 Prior to running each sample the DRI detector and the injector were purged. Flow rate in the apparatus was then increased to 0.5 ml/minute, and the DRI was allowed to stabilize for 8-9 hours before injecting the first sample. The argon ion laser was turned on 1 to 1.5 hours before running samples by running the laser in idle mode for 20-30 minutes and then switching to full power in light
25 regulation mode.

The branching index was measured using *SEC* with an on-line viscometer (*SEC-VIS*) and are reported as g' at each molecular weight in the *SEC* trace. The branching index g' is defined as:

$$g' = \frac{\eta_b}{\eta_l}$$

30 where η_b is the intrinsic viscosity of the branched polymer and η_l is the intrinsic viscosity of a linear polymer of the same viscosity-averaged molecular weight (M_v) as the branched polymer. $\eta_l = KM_v^\alpha$, K and α were measured values for

5 linear polymers and should be obtained on the same SEC-DRI-LS-VIS instrument as the one used for branching index measurement. For polypropylene samples presented in this invention, $K=0.0002288$ and $\alpha=0.705$ were used. The SEC-DRI-LS-VIS method obviates the need to correct for polydispersities, since the intrinsic viscosity and the molecular weight were measured at individual elution volumes, 10 which arguably contain narrowly dispersed polymer. Linear polymers selected as standards for comparison should be of the same viscosity average molecular weight, monomer content and composition distribution. Linear character for polymer containing C2 to C10 monomers is confirmed by Carbon-13 NMR the method of Randall (Rev. Macromol. Chem. Phys., C29 (2&3), p. 285-297). 15 Linear character for C11 and above monomers is confirmed by GPC analysis using a MALLS detector. For example, for a copolymer of propylene, the NMR should not indicate branching greater than that of the co-monomer (i.e. if the comonomer is butene, branches of greater than two carbons should not be present). For a homopolymer of propylene, the GPC should not show branches of 20 more than one carbon atom. When a linear standard is desired for a polymer where the comonomer is C9 or more, one can refer to T. Sun, P. Brant, R. R. Chance, and W. W. Graessley, *Macromolecules*, Volume 34, Number 19, 6812-6820, (2001) for protocols on determining standards for those polymers. In the case of syndiotactic polymers, the standard should have a comparable amount of 25 syndiotacticity as measured by Carbon 13 NMR.

Polymer samples for ^{13}C NMR spectroscopy were dissolved in d_2 -1,1,2,2-tetrachloroethane and the samples were recorded at 125°C . using a NMR spectrometer of 75 or 100 MHz. Polymer resonance peaks are referenced to $\text{mmmm}=21.8$ ppm. Calculations involved in the characterization of polymers by 30 NMR follow the work of F. A. Bovey in "Polymer Conformation and Configuration" Academic Press, New York 1969 and J. Randall in "Polymer Sequence Determination, Carbon-13 NMR Method, Academic Press, New York, 1977. The Bernoullianity index (B) is defined as $B=4[\text{mm}][\text{rr}]/[\text{mr}]^2$. The percent of methylene sequences of two in length, $\%(\text{CH}_2)_2$, were calculated as follows: the 35 integral of the methyl carbons between 14-18 ppm (which are equivalent in

5 concentration to the number of methylenes in sequences of two in length) divided
by the sum of the integral of the methylene sequences of one in length between
45-49 ppm and the integral of the methyl carbons between 14-18 ppm, times 100.
This is a minimum calculation for the amount of methylene groups contained in a
sequence of two or more since methylene sequences of greater than two have been
10 excluded. Assignments were based on H. N. Cheng and J. A. Ewen, Makromol.
Chem. 1989, 190, 1931.

Peak melting point (T_m), peak crystallization temperature (T_c) were
determined according to ASTM E 794-85. The heat of fusion and crystallinity
15 were determined using ASTM D 3417-99. Differential scanning calorimetric
(DSC) data was obtained using a TA Instruments model 2920 machine. Samples
weighing approximately 7-10 mg were sealed in aluminum sample pans. The
DSC data was recorded by first cooling the sample to $-50\text{ }^{\circ}\text{C}$ and then gradually
heating it to $200\text{ }^{\circ}\text{C}$ at a rate of $10\text{ }^{\circ}\text{C}/\text{minute}$. The sample was kept at $200\text{ }^{\circ}\text{C}$ for
20 5 minutes before a second cooling-heating cycle was applied. Both the first and
second cycle thermal events were recorded. Areas under the melting curves were
measured and used to determine the heat of fusion (ΔH) and the degree of
crystallinity. The percent crystallinity was calculated using the formula, [area
under the curve (Joules/gram) / B (Joules/gram)] * 100, where B is the heat of
25 fusion for the homopolymer of the major monomer component. These values for
B are to be obtained from the Polymer Handbook, Fourth Edition, published by
John Wiley and Sons, New York 1999. A value of 189 J/g (B) was used as the
heat of fusion for 100% crystalline polypropylene. For polymers displaying
multiple melting or crystallization peaks, the highest melting peak was taken as
30 peak melting point, and the highest crystallization peak was taken as peak
crystallization temperature.

The ethylene content of ethylene/propylene copolymers was determined
using FTIR according to the following technique. A thin homogeneous film of
polymer, pressed at a temperature of about $150\text{ }^{\circ}\text{C}$, was mounted on a Perkin
35 Elmer Spectrum 2000 infrared spectrophotometer. A full spectrum of the sample

- 5 from 600 cm^{-1} to 4000 cm^{-1} was recorded and the ethylene content in wt.% was calculated according to the following equation:

$$\text{ethylene content (wt.\%)} = 72.698 - 86.495X + 13.696X^2$$

- where $X = \text{AR}/(\text{AR}+1)$. The area under propylene band at $\sim 1165 \text{ cm}^{-1}$ and the area of ethylene band at $\sim 732 \text{ cm}^{-1}$ in the spectrum were calculated. The baseline
10 integration range for the methylene rocking band is nominally from 695 cm^{-1} to the minimum between 745 and 775 cm^{-1} . For the polypropylene band the baseline and integration range is nominally from 1195 to 1126 cm^{-1} . AR is the ratio of the area for the peak at $\sim 1165 \text{ cm}^{-1}$ to the area of the peak at $\sim 732 \text{ cm}^{-1}$.

- 15 The glass transition temperature (T_g) was measured by ASTM E 1356 using a TA Instruments model 2920 machine.

- Melt Viscosity (ASTM D-3236) (also called "viscosity", "Brookfield viscosity") Melt viscosity profiles were typically measured at temperatures from 120 $^{\circ}\text{C}$ to 190 $^{\circ}\text{C}$ using a Brookfield Thermosel viscometer and a number 27
20 spindle.

General Procedure of Polymerization

- Polymerization was performed in a series dual-reactor continuous solution process. Both of the reactors were a 0.5-liter stainless steel autoclave reactor and
25 were equipped with a stirrer, a water-cooling/steam-heating element with a temperature controller, and a pressure controller. Solvents, monomers such as ethylene and propylene, and comonomers (such as butene and hexene), if present, were first purified by passing through a three-column purification system. The purification system consisted of an Oxiclear column (Model # RGP-R1-500 from
30 Labclear) followed by a 5A and a 3A molecular sieve columns. Purification columns were regenerated periodically whenever there was evidence of lower activity of polymerization. Both the 3A and 5A molecular sieve columns were regenerated in-house under nitrogen at a set temperature of 260 $^{\circ}\text{C}$ and 315 $^{\circ}\text{C}$,

5 respectively. The molecular sieve material was purchased from Aldrich. Oxiclear column was regenerated in the original manufacture.

The solvent feed to the reactors was measured by a mass-flow meter. A Pulsafeed pump controlled the solvent flow rate and increased the solvent pressure
10 to the reactors. The compressed, liquefied propylene feed was measured by a mass flow meter and the flow was controlled by a variable speed pump. The monomers were fed into the reactor through pulse pump (>5 ml/minute) or Eldex pump (< 5 ml/minute) and the flow rates were measured using Brookfield mass flow meters or Micro-Motion Coriolis-type flow meters. The solvent, monomers and
15 comonomers were fed into a manifold first. Ethylene from in-house supply was delivered as a gas solubilized in the chilled solvent/monomer mixture in the manifold. The mixture of solvent and monomers were then chilled to about -15 °C by passing through a chiller prior to feeding into the reactor through a single tube. Ethylene flow rate was metered through a Brookfield mass flow controller. A
20 mass flow controller was used to deliver hydrogen into the reactor.

The content of the first reactor flowed into the second reactor. The exit of the first reactor was connected by a heat traced tubing to a second reactor similarly equipped to the first, with a provision for independent catalyst and
25 cocatalyst addition, and additional monomer, hydrogen, solvent addition and reaction temperature control. After the polymer solution stream exited the second reactor, polymerization was stopped with the addition of a small amount of water.

The catalyst compounds used to produce semi-crystalline polypropylene
30 were rac-dimethylsilylbis(2-methyl-4-phenylindenyl)zirconium dimethyl (obtained from Albemarle) and rac-1,2-ethylene-bis(4,7-dimethylindenyl)hafnium dimethyl (obtained from Boulder Scientific Company).

The catalyst compounds used to produce amorphous polypropylene were,
35 dimethylsilyl(tetramethylcyclopentadienyl)(cyclododecylamido)titanium dimethyl (obtained from Albemarle) and [di(p-

- 5 triethylsilylphenyl)methylene](cyclopentadienyl) (3,8-di-t-butylfluorenyl)hafnium dimethyl (obtained from Albemarle).

The catalysts were preactivated with N,N-dimethylanilinium tetrakis(pentafluorophenyl) borate (obtained from Albemarle) at a molar ratio of
10 1:1 to 1:1.1 in 700 ml of toluene at least 10 minutes prior to the polymerization reaction. The catalyst systems were diluted to a concentration of catalyst ranging from 0.2 to 1.4 mg/ml in toluene. All catalyst solutions were kept in an inert atmosphere with <1.5 ppm water content and fed into reactors by metering pumps. The catalyst solution was used for all polymerization runs carried out in the same
15 day. New batch of catalyst solution was prepared when more than 700 ml of catalyst solution was consumed in one day.

