

## TANDEM TETRAMERISATION-POLYMERISATION OF OLEFINS

### Field of the invention

This invention relates to a tandem tetramerisation and polymerisation catalyst system and a process for the production of branched polyolefins wherein the polymerisation catalyst can be homogeneous or supported on a support medium.

### Background to the invention

This invention relates to catalyst systems that facilitate the selective production of 1-octene in combination with a suitable polymerisation catalyst so that the *in situ* polymerisation of the formed 1-octene with the available ethylene feedstock takes place. Preferably, ethylene tetramerisation and polymerisation take place at the same time under the same reaction conditions.

With regard to the oligomerisation catalysts used in this application that are used for the production of 1-octene, the art does not teach a commercially successful process for the tetramerisation of ethylene to produce 1-octene selectively. Conventional ethylene oligomerisation technologies produce a range of  $\alpha$ -olefins following either a Schulz-Flory or Poisson product distribution. By definition, these mathematical distributions limit the mass % of the tetramer that can be formed and make a distribution of products. In this regard, it is known from prior art (US patent 6,184,428) that a nickel catalyst comprising a chelating ligand, preferably 2-diphenyl phosphino benzoic acid (DPPBA), a nickel compound, preferably  $\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$ , and a catalyst activator, preferably sodium tetraphenylborate, catalyse the oligomerisation of ethylene to yield a mixture of linear olefins containing 1-octene. The selectivity towards linear  $\text{C}_8$   $\alpha$ -olefins is claimed to be 19%. Similarly the Shell Higher Olefins Process (SHOP process, US patents 3,676,523 and 3,635,937) using a similar catalyst system is reported to typically yield 11 mass % 1-octene in its product mixture (Chem Systems PERP reports 90-1, 93-6 and 94/95S12).

Ziegler-type technologies based on trialkylaluminium catalysts, independently developed by Gulf Oil Chemicals Company (Chevron, e.g. DE patent 1,443,927) and Ethyl Corporation (BP/Amoco, e.g. US patent 3,906,053), are also commercially used to

oligomerise ethylene to produce mixtures of olefins that reportedly contain 13-25 mass % 1-octene (Chem Systems PERP reports 90-1, 93-6, and 94/95S12).

The prior art also teaches that chromium-based catalysts containing heteroatomic ligands with both phosphorus and nitrogen heteroatoms selectively catalyse the trimerisation of ethylene to 1-hexene. Examples of such heteroatomic ligands for ethylene trimerisation include bis(2-diethylphosphino-ethyl) amine (WO 03/053891) as well as (o-methoxyphenyl)<sub>2</sub>PN(methyl)P(o-methoxyphenyl)<sub>2</sub> (WO 02/04119). Both these catalyst systems and processes are very specific for the production of 1-hexene and only yield 1-octene as an impurity (typically less than 3 mass % of the product mixture as disclosed by WO 02/04119). The coordinating phosphorus heteroatoms in (o-methoxyphenyl)<sub>2</sub>PN(methyl)P(o-methoxyphenyl)<sub>2</sub> (WO 02/04119) are spaced apart by one nitrogen atom. It is believed that the nitrogen atom does not coordinate, at least in the absence of an activator, with the chromium and that without any further electron donating atoms on the ligand that it is a bidentate system. Furthermore it is argued that the polar, or electron donating substituents in the ortho-position of the phenyl groups help form a tridentate system, which is generally believed to enhance selectivity towards 1-hexene formation as reiterated in WO 02/04119 in *Chem. Commun.*, 2002, 858-859: "This has led us to hypothesise that the potential for ortho-methoxy groups to act as pendent donors and increase the coordinative saturation of the chromium centre is an important factor." WO 02/04119 (Example 16) teaches the production of octenes using a trimerisation of olefins process and catalyst system. In this instance, 1-butene was co-trimerised with two ethylene molecules to give 25% octenes. However, the nature of these octenes was not disclosed and the applicant believes that they consist of a mixture of linear and branched octenes.

This application discloses the combination of highly selective tetramerisation cocatalysts for the production of 1-octene and the concomitant polymerisation of the tetramerised olefin with the ethylene feedstock. It has further been found that a chromium-based catalyst containing mixed heteroatomic ligands can be used to selectively tetramerise ethylene to 1-octene often in excess of 60 mass% selectivity. This high 1-octene selectivity cannot be achieved *via* conventional one-step ethylene oligomerisation technologies which at most yield 25 mass% 1-octene and the present invention confers the suitability of a selective 1-octene yielding catalyst for *in situ* polymerisation.

Examples of heterogeneous polymerisation processes that mention the polymerisation of *in situ* formed  $\alpha$ -olefins to yield branched by-products of the polymerisation process are known in the art. However, few examples are available describing the combination of a heterogeneous polymerisation catalyst on a support and a homogeneous oligomerisation catalyst, or the combination of two homogeneous catalyst systems where one system produces an  $\alpha$ -olefin and the other catalyst polymerises the olefin formed *in situ* as a comonomer with ethylene. Specific examples related to tandem oligomerisation and polymerisation catalysis involving a supported polymerisation catalyst include US patent 5,331,070 and EP 0 417 477 in which a metal alkyl, a pyrrole-containing compound and a chromium salt supported on an inorganic support material polymerises an *in situ* generated  $\alpha$ -olefin to yield a branched polyethylene.

Tandem catalysis involving heterogeneous polymerisation catalysts mostly does not involve the combination of a separate oligomerisation catalyst together with a supported polymerisation catalyst as proposed in this application. Due to the nature and geometry of the supported polymerisation catalyst, the formation of  $\alpha$ -olefins through  $\beta$ -hydride elimination is possible to some extent, however the formation of polymers is thermodynamically and kinetically more favoured. After an  $\alpha$ -olefin is produced it has a high probability to be incorporated into other growing polymer chains as the process of chain growth proceeds from the surface of the support material. In this way, polymers containing relatively low amounts of  $\alpha$ -olefins are produced without significant control over the extent of  $\alpha$ -olefin incorporation or the resultant polymer architecture.

Although examples of homogeneous tandem catalysis involving the combination of oligomerisation catalysts in tandem with suitable soluble polymerisation catalysts are rare, the recent patent literature such as US patents 6,586,541, 6,555,631 and 6,586,550 specifically relate to the deliberate combination of oligomerisation catalysts, previously reported for their ability to produce  $\alpha$ -olefins, combined with soluble polymerisation catalysts to produce branched polyolefins.

These patent examples centre on contacting both an oligomerisation catalyst and a polymerisation catalyst with a single feedstock namely ethylene. Both oligomers and

polymers are produced in the same system with concurrent production of  $\alpha$ -olefins and the polymerisation of the resultant  $\alpha$ -olefins with ethylene. Most of the oligomerisation catalysts that convey novelty to these patents are oligomerisation catalysts that produce a distribution of  $\alpha$ -olefins. These distributions are favoured towards a mixture of 1-butene, 1-hexene, 1-octene and 1-decene. Consequently, the structure of the polymers described in these patents exhibit patterns of incorporation according to  $^{13}\text{C}$  NMR analysis that reflect the distribution of  $\alpha$ -olefins.

It is well-known in the open literature that control of reaction conditions such as pressure, temperature, feed gas make-up, comonomer content etc. result in polymers with desirable physical properties such as tear-strength, optical clarity, elasticity and many other physical properties deemed desirable by polymer end-users. In particular, the production of linear low-density polyethylene (LLDPE) sees the polymerisation of individual high purity  $\alpha$ -olefins, such as 1-butene, 1-hexene and 1-octene, and ethylene to produce LLDPE suitable for films and sheets, blow molding, extrusion and wire and cable jacket material as well as rotational molding material. In none of the above examples is there disclosed a process relating to the selective production of 1-octene for *in situ* incorporation into polyethylene.

The combination of the highly selective ethylene tetramerisation catalyst system of the present disclosure with a suitable polymerisation catalyst can therefore result in a tandem process that produces polyethylene grades with high end-user specifications without the need of a separate facility or process to manufacture the 1-octene.

It will be understood that tandem oligomerisation and polymerisation catalysis includes *in situ* catalysis. In-situ catalysis means oligomerisation and polymerisation catalysis in the same reaction medium. The tandem oligomerisation and polymerisation catalysis may be *in situ* concurrent and/or *in situ* consecutive catalysis. The individual components of the oligomerisation and polymerisation catalysts may thus be added together simultaneously or sequentially, in any order, and in the presence or absence of monomer in a suitable solvent. Reaction mixture will be understood to include a reaction medium, reaction products and catalyst components. The reaction medium normally includes a solvent.

## Summary of the invention

This invention generally relates to the need for selectively producing 1-octene from ethylene and concomitant polymerisation of this monomer with ethylene can be satisfied by using a separate distinct transition metal-based ethylene tetramerisation catalyst system containing a heteroatomic ligand in tandem with a separate distinct polymerisation catalyst comprising a transition metal and hetero or homoatomic ligands.

By homoatomic is meant a ligand that consists entirely of similar atoms such as carbon that constitute the skeleton of the ligand such as the cyclopentadienyl ligand for example.

