

Description

Background of the invention

This invention relates to a process for producing olefin polymers and to a high activity catalyst suitable for use in the process.

It is old in the field of olefin polymerization to prepare solid polyolefins by employing catalyst systems comprising a transition metal compound and an organometallic cocatalyst. Until fairly recently most of such catalyst systems were relatively low in activity and as a result generally required the employment of a catalyst removal step in order to obtain a polymer having satisfactory properties. Recently, however, much research has been directed toward the discovery of catalyst systems that are so active that there is no need to employ the catalyst removal step. Such catalyst systems are obviously of significant commercial importance since they allow for appreciable reductions in the costs of producing such polymers.

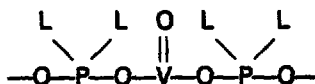
Included among the early relatively low activity catalysts were systems involving the use of a vanadium halide, an aluminum halide, and an organophosphorus compound. See for example the catalyst systems disclosed in U.S. Patent 2,969,345 and U.S. Patent 3,186,977. It has been noted that when a vanadium halide, an organoaluminum halide, and an organophosphorus compound are combined in the polymerization diluent as taught by those patents, an intractable oily or gummy form of catalyst is produced. While this might not present too much of a problem if the polymerization were conducted at temperatures under which the polymer is substantially dissolved, for modern particle form polymerization wherein the polymer is produced in solid form, it would lead to serious fouling of the reactor. For a particle form polymerization, it would obviously be more desirable, if not absolutely necessary, to have the catalyst in the form of individual solid particles. It was further noted that the activity of the catalysts of the two aforementioned patents was not nearly high enough to produce polymer in such quantity that subsequent catalyst removal could be eliminated.

The present invention is based upon the discovery of a way to produce a solid vanadium-containing catalyst capable of producing ethylene polymers in such amounts that the catalyst removal step is generally unnecessary.

Summary of the invention

In accordance with one aspect of the present invention, a solid particulate vanadium-containing catalyst component is produced by reacting reactants comprising a vanadium halide compound, an organophosphorus compound, and an organoaluminum compound or an organoaluminum compound and a vanadium organophosphate under such conditions that a particulate solid is obtained.

Accordingly, the invention relates to a process for preparing a catalyst component comprising reacting in a liquid components (A) and (B), component (A) being comprised of (A1) at least one vanadium halide having a V—X linkage where X is halogen and at least one organophosphorus compound having a P—L linkage where L is selected from alkyl, alkoxy, aryl, aryloxy, cycloalkyl, cycloalkyloxy, cycloalkylaryl, alkylamino and haloalkyloxy radicals, or (A2) at least one compound having functionality of the formula



wherein each L is individually as defined hereinbefore, and (B) being at least one organoaluminum compound having the formula AlR'_nX_{3-n} wherein X is halogen, each R' is individually selected from alkyl, aryl, cycloalkyl and alkoxy radicals, and n is a number from 1 to 3, and recovering the reaction product from said liquid, which process is characterized in that said reaction is carried out in a liquid consisting substantially of nonaromatic halohydrocarbon.

In a specific embodiment said organophosphorus compound is selected from triethylphosphite, tributylphosphate, tributylphosphite, triethylphosphate, trioctylphosphate, trimethylphosphite, hexamethylphosphoramide, diethylphosphite, tri(2-chloro-ethyl)phosphite, trimethylphosphate, tri(2-ethylhexyl)phosphate, triisopropylphosphite, triphenylphosphine, and triphenylphosphine oxide.

In a further specific embodiment said organophosphorus compound is tri-n-butylphosphate or triethylphosphite.

In a further specific embodiment said organoaluminum compound is selected from ethylaluminum dichloride, diethylaluminum chloride, trimethylaluminum, triisobutylaluminum, ethylaluminum sesquichloride and diethylaluminum ethoxide.

In yet a further specific embodiment said organoaluminum compound is ethylaluminum dichloride.

The catalyst compound of the invention is used for the polymerization or copolymerization of α -olefins, e.g. the polymerization of ethylene.

In a specific embodiment said polymerization is carried out in the presence of a cocatalyst system comprising at least one organoaluminum compound and at least one halogen-substituted alkane having from 1 to 10 carbon atoms as halohydrocarbon activator.