When multiple catalysts were utilized in one reactor, each catalyst solution was pumped through separate lines, and then mixed in a manifold, and
20 fed into the reactor through a single line. The connecting tube between the catalyst manifold and reactor inlet was about one meter long. The contact of catalyst, solvent and monomers took place in the reactor. Catalyst pumps were calibrated periodically using toluene as the calibrating medium. Catalyst concentration in the feed was controlled through changing the catalyst
25 concentration in catalyst solution and/or changing in the feed rate of catalyst solution. The feed rate of catalyst solution varied in a range of 0.2 to 5 ml/minute.

As an impurity scavenger, 250 ml of tri-n-octylaluminum (TNOA) (25 wt.% in toluene, Akzo Noble) was diluted in 22.83 kilogram of hexane. The
30 diluted TNOA solution was stored in a 37.9-liter cylinder under nitrogen blanket. The solution was used for all polymerization runs until about 90% of consumption, and then a new batch was prepared. Feed rates of the tri-n-octylaluminum (TNOA) solution varied from polymerization reaction to reaction, ranging from 0 (no scavenger) to 4 ml per minutes in order to optimize the overall
35 catalyst efficiency.

5 For polymerization reactions involving alpha, omega-dienes, 1,9-decadiene was diluted to a concentration ranging from 4.8 to 9.5 vol.% in toluene. The diluted solution was then fed into the reactor by a metering pump through a comonomer line. The 1,9-decadiene was obtained from Aldrich and was purified by first passing through alumina activated at high temperature under nitrogen,
10 followed by molecular sieve activated at high temperature under nitrogen.

The reactors were first cleaned by continuously pumping solvent (e.g., hexane) and scavenger through the reactor system for at least one hour at a maximum allowed temperature (about 150°C). After cleaning, the reactors were
15 heated/cooled to the desired temperature using water/steam mixture flowing through the reactor jacket and controlled at a set pressure with controlled solvent flow. Monomers and catalyst solutions were then fed into the reactor. An automatic temperature control system was used to control and to maintain the reactors at set temperatures. Onset of polymerization activity was determined by
20 observations of a viscous product and lower temperature of water-steam mixture. Once the activity was established and system reached steady state, the reactors were lined out by continuing operating the system under the established condition for a time period of at least five times of mean residence time prior to sample collection. The resulting mixture from the second reactor, containing mostly
25 solvent, polymer and unreacted monomers, was collected in a collection box. The collected samples were first air-dried in a hood to evaporate most of the solvent, and then dried in a vacuum oven at a temperature of about 90 °C for about 12 hours. The vacuum oven dried samples were weighed to obtain yields. All the reactions were carried out at a pressure of about 2.41 MPa-g and in the
30 temperature range of 70 to 130 °C.

EXAMPLE 1a-1i

These examples demonstrate a series dual-reactor continuous solution
35 process with the use of rac-dimethylsilyl bis(2-methyl-4-phenylindenyl)zirconium dimethyl catalyst (Catalyst A) (obtained from Albemarle) in the first reactor to

- 5 produce isotactic polypropylene and [di(p-triethylsilylphenyl) methylene]
(cyclopentadienyl) (3,8-di-t-butylfluorenyl)hafnium dimethyl catalyst (Catalyst B)
(obtained from Albemarle) in the second reactor to produce amorphous
polypropylene. Molecular weight of the isotactic polypropylene was controlled
using reaction temperature and molecular weight of the amorphous polypropylene
10 was adjusted through the addition of hydrogen as a molecular weight control
agent. Propylene, solvent, catalyst A solution and scavenger were fed into the first
reactor. The content of the first reactor flows into the second reactor. Catalyst B
solution and hydrogen were fed into the second reactor. The general procedure
described above was followed, and the detailed reaction condition and polymer
15 properties are listed in Table 1. The catalyst A in Example 1i was rac-
dimethylsilyl bis (indenyl) hafnium dimethyl (obtained from Albemarle).

Table 1					
Example	1a	1b	1c	1d	1e
Reactor 1 Temperature (°C)	125	120	115	125	120
Reactor 2 Temperature (°C)	125	125	125	125	125
Hexane feed rate to reactor 1 (ml/min)	90	90	90	90	90
Propylene feed rate to reactor 1 (g/min)	28.3	28.3	28.3	28.3	28.3
Catalyst B feed rate to reactor 2 (mole/min)	9.42E-07	9.42E-07	9.42E-07	9.42E-07	9.42E-07
Catalyst A feed rate to reactor 1 (mole/min)	4.89E-08	4.89E-08	4.89E-08	5.59E-08	5.59E-08
H ₂ feed rate to reactor 2 (cc/min)	20	20	20	20	20
T _c (°C)	90.4	99.7	99.7	98.1	101.1
T _m (°C)	131.6	150.0	152.4	145.3	149.9
T _g (°C)	-10.9	-7.1	-7.0	-2.4	-8.3
delta H (J/g)	21.8	44.4	52.9	42.5	52.9
M _n (kg/mol)	15.7	12.8	12.3	16.9	12.0
M _w (kg/mol)	37.0	34.4	37.9	49.3	32.9
M _z (kg/mol)	79.8	66.1	72.7	186.0	63.5
g' @M _z	0.90	0.88	0.88	0.81	0.86
Viscosity at 190 °C (cp)	2690	2000	2560	5980	1470

5

Table 1 (continued)				
Example	1f	1g	1h	1i
Reactor 1 Temperature (°C)	115	120	115	120
Reactor 2 Temperature (°C)	125	125	125	90
Hexane feed rate to reactor 1 (ml/min)	90	90	90	80
Propylene feed rate to reactor 1 (g/min)	28.3	28.3	28.3	14
Catalyst B feed rate to reactor 2 (mole/min)	9.42E-07	9.42E-07	9.42E-07	8.24E-08
Catalyst A feed rate to reactor 1 (mole/min)	5.59E-08	4.89E-08	4.89E-08	1.11E-06
H ₂ feed rate to reactor 2 (cc/min)	20	20	20	0
T _c (°C)	101.1	100.3	101.3	69.0
T _m (°C)	151.6	150.5	153.2	106.4
T _g (°C)	-8.1	-7.3	-4.5	-12.9
delta H (J/g)	53.8	50.0	58.8	46.8
M _n (kg/mol)	13.3	16.0	21.7	
M _w (kg/mol)	37.4	37.8	48.4	
M _z (kg/mol)	73.8	70.5	89.8	
g' @M _z	0.88	0.92	0.93	
Viscosity at 190 °C (cp)	2350	2760	6310	718

EXAMPLE 2a-2f

These examples demonstrate a series dual-reactor continuous solution process with the use of rac-dimethylsilyl bis(2-methyl-4-phenylindenyl)zirconium dimethyl catalyst (Catalyst A) in the first reactor to produce isotactic polypropylene and [di(p-triethylsilylphenyl) methylene] (cyclopentadienyl) (3,8-di-t-butylfluorenyl)hafnium dimethyl catalyst (Catalyst B) in the second reactor to produce propylene/ethylene copolymer. Propylene, solvent, catalyst A solution and scavenger were fed into the first reactor. The content of the first reactor flows into the second reactor. Ethylene and catalyst B solution were fed into the second reactor. The crystallinity of ethylene/propylene copolymers was adjusted through propylene conversion in the first reactor and amount of ethylene fed into the second reactor. Sufficient ethylene feed rate is required in order to produce amorphous ethylene/propylene copolymer. The general procedure described above

- 5 was followed, and the detailed reaction condition and polymer properties are listed in Table 2.

Table 2						
Example	2a	2b	2c	2d	2e	2f
Reactor 1 Temperature (°C)	118	125	125	125	125	125
Reactor 2 Temperature (°C)	120	125	125	125	125	125
Propylene feed rate to reactor 1 (g/min)	14	28.3	28.3	28.3	28.3	28.3
Ethylene feed rate to reactor 2 (SLPM)	0.6	0.6	0.6	0.6	0.6	0.6
Hexane feed rate to reactor 1 (ml/min)	90	90	90	90	90	90
Catalyst B feed rate to reactor 2 (mole/min)	8.83E-07	5.89E-07	5.89E-07	5.89E-07	5.89E-07	5.89E-07
Catalyst A feed rate to reactor 1 (mole/min)	1.40E-07	1.40E-07	6.99E-08	6.99E-08	6.99E-08	4.89E-08
H ₂ feed rate to reactor 2 (cc/min)	0	20	30	10	20	30
T _c (°C)	98.9	97.3	82.2	92.5	61.1	60.6
T _m (°C)	150.5	138.0	144.3	147.9	110.3	108.1
T _g (°C)	-15.8	-16.5	-16.1	-16.5	-13.5	-13.7
delta H (J/g)	47.9	68.5	21.6	37.1	19.3	15.1
M _n	17.8	9.7	10.0	11.1	13.5	11.7
M _w	42.1	27.0	30.0	31.6	35.2	30.4
M _z	76.8	50.4	59.7	64.2	69.0	57.6
g' @M _z	0.79	0.76	0.87	0.84	0.88	0.89
Viscosity @190 °C (cp)	2620	518	1290	1390	3040	1800
Ethylene content (wt %)	5.4	4.11	5.61	5.11	5.12	5.07

EXAMPLE 3a-3e

- 5 These examples demonstrate a series dual-reactor continuous solution process with the use of rac-dimethylsilyl bis(2-methyl-4-phenylindenyl)zirconium dimethyl catalyst (Catalyst A) in the first reactor to produce isotactic polypropylene and [di(p-triethylsilylphenyl) methylene] (cyclopentadienyl) (3,8-di-t-butylfluorenyl)hafnium dimethyl catalyst (Catalyst B) in the second reactor to
- 10 produce propylene/octene copolymer. Propylene, solvent, catalyst A solution and scavenger were fed into the first reactor. The content of the first reactor flows into the second reactor. Octene and catalyst B solution were fed into the second reactor. The crystallinity of propylene/hexene copolymers was adjusted through propylene conversion in the first reactor and amount of hexene fed into the second
- 15 reactor. For most samples, the propylene/hexene copolymers were amorphous. The general procedure described above was followed, and the detailed reaction condition and polymer properties are listed in Table 3.