Thus, according to a first aspect of the invention, there is provided a process for polymerising olefins to produce branched polyolefins in the presence of a polymerisation catalyst and a cocatalyst, wherein the cocatalyst produces 1-octene in a selectivity greater than 30% and the 1-octene produced is at least partially incorporated into the polyolefin chain.

The invention allows the cocatalyst and/or process conditions to be selected to produce more than 40%, 50%, 60% or 70% 1-octene and the said 1-octene is at least partially incorporated into the polyolefin chain.

In this specification, % will be understood to be a mass %.

The branched polyolefin may be a branched polyethylene.

The branched polyolefin may be linear low density polyethylene (LLDPE).

The cocatalyst may be an ethylene tetramerisation catalyst, which includes a transition metal precursor and a heteroatomic ligand.

In this specification the term "tetramerisation" refers to a catalytic reaction of a single olefinic monomer or a mixture of olefinic monomers giving products enriched in

tetramers derived from those olefinic monomers. The tetramerisation product stream may consist of linear or branched olefins.

By heteroatomic tetramerisation ligand is meant a ligand that contains at least two heteroatoms, which can be the same or different, where the heteroatoms may be selected from phosphorus, arsenic, antimony, sulphur, oxygen, bismuth, selenium or nitrogen.

The heteroatomic ligand for the ethylene tetramerisation catalyst may be described by the following general formula  $(R)_nA-B-C(R)_m$  where A and C are independently selected from a group which comprises phosphorus, arsenic, antimony, oxygen, bismuth, sulphur, selenium, and nitrogen, and B is a linking group between A and C, and R is independently selected from any homo or heterohydrocarbyl group and n and m are determined by the respective valence and oxidation state of A and C.

A and/or C may be a potential electron donor for coordination with the transition metal.

An electron donor is defined as that entity that donates electrons used in chemical, including dative covalent, bond formation.

The heteroatomic ligand for the ethylene tetramerisation catalyst may be described by the following general formula  $(R^1)(R^2)A-B-C(R^3)(R^4)$  where A and C are independently selected from a group which comprises phosphorus, arsenic, antimony, bismuth and nitrogen and B is a linking group between A and C, and  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are independently selected from hydrocarbyl or heterohydrocarbyl or substituted hydrocarbyl or substituted heterohydrocarbyl groups.

The heteroatomic ligand for the ethylene tetramerisation catalyst may be described by the following general formula  $(R^1)(R^2)A-B-C(R^3)(R^4)$  where A and C are independently selected from a group which comprises phosphorus, arsenic, antimony, bismuth and nitrogen and B is a linking group between A and C, and  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are independently non-aromatic or aromatic, including heteroaromatic, groups.

R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> may be aromatic, including heteroaromatic, groups and preferably not all the groups R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup>, if aromatic, have a substituent on the atom adjacent to the atom bound to A or C.

R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> may have substituents on the atom adjacent to the atom bound to A or C.

At least one or more of R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> may be substituted with a polar substituent on a second or further atom from the atom bound to A or C.

Any polar substituents on R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> may preferably not be on the atom adjacent to the atom bound to A or C.

Any polar substituent on one or more of R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> may be electron-donating.

Any substituents on one or more of R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> may be not electron-donating.

In another embodiment, each non-electron donating substituent may be non-polar.

Each of R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> may be aromatic, including heteroaromatic, but not all of R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> are substituted by any substituent on an atom adjacent to the atom bound to A or C.

Preferably, not more than two of R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> may have substituents on the atom adjacent to the atom bound to A or C.

Polar is defined by IUPAC as an entity with a permanent electric dipole moment. Polar substituents include methoxy, ethoxy, isopropoxy, C<sub>3</sub>-C<sub>20</sub> alkoxy, phenoxy, pentafluorophenoxy, trimethylsiloxy, dimethylamino, methylsulfanyl, tosyl, methoxymethyl, methylthiomethyl, 1,3-oxazolyl, methoxymethoxy, hydroxyl, amino, phosphino, arsino, stibino, sulphate, nitro and the like.

Any of the groups in the ligand for the tetramerisation catalyst,  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  may independently be linked to one or more of each other or to the linking group B to form a cyclic structure together with A and C, A and B or B and C.

$R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  in the tetramerisation ligand may be independently selected from a group comprising a benzyl, phenyl, tolyl, xylyl, mesityl, biphenyl, naphthyl, anthracenyl, methoxy, ethoxy, phenoxy, tolyloxy, dimethylamino, diethylamino, methylethylamino, thiophenyl, pyridyl, thioethyl, thiophenoxy, trimethylsilyl, dimethylhydrazyl, methyl, ethyl, ethenyl, propyl, butyl, propenyl, propynyl, cyclopentyl, cyclohexyl, ferrocenyl and tetrahydrofuranlyl group. Preferably,  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  may independently be selected from a group comprising a phenyl, tolyl, biphenyl, naphthyl, thiophenyl and ethyl group.

B may be selected from any one of a group comprising: organic linking groups comprising a hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl and a substituted heterohydrocarbyl; inorganic linking groups comprising single atom links; ionic links and a group comprising methylene, dimethylmethylene, 1,2-ethane, 1,2-phenylene, 1,2-propane, 1,2-catechol, 1,2-dimethylhydrazine,  $-B(R^5)-$ ,  $-Si(R^5)_2-$ ,  $-P(R^5)-$  and  $-N(R^5)-$  where  $R^5$  is hydrogen, a hydrocarbyl or substituted hydrocarbyl, a substituted heteroatom or a halogen. Preferably, B may be  $-N(R^5)-$  and  $R^5$  is a hydrocarbyl or a substituted hydrocarbyl group.  $R^5$  may be hydrogen or may be selected from the groups consisting of alkyl, substituted alkyl, aryl, substituted aryl, aryloxy, substituted aryloxy, halogen, nitro, alkoxycarbonyl, carbonyloxy, alkoxy, aminocarbonyl, carbonylamino, dialkylamino, silyl groups or derivatives thereof, and aryl substituted with any of these substituents. Preferably  $R^5$  may be an isopropyl, a 1-cyclohexylethyl, a 2-methylcyclohexyl or a 2-octyl group.

In another embodiment of the invention any of the groups in the ligand for the tetramerisation catalyst,  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$  or  $R^5$  may include any cyclic heteroatomic group such as a cyclopentadienyl-dimethylsilyl-t-butyl group or a cyclic homoatomic group such as a cyclopentadienyl, indenyl or fluorene group.

B may be selected to be a single atom spacer. A single atom linking spacer is defined as a substituted or non-substituted atom that is bound directly to A and C.

A and/or C may be independently oxidised by S, Se, N or O.

A and C may be independently phosphorus or phosphorus oxidised by S or Se or N or O.

The tetramerisation ligand may also contain multiple  $(R)_nA-B-C(R)_m$  units. Not limiting examples of such ligands include dendrimeric ligands as well as ligands where the individual units are coupled either *via* one or more of the R groups or *via* the linking group B. More specific, but not limiting, examples of such ligands may include 1,2-di-(N(P(phenyl)<sub>2</sub>)<sub>2</sub>)-benzene, 1,4-di-(N(P(phenyl)<sub>2</sub>)<sub>2</sub>)-benzene, N(CH<sub>2</sub>CH<sub>2</sub>N(P(phenyl)<sub>2</sub>)<sub>2</sub>)<sub>3</sub>, 1,4-di-(P(phenyl)N(methyl)P(phenyl)<sub>2</sub>)-benzene, 1,2-di-(N(P(*p*-methoxyphenyl)<sub>2</sub>)<sub>2</sub>)-benzene, 1,4-di-(N(P(*p*-methoxyphenyl)<sub>2</sub>)<sub>2</sub>)-benzene, N(CH<sub>2</sub>CH<sub>2</sub>N(P(*p*-methoxyphenyl)<sub>2</sub>)<sub>2</sub>)<sub>3</sub> and 1,4-di-(P(*p*-methoxyphenyl)N(methyl)P(*p*-methoxyphenyl)<sub>2</sub>)-benzene.