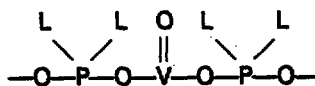
Detailed description of the invention

The vanadium compound used in making the particulate catalyst component is selected from those vanadium compounds having a V—X linkage wherein X is a halogen. Typical examples of such compounds are the halides, oxyhalides, alkylhalides, and alkoxyhalides. Especially notable halo-vanadium compounds are those compounds of the formula $VX_{4-n}Y_n$, wherein X is a halogen atom, Y is an alkoxy group with 1 to 12, preferably 1 to 6 carbon atoms or trialkylsiloxy group ($-\text{OSiR}^3$, group where R^3 is an alkyl group with 1 to 12 carbon atoms), and n is a number defined by: $0 \leq n < 4$, and the compounds of the formula: $\text{VOX}_{3-m}Y_m$ (wherein X is a halogen atom, Y is an alkoxy group with 1 to 12, preferably 1 to 6 carbon atoms or trialkylsiloxy group ($-\text{OSiR}^3$, group wherein R^3 is an alkyl group with 1 to 12 carbon atoms), and m is a number defined by: $0 \leq m < 3$). More specific examples of compounds having such a V—X linkage are vanadium tetrachloride, vanadium tetrabromide, vanadium tetraiodide, mono-n-butoxy-vanadium trichloride, vanadyl trichloride, vanadyl tribromide, di-n-butoxy-chlorovanadyl, diisopropoxymono-chlorovanadyl, mono-n-butoxydichlorovanadyl, monoisopropoxydichlorovanadyl, and monotrimethyl-siloxydichlorovanadyl. The currently preferred vanadium compounds are vanadium tetrachloride and vanadyl trichloride, which is also sometimes referred to hereafter as vanadium oxychloride or as VOCl_3 .

The organoaluminum compound used in preparing the novel particulate catalyst is selected from those having the formula $\text{AlR}'_n\text{X}_{3-n}$, wherein X is halogen, each R' is individually selected from the group consisting of alkyl radicals, aryl radicals, cycloalkyl radicals, and alkoxy radicals, and n is a number from 1 to 3. Representative types of compounds having such formulas include dialkylaluminum monohalides, alkylaluminum dihalides, trialkyl aluminums, alkyl, cycloalkyl, or arylaluminum sesquihalides, and dialkylaluminum alkyloxides. Particularly preferred organoaluminum compounds are diethylaluminum chloride, ethylaluminum dichloride, ethylaluminum sesquichloride, and diethylaluminum ethoxide. The best organoaluminum compound to use varies somewhat depending upon the particular organophosphorus compound being used.

The organophosphorus compound used in preparing the novel particulate catalyst is selected from those having a P—L linkage wherein L is selected from the group consisting of alkyl radicals, alkoxy radicals, aryl radicals, aryloxy radicals, cycloalkyl radicals, cycloalkyloxy radicals, cycloalkylaryl radicals, alkylamino radicals, and haloalkyloxy radicals. Typical examples of such compounds include tris-N,N-dialkyl phosphoramides, dialkyl phosphites, trialkyl phosphites, trialkyl phosphates, trialkyl phosphines, triaryl phosphines, triaryl phosphine oxides, triaryl phosphites, and haloalkyl phosphites. Generally, the radicals of the L group will have only 1 to 10 carbon atoms.

In an alternative embodiment of the present invention instead of reacting the organoaluminum compound, and the vanadium and organophosphorus compound, one can merely react the organoaluminum compound and a vanadium organophosphate containing functionality of the formula



wherein each L is individually selected from the group of L's described above. The preparation of such compounds disclosed in U.S. Patent No. 3,595,890. For best results a vanadium organophosphate must be selected that is soluble in the reaction liquid. A particularly preferred type of vanadium organophosphate are the vanadium oxy bis(dialkyl phosphates).

The catalyst preparation is carried out in a liquid consisting substantially of nonaromatic halohydrocarbon. Examples of halohydrocarbons which have been found suitable are 1,1,2-trichlorotrifluoroethane, dichloromethane, 1-chlorobutane, dibromomethane, and tetrachloroethylene. The currently most favored liquid is dichloromethane. The amount of liquid employed can vary widely, however, if some of the reactants are added in a liquid which is not suitable for the formation of the particulate product, the quantity of such liquid should preferably be less than that of the liquid in which the particulate product can be obtained.