Table 3					
Example	3a	3b	3c	3d	3e
Reactor 1 Temperature (°C)	125	120	115	125	120
Reactor 2 Temperature (°C)	125	125	125	125	125
Propylene feed rate to reactor 1 (g/min)	28.3	28.3	28.3	28.3	28.3
Octene feed rate to reactor 2 (ml/min)	1	1	1	1	1
Hexane feed rate to reactor 1 (ml/min)	90	90	90	90	90
Catalyst B feed rate to reactor 2 (mole/min)	9.42E-07	9.42E-07	9.42E-07	9.42E-07	9.42E-07
Catalyst A feed rate to reactor 1 (mole/min)	3.64E-08	3.64E-08	3.64E-08	4.20E-08	4.20E-08
H ₂ feed rate to reactor 2 (cc/min)	10.0	20.0	30.0	10.0	20.0
T _c (°C)				70.0	62.5
T _m (°C)				132.9	129.2
T _g (°C)				-6.8	-10.6
delta H (J/g)				27.0	17.3
Viscosity @190°C (cp)	7950	2970	1420	3940	2140

20

EXAMPLE 4a-4e

5

These examples demonstrated a series dual-reactor continuous solution process with the use of [di(p-triethylsilylphenyl) methylene] (cyclopentadienyl) (3,8-di-t-butylfluorenyl)hafnium dimethyl catalyst (Catalyst B) in the first reactor to produce ethylene/propylene copolymers and rac-dimethylsilyl bis(2-methyl-4-phenylindenyl)zirconium dimethyl catalyst (Catalyst A) in the second reactor to produce isotactic polypropylene. Catalyst feed rate to the first reactor was adjusted to ensure over 50% propylene conversion in the first reactor. At this propylene conversion, the ethylene conversion was about over 90 %. High conversion of ethylene in the first reactor allowed for a more crystalline product to be made in the second reactor. Propylene, ethylene, solvent, catalyst A solution and scavenger were fed into the first reactor. The content of the first reactor flows into the second reactor. Catalyst B solution was fed into the second reactor. The general procedure described above was followed, and the detailed reaction condition and polymer properties are listed in Table 4.

20

Table 4					
Example	4a	4b	4c	4d	4e
Reactor 1 Temperature (°C)	125	125	125	125	125
Reactor 2 Temperature (°C)	110	100	90	110	100
Propylene feed rate to reactor 1 (g/min)	14	14	14	28.3	28.3
Ethylene feed rate to reactor 1 (SLPM)	0.6	0.6	0.6	0.6	0.6
Hexane feed rate to reactor 1 (ml/min)	90	90	90	90	90
Catalyst B feed rate to reactor 1 (mole/min)	1.47E-07	1.47E-07	1.47E-07	5.89E-07	5.89E-07
Catalyst A feed rate to reactor 2 (mole/min)	2.83E-07	2.83E-07	2.83E-07	1.89E-07	1.89E-07
T _c (°C)	39.3	49.8	57.3	62.6	75.8
T _m (°C)	81.8	91.3	99.7	111.7	122.8
T _g (°C)	-23.7	-26.5	-22.7	-12.9	-11.6
delta H (J/g)	17.9	24.7	30.1	17.9	25.1
M _n (kg/mol)	3.8	5.1	6.5	14.3	15.8
M _w (kg/mol)	30.7	35.2	34.7	60.6	66.9

Mz (kg/mol)	93.5	100.2	83.5	121.1	133.5
Viscosity @190 °C (cp)	2790	3360	2890	25600	28200
Ethylene content (wt %)	9.11	9.22	8.58	4.7	4.65

5

EXAMPLE 5a-5c

These examples demonstrate a series dual-reactor continuous solution process with the use of rac-dimethylsilyl bis(2-methyl-4-phenylindenyl)zirconium dimethyl catalyst (Catalyst A) to produce isotactic polypropylene in the first reactor and propylene/ethylene copolymer in the second reactor. Propylene, solvent, catalyst A solution and scavenger were fed into the first reactor. The content of the first reactor flows into the second reactor. Ethylene was fed into the second reactor. The crystallinity of ethylene/propylene copolymers was adjusted through propylene conversion in the first reactor and amount of ethylene fed into the second reactor. Sufficient ethylene fed rate is required in order to produce amorphous ethylene/propylene copolymer in the second reactor. The general procedure described above was followed, and the detailed reaction condition and polymer properties are listed in Table 5.

20

Table 5			
Example	5a	5b	5c
Reactor 1 Temperature (°C)	118	90	90
Reactor 2 Temperature (°C)	100	90	100
Propylene feed rate to reactor 1 (g/min)	14	14	14
Ethylene feed rate to reactor 2 (SLPM)	0.6	0.6	0.6
Hexane feed rate to reactor 1 (ml/min)	90	90	90
Catalyst A feed rate to reactor 1 (mole/min)	6.99E-08	6.99E-08	6.99E-08
Catalyst A feed rate to reactor 2 (mole/min)	2.10E-07	2.10E-07	2.10E-07
Tc (°C)	74.6	52.5	87.4
Tm (°C)	145.4	99.1	151.6
Tg (°C)	-19.1	-11.9	-21.4
delta H (J/g)	45.1	42.9	29.0
Mn (kg/mol)	6.7	12.8	5.3
Mw (kg/mol)	26.2	33.3	23.8
Mz (kg/mol)	49.7	58.6	94.1

Viscosity @190 °C (cp)	455	1240	320
Ethylene content (wt.%)	6.47	8.23	6.82

5

EXAMPLE 6a-6c

- These examples demonstrate a series dual-reactor continuous process with the use of rac-dimethylsilyl bis(2-methyl-4-phenylindenyl)zirconium dimethyl catalyst (Catalyst A) in the first reactor to produce low molecular weight polyethylene and a mixture of rac-dimethylsilyl bis(2-methyl-4-phenylindenyl)zirconium dimethyl catalyst (Catalyst A) and [di(p-triethylsilylphenyl) methylene] (cyclopentadienyl) (3,8-di-t-butylfluorenyl)hafnium dimethyl catalyst (Catalyst B) in the second reactor to produce atactic (aPP) and isotactic polypropylene (iPP) as well as aPP/iPP branch-blocks. Ethylene, solvent, catalyst A solution and scavenger were fed into the first reactor. The content of the first reactor flows into the second reactor. Propylene and catalyst B solution were fed into the second reactor. The catalyst A fed rate was high enough that over 90% of ethylene were converted in the first reactor.
- The ratio of catalyst A and catalyst B was adjusted to control and aPP/iPP ratio in the second reactor. The general procedure described above was followed, and the detailed reaction condition and polymer properties are listed in Table 6.

Table 6			
Example	6a	6b	6c
Reactor 1 Temperature (°C)	125	120	120
Reactor 2 Temperature (°C)	125	110	100
Ethylene feed rate reactor 1 (SLPM)	0.2	0.2	0.2
Propylene feed rate to reactor 2 (g/min)	14	14	14
Hexane fed rate to reactor 1 (ml/min)	90	90	90
Catalyst B feed rate to reactor 2 (mole/min)	8.83E-07	8.83E-07	8.83E-07
Catalyst A feed rate to reactor 1 (mole/min)	2.10E-07	2.10E-07	2.10E-07
Tc (°C)	91.2	103.6	101.6
Tm (°C)	128.0	138.1	139.9
Tg (°C)	-13.3	-8.8	-6.1
delta H (J/g)	26.7	59.3	42.6

Mn (kg/mol)	4.9	7.1	12.7
Mw (kg/mol)	37.3	59.6	127.9
Mz (kg/mol)	112.7	218.5	320.9
Viscosity @190 °C (cp)	2435	28400	89800

5

Adhesive Testing

A number of hot melt adhesives were prepared by using the neat polymers or blending the neat polymer, functionalized additives, tackifier, wax, antioxidant, and other ingredients under low shear mixing at elevated temperatures to form fluid melt. The mixing temperature varied from about 130 to about 190 °C. Adhesive test specimens were created by bonding the substrates together with a dot of molten adhesive and compressing the bond with a 500-gram weight until cooled to room temperature. The dot size was controlled by the adhesive volume such that in most cases the compressed disk which formed gave a uniform circle just inside the dimensions of the substrate.

Once a construct has been produced it can be subjected to various insults in order to assess the effectiveness of the bond. Once a bond fails to a paper substrate a simple way to quantify the effectiveness is to estimate the area of the adhesive dot that retained paper fibers as the construct failed along the bond line. This estimate was called percent substrate fiber tear. An example of good fiber, after conditioning a sample for 15 hours at -12 °C and attempting to destroy the bond, would be an estimate of 80-100% substrate fiber tear. It is likely that 0% substrate fiber tear under those conditions would signal a loss of adhesion.

Substrate fiber tear: The specimens were prepared using the same procedure as that described above. For low temperature fiber tear test, the bond specimens were placed in a freezer or refrigerator to obtain the desired test temperature. For substrate fiber tear at room temperature, the specimens were aged at ambient conditions. The bonds were separated by hand and a determination made as to the type of failure observed. The amount of substrate fiber tear was expressed in percentage.

5 Dot T-Peel was determined according to ASTM D 1876, except that the specimen was produced by combining two 1 inch by 3 inch (2.54 cm x 7.62 cm) substrate cut outs with a dot of adhesive with a volume that, when compressed under a 500-gram weight occupied about 1 square inch of area (1 inch = 2.54 cm). Once made all the specimens were pulled apart in side by side testing at a rate of 2
10 inches per minute by a machine that records the destructive force of the insult being applied. The maximum force achieved for each sample tested was recorded and averaged, thus producing the average maximum force which is reported as the Dot T-Peel.