The heteroatomic ligands for tetramerisation can be prepared using procedures known to one skilled in the art and procedures disclosed in published literature. Examples of ligands are: (phenyl)<sub>2</sub>PN(methyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(pentyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(phenyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(*p*-methoxyphenyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(*p*-*t*butylphenyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN((CH<sub>2</sub>)<sub>3</sub>-N-morpholine)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(Si(CH<sub>3</sub>)<sub>3</sub>)P(phenyl)<sub>2</sub>, (((phenyl)<sub>2</sub>P)<sub>2</sub>NCH<sub>2</sub>CH<sub>2</sub>)N, (ethyl)<sub>2</sub>PN(methyl)P(ethyl)<sub>2</sub>, (ethyl)<sub>2</sub>PN(isopropyl)P(phenyl)<sub>2</sub>, (ethyl)(phenyl)PN(methyl)P(ethyl)(phenyl), (ethyl)(phenyl)PN(isopropyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>P(=Se)N(isopropyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PCH<sub>2</sub>CH<sub>2</sub>P(phenyl)<sub>2</sub>, (*o*-ethylphenyl)(phenyl)PN(isopropyl)P(phenyl)<sub>2</sub>, (*o*-methylphenyl)<sub>2</sub>PN(isopropyl)P(*o*-methylphenyl)(phenyl), (phenyl)<sub>2</sub>PN(benzyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(1-cyclohexylethyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN[CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>Si(OMe)<sub>3</sub>]P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(cyclohexyl)P(phenyl)<sub>2</sub>, phenyl)<sub>2</sub>PN(2-methylcyclohexyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(allyl)P(phenyl)<sub>2</sub>, (2-naphthyl)<sub>2</sub>PN(methyl)P(2-naphthyl)<sub>2</sub>, (*p*-biphenyl)<sub>2</sub>PN(methyl)P(*p*-biphenyl)<sub>2</sub>, (*p*-methylphenyl)<sub>2</sub>PN(methyl)P(*p*-methylphenyl)<sub>2</sub>, (2-thiophenyl)<sub>2</sub>PN(methyl)P(2-thiophenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(methyl)N(methyl)P(phenyl)<sub>2</sub>, (*m*-methylphenyl)<sub>2</sub>PN(methyl)P(*m*-methylphenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(isopropyl)P(phenyl)<sub>2</sub> and (phenyl)<sub>2</sub>P(=S)N(isopropyl)P(phenyl)<sub>2</sub>.

Examples of ligands with polar substituents on R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> include: (*m*-methoxyphenyl)<sub>2</sub>PN(methyl)P(*m*-methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(methyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*m*-methoxyphenyl)<sub>2</sub>PN(isopropyl)P(*m*-methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(isopropyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(2-ethylhexyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*m*-methoxyphenyl)(phenyl)PN(methyl)P(phenyl)<sub>2</sub> and (*p*-methoxyphenyl)(phenyl)PN(methyl)P(phenyl)<sub>2</sub>, (*m*-methoxyphenyl)(phenyl)PN(methyl)P(*m*-methoxyphenyl)(phenyl), (*p*-methoxyphenyl)(phenyl)PN(methyl)P(*p*-methoxyphenyl)(phenyl), (*m*-methoxyphenyl)<sub>2</sub>PN(methyl)P(phenyl)<sub>2</sub> and (*p*-methoxyphenyl)<sub>2</sub>PN(methyl)P(phenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(1-cyclohexylethyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(2-methylcyclohexyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(decyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(pentyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(benzyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(phenyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-fluorophenyl)<sub>2</sub>PN(methyl)P(*p*-fluorophenyl)<sub>2</sub>, (*o*-fluorophenyl)<sub>2</sub>PN(methyl)P(*o*-fluorophenyl)<sub>2</sub>, (*p*-dimethylamino-phenyl)<sub>2</sub>PN(methyl)P(*p*-dimethylamino-phenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(allyl)P(*p*-methoxyphenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(isopropyl)P(*o*-methoxyphenyl)<sub>2</sub>, (*p*-(*p*-methoxyphenyl)-phenyl)<sub>2</sub>PN(isopropyl)P(*p*-(*p*-methoxyphenyl)-phenyl)<sub>2</sub> and (*p*-methoxyphenyl)(phenyl)PN(isopropyl)P(phenyl)<sub>2</sub>.

The heteroatomic tetramerisation ligand can be modified to be attached to a polymer chain so that the resulting heteroatomic coordination complex of the transition metal is soluble at elevated temperatures, but becomes insoluble at 25°C. This approach would enable the recovery of the complex from the reaction mixture for reuse and has been used for other catalyst as described by D.E. Bergbreiter *et al.*, *J. Am. Chem. Soc.*, 1987, **109**, 177-179. In a similar vein these transition metal complexes can also be immobilised by binding the heteroatomic ligands to silica, silica gel, polysiloxane or alumina backbone as, for example, demonstrated by C. Yuanyin *et al.*, *Chinese J. React. Pol.*, 1992, **1**(2), 152-159 for immobilising platinum complexes.

With regard to the tetramerisation process, the process may include the step of generating a heteroatomic tetramerisation coordination complex from a transition metal precursor and a heteroatomic ligand and adding a polymerisation catalyst either at the same time or over a period of time. The process may include the step of adding a pre-

formed coordination complex, prepared using a heteroatomic ligand and a transition metal precursor, to a reaction mixture, or the step of adding separately to the reactor, a heteroatomic ligand and a transition metal precursor such that a heteroatomic coordination complex of a transition metal is generated *in situ*. In all cases, the polymerisation catalyst may be added to the reactor at the same time as the tetramerisation catalyst or the polymerisation catalyst may be fed sequentially into the reactor over a period of time. By generating a heteroatomic coordination complex *in situ* is meant that the complex is generated in the medium in which catalysis takes place. Typically, the heteroatomic coordination complex is generated *in situ*. Typically, the transition metal precursor, and heteroatomic ligand are combined (both *in situ* and *ex situ*) to provide metal/ligand ratios from about 0.01:100 to 10 000:1, and preferably, from about 0.1:1 to 10:1.

The tetramerisation cocatalyst, as defined previously to include a transition metal precursor and a heteroatomic ligand, can be combined with the polymerisation catalyst in the ratio 0.01:100 to 10 000:1, and preferably, from about 1:1 to 100:1.

With regard to the tetramerisation transition metal precursor, the transition metal may be selected from any one of a group comprising chromium, molybdenum, tungsten, titanium, tantalum, vanadium and zirconium, preferably chromium.

With regard to the tetramerisation catalyst, the transition metal precursor which, upon mixing with the heteroatomic ligand and an activator, catalyses ethylene tetramerisation in accordance with the invention, may be a simple inorganic or organic salt, a coordination or organometallic complex and may be selected from any one of a group comprising chromium trichloride tris-tetrahydrofuran complex, (benzene)-tricarbonyl chromium, chromium (III) octanoate, chromium hexacarbonyl, chromium (III) acetylacetonate and chromium (III) 2-ethylhexanoate. The preferred transition metal precursors include chromium (III) acetylacetonate and chromium (III) 2-ethylhexanoate.

The process may include the step of combining in any order a heteroatomic ligand with a transition metal precursor, a polymerisation catalyst and an activator.

Examples of suitable polymerisation catalysts for the polymerisation of ethylene and the *in situ* synthesised 1-octene include but are not limited to the following:

### ZIEGLER-NATTA CATALYSTS

$\text{TiCl}_3\text{-Et}_2\text{AlCl}$ ,  $\text{AlR}_3\text{-TiCl}_4$ , wherein R belongs to the group comprising of alkyl, substituted alkyl, cyclic alkyl, aryl, substituted aryl, alkene and substituted alkene, and the like.

### UNBRIDGED METALLOCENES

Bis(cyclopentadienyl)-chromium(II), bis(cyclopentadienyl)-zirconium chloride hydride, bis(cyclopentadienyl)-titanium dichloride, bis(cyclopentadienyl)-zirconium dichloride, bis(cyclopentadienyl)-zirconium dimethyl, bis(*n*-butylcyclopentadienyl)-zirconium dichloride, bis(*n*-dodecylcyclopentadienyl)-zirconium dichloride, bis(ethylcyclopentadienyl)-zirconium dichloride, bis(iso-butylcyclopentadienyl)-zirconium dichloride, bis(isopropylcyclopentadienyl)-zirconium dichloride, bis(methylcyclopentadienyl)-zirconium dichloride, bis(*n*-octylcyclopentadienyl)-zirconium dichloride, bis(*n*-pentylcyclopentadienyl)-zirconium dichloride, bis(*n*-propylcyclopentadienyl)-zirconium dichloride, bis(trimethylsilylcyclopentadienyl)-zirconium dichloride, bis(1,3-bis(trimethylsilyl)cyclopentadienyl)-zirconium dichloride, bis(1-ethyl-3-methylcyclopentadienyl)-zirconium dichloride, bis(pentamethylcyclopentadienyl)-zirconium dichloride, bis(pentamethylcyclopentadienyl)-zirconium dimethyl, bis(1-propyl-3-methylcyclopentadienyl)-zirconium dichloride, bis(4,7-dimethylindenyl)-zirconium dichloride, bis(indenyl)-zirconium dichloride, bis(2-methylindenyl)-zirconium dichloride, bis(2-methylindenyl)-zirconium dichloride, cyclopentadienylindenyl-zirconium dichloride

### HALF SANDWICH METALLOCENES

Cyclopentadienyl-zirconium trichloride, pentamethylcyclopentadienyl titanium trichloride, pentamethylcyclopentadienyl-titanium trimethoxide, trimethyl-titanium-pentamethylcyclopentadienide, pentamethylcyclopentadienyl-zirconium trichloride, tetramethylcyclopentadienyl-zirconium trichloride, 1,2,4-trimethylcyclopentadienyl-zirconium trichloride