In the embodiment of the invention involving three reactants, the reactants can be combined in any order. It has, however, been noted that the most active catalysts have been obtained from those reactions in which the vanadium compound is first reacted with the organophosphorus compound and then the resulting product is reacted with the organoaluminum compound.

The reaction of the reactants can be carried out under any suitable conditions. Generally, the reaction will be conducted at a temperature in the range of about 0°C to about 100°C and a pressure in the range of 69—690 kPa (10 to 100 psia). Preferably, the reaction is conducted without the addition of heat and when the addition of a reactant produces an exothermic reaction, the reactant addition is preferably spread out over a period of time such that large temperature increases are avoided.

The particulate solid can be recovered from the reaction mixture by any suitable means. Most typically the solid is separated from the liquid by filtration. In one particularly preferred embodiment, the solid is washed with a hydrocarbon liquid until the solid is substantially free of compounds soluble in said hydrocarbon liquid. Typical suitable wash liquid include liquids of the type generally used as liquid diluent

in the polymerization of ethylene. Examples of such liquids include heptane, hexane, pentane, benzene, toluene and the like.

The reactants can be combined in any amounts that will allow the production of recoverable particulate solid. Generally, the P/V atom ratio of the reactants is in the range of about 0.2/1 to about 4/1, more preferably in the range of about 0.3/1 to 3/1. Generally, the Al/P atom ratio of the reactants is in the range of about 0.5/1 to about 2/1, more preferably about 1/1.

The solid particulate catalyst of this invention is useful in the polymerization of olefins. It provides particularly surprising results in the production of homopolymers of ethylene and copolymers containing at least 90 mole percent and preferably 95 mole percent ethylene.

Examples of typical olefin monomers include alpha-olefins containing 2 to 20 carbon atoms per molecule, the most preferred being usually those with 2 to 6 carbon atoms per molecule, such as ethylene, propylene, butene-1, 4-methylpentene-1, and hexene-1. Such mono-olefins can also be copolymerized with each other and with diolefins, preferably diolefins containing 4 to 20 carbon atoms. Examples of such diolefins include 1,3-butadiene, 1,5-hexadiene, 4-vinylcyclohexene, 1,3-divinylcyclohexane, cyclopentadiene, and the like.

The polymerization may be carried out in any suitable manner. Typically, the polymerization is carried out in a liquid diluent. Typical diluents include hydrocarbons, such as for example, alkanes or cycloalkanes, such as n-butane, n-hexane, n-pentane, n-heptane, cyclohexane, isobutane, methylcyclohexane, or mixtures thereof.

The polymerization pressure is generally in the range of about 15 psia (0.103 MPa) to about 1000 psia (6.89 MPa), more preferably no higher than about 700 psia (4.83 MPa). The temperature is generally selected between 20°C and 200°C, more preferably between 60°C and 120°C. The catalyst is suitable for continuous or batch polymerization.

The catalyst is used with a cocatalyst comprising an organometallic reducing agent. Typical examples of such reducing agents are the organometallic compounds of the metals of Groups Ia, IIa, IIb, IIIa, and IVb of the Periodic Table, such as organic compounds of lithium, magnesium, zinc, aluminum and tin. More specific examples of such compounds include the completely alkylated compounds whose alkyl radicals contain 1 to 20 carbon atoms, either straight or branched, such as for example, n-butyl lithium, diethylmagnesium, diethylzinc, trimethylaluminum, triethylaluminum, triisobutylaluminum, tri-n-butylaluminum, tri-n-decylaluminum, tetraethyl tin and tetrabutyl tin. Also included are the alkyl metal hydrides, for example, those containing alkyl radicals having 1 to 20 carbon atoms, such as diisobutylaluminum hydride and trimethyl tin hydride. Also suitable are the alkyl halides in which the alkyl radicals contain 1 to 20 carbon atoms such as ethylaluminum sesquichloride, diethylaluminum chloride, and diisobutylaluminum dichloride.