 Peel Strength (modified ASTM D1876): Substrates (1 x 3 inches (25 x 76
15 mm)) were heat sealed with adhesive film (5 mils (130 μ m) thickness) at 135 °C for 1 to 2 seconds and 40 psi (0.28 MPa) pressure. Bond specimens were peeled back in a tensile tester at a constant crosshead speed of 2 in/min (51 mm/min). The average force required to peel the bond (5 specimens) apart is recorded.

 Set time is defined as the time it takes for a compressed adhesive substrate
20 construct to fasten together enough to give substrate fiber tear when pulled apart, and thus the bond is sufficiently strong to remove the compression. The bond will likely still strengthen upon further cooling, however, it no longer requires compression. These set times were measured by placing a molten dot of adhesive on to a file folder substrate taped to a flat table. A file folder tab (1 inch by 3 inch
25 (2.5 cm x 7.6 cm)) was placed upon the dot 3 seconds later and compressed with a 500 gram weight. The weight was allowed to sit for about 0.5 to about 10 seconds. The construct thus formed was pulled apart to check for a bonding level good enough to produce substrate fiber tear. The set time was recorded as the minimum time required for this good bonding to occur. Standards were used to
30 calibrate the process.

 SAFT (modified D 4498-00) measures the ability of a bond to withstand an elevated temperature rising at 10 °F (5.5 °C) /15 min., under a constant force that pulls the bond in the shear mode. Bonds were formed in the manner described above on Kraft paper (1 inch by 3 inch (2.5 cm x 7.6 cm)). The test

- 5 specimens were suspended vertically in an oven at room temperature with a 500-gram load attached to the bottom. The temperatures at which the weight fell was recorded (when the occasional sample reached temperatures above the oven capacity >265°F (129°C) it was terminated and averaged in with the other samples at termination temperature).

- 10 Shore A hardness was measured according to ASTM D 2240. An air-cooled dot of adhesive was subjected to the needle and the deflection was recorded from the scale.

Examples

- 15 The following materials were used in examples of adhesive performance testing.

Trade name	Description	Source
Tackifiers		
Escorez® 5637	Hydrogenated aromatic modified resin produced from dicyclopentadiene feedstock, exhibiting a ring and ball softening point of 130 °C	ExxonMobil Chemical Company
Escorez® 5690	Hydrogenated aromatic modified resin produced from dicyclopentadiene feedstock, exhibiting a ring and ball softening point of 90 °C	ExxonMobil Chemical Company
Escorez® 2203	A low aromatic modified hydrocarbon resin having a narrow molecular weight distribution produced from a feed of C5, C6 and C9 olefins and di-olefins, having a ring and ball softening point of about 95°C	ExxonMobil Chemical Company
Oils		
Kadol oil	Refined white mineral oil	Witco
Polymers / Adhesives		
Rextac RT 2715	Propylene/butene copolymer, Ziegler Natta APAO	Huntsman, Odessa Texas
Henkel hot melt 80-8368	Commercial blend of EVA, tackifier, and wax	Henkel Corporation
Advantra 9250	Commercial blend of ethylene/octene metallocene polymers, tackifiers, and wax	Fuller
Tite bond wood glue	Water based adhesive	Home Depot, Houston, Texas

Waxes/Funtionalized additives		
AC395A	Oxidized polyethylene with density of 1.0 g/cc, viscosity of 2500 cP at 150 °C and acid number of 45~50 mg/KOH/g.	Honeywell, Morristown, New Jersey
AC 596P	Polypropylene-maleic anhydride copolymer with viscosity 189 cps at 190 °C and Saponification number of 40 mg/KOH/g	Honeywell, Morristown, New Jersey
AC 597	Polypropylene-maleic anhydride copolymer with viscosity 374 cps at 190 °C and Saponification number of 80 mg/KOH/g	Honeywell, Morristown, New Jersey
AC X1325	Polypropylene-maleic anhydride copolymer with viscosity 1490 cps at 190 °C and acid number of 16 mg/KOH/g	Honeywell, Morristown, New Jersey
A-C-1302P	Ethylene-maleic anhydride copolymer with viscosity of 248 cP at 190 °C and Saponification number of 5 mg KOH/g	Honeywell, Morristown, New Jersey
PP-grafted maleic anhydride	PP-grafted maleic anhydride with averaged Mw of ~9,100, Mn of ~3,900 by GPC, viscosity of 400 cP at 190 °C, acid number of 47 mg KOH/g, softening point of 157 °C (ring and ball) and density of 0.934 g/ml	Sigma-Aldrich, Product number: 42784-5
MAPP 40	Maleated polypropylene with acid value of 45 ~50, viscosity at 190 °C of 400 - 425 cp, and soft point of 143~155 °C	Chusei, Pasadena, Texas
Paraflint H-1	Fisher-Tropsch wax, 10 mPa @ 250 °F	Moore and Munger
C80 wax	Fischer Tropsch fractionated wax	Moore and Munger
Antioxidants and other additives		
Irganox 1010	Phenolic antioxidant	Ciba-Geigy
Test surfaces (substrates)		
Paperboard 84B	Generic poster board clay coated newsprint	Huckster Packaging and Supply, Houston, Texas
Paperboard 84C	Generic corrugated cardboard 200# stock	Huckster Packaging and Supply, Houston, Texas
Inland paper board	High Performance box board	Inland Paper Board and Packaging Company of Rome
Black white fabric	Printed stretch 100% Cotton with a Thread Count of 17 by 13 per square cm, a more loosely woven fabric	High Fashion Fabrics, Houston, Texas
Formica	Tab were made from standard sheet Formica	Lowe's Hardware, Houston, Texas
Blue fabric	Tab were made from Blue Stock	High Fashion Fabrics,

	038C0TP 100% Cotton, Thread Count 21by 45 per square cm with a weight of 0.022 grams per square cm, a tightly woven cotton fabric	Houston, Texas.
Seton catalog paper	Book paper bound by a hot melt process as determined from examination	Seton Catalog
PET	Polyester (PET), Commonly called Mylar	Several Sources
Kraft paper	Kraft paper	Georgia Pacific, Atlanta, Georgia
File folder	File folder is a typical manila letter size (1/3 cut) stock having a minimum of 10% post consumer recycle paper content	Smead Paper, stock number 153L, UPC number 10330
PP cast film	Polypropylene cast film PP4772.	ExxonMobil Chemical Company

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As examples, the adhesive formulation and adhesive performance for selected polymers produced using current invention are listed in the following tables.

Table 7. Adhesion tests of substrate fiber tear (Dot-T peel)

	HMA1	HMA2	HMA3	HMA4	HMA5	HMA6	HMA7
Polymer	1b	1b	1i	1i	2a	2a	2a
Polymer (wt.%)	100	87	97.5	92.5	100	89.5	89.5
Irganox 1010 (wt.%)	0	0				0.5	0.5
MAPP 40 (wt.%)	0	2.5	2.5	2.5			
AC1302P (wt.%)							10
C80 wax (wt.%)	0	7		5		5	
Escorez 5690 (wt.%)	0	3.5					
Escorez 2203 (wt.%)						5	
Shore A hardness	95	94	98	97	94	93	96
Set time (sec)	6+	3.5~4	6+	3.5	3.5	3.5	3.5
Viscosity @ 177 °C (cps)		1490					
Viscosity @ 190 °C (cps)						1790	2190
Fiber tear at low temperatures							
Paperboard 84C overnight at 2 °C (%)	11	97					
Paperboard 84C					97	48	97

overnight at -10 °C (%)							
Paperboard 84C overnight at -30 °C (%)					98	93	95
Inland Paper Board overnight at 2 °C (%)	1	84	89	85			
Inland paper board overnight at 6 °C (%)	0, af	90					
Inland Paper Board overnight at -10 °C (%)	0, af, ab	45			49	19	5
Inland Paper Board overnight at -25 ~ -30 °C (%)			93 (tested @ -18 C)	65 (tested @ -18 C)	72	0, af, ab	0, af, ab
Fiber tear at ambient condition							
Inland paper board	0, af, ab	97	79	94	0, af	0, af	0, af
Paperboard 84C	55	97			6	9	0, af
af -- adhesive failure, ab --adhesive break, set time: 6+ --set time was longer than 6 seconds.							

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Table 8. Adhesion tests of substrate fiber tear (Dot-T peel)							
	HMA8	HMA9	HMA10	HMA11	HMA12	HMA13	HMA14
Polymer	5a	5a	5a	5b	5b	5c	5c
Polymer (wt.%)	100	89.5	89.5	100	87	100	87
Irganox 1010 (wt.%)		0.5	0.5				
MAPP 40 (wt.%)					2.5		2.5
AC1302P (wt.%)			10				
C80 wax (wt.%)		5			7		7
Escorez 5690 (wt.%)					3.5		3.5
Escorez 2203 (wt.%)		5					
Shore A hardness	97	95	95	94	92	93	87
Set time (sec)	5	2	2	6+	3	6+	2.5
Viscosity @ 177 °C (cps)		482.5	585	1610	1050	433	315
Viscosity @ 190 °C (cps)		351	448				

Fiber tear at low temperatures							
Paperboard 84C overnight at -10 °C (%)	10	0, af, ab	2				
Paperboard 84C overnight at -30 °C (%)	69	5	75				
Inland Paper Board overnight at 0 °C (%)				2	74	0, af, ab	74
Inland Paper Board overnight at -10 °C (%)	0, af, ab	0, af, ab	0, af, ab	10	74	0, af, ab	58
Inland Paper Board overnight at -25 ~ -30 °C (%)	0, af, ab	0, af, ab	0, af, ab	5	55	0, af, ab	0, af, ab
Fiber tear at ambient condition							
Inland paper board	0, af	0, af	0, af	3	10	0, af	90
Paperboard 84C	0, af	0, af	0, af				
Af -- adhesive failure, ab --adhesive break, set time: 6+ --set time was longer than 6 seconds.							