## CARBON-BRIDGED METALLOCENES

Diphenylmethylenedicyclopentadienyl-(9-fluorenyl)-zirconium dichloride, diphenylmethylenedicyclopentadienyl-(indenyl)-zirconium dichloride, isopropylidene-bis(cyclopentadienyl)-zirconium dichloride, isopropylidene-(cyclopentadienyl)(9-fluorenyl)-zirconium dichloride, isopropylidene-(3-methylcyclopentadienyl)-(9-fluorenyl)-zirconium dichloride, ethylene-bis(9-fluorenyl)-zirconium dichloride, *meso*-ethylene-bis(1-indenyl)-zirconium dichloride, *rac*-ethylene-bis(1-indenyl)-zirconium dichloride, *rac*-ethylene-bis(1-indenyl)-zirconium dimethyl, *rac*-ethylene-bis(2-methyl-1-indenyl)-zirconium dichloride, *rac*-ethylene-bis(4,5,6,7-tetrahydro-1-indenyl)-zirconium dichloride

## SILYL-BRIDGED METALLOCENES

Dimethylsilyl-bis(cyclopentadienyl)-zirconium dichloride, dimethylsilyl-bis(9-fluorenyl)-zirconium dichloride, *rac*-dimethylsilyl-bis(1-indenyl)-zirconium dichloride, *meso*-dimethylsilyl-bis(2-methylindenyl)-zirconium dichloride, *rac*-dimethylsilyl-bis(2-methylindenyl)-zirconium dichloride, *rac*-dimethylsilyl-bis(tetrahydroindenyl)-zirconium dichloride, dimethylsilyl-bis(tetramethylcyclopentadienyl)-zirconium dichloride, diphenylsilyl(cyclopentadienyl)(9-fluorenyl)-zirconium dichloride, diphenylsilyl-bis(indenyl)hafnium dichloride.

The activator for use in the process may in principle be any compound that generates an active catalyst when combined with the heteroatomic ligand and the transition metal precursor. Mixtures of activators may also be used. Suitable compounds include organoaluminium compounds, organoboron compounds, organic salts, such as methyl lithium and methylmagnesium bromide, inorganic acids and salts, such as tetrafluoroboric acid etherate, silver tetrafluoroborate, sodium hexafluoroantimonate and the like.

Suitable organoaluminium compounds include compounds of the formula  $\text{AlR}_3$ , where each R is independently a  $\text{C}_1\text{-C}_{12}$  alkyl, an oxygen containing moiety or a halide, and compounds such as  $\text{LiAlH}_4$  and the like. Examples include trimethylaluminium (TMA),

triethylaluminium (TEA), tri-isobutylaluminium (TIBA), tri-*n*-octylaluminium, methylaluminium dichloride, ethylaluminium dichloride, dimethylaluminium chloride, diethylaluminium chloride, aluminium isopropoxide, ethylaluminiumsesquichloride, methylaluminiumsesquichloride, and aluminoxanes. Aluminoxanes are well known in the art as oligomeric compounds which can be prepared by the controlled addition of water to an alkylaluminium compound, for example trimethylaluminium. Such compounds can be linear, cyclic, cages or mixtures thereof. Mixtures of different aluminoxanes may also be used in the process.

Examples of suitable organoboron compounds are boroxines,  $\text{NaBH}_4$ , triethylborane, tris(pentafluorophenyl)borane, tributyl borate and the like.

The activator may also be or contain a compound that acts as a reducing or oxidising agent, such as sodium or zinc metal and the like, or oxygen and the like.

The activator may be selected from alkylaluminoxanes such as methylaluminoxane (MAO) and ethylaluminoxane (EAO) as well as modified alkylaluminoxanes such as modified methylaluminoxane (MMAO). Modified methylaluminoxane (a commercial product from Akzo Nobel) contains modifier groups such as isobutyl or *n*-octyl groups, in addition to methyl groups.

The transition metal precursor of the tetramerisation catalyst and the aluminoxane may be combined in proportions to provide Al/metal ratios from about 1:1 to 10 000:1, preferably from about 1:1 to 1000:1, and more preferably from 1:1 to 300:1.

The process may include the step of adding to the catalyst system (including both the polymerisation and olefin tetramerisation catalysts) a trialkylaluminium compound in amounts of between 0.01 to 1000 mol per mol of alkylaluminoxane.

It should be noted that aluminoxanes generally also contain considerable quantities of the corresponding trialkylaluminium compounds used in their preparation. The presence of these trialkylaluminium compounds in aluminoxanes can be attributed to their incomplete hydrolysis with water. Any quantity of a trialkylaluminium compound quoted

in this disclosure is additional to alkylaluminium compounds contained within the aluminoxanes.

The activator may also be a binuclear cocatalyst such as a bisborane 1,4-(C<sub>6</sub>F<sub>5</sub>)<sub>2</sub>B(C<sub>6</sub>F<sub>4</sub>)B-(C<sub>6</sub>F<sub>5</sub>)<sub>2</sub>. This binuclear cocatalyst may be combined with the tetramerisation and polymerisation catalysts in such a way that a bimolecular species may be formed that has metal centres in close molecular proximity.

The catalyst systems or its individual components, in accordance with the invention, or its individual components, may also be immobilised by supporting it on a support material, for example, silica, alumina, MgCl<sub>2</sub>, zirconia, artificial hectorite or smectorite clays such as Laponite<sup>TM</sup> RD or mixtures thereof, or on a polymer, for example polyethylene, polypropylene, polystyrene, or poly(aminostyrene). The catalyst can be formed *in situ* in the presence of the support material, or the support can be pre-impregnated or premixed, simultaneously or sequentially, with one or more of the catalyst components or the polymerisation catalyst. In some cases, the support material can also act as a component of the activator. This approach would also facilitate the recovery of the catalyst from the reaction mixture for reuse. The concept was, for example, successfully demonstrated with a chromium-based ethylene trimerisation catalyst by T. Monoi and Y. Sasaki, *J. Mol. Cat.A: Chem.*, 1987, **109**, 177-179. In some cases, the support can also act as a catalyst component, for example where such supports contain aluminoxane functionalities or where the support is capable of performing similar chemical functions as an aluminoxane, which is for instance the case with IOLA<sup>TM</sup> (a commercial product from Grace Davison).

According to a further aspect of the invention, there is provided a tandem oligomerisation and polymerisation process for the oligomerisation of ethylene to produce an olefinic product stream and the *in situ* polymerisation of the oligomeric olefin with ethylene, the process including the step of contacting the olefins with an oligomerisation catalyst, which catalyst includes a heteroatomic ligand and a transition metal precursor, and a polymerisation catalyst, which catalyst includes a transition metal and and/or a homo- or heteroatomic ligand.

The oligomerisation catalyst may be an ethylene tetramerisation catalyst.

According to yet another aspect of the invention the tetramerisation process includes the step of contacting an olefinic feedstream with a catalyst system which includes a transition metal and a heteroatomic ligand and wherein the product of the tetramerisation process is an olefin and the olefin is concomitantly copolymerised with the olefinic feedstream by contacting the olefins produced in the first step and the olefinic feedstream with a catalyst comprising of a transition metal and/or a homo- or heteroatomic ligand.

By homoatomic is meant a ligand that consists entirely of similar atoms such as carbon that constitute the skeleton of the ligand such as the cyclopentadienyl ligand for example.

The ethylene to be tetramerised and polymerised can be introduced into the process according to the invention in a continuous or batch fashion.

The tandem catalysis product stream will be understood to include a polymer, which polymer is produced according to the invention in a continuous or batch fashion.

The process may include a process for tetramerisation of ethylene to selectively yield 1-octene.

The process may be a process for tetramerisation of ethylene and concomitant polymerisation of the produced 1-octene with ethylene .

The ethylene may be contacted with the catalyst system comprising of the tetramerisation catalyst and the polymerisation catalyst at a pressure of 1 barg, preferably greater than 10 barg, more preferably greater than 30 barg.

The tandem catalysis process may include the step of mixing the components of the catalyst systems (both the tetramerisation catalyst and the polymerisation catalyst) at any temperature between -20°C and 250°C in the presence of an olefin. The preferred temperature range being 20 °C - 100 °C.

The individual components of both catalyst systems described herein may be combined simultaneously or sequentially in any order, and in the presence or absence of a solvent, in order to give active catalysts. The presence of an olefin during the mixing of the catalyst components generally provides a protective effect which may result in improved catalyst performance. The preferred temperature range may be between 20°C and 100°C.

The reaction products derived from the tandem catalysis process as described herein, may be prepared using the disclosed catalyst system by a homogeneous liquid phase reaction in the presence or absence of an inert solvent, and/or by slurry reaction where the catalyst system is in a form that displays little or no solubility, and/or a two-phase liquid/liquid reaction, and/or a bulk phase reaction in which neat reagent and/or product olefins serve as the dominant medium, and/or gas phase reaction, using conventional equipment and contacting techniques.

The tandem catalysis process may also be carried out in an inert solvent. Any inert solvent that does not react with the activator can be used. These inert solvents may include, saturated aliphatic, unsaturated aliphatic, aromatic hydrocarbon and halogenated hydrocarbon. Typical solvents include, but are not limited to, benzene, toluene, xylene, cumene, heptane, methylcyclohexane, methylcyclopentane, cyclohexane, ionic liquids and the like.

The tandem catalysis process may be carried out at pressures from atmospheric to 500 barg. Ethylene pressures in the range of 10-70 barg are preferred. Particularly preferred pressures range from 30-50 barg.