An especially preferred embodiment of the invention employs an organoaluminum cocatalyst and a halohydrocarbon activator, selected from halogen-substituted alkanes having 1 to 10 carbon atoms, more preferably 1 to 5 carbon atoms per molecule. The especially preferred activators are chlorine-substituted alkanes having no more than two chlorines on any primary carbon, no more than one chlorine on any secondary carbon atom, and no chlorine on any tertiary carbon atom. Examples of typical activators include dichloromethane, bromo-chloromethane, 2-bromopropane, 1,2-dibromoethane, dibromomethane, fluorotrichloromethane, 1,1,2-trichloro-1,2,2-trifluoroethane, 1,1,1-trichloroethane, 1,1,2-trichloroethane, 1,2-dichloroethane, 1-fluoro-1,1-dichloro-2,2,2-trichloroethane, 1,2-difluoro-1,2-dichloroethane, 1,2-dichlorotetrafluoroethane, difluorodichloromethane, 1,2-difluorotetrachloroethane and 1-chlorobutane. The currently preferred activators are dichloromethane and because of its effectiveness, low toxicity, and moderate boiling point, 1,2-difluorotetrachloroethane.

In the practice of this invention, the particulate solid catalyst is effective when used in amounts conventionally used in such reactions, i.e., about 0.03 to 2 g per liter of diluent. However, it is not necessary to use such large amounts of the catalyst and such would in fact defeat one of the advantages of the catalyst which is the production of such large levels of polymer per unit of catalyst that one can dispense with removal of catalyst from the polymer. It is, therefore, preferred to use an amount of catalyst which provides about 0.0001 to 0.05 g of vanadium-containing solid catalyst per liter of diluent, more preferably about 0.001 to 0.01 g per liter of diluent.

The amount of cocatalyst employed can vary over a wide range depending upon the particular cocatalyst selected and the results desired. With the preferred organoaluminum cocatalysts enough is used to provide at least 0.3 mole of the cocatalyst per gram atom of vanadium in the particulate solid catalyst employed. More generally the ratio of Al in the cocatalyst to vanadium in the catalyst is in the range of about 0.3/1 to 1000/1, more preferably about 1/1 to 500/1.

The ratio of the activator to the cocatalyst can also vary over a wide range. If desired the activator can be used in excess of that needed for maximum activation. Typical ratios for the moles of activator to mg atoms of Al would be about 0.5/1 to about 160/1, more preferably in the range of about 3/1 to about 100/1. When triethylaluminum is the cocatalyst particularly good results are obtained when the ratio is about 20/1.

The order of addition of the polymerization reaction components is not considered to be particularly critical. Typically in a batch process cocatalyst is added, when activator, then catalyst, then hydrogen, then hydrocarbon diluents, and then monomer. In view of the fact that many halohydrocarbons which permit the formation of the particulate solid are also activators, it is possible to prepare the particulate solid

directly in the reaction vessel eliminating the need for recovering the catalyst. For example, a vanadium compound such as VCl_4 and a phosphorus compound such as triethylphosphite can be added to CH_2Cl_2 in the polymerization vessel to form a solution and then an organoaluminum compound can be added to cause formation of the particulate solid. Subsequently, hydrocarbon diluent and cocatalyst can be added to the polymerization vessel.

The mean molecular weight based on the melt index of the polymers produced using the inventive catalyst can be regulated by the employment of molecular weight modifier, such as for example hydrogen. The high hydrogen response of the vanadium-containing catalyst leads to very low molecular weight polyethylenes when the ethylene pressure is reduced by hydrogen. Thus, if an attempt is being made to produce high molecular weight polyethylenes hydrogen control would have to be closely monitored.

It has also been noted that by varying the P/V ratio in the catalyst, it is possible to control to some extent the molecular weight distribution of polyethylene.

The density of polymers obtained from ethylene can be varied by copolymerization with butene-1 or other 1-olefin as known in the art.

The inventive catalysts have provided ethylene polymers having densities varying from 0.968 to 0.901 g/ml, melt indexes varying from 0.1 HLMI to over 600 MI, and Mw/Mn varying from 6.4 to 21. The polymers have many physical characteristics in common with polymers obtained using the more conventional chromium on silica catalysts. In addition, the inventive polymerization catalysts allow one to obtain substantially saturated ethylene polymers like those generally produced using titanium catalysts. For example, with the inventive vanadium catalyst, it is possible to obtain ethylene polymers wherein the number of vinyl groups per 1000 carbon atoms as determined by infrared analysis is less than about 0.1. In view of the fact that such polymers are substantially free of vinyl unsaturation they are particularly suitable for applications requiring superior environmental stress crack resistance.