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Table 9. Adhesion tests of substrate fiber tear (Dot-T peel)						
	HMA15	HMA16	HMA17	HMA18	HMA19	HMA20
Polymer	2c	2c	2c	2d	2d	2d
Polymer (wt.%)	100	89.5	87	100	89.5	87
MAPP 40 (wt.%)			2.5			2.5
C80 wax (wt.%)		7	7		7	7
Escorez 5690 (wt.%)		3.5	3.5		3.5	3.5
Shore A hardness	59	60	58	81	82	87
Set time (sec)	6+	3~3.5	3.5~4	6	3	2.5
Low temperature fiber tear						
Paperboard 84C overnight at 6 °C (%)	94	99	100	88	30	98
Inland paper board overnight at 6 °C (%)	95	82	100	90	49	100
Inland Paper Board overnight at -5 °C (%)	90	93	99	73	47	100
Inland Paper Board overnight at -25 ~ -30 °C (%)	66	30	81	35	27	93
Fiber tear at ambient condition						

Inland paper board	97	99	100	70	69	100
Paperboard 84C	100	99	100	65	25	100

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Table 10. Dot T-peel strength to Mylar and failure types

	HMA21	HMA22	HMA23	HMA24	HMA25	HMA26
Polymer	2c	2d	6a	1a	4c	3d
Polymer (wt.%)	100	100	100	100	100	100
Shore A hardness	51	81	38	29	79	35
Dot T-Peel failure type	CF, AF, SS	AF, SS	AF, CF	CF	AF	CF
Peel force to Mylar (aged for three days) (lbf)	1.954	0.017	2.549	2.819	0.0083	3.84
cf -- cohesive failure; af -- adhesive failure; ab--adhesive break, sf -- substrate failure; ss --slip stick						

Table 11. Dot T-peel strength to Mylar and failure types

	HMA27	HMA28	HMA29	HMA30	HMA31	HMA32
Polymer	2c	2d	6a	1a	4c	3d
Polymer (wt.%)	70	70	70	70	70	70
Escorez 5637 (wt.%)	30	30	30	30	30	30
Viscosity @ 190 °C (cp)	1032	1032	2770	1830	2305	3230
Shore A hardness	42	71	56	31	73	33
Dot T-Peel failure type	CF	AF, CF	CF, AF	CF	CF, AF	CF
Peel force to Mylar (aged for three days) (lbf)	19.47	19.71	13.31	17.82	8.506	21.98
cf -- cohesive failure; af -- adhesive failure; ab--adhesive break; sf -- substrate failure; ss --slip stick.						

While the invention has been described with reference to an exemplary embodiment, it will be understood by those skilled in the art that various changes may be made and equivalents may be substituted for elements thereof without departing from the scope of the invention. In addition, many modifications may be made to adapt a particular situation or material to the teachings of the invention without departing from the essential scope thereof. Therefore, it is intended that

- 5 the invention not be limited to the particular embodiment disclosed as the best mode contemplated for carrying out this invention, but that the invention will include all embodiments falling within the scope of the appended claims.

CLAIMS

1. A continuous process to produce a branched olefin polymer comprising:
 - 1) selecting a first catalyst component capable of producing a polymer having a weight average molecular weight of 100,000 or less and a crystallinity of 20% or less under selected polymerization conditions;
 - 2) selecting a second catalyst component capable of producing a polymer having a weight average molecular weight of 100,000 or less and a crystallinity of 20% or more at the selected polymerization conditions;
 - 3) contacting said first catalyst component or said second catalyst component, one or more activators, and one or more C₂ to C₄₀ olefins in a first reaction zone, at a temperature of greater than 70°C, and at a residence time of 120 minutes or less; and
 - 4) transferring the contents of the first reaction zone directly to a second reaction zone and further contacting the contents with said first catalyst component or said second catalyst component, one or more activators, and one or more C₂ to C₄₀ olefins, at a temperature of greater than 70°C, and at a residence time of 120 minutes or less; and
 - 5) optionally, transferring the contents of the second reaction zone directly to a third reaction zone and further contacting the contents with said first catalyst component or said second catalyst component, one or more activators, and one or more C₂ to C₄₀ olefins, at a temperature of greater than 70°C, and at a residence time of 120 minutes or less; and
 - 6) recovering a branched olefin polymer comprising at least 50 mole% of one or more C₃ to C₄₀ olefins,
wherein:
 - (a) both catalyst components are present in at least one reaction zone, selected from the first and second reaction zones; and
 - (b) in at least one of the first and the second reaction zones, the C₂ to C₄₀ olefin is a C₃ to C₄₀ alpha-olefin.
2. The process of claim 1 wherein the olefin polymer comprises from 50 to 100 mole% of propylene.
3. The process of claim 1 wherein the olefin polymer is homopolypropylene.

4. The process of claim 1 wherein the first catalyst component comprises a non-stereospecific metallocene catalyst compound.
5. The process of claim 1 wherein the second catalyst component comprises a stereospecific metallocene catalyst compound.
6. The process of claim 1 wherein propylene is present in the first reaction zone.
7. The process of claim 1 wherein propylene is present in the second reaction zone.
8. The process of claim 1 wherein propylene is present in a third reaction zone.
9. The process of claim 1 wherein ethylene is present in the first reaction zone.
10. The process of claim 1 wherein ethylene is present in the second reaction zone.
11. The process of claim 1 wherein ethylene is present in a third reaction zone.
12. The process of claim 1 wherein propylene and ethylene are present in the first reaction zone.
13. The process of claim 1 wherein propylene and ethylene are present in the second reaction zone.
14. The process of claim 1 wherein propylene is present in the first reaction zone at 100 weight %, based upon the weight of the monomers present.
15. The process of claim 1 wherein propylene is present in the second reaction zone at 100 weight %, based upon the weight of the monomers present.
16. The process of claim 1 wherein propylene is present in a third reaction zone at 100 weight %, based upon the weight of the monomers present.
17. The process of claim 1 wherein propylene is present in the first reaction zone at 100 weight %, based upon the weight of the monomers present in the first reaction zone and ethylene

is present in the second reaction zone at up to 50 weight%, based upon the weight of the monomers present in the second reaction zone.

18. The process of claim 1 wherein propylene and ethylene are present in the first reaction zone and no ethylene, other than residual ethylene monomer present in the contents of the first reaction zone, is introduced into the second reaction zone.

19. The process of claim 1 wherein ethylene is intermittently introduced into one or more reaction zones.

20. The process of claim 1 wherein ethylene is present in a reaction zone at less than 10 weight%, based upon the weight of the monomers in the reaction zone..

21. The process of claim 1 wherein propylene is present in the first reaction zone, ethylene is present in the second reaction zone, the second catalyst component is present in the first reaction zone, and the first catalyst component is present in the second reaction zone.

22. The process of claim 1 wherein propylene is present in the first reaction zone, propylene and ethylene are present in the second reaction zone, the second catalyst component is present in the first reaction zone, and the first catalyst component is present in the second reaction zone.

23. The process of claim 1 wherein propylene and ethylene are present in the first reaction zone, propylene is present in the second reaction zone, the first catalyst component is present in the first reaction zone, and the second catalyst component is present in the second reaction zone.

24. The process of claim 1 wherein propylene is present in the first reaction zone, propylene and ethylene are present in the second reaction zone, the second catalyst component is present in the first reaction zone, and the second catalyst component is present in the second reaction zone.

25. The process of claim 1 wherein ethylene is present in the first reaction zone, propylene is present in the second reaction zone, propylene is present in the third reaction zone, the first catalyst component is present in the second reaction zone, and the second catalyst component is present in the third reaction zone, and the catalyst component present in the first reaction zone is

capable of producing polymer having a weight average molecular weight of 20,000 or less and a crystallinity of 10% or less at the selected polymerization conditions.

26. The process of claim 1 wherein

- 1) the first catalyst component comprises a non-stereospecific metallocene catalyst compound;
- 2) the second catalyst component comprises a stereospecific metallocene catalyst compound;
- 3) the first reaction zone is a reactor comprising solvent, monomer, said first catalyst component or said second catalyst component, and activator at a temperature of greater than 70°C; and
- 4) The second reaction zone is a reactor comprising solvent, monomer, said first catalyst component or said second catalyst component, and activator at a temperature of greater than 70°C.

27. The process of claim 1 wherein the first catalyst component capable of producing polymer having a weight average molecular weight of 100,000 or less and a crystallinity of 20% or less is selected from the group consisting of: rac-dimethylsilylbis(4,7-dimethylindenyl) hafnium dichloride, rac-dimethylsilylbis(4,7-dimethylindenyl) hafnium dimethyl, rac-dimethylsilylbis(4,7-dimethylindenyl) zirconium dichloride, rac-dimethylsilylbis(4,7-dimethylindenyl) zirconium dimethyl, rac-dimethylsilylbis(indenyl) hafnium dichloride, rac-dimethylsilylbis(indenyl) hafnium dimethyl, rac-dimethylsilylbis(indenyl) zirconium dichloride, rac-dimethylsilylbis(indenyl) zirconium dimethyl, rac-dimethylsilylbis(tetrahydroindenyl) hafnium dichloride, rac-dimethylsilylbis(tetrahydroindenyl) hafnium dimethyl, rac-dimethylsilylbis(tetrahydroindenyl) zirconium dichloride, rac-dimethylsilylbis(tetrahydroindenyl) zirconium dimethyl, rac-diphenylsilylbis(4,7-dimethylindenyl) hafnium dichloride, rac-diphenylsilylbis(4,7-dimethylindenyl) hafnium dimethyl, rac-diphenylsilylbis(4,7-dimethylindenyl) zirconium dichloride, rac-diphenylsilylbis(4,7-dimethylindenyl) zirconium dimethyl, rac-diphenylsilylbis(indenyl) hafnium dichloride, rac-diphenylsilylbis(indenyl) hafnium dimethyl, rac-diphenylsilylbis(indenyl) zirconium dichloride, rac-diphenylsilylbis(indenyl) zirconium dimethyl, rac-diphenylsilylbis(tetrahydroindenyl) hafnium dichloride, rac-diphenylsilylbis(tetrahydroindenyl) hafnium dimethyl, rac-diphenylsilylbis(tetrahydroindenyl) zirconium dichloride, rac-diphenylsilylbis(tetrahydroindenyl)