The tandem catalysis process may be carried out at temperatures from -20 °C - 250 °C. Temperatures in the range of 15-130 °C are preferred. Particularly preferred temperatures range from 35-150°C.

The tandem catalysis process may be carried out in a plant which includes any type of reactor. Examples of such reactors include, but are not limited to, batch reactors, semi-batch reactors and continuous reactors. The plant may include, in combination a) a reactor, b) at least one inlet line into this reactor for olefin reactant and the catalyst

system, c) effluent lines from this reactor for polymerisation reaction products, and d) at least one separator to separate the desired polymerisation reaction products, wherein the catalyst system may include a heteroatomic ligand, a transition metal precursor, a polymerisation catalyst and an activator, as described herein.

In another embodiment of this invention, a combination of reactors may be preferred where the first reactor may be a continuously stirred tank reactor and the second reactor may be a batch, continuously stirred tank reactor or a plug-flow reactor.

### **Examples of performing the invention**

The invention will now be described with reference to the following non-limiting examples. The individual components of the examples may conceivably be omitted or substituted and, although not necessarily ideal, the invention may conceivably still be performed and these components are not to be taken as essential to the working of the invention.

In the examples that follow all procedures were carried out under inert conditions, using pre-dried reagents. Chemicals were obtained from Sigma-Aldrich or Strem Chemicals unless stated otherwise. All trialkylaluminium and aluminoxane compounds and solutions thereof were obtained from Crompton GmbH, Akzo Nobel and Albemarle Corporation. In all the examples, the molar mass of methylaluminoxane (MAO) was taken to be 58.016 g/mol, corresponding to the (CH<sub>3</sub>-Al-O) unit, in order to calculate the molar quantities of MAO used in the preparation of the catalysts described in the examples below. Ethylene oligomerisation products were analysed by GC-MS and GC-FID. Polymer samples were dried under vacuum for 12 hours prior to characterisation. Polymers were characterised using differential scanning calorimetry (DSC) to determine melting points and <sup>13</sup>C NMR for the determination of the mol% incorporation of  $\alpha$ -olefins. Where possible the remaining liquid after tandem polymerisation was analysed by GC-FID for residual olefin composition.

**Example 1: Ethylene tetramerisation reaction using CrCl<sub>3</sub>(THF)<sub>3</sub>, (p-methoxyphenyl)<sub>2</sub>PN(isopropyl)P(p-methoxyphenyl)<sub>2</sub> and MAO**

A solution of 9.8 mg of (*p*-methoxyphenyl)<sub>2</sub>PN(isopropyl)P(*p*-methoxyphenyl)<sub>2</sub> (0.018 mmol) in 10 ml of toluene was added to a solution of 5.6 mg CrCl<sub>3</sub>(THF)<sub>3</sub> (0.015 mmol) in 10 ml toluene in a Schlenk vessel. The mixture was stirred for 5 min at ambient temperature and was then transferred to a 300 ml pressure reactor (autoclave) containing a mixture of toluene (80ml) and MAO (methylaluminoxane, 4.5 mmol) at 40°C. The pressure reactor was charged with ethylene after which the reactor temperature was maintained at 45°C, while the ethylene pressure was kept at 45 barg. Thorough mixing was ensured throughout by mixing speeds of 1100 RPM's using a gas entraining stirrer. The reaction was terminated after 30 minutes by discontinuing the ethylene feed to the reactor and cooling the reactor to below 10°C. After releasing the excess ethylene from the autoclave, the liquid contained in the autoclave was quenched with ethanol followed by 10% hydrochloric acid in water. Nonane was added as an internal standard for the analysis of the liquid phase by GC-FID. A small sample of the organic layer was dried over anhydrous sodium sulfate and then analysed by GC-FID. The remainder of the organic layer was filtered to isolate the solid wax/polymeric products. These solid products were dried overnight in an oven at 100°C and then weighed to yield 1.0831 g of polyethylene. The GC analyses indicated that the reaction mixture contained 42.72 g oligomers. The oligomers comprised 72% 1-octene (99% purity).

**Example 2: Ethylene tetramerisation reaction using Cr(acetylacetonate)<sub>3</sub>, (phenyl)<sub>2</sub>PN(isopropyl)P(phenyl)<sub>2</sub> and MAO**

A solution of 18.8 mg of (phenyl)<sub>2</sub>PN(isopropyl)P(phenyl)<sub>2</sub> (0.044 mmol) in 6.4 ml of cumene was added to a solution of 7.7 mg Cr(acetylacetonate)<sub>3</sub> (0.022 mmol) in 8 ml cumene in a Schlenk vessel. The mixture was stirred for 5 min at ambient temperature and was then transferred to a 1000 ml pressure reactor (autoclave) containing a mixture of cumene (180 ml) and MAO (methylaluminoxane, 4.4 mmol, 10 % solution in toluene) at 40°C. The pressure reactor was charged with ethylene after which the reactor temperature was controlled at 45°C, while the ethylene pressure was maintained at 45 barg. The reaction was terminated after 25 min, and the procedure of Example 2 above was employed. The product mass was 118.78 g. The product comprised 69.5% 1-octene (98.9% purity).

**Example 3: Tandem catalysis reaction using  $\text{Cr}(\text{acetylacetonate})_3$ ,  $(\text{phenyl})_2\text{PN}(\text{isopropyl})\text{P}(\text{phenyl})_2$ , MAO and dimethylsilyl-bis(2-methyl{4,5}benzoindenyl) zirconium dichloride**

In this example, 0.011 mmol Chromium acetylacetonate (15.2 mg) was combined with 0.022 mmol  $(\text{phenyl})_2\text{PN}(\text{isopropyl})\text{P}(\text{phenyl})_2$  (36 mg) in a Schlenk tube under an argon atmosphere with 10 ml anhydrous toluene as solvent. The metal salt and the ligand was stirred for 5 minutes and then added under inert conditions to a Parr autoclave (300 ml capacity). The autoclave was filled *a priori* with 70 ml anhydrous toluene as solvent. Separately, 2 ml of a solution of dimethylsilyl-bis(2-methyl(4,5)benzoindenyl) zirconium dichloride ( 0.0052 mmol, 0.0022 mg) was added to 18 ml anhydrous toluene in a separate reservoir under inert conditions connected to the inlet of an HPLC pump. Approximately 1200 eq (on Cr) methylaluminoxane (MAO) was added to the autoclave under argon atmosphere. The autoclave was heated to 45 °C, sealed and pressurised to 35 barg with ethylene whilst stirring was commenced at 1200 rpm. At the same time the polymerisation catalyst solution was added to the autoclave through an inlet on the Parr reactor that was connected to the outlet of the HPLC pump. The polymerisation catalyst was added at a rate of 0.66 ml/min over a period of 30 minutes after which the reaction vessel was cooled down and quenched with ethanol. After 30 minutes, the autoclave was opened and the contents collected for analysis. The amount of polymer collected was found to be 23.16 g with a melting point of 126 °C and an amount of 1-octene incorporation as determined by  $^{13}\text{C}$  nuclear magnetic resonance spectroscopy of 3.83 %.

**Example 4: Tandem catalysis reaction using  $\text{Cr}(\text{acetylacetonate})_3$ ,  $(\text{phenyl})_2\text{PN}(\text{isopropyl})\text{P}(\text{phenyl})_2$ , MAO and tetramethylcyclopentadienyl dimethylsilyl t-butylamidato titanium dichloride**

In this example, a solution of 20 ml  $(\text{phenyl})_2\text{PN}(\text{iso-propyl})\text{P}(\text{phenyl})_2$  (0.044 mmol) in anhydrous toluene and 20 ml of a solution of  $\text{Cr}(\text{acetylacetonate})_3$  (0.022 mmol) in 30 ml

anhydrous toluene was added to a Parr autoclave (300 ml capacity). At the same time, 10 ml of a solution of tetramethylcyclopentadienyl-dimethylsilyl-*t*-butylamidato titanium dichloride (0.026 mmol, 0.01 mg) was added to an external reservoir along with 10 ml anhydrous toluene. The external reservoir (under argon atmosphere) was connected to the inlet of an HPLC pump. Approximately 1200 eq (on Cr) methylaluminoxane (MAO) was added to the autoclave under inert conditions. The autoclave was heated to 45 °C, sealed and pressurised to 43 barg with ethylene whilst stirring was commenced at 1200 rpm. At the same time the polymerisation catalyst solution was added to the autoclave through a separate inlet on the Parr reactor that was connected to the outlet of the HPLC pump. The polymerisation catalyst was added at a rate of 0.66 ml/min over a period of 30 minutes after which the reaction vessel was cooled down and quenched with ethanol. The autoclave was opened and the contents collected for analysis. Analysis of the polymer after drying in vacuum for 12 h at 60 °C revealed a melting point of 99 °C and 2.39 % incorporation of 1-octene as determined using  $^{13}\text{C}$  NMR. The amount of polymer collected was 24.53g. The density of the polymer was determined to be  $0.9202 \text{ g.cm}^{-3}$ .