The polyolefins obtained by the process of this invention may be used according to all who know fabricating techniques such as extrusion, injection, blow extrusion or rolling.

A further understanding of the present invention and its advantages will be provided by the following examples resulting from the applicant's research.

In the following examples unless indicated otherwise the following procedures were used:

All phosphorus compounds as purchased were dried over Davison H-151[®] alumina. The dichloromethane used was Fisher certified and stored over Davison H-151[®] alumina. The aluminum alkyls were used as the hydrocarbon solutions as received from Texas Alkyls. The vanadium oxytrichloride and vanadium tetrachloride was used as received from Stauffer Chemical Company.

All ethylene polymerizations were carried out in a one gallon packed, Autoclave Engineers, stirred reactor. The autoclave was partly filled with heptane and heated to above 150°C. Hot heptane was flushed out through a bottom valve and then the reactor was flushed with ethylene and then three times with isobutane. Cooling water was run through the reactor jacket to bring the temperature below 40°C before charging the reactants. The order of charging was cocatalyst, haloalkane activator, catalyst, hydrogen and isobutane. The volume of isobutane was always 2 l while hydrogen was charged as the pressure drop from either a 2.4 l or a 0.325 l vessel. The reactor was then raised to the polymerization temperature and ethylene was added so as to raise the reactor pressure by 1586 kPa (230 psi). Ethylene pressure was maintained at this value for 1 hour and then volatiles were vented from the reactor to terminate the run. Productivities of (g/g) refer to grams of polymer obtained per gram of catalyst per hour of polymerization.

Polymer evaluations reported in the following examples used the following procedures:

MI (Melt index, g/10 min 190°C), ASTM D 1238, Condition E.

HLMI (High load melt index, g/10 min, 190°C) ASTM D 1238, Condition F.

HLMI/MI=Shear response where larger value indicates a broader molecular weight distribution.

Mw (Weight average molecular weight), Determined by size exclusion chromatography.

Mn (Number average molecular weight), Determined by size exclusion chromatography.

HI (Heterogeneity Index=Mw/Mn), The higher the number the broader the molecular weight distribution.

Flexural modulus, MPa (psi), ASTM D 790.

ESCR (Environmental Stress Cracking Resistance), F_{50} in hours, ASTM D 1693, Condition A.

RDI (Rheological Distribution Index), Another way of evaluating the molecular weight distribution involving subjecting a molten polymer sample to a sinusoidal oscillating shear strain between two parallel circular plates at 190°C. The frequency of oscillation is varied from 0.1 to 500 radians per second. A log-log plot of frequency vs. dynamic loss modulus is made and the slope of the curve determined when the loss modulus is equal to 10,000 Pa (100,000 (10⁵) dynes/cm²). RDI equals the reciprocal of that slope.

Density, g/ml, ASTM D 1505.

Example I

When ethylaluminum dichloride (EADC) is added to a mixture of tri-n-butylphosphate and VCl_4 in heptane as taught in prior art patents such as U.S. Patent No. 2,969,345, an oily mass was produced. No particulate solid was recovered. Such catalyst systems when used to polymerize ethylene produce only relatively small amounts of polyethylene.

Example II

A 4 ml CH_2Cl_2 solution of 16.6 mmoles VCl_4 was added to 20 ml of CH_2Cl_2 . Then 4.5 ml of tri-n-butylphosphate (16.6 mmoles) was added to give a dark red solution. The solution was heated for 3 hours at 100°C then filtered in a glove box. No solid was obtained. Over a 15 minute period, 11 ml of 25 weight percent heptane solution of ethylaluminum dichloride (16.6 mmoles) was added to the red solution. The resulting slurry was stirred for 6 hours and then filtered. A violet solid was recovered and washed with about 20 ml of hexane and dried. The weight of solid recovered was 5.91 g. The solid thus theoretically contained 14.3 weight percent vanadium based upon the assumption of a complete precipitation of all the vanadium.

The activity of the solid as a catalyst component for ethylene polymerization was then determined by using the respective solid catalyst, 0.5 mmole (0.5 ml) of triethylaluminum as a 1 M solution in n-heptane and 3 ml of dichloromethane. The hydrogen was measured as the pressure drop from a 2.4 l vessel. The results at three different hydrogen levels are shown in Table I.