zirconium dimethyl, rac-methylphenylsilylbis(4,7-dimethylindenyl) hafnium dichloride, rac-methylphenylsilylbis(4,7-dimethylindenyl) hafnium dimethyl, rac-methylphenylsilylbis(4,7-dimethylindenyl) zirconium dichloride, rac-methylphenylsilylbis(4,7-dimethylindenyl) zirconium dimethyl, rac-methylphenylsilylbis(indenyl) hafnium dichloride, rac-methylphenylsilylbis(indenyl) hafnium dimethyl, rac-methylphenylsilylbis(indenyl) zirconium dichloride, rac-methylphenylsilylbis(indenyl) zirconium dimethyl, rac-methylphenylsilylbis(tetrahydroindenyl) hafnium dichloride, rac-methylphenylsilylbis(tetrahydroindenyl) hafnium dimethyl, rac-methylphenylsilylbis(tetrahydroindenyl) zirconium dichloride, rac-methylphenylsilylbis(tetrahydroindenyl) zirconium dimethyl, rac-ethylenebis(4,7-dimethylindenyl) hafnium dichloride, rac-ethylenebis(4,7-dimethylindenyl) hafnium dimethyl, rac-ethylenebis(4,7-dimethylindenyl) zirconium dichloride, rac-ethylenebis(4,7-dimethylindenyl) zirconium dimethyl, rac-ethylenebis(indenyl) hafnium dichloride, rac-ethylenebis(indenyl) hafnium dimethyl, rac-ethylenebis(indenyl) zirconium dichloride, rac-ethylenebis(indenyl) zirconium dimethyl, rac-ethylenebis(tetrahydroindenyl) hafnium dichloride, rac-ethylenebis(tetrahydroindenyl) hafnium dimethyl, rac-ethylenebis(tetrahydroindenyl) zirconium dichloride, and rac-ethylenebis(tetrahydroindenyl) zirconium dimethyl.

28. The process of claim 1 wherein the first catalyst component comprises one or more of dimethylsilyl(tetramethylcyclopentadienyl)(cyclododecylamido) titanium dichloride, dimethylsilyl(tetramethylcyclopentadienyl)(cyclohexyl-amido) titanium dichloride, dimethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dichloride, dimethylsilyl(tetramethylcyclopentadienyl)(t-butylamido) titanium dichloride, dimethylsilyl(tetramethylcyclopentadienyl)(s-butylamido) titanium dichloride, dimethylsilyl(tetramethylcyclopentadienyl)(n-butylamido) titanium dichloride, dimethylsilyl(tetramethylcyclopentadienyl)(exo-2-norbomylamido) titanium dichloride, diethylsilyl(tetramethylcyclopentadienyl)(cyclododecyl-amido) titanium dichloride, diethylsilyl(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dichloride, diethylsilyl(tetramethylcyclopentadienyl)(cyclohexyl-amido) titanium dichloride, diethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dichloride, methylene(tetramethylcyclopentadienyl)(cyclododecyl-amido) titanium dichloride, methylene(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dichloride, methylene(tetramethylcyclopentadienyl)(cyclohexylamido) titanium dichloride, methylene(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dichloride,

dimethylsilyl(tetramethylcyclopentadienyl)(cyclododecylamido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(cyclohexyl-amido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,
 dimethylsilyl(2,5-dimethylcyclopentadienyl)(cyclododecylamido) titanium dichloride,
 dimethylsilyl(2,5-dimethylcyclopentadienyl)(exo-2-norbornylamido) titanium dichloride,
 dimethylsilyl(2,5-dimethylcyclopentadienyl)(cyclohexylamido) titanium dichloride,
 dimethylsilyl(2,5-dimethylcyclopentadienyl)(1-adamantylamido) titanium dichloride,
 dimethylsilyl(3,4-dimethylcyclopentadienyl)(cyclododecylamido) titanium dichloride,
 dimethylsilyl(3,4-dimethylcyclopentadienyl)(exo-2-norbornylamido) titanium dichloride,
 dimethylsilyl(3,4-dimethylcyclopentadienyl)(cyclohexylamido) titanium dichloride,
 dimethylsilyl(3,4-dimethylcyclopentadienyl)(1-adamantylamido) titanium dichloride,
 dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(cyclododecylamido) titanium dichloride,
 dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(exo-2-norbornylamido) titanium dichloride,
 dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(cyclohexylamido) titanium dichloride,
 dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(1-adamantylamido) titanium dichloride,
 dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(cyclododecylamido) titanium dichloride,
 dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(exo-2-norbornylamido) titanium dichloride,
 dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(cyclohexylamido) titanium dichloride,
 dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(1-adamantylamido) titanium dichloride,
 dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(cyclododecylamido) titanium
 dichloride,
 dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(exo-2-norbornylamido) titanium
 dichloride,
 dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(cyclohexylamido)
 titanium dichloride,
 dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(1-adamantylamido) titanium
 dichloride,
 dimethylsilyl(2-tetrahydroindenyl)(cyclododecylamido) titanium dichloride,
 dimethylsilyl(2-tetrahydroindenyl)(cyclohexylamido) titanium dichloride,
 dimethylsilyl(2-tetrahydroindenyl)(1-adamantylamido) titanium dichloride,
 dimethylsilyl(2-tetrahydroindenyl)(exo-2-norbornylamido) titanium dichloride,
 dimethylsilyl(tetramethylcyclopentadienyl)(cyclododecylamido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(cyclohexyl-amido) titanium dimethyl,

dimethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(t-butylamido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(s-butylamido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(n-butylamido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(exo-2-norbomylamido) titanium dimethyl,
 diethylsilyl(tetramethylcyclopentadienyl)(cyclododecyl-amido) titanium dimethyl,
 diethylsilyl(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
 diethylsilyl(tetramethylcyclopentadienyl)(cyclohexyl-amido) titanium dimethyl,
 diethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,
 methylene(tetramethylcyclopentadienyl)(cyclododecyl-amido) titanium dimethyl,
 methylene(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
 methylene(tetramethylcyclopentadienyl)(cyclohexylamido) titanium dimethyl,
 methylene(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(cyclododecylamido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(cyclohexyl-amido) titanium dimethyl,
 dimethylsilyl(tetramethylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,
 dimethylsilyl(2,5-dimethylcyclopentadienyl)(cyclododecylamido) titanium dimethyl,
 dimethylsilyl(2,5-dimethylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
 dimethylsilyl(2,5-dimethylcyclopentadienyl)(cyclohexylamido) titanium dimethyl,
 dimethylsilyl(2,5-dimethylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,
 dimethylsilyl(3,4-dimethylcyclopentadienyl)(cyclododecylamido) titanium dimethyl,
 dimethylsilyl(3,4-dimethylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
 dimethylsilyl(3,4-dimethylcyclopentadienyl)(cyclohexylamido) titanium dimethyl,
 dimethylsilyl(3,4-dimethylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,
 dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(cyclododecylamido)titanium dimethyl,
 dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
 dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(cyclohexylamido) titanium dimethyl,
 dimethylsilyl(2-ethyl-5-methylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,
 dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(cyclododecylamido)titanium dimethyl,
 dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
 dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(cyclohexylamido) titanium dimethyl,
 dimethylsilyl(3-ethyl-4-methylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,

dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(cyclododecylamido) titanium dimethyl,
 dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(exo-2-norbornylamido) titanium dimethyl,
 dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(cyclohexylamido) titanium dimethyl,
 dimethylsilyl(2-ethyl-3-hexyl-5-methyl-4-octylcyclopentadienyl)(1-adamantylamido) titanium dimethyl,
 dimethylsilyl(2-tetrahydroindenyl)(cyclododecylamido) titanium dimethyl,
 dimethylsilyl(2-tetrahydroindenyl)(cyclohexylamido) titanium dimethyl,
 dimethylsilyl(2-tetrahydroindenyl)(1-adamantylamido) titanium dimethyl, and
 dimethylsilyl(2-tetrahydroindenyl)(exo-2-norbornylamido) titanium dimethyl.

29. The process of claim 1 wherein the second catalyst component comprises one or more of the racemic versions of:

dimethylsilyl (2-methyl-4-phenylindenyl) zirconium dichloride,
 dimethylsilyl (2-methyl-4-phenylindenyl) zirconium dimethyl,
 dimethylsilyl (2-methyl-4-phenylindenyl) hafnium dichloride,
 dimethylsilyl (2-methyl-4-phenylindenyl) hafnium dimethyl,
 dimethylsilyl bis(indenyl)hafnium dimethyl,
 dimethylsilyl bis(indenyl)hafnium dichloride,
 dimethylsilyl bis(indenyl)zirconium dimethyl,
 dimethylsilyl bis(indenyl)zirconium dichloride,

the racemic isomers of:

dimethylsilanediylbis(2-methyl)metal dichloride;
 dimethylsilanediylbis(indenyl)metal dichloride;
 dimethylsilanediylbis(indenyl)metal dimethyl;
 dimethylsilanediylbis(tetrahydroindenyl)metal dichloride;
 dimethylsilanediylbis(tetrahydroindenyl)metal dimethyl;
 dimethylsilanediylbis(indenyl)metal diethyl; and
 dibenzylsilanediylbis(indenyl)metal dimethyl;

wherein the metal can be chosen from Zr, Hf, or Ti.