**Example 5: Tandem catalysis reaction using (p-methoxyphenyl)<sub>2</sub>PN(isopropyl)P(p-methoxyphenyl)<sub>2</sub>, Cr(acetylacetonate)<sub>3</sub>, MAO and dimethylsilyl bisindenyl zirconium dichloride**

In this example, 10 ml of a toluene solution containing 0.03 mmol chromium acetylacetonate was combined with 0.06 mmol (p-methoxyphenyl)<sub>2</sub>PN(isopropyl)P(p-methoxyphenyl)<sub>2</sub> [30 mg] in a Schlenk tube under an argon atmosphere. The metal salt and the ligand was stirred for 5 minutes and then added to a Parr autoclave (300 ml capacity) under inert conditions. The autoclave was filled *a priori* with 60 ml anhydrous toluene as solvent. Separately, 5 ml of a solution of dimethylsilyl bisindenyl zirconium dichloride (0.015 mmol, 0.0067 mg), was added to 15 ml anhydrous toluene in a separate reservoir under argon that was connected to the inlet of an HPLC pump. Approximately 600 eq (on Cr) methylaluminoxane (MAO) was added to the autoclave under inert conditions. The autoclave was heated to 45 °C, sealed and pressurised to a pressure of 41 barg with ethylene whilst stirring was commenced at 1200 rpm. At the same time the polymerisation catalyst solution was added to the autoclave through a separate inlet connected to the outlet of the HPLC pump. The polymerisation catalyst

was added at a rate of 0.33 ml/min over a period of 60 minutes after which the reaction vessel was cooled down and quenched with ethanol. The autoclave was opened and the contents collected for analysis. The amount of polymer collected was first dried in a vacuum oven at 60 °C for 12 h yielding an amount of 51.1 g with a melting point of 106 °C and an amount of 1-octene incorporated as determined by  $^{13}\text{C}$  nuclear magnetic resonance spectroscopy of 5.55 %. The density of the polymer was determined to be  $0.8155 \text{ g.cm}^{-3}$ .

**Example 6: Tandem catalysis reaction using (p-methoxyphenyl)<sub>2</sub>-PN(isopropyl)P(p-methoxyphenyl)<sub>2</sub>, Cr(acetylacetonate)<sub>3</sub>, MAO and cyclopentadienyl dimethylsilyl titanium dichloride**

In this example, 0.025 g of (p-methoxyphenyl)<sub>2</sub>-PN(isopropyl)P(p-methoxyphenyl)<sub>2</sub> [0.06 mmol] was added to 10 ml of a toluene solution of Cr(acetylacetonate)<sub>3</sub> (0.03 mmol) in a Schlenk tube under an argon atmosphere and allowed to stir for 5 minutes until fully dissolved. At the same time, 10 ml of a cyclopentadienyl dimethylsilyl titanium dichloride (0.03 mmol, 0.009 mg) solution in toluene was added to an external reservoir along with 10 ml anhydrous toluene. The external reservoir was connected to the inlet of an HPLC pump. After this, the 300 ml Parr autoclave was charged with 60 ml anhydrous toluene as well as the previously stirred Cr(acetylacetonate)<sub>3</sub>/(p-methoxyphenyl)<sub>2</sub>-PN(isopropyl)P(p-methoxyphenyl)<sub>2</sub> ligand solution under inert conditions. Approximately 600 eq (on Cr) methylaluminoxane (MAO) was added to the autoclave under inert conditions. The autoclave was connected to the HPLC pump outlet and heated to 45 °C, sealed and pressurised to a pressure of 53 barg with ethylene for the duration of the reaction (60 minutes) whilst stirring was commenced at 1200 rpm. At the same time the polymerisation catalyst solution was added to the autoclave through the HPLC pump. The polymerisation catalyst was added at a rate of 0.33 ml/min over a period of 60 minutes after which the reaction vessel was cooled down and quenched with ethanol. The autoclave was opened and the contents collected for analysis. The polymer collected was washed in acetone and dried in a vacuum oven for 12h at 60 °C. The dried polymer amounted to 3.35 g. Analyses of the polymer revealed a melting point of 128 °C and 1.02 % incorporation of 1-octene as determined by  $^{13}\text{C}$  NMR.

## Claims

1. A process for polymerising olefins to branched polyolefins in the presence of a polymerisation catalyst and a cocatalyst, wherein the cocatalyst produces 1-octene in a selectivity greater than 30%.
2. A process as claimed in Claim 1 wherein the branched polyolefin is a branched polyethylene.
3. A process as claimed in Claim 1 or Claim 2 wherein the branched polyolefin is linear low density polyethylene.
4. A process as claimed in any one of the preceding claims wherein the cocatalyst is an ethylene tetramerisation catalyst.
5. A process as claimed in Claim 4 wherein the ethylene tetramerisation catalyst includes a transition metal and a heteroatomic ligand.
6. A process as claimed in Claim 5 wherein the heteroatomic ligand is described by a general formula  $(R)_nA-B-C(R)_m$  where A and C are independently selected from a group which comprises phosphorus, arsenic, antimony, oxygen, bismuth, sulphur, selenium, and nitrogen, and B is a linking group between A and C, and the R groups are the same or different and each R is independently selected from any homo or heterohydrocarbyl group and n and m for each R is independently determined by the respective valence and oxidation state of A and C.
7. A process as claimed in Claim 5 or Claim 6 wherein the heteroatomic ligand is described by a general formula  $(R^1)(R^2)A-B-C(R^3)(R^4)$  where A and C are independently selected from a group which comprises phosphorus, arsenic, antimony, bismuth and nitrogen and B is a linking group between A and C, and  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are independently selected from hydrocarbyl or

heterohydrocarbyl or substituted hydrocarbyl or substituted heterohydrocarbyl groups.

8. A process as claimed in any one of claims 5 to 7, wherein the heteroatomic ligand is described by the following general formula  $(R^1)(R^2)A-B-C(R^3)(R^4)$  where A and C are independently selected from a group which comprises phosphorus, arsenic, antimony, bismuth, and nitrogen and B is a linking group between A and C, and  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are independently non-aromatic or aromatic, including heteroaromatic, groups.
9. A process as claimed in Claim 8, wherein  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are independently aromatic, including heteroaromatic groups and not all the groups  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  have a substituent on the atom adjacent to the atom bound to A or C.
10. A process as claimed in Claim 9, wherein not more than two of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  have substituents on the atom adjacent to the atom bound to A or C.
11. A process as claimed in Claim 9 or Claim 10 wherein at least one or more of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are substituted with a polar substituent on a second or further atom from the atom bound to A or C.
12. A process as claimed in any one of claims 9 to 11, wherein any polar substituents on  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are not on the atom adjacent to the atom bound to A or C.
13. A process as claimed in Claim 11 or Claim 12 wherein any polar substituent on one or more of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are electron-donating.
14. A process as claimed in Claim 9 or Claim 10, wherein any substituents on one or more of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are not electron-donating.
15. A process as claimed in Claim 14 wherein each non-electron donating substituent is non-polar.

16. A process as claimed in any one of claims 10 to 15 wherein each of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  is aromatic, including heteroaromatic, but not all of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are substituted by any substituent on an atom adjacent to the atom bound to A or C.
17. A process as claimed in any one of claims 6 to 16, wherein B is selected from any one of a group comprising: organic linking groups comprising a hydrocarbyl, substituted hydrocarbyl, heterohydrocarbyl and a substituted heterohydrocarbyl; inorganic linking groups comprising single atom links; ionic links and a group comprising methylene, dimethylmethylene, 1,2-ethane, 1,2-phenylene, 1,2-propane, 1,2-catechol, 1,2-dimethylhydrazine,  $-B(R^5)-$ ,  $-Si(R^5)_2-$ ,  $-P(R^5)-$  and  $-N(R^5)-$  where  $R^5$  is hydrogen, a hydrocarbyl or substituted hydrocarbyl, a substituted heteroatom, a halogen, a cyclic heteroatomic group or a cyclic homoatomic group.
18. A process as claimed in Claim 17 wherein B is selected to be a single atom spacer.
19. A process as claimed in Claim 17 or Claim 18, wherein B is selected to be  $-N(R^5)-$ , wherein  $R^5$  is hydrogen or selected from the groups consisting of alkyl, aryl, aryloxy, halogen, nitro, alkoxycarbonyl, carbonyloxy, alkoxy, aminocarbonyl, carbonylamino, dialkylamino, or derivatives thereof, aryl substituted with any of these substituents, and any cyclic heteroatomic group including cyclopentadienyl dimethylsilyl-t-butylamidato or a cyclic homoatomic group including cyclopentadienyl, indenyl and fluorene.
20. A process as claimed in any one of claims 6 to 19, wherein A and/or C is independently oxidised by S, Se, N or O, where the valence of A and/or C allows for such oxidation.
21. A process as claimed in Claim 20, wherein A and/or C is independently phosphorus or phosphorus oxidised by S or Se or N or O.
22. A process as claimed in any one of claims 7 to 21, wherein  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are independently selected from a group comprising a benzyl, phenyl, tolyl, xylyl,

mesityl, biphenyl, naphthyl, anthracenyl, methoxy, ethoxy, phenoxy, tolyloxy, dimethylamino, diethylamino, methylethylamino, thiophenyl, pyridyl, thioethyl, thiophenoxy, trimethylsilyl, dimethylhydrazyl, methyl, ethyl, ethenyl, propyl, butyl, propenyl, propynyl, cyclopentyl, cyclohexyl, ferrocenyl and tetrahydrofuranyl group.

23. A process as claimed in any one of claims 7 to 21 wherein  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are independently selected from a group comprising a phenyl, tolyl, biphenyl, naphthyl, thiophenyl and ethyl group.
24. A process as claimed in any one of claims 5 to 9, Claim 14 and Claim 15 wherein the heteroatomic ligand is selected from any one of a group comprising
- (phenyl)<sub>2</sub>PN(methyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(pentyl)P(phenyl)<sub>2</sub>,  
 (phenyl)<sub>2</sub>PN(phenyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(p-methoxyphenyl)P(phenyl)<sub>2</sub>,  
 (phenyl)<sub>2</sub>PN(*p*-*t*-butylphenyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN((CH<sub>2</sub>)<sub>3</sub>-N-morpholine)P(phenyl)<sub>2</sub>,  
 (phenyl)<sub>2</sub>PN(Si(CH<sub>3</sub>)<sub>3</sub>)P(phenyl)<sub>2</sub>,  
 (((phenyl)<sub>2</sub>P)<sub>2</sub>NCH<sub>2</sub>CH<sub>2</sub>)N, (ethyl)<sub>2</sub>PN(methyl)P(ethyl)<sub>2</sub>,  
 (ethyl)<sub>2</sub>PN(isopropyl)P(phenyl)<sub>2</sub>, (ethyl)(phenyl)PN(methyl)P(ethyl)(phenyl),  
 (ethyl)(phenyl)PN(isopropyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>P(=Se)N(isopropyl)P(phenyl)<sub>2</sub>,  
 (phenyl)<sub>2</sub>PCH<sub>2</sub>CH<sub>2</sub>P(phenyl)<sub>2</sub>, (*o*-ethylphenyl)(phenyl)PN(isopropyl)P(phenyl)<sub>2</sub>, (*o*-methylphenyl)<sub>2</sub>PN(isopropyl)P(*o*-methylphenyl)(phenyl), (phenyl)<sub>2</sub>PN(benzyl)-P(phenyl)<sub>2</sub>,  
 (phenyl)<sub>2</sub>PN(1-cyclohexylethyl)P(phenyl)<sub>2</sub>,  
 (phenyl)<sub>2</sub>PN[CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>Si(OMe)<sub>3</sub>]P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(cyclohexyl)P(phenyl)<sub>2</sub>,  
 phenyl)<sub>2</sub>PN(2-methylcyclohexyl)P(phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(allyl)P(phenyl)<sub>2</sub>, (*o*-naphthyl)<sub>2</sub>PN(methyl)P(*o*-naphthyl)<sub>2</sub>, (*p*-biphenyl)<sub>2</sub>PN(methyl)P(*p*-biphenyl)<sub>2</sub>, (*p*-Me-phenyl)<sub>2</sub>PN(methyl)P(*p*-Me-phenyl)<sub>2</sub>, (*o*-thiophenyl)<sub>2</sub>PN(methyl)P(*o*-thiophenyl)<sub>2</sub>,  
 (phenyl)<sub>2</sub>PN(methyl)N(methyl)P(phenyl)<sub>2</sub>, (*m*-Me-phenyl)<sub>2</sub>PN(methyl)P(*m*-Me-phenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(isopropyl)P(phenyl)<sub>2</sub>,  
 (phenyl)<sub>2</sub>P(=S)N(isopropyl)P(phenyl)<sub>2</sub>, and  
 (phenyl)<sub>2</sub>P(=S)N(isopropyl)P(=S)(phenyl)<sub>2</sub>.
25. A process as claimed in any one of claims 5 to 13, wherein the heteroatomic ligand is selected from any one of a group comprising ((*m*-methoxyphenyl)<sub>2</sub>PN(methyl)P(*m*-methoxyphenyl)<sub>2</sub>, (*p*-

methoxyphenyl)<sub>2</sub>PN(methyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*m*-  
 methoxyphenyl)<sub>2</sub>PN(isopropyl)P(*m*-methoxyphenyl)<sub>2</sub>, (*p*-  
 methoxyphenyl)<sub>2</sub>PN(isopropyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(2-  
 ethylhexyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*m*-  
 methoxyphenyl)(phenyl)PN(methyl)P(phenyl)<sub>2</sub> and (*p*-  
 methoxyphenyl)(phenyl)PN(methyl)P(phenyl)<sub>2</sub>, (*m*-  
 methoxyphenyl)(phenyl)PN(methyl)P(*m*-methoxyphenyl)(phenyl), (*p*-  
 methoxyphenyl)(phenyl)PN(methyl)P(*p*-methoxyphenyl)(phenyl), (*m*-  
 methoxyphenyl)<sub>2</sub>PN(methyl)P(phenyl)<sub>2</sub> and (*p*-  
 methoxyphenyl)<sub>2</sub>PN(methyl)P(phenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(1-  
 cyclohexylethyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(2-  
 methylcyclohexyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(decyl)P(*p*-  
 methoxyphenyl)<sub>2</sub>, (*p*-methoxyphenyl)<sub>2</sub>PN(pentyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-  
 methoxyphenyl)<sub>2</sub>PN(benzyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-  
 methoxyphenyl)<sub>2</sub>PN(phenyl)P(*p*-methoxyphenyl)<sub>2</sub>, (*p*-  
 fluorophenyl)<sub>2</sub>PN(methyl)P(*p*-fluorophenyl)<sub>2</sub>, (*o*-fluorophenyl)<sub>2</sub>PN(methyl)P(*o*-  
 fluorophenyl)<sub>2</sub>, (*p*-dimethylamino-phenyl)<sub>2</sub>PN(methyl)P(*p*-dimethylamino-phenyl)<sub>2</sub>,  
 (*p*-methoxyphenyl)<sub>2</sub>PN(allyl)P(*p*-methoxyphenyl)<sub>2</sub>, (phenyl)<sub>2</sub>PN(isopropyl)P(*o*-  
 methoxyphenyl)<sub>2</sub>, (*p*-(*p*-methoxyphenyl)-phenyl)<sub>2</sub>PN(isopropyl)P(*p*-(*p*-  
 methoxyphenyl)-phenyl)<sub>2</sub> and (*p*-  
 methoxyphenyl)(phenyl)PN(isopropyl)P(phenyl)<sub>2</sub>.

26. A process as claimed in any one of the preceding claims wherein the polymerisation catalyst is selected from a group of catalysts consisting of Ziegler-Natta catalysts, unbridged metallocenes, half sandwich metallocenes, methyl-bridged metallocenes, ethyl-bridged metallocenes and silyl-bridged metallocenes.
27. A process as claimed in Claim 26, wherein the Ziegler-Natta catalyst is selected from TiCl<sub>3</sub>-Et<sub>2</sub>AlCl, AlR<sub>3</sub>-TiCl<sub>4</sub>.
28. A process as claimed in Claim 26 wherein the unbridged metallocene is selected from:
  - bis(cyclopentadienyl)chromium(II),

bis(cyclopentadienyl)-zirconium Chloride hydride,  
 bis(cyclopentadienyl)-titanium dichloride,  
 bis(cyclopentadienyl)-zirconium dichloride,  
 bis(cyclopentadienyl)-zirconium dimethyl,  
 bis(*n*-butylcyclopentadienyl)-zirconium dichloride,  
 bis(*n*-dodecylcyclopentadienyl)-zirconium dichloride,  
 bis(ethylcyclopentadienyl)-zirconium dichloride,  
 bis(iso-butylcyclopentadienyl)-zirconium dichloride,  
 bis(isopropylcyclopentadienyl)-zirconium dichloride,  
 bis(methylcyclopentadienyl)-zirconium dichloride,  
 bis(*n*-octylcyclopentadienyl)-zirconium dichloride,  
 bis(*n*-pentylcyclopentadienyl)-zirconium dichloride,  
 bis(*n*-propylcyclopentadienyl)-zirconium dichloride,  
 bis(trimethylsilylcyclopentadienyl)-zirconium dichloride,  
 bis(1,3-bis(trimethylsilyl)cyclopentadienyl)-zirconium dichloride,  
 bis(1-ethyl-3-methylcyclopentadienyl)-zirconium dichloride,  
 bis(pentamethylcyclopentadienyl)-zirconium dichloride,  
 bis(pentamethylcyclopentadienyl)-zirconium dimethyl,  
 bis(1-propyl-3-methylcyclopentadienyl)-zirconium dichloride,  
 bis(4,7-dimethylindenyl)-zirconium dichloride,  
 bis(indenyl)-zirconium dichloride,  
 bis(2-methylindenyl)-zirconium dichloride,  
 bis(2-methylindenyl)-zirconium dichloride, and  
 cyclopentadienylindenyl-zirconium dichloride.