30. The process of claim 1 wherein the first catalyst component comprises 1, 1'-bis(4-triethylsilylphenyl)methylene-(cyclopentadienyl)(2,7-di-tertiary-butyl-9-fluorenyl)hafnium dimethyl and or 1, 1'-bis(4-triethylsilylphenyl)methylene-(cyclopentadienyl)(3,8-di-tertiary-butyl-fluorenyl)hafnium dimethyl.
31. The process of claim 1 wherein the second catalyst component comprises dimethylsilyl bis(2-methyl-5-phenylindenyl) zirconium dimethyl and/or dimethylsilyl bis(2-methyl-4-phenylindenyl) zirconium dimethyl.
32. The process of claim 1 wherein the activator comprises dimethylaniliniumtetrakis(pentafluorophenyl) borate and or trityl tetrakis(pentafluorophenyl) borate.
33. The process of claim 1 wherein the activator comprises an alumoxane.
34. The process of claim 1 wherein the activator comprises an ionizing compound.
35. The process of claim 1 wherein the activator comprises a non-coordinating anion.
36. The process of claim 1 wherein the activator comprises one or more of methylalumoxane, trimethylammonium tetraphenylborate, triethylammonium tetraphenylborate, tripropylammonium tetraphenylborate, tri(n-butyl)ammonium tetraphenylborate, tri(t-butyl)ammonium tetraphenylborate, N,N-dimethylanilinium tetraphenylborate, N,N-diethylanilinium tetraphenylborate, N,N-dimethyl-(2,4,6-trimethylanilinium) tetraphenylborate, trimethylammonium tetrakis(pentafluorophenyl)borate, triethylammonium tetrakis(pentafluorophenyl)borate, tripropylammonium tetrakis(pentafluorophenyl)borate, tri(n-butyl)ammonium tetrakis(pentafluorophenyl)borate, tri(sec-butyl)ammonium tetrakis(pentafluorophenyl) borate, N,N-dimethylanilinium tetrakis(pentafluorophenyl) borate,

N,N-diethylanilinium tetrakis(pentafluorophenyl) borate,
 N,N-dimethyl-(2,4,6-trimethylanilinium) tetrakis(pentafluorophenyl) borate, trimethylammonium tetrakis-(2,3,4,6-tetrafluorophenyl)borate,
 triethylammonium tetrakis-(2,3,4,6-tetrafluorophenyl) borate,
 tripropylammonium tetrakis-(2,3,4,6-tetrafluorophenyl) borate,
 tri(n-butyl)ammonium tetrakis-(2,3,4,6-tetrafluoro-phenyl) borate,
 dimethyl(t-butyl)ammonium tetrakis-(2,3,4,6-tetrafluorophenyl) borate,
 N,N-dimethylanilinium tetrakis-(2,3,4,6-tetrafluorophenyl) borate,
 N,N-diethylanilinium tetrakis-(2,3,4,6-tetrafluorophenyl) borate, and
 N,N-dimethyl-(2,4,6-trimethylanilinium)tetrakis-(2,3,4,6-tetrafluorophenyl) borate;
 di-(i-propyl)ammonium tetrakis(pentafluorophenyl) borate;
 dicyclohexylammonium tetrakis(pentafluorophenyl) borate;
 triphenylphosphonium tetrakis(pentafluorophenyl) borate;
 tri(o-tolyl)phosphonium tetrakis(pentafluorophenyl) borate; and
 tri(2,6-dimethylphenyl)phosphonium tetrakis(pentafluorophenyl) borate.

37. The process of claim 1 wherein the first catalyst component is capable of polymerizing macromonomers having reactive termini; and the second catalyst is capable of producing macromonomers having reactive termini.

38. The process of claim 1 wherein one or more reaction zones further comprise diolefin.

39. The process of claim 1 wherein one or more reaction zones further comprise one or more C4 to C40 dienes.

40. The process of claim 1 wherein one or more reaction zones further comprise one or more dienes selected from the group consisting of 1,6-heptadiene, 1,7-octadiene, 1,8-nonadiene, 1,9-decadiene, 1,10-undecadiene, 1,11-dodecadiene, 1,12-tridecadiene, 1,13-tetradecadiene, cyclopentadiene, vinylnorbornene, norbornadiene, ethylidene norbornene, divinylbenzene, dicyclopentadiene, polybutadienes having an Mw less than 1000 g/mol, or combinations thereof.

41. The process of claim 1 wherein at least one reaction zone is a gas phase reactor.

42. The process of claim 1 wherein at least one reaction zone is a solution phase reactor.

43. The process of claim 1 wherein at least one reaction zone is a slurry phase reactor.
44. The process of claim 1 wherein all of the reaction zones are a solution phase reactor.
45. The process of claim 1 wherein the first catalyst component, the second catalyst component and the activator comprise one or more of the following combinations (where Me equals methyl, Ph equals phenyl, Et equals ethyl, Cp equals cyclopentadienyl, 3,8-di-t-BuFlu equals 3,8-di-tert-butylfluorenyl, 2-Me-4-PhInd equals 2-methyl-4-phenylindenyl, 2-MeInd means 2-methylindenyl, c-C₁₂H₂₃ equals cyclododecyl, Me₄C₅ - tetramethylcyclopentadienyl, H₄Ind equals tetrahydroindenyl, and Ind equals indenyl):
- (1) Me₂Si(Me₄C₅)(N-c-C₁₂H₂₃)TiCl₂ and rac-Me₂Si(2-Me-4-PhInd)₂ZrCl₂ activated with an alumoxane;
- (2) Me₂Si(Me₄C₅)(N-c-C₁₂H₂₃)TiMe₂ and rac-Me₂Si(2-Me-4-PhInd)₂ZrMe₂ activated with a non-coordinating anion activator,
- (2a) Me₂Si(Me₄C₅)(N-c-C₁₂H₂₃)TiMe₂ and rac-Me₂Si(2-Me-4-PhInd)₂ZrMe₂ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;
- (3) Me₂Si(Me₄C₅)(N-c-C₁₂H₂₃)TiCl₂ and rac-Me₂Si(2-MeInd)₂ZrCl₂ activated with an alumoxane;
- (4) Me₂Si(Me₄C₅)(N-c-C₁₂H₂₃)TiMe₂ and rac-Me₂Si(2-MeInd)₂ZrMe₂ activated with a non-coordinating anion activator;
- (4a) Me₂Si(Me₄C₅)(N-c-C₁₂H₂₃)TiMe₂ and rac-Me₂Si(2-MeInd)₂ZrMe₂ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;
- (5) Me₂Si(Me₄C₅)(N-1-adamantyl)TiCl₂ and rac-Me₂Si(2-Me-4-PhInd)₂ZrCl₂ activated with an alumoxane;

(6) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-1-adamantyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(6a) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-1-adamantyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(7) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-1-adamantyl})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrCl}_2$ activated with an alumoxane;

(8) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-1-adamantyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(8a) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-1-adamantyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(9) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-butyl})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrCl}_2$ activated with an alumoxane;

(10) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-butyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(10a) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-butyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(11) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-butyl})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})$ activated with an alumoxane;

(12) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-butyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(12a) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-t-butyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(13) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-exo-norbornyl})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrCl}_2$ activated with an alumoxane;

(14) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-exo-norbornyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(14a) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-exo-norbornyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(15) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-exo-norbornyl})\text{TiCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrCl}_2$ activated with an alumoxane;

(16) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-exo-norbornyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(16a) $\text{Me}_2\text{Si}(\text{Me}_4\text{C}_5)(\text{N-exo-norbornyl})\text{TiMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(17) $(\text{p-Et}_3\text{SiPh})_2\text{C}(\text{Cp})(3,8\text{-di-t-BuFlu})\text{HfCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrCl}_2$ activated with an alumoxane;

(18) $(\text{p-Et}_3\text{SiPh})_2\text{C}(\text{Cp})(3,8\text{-di-t-BuFlu})\text{HfMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(18a) $(\text{p-Et}_3\text{SiPh})_2\text{C}(\text{Cp})(3,8\text{-di-t-BuFlu})\text{HfMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(19) $(p\text{-Et}_3\text{SiPh})_2\text{C}(\text{Cp})(3,8\text{-di-}t\text{-BuFlu})\text{HfCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrCl}_2$ activated with an alumoxane;

(20) $(p\text{-Et}_3\text{SiPh})_2\text{C}(\text{Cp})(3,8\text{-di-}t\text{-BuFlu})\text{HfMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(20a) $(p\text{-Et}_3\text{SiPh})_2\text{C}(\text{Cp})(3,8\text{-di-}t\text{-BuFlu})\text{HfMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(21) $\text{meso-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si}(\text{H}_4\text{Ind})_2\text{ZrCl}_2$ activated with an alumoxane;

(22) $\text{meso-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si}(\text{H}_4\text{Ind})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(22a) $\text{meso-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si}(\text{H}_4\text{Ind})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(23) $\text{meso-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrCl}_2$ activated with an alumoxane;

(24) $\text{meso-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(24a) $\text{meso-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si}(2\text{-MeInd})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(25) $\text{meso-Me}_2\text{Si}(\text{Ind})_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si}(\text{H}_4\text{Ind})_2\text{ZrCl}_2$ activated with an alumoxane;

(26) $\text{meso-Me}_2\text{Si}(\text{Ind})_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si}(\text{H}_4\text{Ind})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(26a) $\text{meso-Me}_2\text{Si(Ind)}_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si(H}_4\text{Ind)}_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(27) $\text{meso-Me}_2\text{Si(Ind)}_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si(2-MeInd)}_2\text{ZrCl}_2$ activated with an alumoxane;

(28) $\text{meso-Me}_2\text{Si(Ind)}_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(28a) $\text{meso-Me}_2\text{Si(Ind)}_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(29) $\text{meso-Me}_2\text{Si(2-MeInd)}_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si(2-Me-4-PhInd)}_2\text{ZrCl}_2$ activated with an alumoxane;

(30) $\text{meso-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si(2-Me-4-PhInd)}_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(30a) $\text{meso-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si(2-Me-4-PhInd)}_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(31) $\text{meso-Me}_2\text{Si(2-MeInd)}_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si(2-MeInd)}_2\text{ZrCl}_2$ activated with an alumoxane;

(32) $\text{meso-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(32a) $\text{meso-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(33) $\text{meso-CH}_2\text{CH}_2\text{(2-MeInd)}_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si(2-Me-4-PhInd)}_2\text{ZrCl}_2$ activated with an alumoxane;

(34) meso-CH₂CH₂(2-MeInd)₂ZrMe₂ and rac-Me₂Si(2-Me-4-PhInd)₂ZrMe₂ activated with a non-coordinating anion activator;

(34a) meso-CH₂CH₂(2-MeInd)₂ZrMe₂ and rac-Me₂Si(2-Me-4-PhInd)₂ZrMe₂ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(35) meso-CH₂CH₂(2-MeInd)₂ZrCl₂ and rac-Me₂Si(2-MeInd)₂ZrCl₂ activated with an alumoxane;