29. A process as claimed in Claim 26, wherein the half sandwich metallocene is selected from:

Cyclopentadienyl-zirconium trichloride,  
 pentamethylcyclopentadienyl titanium trichloride,  
 pentamethylcyclopentadienyl-titanium Trimethoxide,  
 pentamethylcyclopentadienyl-titanium Trimethyl,  
 pentamethylcyclopentadienyl-zirconium trichloride,  
 tetramethylcyclopentadienyl-zirconium trichloride, and  
 1,2,4-trimethylcyclopentadienyl-zirconium trichloride.

30. A process as claimed in Claim 26, wherein the carbon-bridged metallocene is selected from:  
Diphenylmethylenedicyclopentadienyl-(9-fluorenyl)-zirconium dichloride,  
diphenylmethylenedicyclopentadienyl-(indenyl)-zirconium dichloride,  
iso-propylidenebis(cyclopentadienyl)-zirconium dichloride,  
iso-propylidene(cyclopentadienyl)(9-fluorenyl)-zirconium dichloride, and  
iso-propylidene(3-methylcyclopentadienyl)-(9-fluorenyl)-zirconium dichloride.  
ethylene-bis(9-fluorenyl)-zirconium dichloride,  
*meso*-ethylene-bis(1-indenyl)-zirconium dichloride,  
*rac*-ethylene-bis(1-indenyl)-zirconium dichloride,  
*rac*-ethylene-bis(1-indenyl)-zirconium dimethyl,  
*rac*-ethylene-bis(2-methyl-1-indenyl)-zirconium dichloride, and  
*rac*-ethylene-bis(4,5,6,7-tetrahydro-1-indenyl)-zirconium dichloride
32. A process as claimed in Claim 26, wherein the silyl-bridged metallocene is selected from:  
Dimethylsilyl-bis(cyclopentadienyl)-zirconium dichloride,  
dimethylsilyl-bis(9-fluorenyl)-zirconium dichloride,  
*rac*-dimethylsilyl-bis(1-indenyl)-zirconium dichloride,  
*meso*-dimethylsilyl-bis(2-methylindenyl)-zirconium dichloride,  
*rac*-dimethylsilyl-bis(2-methylindenyl)-zirconium dichloride,  
*rac*-dimethylsilyl-bis(tetrahydroindenyl)-zirconium dichloride,  
dimethylsilyl-bis(tetramethylcyclopentadienyl)-zirconium dichloride,  
diphenylsilyl(cyclopentadienyl)(9-fluorenyl)-zirconium dichloride, and  
diphenylsilyl-bis(indenyl)hafnium dichloride.
33. A process as claimed in any one of the preceding claims, wherein branched polymers are formed by means of a tandem tetramerisation and polymerisation process.
34. A process as claimed in Claim 33, the tandem tetramerisation and polymerisation process being an *in situ* catalysis process wherein ethylene tetramerisation and polymerisation takes place in the same reaction medium.

35. A process as claimed in Claim 34, wherein the *in situ* catalysis process is an in-situ concurrent catalysis process.
36. A process as claimed in Claim 34, wherein the in-situ catalysis process is an *in situ* consecutive catalysis process.
37. A process as claimed in Claim 34, wherein the process includes the steps of tetramerising ethylene using a tetramerisation catalyst to produce 1-octene selectively and copolymerising *in situ* the 1-octene with ethylene using a polymerisation catalyst wherein the tetramerisation catalyst produces 1-octene in more than 30% selectivity.
38. A process as claimed in Claim 37, wherein the process includes a step of contacting the ethylene with a tetramerisation catalyst, and a step of contacting ethylene and the 1-octene derived from tetramerisation with a polymerisation catalyst, which catalyst includes a transition metal, and wherein the tetramerisation catalyst includes a transition metal and a heteroatomic ligand.
39. A process as claimed in any one of claims 5 to 38, which includes a step of combining the heteroatomic ligand with a transition metal precursor and an activator in the presence of the polymerisation catalyst.
40. A process as claimed in Claim 39, wherein the transition metal for the transition metal precursor is selected from a group consisting of chromium, molybdenum, tungsten, titanium, tantalum and nickel.
41. A process as claimed in Claim 39 or Claim 40, wherein the transition metal for the transition metal precursor is chromium.
42. A process as claimed in any one of claims 39 to 41, wherein the transition metal precursor is selected from a group comprising of an inorganic salt, organic salt, a coordination complex and organometallic complex.

43. A process as claimed in Claim 42, wherein the transition metal precursor is selected from any one of a group comprising chromium trichloride tris-tetrahydrofuran complex, (benzene)-tricarbonyl chromium, chromium (III) octanoate, chromium (III) acetylacetonate, chromium hexacarbonyl and chromium (III) 2-ethylhexanoate.
44. A process as claimed in Claim 43, wherein the transition metal precursor is selected from chromium (III) acetylacetonate and chromium (III) 2-ethylhexanoate.
45. A process as claimed in any one of claims 39 to 44, wherein the transition metal from the transition metal precursor and the heteroatomic ligand are combined to provide metal/ligand ratios from about 0.01:100 to 10 000:1.
46. A process as claimed in Claim 45, wherein the transition metal precursor and the heteroatomic ligand are combined to provide metal/ligand ratios from about 0.1:1 to 10:1.
47. A process as claimed in any one of claims 39 to 46, wherein the catalyst system includes an activator selected from any one of a group consisting of organoaluminium compounds, organoboron compounds, organic salts, inorganic acids and inorganic salts.
48. A process as claimed in Claim 47 wherein the organic salts are selected from a group consisting of methyllithium bromide and methylmagnesium bromide.
49. A process as claimed in Claim 48 wherein inorganic acids and inorganic salts are selected from a group consisting of as tetrafluoroboric acid etherate, silver tetrafluoroborate and sodium hexafluoroantimonate.
50. A process as claimed in any one of claims 39 to 49, wherein the activator is selected from alkylaluminoxanes.

51. A process as claimed in Claim 50, wherein the alkylaluminumoxane, or mixtures thereof, are selected from a group which consists of methylaluminumoxane (MAO), ethylaluminumoxane (EAO) and modified alkylaluminumoxanes (MMAO).
52. A process as claimed in Claim 51, wherein the transition metal and the aluminumoxane are combined in proportions to provide Al/metal ratios from about 1:1 to 10 000:1.
53. A process as claimed in Claim 52, wherein the transition metal and the aluminumoxane are combined in proportions to provide Al/metal ratios from about 1:1 to 1000:1.
54. A process as claimed in Claim 53, wherein the transition metal and the aluminumoxane are combined in proportions to provide Al/metal ratios from about 1:1 to 300:1.
55. A process as claimed in any one of claims 50 to 54, which process includes the step of adding to the catalyst system a trialkylaluminum compound in amounts of between 0.01 to 100 mol per mol of alkylaluminumoxane.
56. A process as claimed in any one of claims 39 to 55, wherein the heteroatomic ligand and the transition metal precursor are combined at any temperature between -20°C and 250°C in the presence of an olefin.
57. A process as claimed in any one of claims 39 to 56, wherein the activator is a binuclear activator such as bisborane 1,4-(C<sub>6</sub>F<sub>5</sub>)<sub>2</sub>B(C<sub>6</sub>F<sub>4</sub>)B-(C<sub>6</sub>F<sub>5</sub>)<sub>2</sub>.
58. A process as claimed in any one of claims 4 to 57, which process includes a step of combining the polymerisation catalyst and the tetramerisation catalyst in a ratio of 0.1:1 to 1:1000 respectively.
59. A process as claimed in any one of the preceding claims 1 to 58, which process is carried out in an inert solvent.

60. A process as claimed in Claim 59 wherein ethylene is contacted with the tetramerisation catalyst system at a pressure of more than 1 barg.
61. A process as claimed in any one of the preceding 1 to 60, which process includes the step of mixing the polymerisation catalyst and the cocatalyst at any temperature between -20°C and 250°C in the presence of an olefin.
62. A process as claimed in Claim 61, wherein the temperature range is between 20°C and 100°C.
63. A process as claimed in any one of the proceeding claims where a combination of reactors is preferred where the first reactor may be a continuous stirred tank reactor and the second reactor may be a batch, continuous stirred tank reactor or a plug flow reactor.
64. A process as claimed in any one of claims 4 to 63, wherein the cocatalyst and polymerisation catalyst is combined in the ratio 0.01:100 to 10 000:1.
65. A process as claimed in any one of claims 4 to 63, wherein the cocatalyst and polymerisation catalyst is combined in the ratio of 1:1 to 100:1.
66. A process as claimed in any one of claims 1 to 65, wherein the conditions are selected such that the cocatalyst produces 1-octene in a selectivity greater than 40%.
67. A process as claimed in any one of claims 1 to 65, wherein the conditions are selected such that the cocatalyst produces 1-octene in a selectivity greater than 50%.
68. A process as claimed in any one of claims 1 to 65, wherein the conditions are selected such that the cocatalyst produces 1-octene in a selectivity greater than 60%.

69. A process as claimed in any one of claims 1 to 65, wherein the conditions are selected such that the cocatalyst produces 1-octene in a selectivity greater than 70%.
70. A process substantially as described herein.