(36) meso-CH₂CH₂(2-MeInd)₂ZrMe₂ and rac-Me₂Si(2-MeInd)₂ZrMe₂ activated with a non-coordinating anion activator;

(36a) meso-CH₂CH₂(2-MeInd)₂ZrMe₂ and rac-Me₂Si(2-MeInd)₂ZrMe₂ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(37) meso-Me₂Si(2-Me-4-PhInd)₂ZrCl₂ and rac-Me₂Si(2-Me-4-PhInd)₂ZrCl₂ activated with an alumoxane;

(38) meso-Me₂Si(2-Me-4-PhInd)₂ZrMe₂ and rac-Me₂Si(2-Me-4-PhInd)₂ZrMe₂ activated with a non-coordinating anion activator;

(38a) meso-Me₂Si(2-Me-4-PhInd)₂ZrMe₂ and rac-Me₂Si(2-Me-4-PhInd)₂ZrMe₂ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(39) meso-CH₂CH₂(2-Me-4-PhInd)₂ZrCl₂ and rac-CH₂CH₂(2-Me-4-PhInd)₂ZrCl₂ activated with an alumoxane;

(40) meso-CH₂CH₂(2-Me-4-PhInd)₂ZrMe₂ and rac-CH₂CH₂(2-Me-4-PhInd)₂ZrMe₂ activated with a non-coordinating anion activator;

(40a) $\text{meso-CH}_2\text{CH}_2(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$ and $\text{rac-CH}_2\text{CH}_2(2\text{-Me-4-PhInd})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(41) $\text{meso-CH}_2\text{CH}_2(2\text{-MeInd})_2\text{ZrCl}_2$ and $\text{rac-CH}_2\text{CH}_2(2\text{-MePhInd})_2\text{ZrCl}_2$ activated with an alumoxane;

(42) $\text{meso-CH}_2\text{CH}_2(2\text{-MeInd})_2\text{ZrMe}_2$ and $\text{rac-CH}_2\text{CH}_2(2\text{-MeInd})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(42a) $\text{meso-CH}_2\text{CH}_2(2\text{-MeInd})_2\text{ZrMe}_2$ and $\text{rac-CH}_2\text{CH}_2(2\text{-MeInd})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(43) $\text{meso-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrCl}_2$ and $\text{rac-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrCl}_2$ activated with an alumoxane;

(44) $\text{meso-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrMe}_2$ and $\text{rac-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(44a) $\text{meso-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrMe}_2$ and $\text{rac-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(45) $\text{meso-Me}_2\text{Si}(\text{Ind})_2\text{ZrCl}_2$ and $\text{rac-Me}_2\text{Si}(\text{Ind})_2\text{ZrCl}_2$ activated with an alumoxane;

(46) $\text{meso-Me}_2\text{Si}(\text{Ind})_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si}(\text{Ind})_2\text{ZrMe}_2$ activated with a non-coordinating anion activator;

(46a) $\text{meso-Me}_2\text{Si}(\text{Ind})_2\text{ZrMe}_2$ and $\text{rac-Me}_2\text{Si}(\text{Ind})_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(47) $\text{meso-CH}_2\text{CH}_2(\text{Ind})_2\text{ZrCl}_2$ and $\text{rac-CH}_2\text{CH}_2(4,7\text{-Me}_2\text{Ind})_2\text{ZrCl}_2$ (4,7-Me₂Ind = 4,7-dimethylindenyl) activated with an alumoxane;

(48) meso-CH₂CH₂(Ind)₂ZrMe₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrMe₂ activated with a non-coordinating anion activator;

(48a) meso-CH₂CH₂(Ind)₂ZrMe₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrMe₂ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(49) meso-Me₂Si(Ind)₂ZrCl₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrCl₂ activated with an alumoxane;

(50) meso-Me₂Si(Ind)₂ZrMe₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrMe₂ activated with a non-coordinating anion activator;

(50a) meso-Me₂Si(Ind)₂ZrMe₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrMe₂ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(51) meso-CH₂CH₂(2-MeInd)₂ZrCl₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrCl₂ (4,7-Me₂Ind = 4,7-dimethylindenyl) activated with an alumoxane;

(52) meso-CH₂CH₂(2-MeInd)₂ZrMe₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrMe₂ activated with a non-coordinating anion activator;

(52a) meso-CH₂CH₂(2-MeInd)₂ZrMe₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrMe₂ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron;

(53) meso-Me₂Si(2-MeInd)₂ZrCl₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrCl₂ activated with an alumoxane;

(54) meso-Me₂Si(2-MeInd)₂ZrMe₂ and rac-CH₂CH₂(4,7-Me₂Ind)₂ZrMe₂ activated with a non-coordinating anion activator; or

(54a) $\text{meso-Me}_2\text{Si(2-MeInd)}_2\text{ZrMe}_2$ and $\text{rac-CH}_2\text{CH}_2\text{(4,7-Me}_2\text{Ind)}_2\text{ZrMe}_2$ activated with N,N-dimethylanilinium tetrakis(pentafluorophenyl)boron and or triphenylcarbonium tetrakis(pentafluorophenyl)boron.

46. The process of claim 1 wherein:

- 1) the first catalyst component is capable of producing a polymer having a weight average molecular weight of 80,000 or less and a crystallinity of 15% or less under selected polymerization conditions;
- 2) the a second catalyst component is capable of producing polymer having a weight average molecular weight of 80,000 or less and a crystallinity of 50% or more at the selected polymerization conditions;
- 3) the temperature in the reaction zones is greater than 105°C;
- 4) the residence time in the reaction zones is 10 minutes or less;
- 5) the ratio of the first catalyst to the second catalyst is from 1:1 to 20:1;
- 6) wherein the activity of the catalyst components is at least 100 kilograms of polymer per gram of the catalyst components; and wherein at least 80% of the olefins are converted to polymer.

47. The process of claim 46 wherein:

- a) the olefins comprise propylene and one or more of butene, pentene, hexene, heptene, octene; nonene, decene, dodecene; and
- b) the temperature is greater than 110°C; and
- c) the residence time is 5 minutes or less; and
- d) the ratio of the first catalyst to the second catalyst is from 1:1 to 1:10.

48. The process of claim 1 wherein the step of recovering a branched olefin polymer comprising at least 50 mole% of one or more C3 to C40 olefins comprises:

- 1) withdrawing polymer solution from the reaction zone;
- 2) removing at least 10% solvent from the polymer solution;
- 3) quenching the reaction;
- 4) devolatilizing the polymer solution to form molten polymer;
- 5) combining the molten polymer and one or more additives in a static mixer;
- 6) removing the polymer combination from the static mixer; and
- 7) pelletizing or drumming the polymer combination.

49. The process of claim 48 wherein the additives in step 5) comprise nucleating agent.

50. The process of claim 1 wherein the second catalyst component comprises one or more of:

dimethylsiladiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;

dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;

dimethylsiladiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂
 hafnium dichloride;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂
 hafnium dichloride;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;

dimethylsiladiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-methyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂
 hafnium dimethyl;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂
 hafnium dimethyl;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;

dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-isobutyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;

dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylsiladiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;

9-silafluorendiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 9-silafluorendiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 dimethylsiladiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylsiladiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;

9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 9-silafluorendiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dichloride;
 9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂hafnium dimethyl;
 dimethylsiladiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylsiladiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylsiladiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

9-silafluorendiyl(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;
 9-silafluorendiyl(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 dimethylamidoborane(2-iso-propyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 dimethylamidoborane(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 dimethylamidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 dimethylamidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 dimethylamidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 dimethylamidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 dimethylamidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;
 dimethylamidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

dimethylamidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ zirconium dimethyl;

dimethylamidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ zirconium dimethyl;

dimethylamidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ zirconium dimethyl;

dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ zirconium dimethyl;

dimethylamidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dimethyl;
 dimethylamidoborane(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dimethyl;
 dimethylamidoborane(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dimethyl;
 dimethylamidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dimethyl;
 dimethylamidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dimethyl;
 dimethylamidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl
 dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 dimethylamidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;

diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 diisopropylamidoborane(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 diisopropylamidoborane(2-iso-propyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 diisopropylamidoborane(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 diisopropylamidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 diisopropylamidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 diisopropylamidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;
 diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

diisopropylamidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl) η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η^4 -1,4-diphenyl-1,3-butadiene;

diisopropylamidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ zirconium dimethyl;

diisopropylamidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ zirconium dimethyl;

diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ zirconium dimethyl;

diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ zirconium dimethyl;

diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ zirconium dimethyl;

diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 diisopropylamidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;
 bis(trimethylsilyl)amidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dichloride;
 bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-bis-
 trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-bis-
 trifluoromethylphenyl]indenyl)₂zirconium dichloride;
 bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
 dichloride;
 bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium
 dichloride;

bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dichloride;

bis(trimethylsilyl)amidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-*t*-butylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-*tert*-butyl, 4-[3',5'-di-*t*-butylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-*n*-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-bis- trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-*n*-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-*tert*-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-*n*-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-*n*-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-*tert*-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂ η⁴-1,4-diphenyl-1,3-butadiene;

bis(trimethylsilyl)amidoborane(2-methyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-tbutylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-bis-trifluoromethylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl

bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-iso-propylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-methyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-ethyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-n-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-iso-propyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-n-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-iso-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl;

bis(trimethylsilyl)amidoborane(2-sec-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl; or

bis(trimethylsilyl)amidoborane(2-tert-butyl, 4-[3',5'-di-phenylphenyl]indenyl)₂zirconium dimethyl.

51. The process of claim 1 wherein the first catalyst component is also present in the second reaction zone.
52. The process of claim 1 wherein one catalyst component is present in at least one reaction zone and the other catalyst component is present in the second reaction zone.
53. The process of claim 1 wherein 3% or more of the combined catalyst components is not injected into the first reaction zone.
54. The process of claim 1 wherein hydrogen is introduced into a reaction zone.
55. The process of claim 1 wherein the temperature is different in the reaction zones.
56. The process of claim 1 wherein the temperature in a reaction zone is varied.