TABLE I

Run No.	Catalyst, g	Productivity, (g/g)	H_2 , kPa (psi)	Melt index
1	.0168	17,600	276 (40)	126
2	.0119	23,100	103 (15)	0.13
3	.0143	28,500	69 (10)	0.04

The polymer resulting from Run 2 had a heterogeneity index as determined by Mw/Mn of 13.4, flexural modulus of 1545 MPa, environmental stress crack resistance of 131 hours, a rheological distribution index of 1.66, and a density of 0.9599 g/ml. These properties are consistent with those of many polyethylenes having significant commercial applications. (The term "catalyst" as used in the regard to productivity refers to the solid catalyst component only and does not include the cocatalyst and activator.)

Example III

A series of catalysts were prepared using VCl_4 , ethylaluminum dichloride, and various phosphorus compounds in the same manner as generally employed in Example II. The resulting solid catalysts were evaluated for ethylene polymerization as in Example II, the only exception being that in some runs, viz. Runs 4—14, the hydrogen was 414 kPa (60 psi) measured from the pressure drop of a 0.325 l vessel and in Runs 15—20, it was 138 kPa (20 psi) as measured from a 2.4 l vessel. The results of the polymerizations are set forth in Table II.

TABLE II
Additional catalysts from VCl_4 and EADC

Run No.	Phosphorus compound	Productivity (g/g)	MI	HLM/MI	Density (g/ml)
4	triethylphosphite	54,700	1.9	47	
5	tributylphosphate	18,200	0.36	60	0.9616
6	triethylphosphate	44,100	0.16	87	
7	triethylphosphate	27,900	1.5	46	
8	triethylphosphate	41,000	0.68	61	
9	trioctylphosphate	6,460	0.49	55	
10	trimethylphosphite	43,600	1.2	31	
11	hexamethylphosphoramide	45,300	5.0	37	
12	diethylphosphite	18,500	0.14	86	
13	tri(2-chloroethyl)phosphite	27,900	1.7	36	
14	trimethylphosphate	16,300	1.6	74	
15	tri-2-ethyl hexylphosphate	13,070	1.1	61	
16	triethylphosphite	70,200	1.4	49	0.9628
17	triethylphosphite	49,400	4.2	49	0.9659
18	triisopropylphosphite	22,700	8.0	66	
19	triphenylphosphine	33,700	8.5	74	
20	triphenylphosphine oxide	23,800	4.6	47	

Most of the phosphorus compounds produced precipitates in substantially the same way as observed in the preparation of Example II, namely the formation of a solution when the VCl_4 and the phosphorus compound were combined in CH_2Cl_2 and then the formation of a particulate solid when the EADC was added. With trioctylphosphate all the vanadium was not precipitated. However, as shown by Run 9, the recovered particulate solid was a moderately active catalyst. The combination of trimethylphosphite with VCl_4 in a 1/1 molar ratio gave a sticky solid in CH_2Cl_2 . Doubling the molar amount of trimethylphosphite resulted in a solution which required an EADC/ VCl_4 molar ratio of 2/1 in order to obtain a precipitate. The resulting precipitate was employed in Run 10 and is clearly a highly active catalyst.

Example IV

Vanadium oxytrichloride, 6.30 mmoles, was added to 20 ml of CH_2Cl_2 . Then 1.70 ml (6.30 mmoles) of tri-n-butylphosphate was added to give a solution black to reflected light and dark red to transmitted light. Addition of 6.30 mmoles of ethylaluminum dichloride caused a gradual color change to orange-black, to dark green, and then to dark blue as a solid was formed. The yield of blue solid was 1.70 g. This calculates to a theoretical maximum vanadium content of 18.8 weight percent.

Another catalyst was prepared in the same manner using VOCl_3 and triethylphosphite rather than the tri-n-butylphosphate. A violet solid in a yield of 1.82 g was obtained. The calculated vanadium content of the solid was 17.7 weight percent.

Polymerizations were carried out with these two catalysts at 100°C in 2 l of isobutane with an ethylene pressure of 1586 kPa (230 psi) and hydrogen measured from a 2.4 l vessel. A mixture of 0.5 mmole of triethylaluminum and 3 ml of CH_2Cl_2 was used as cocatalyst. The results are summarized in Table III.

TABLE III
Catalysts from VOCl_3

Run No.	Phosphorus compound	Productivity (g/g)	H_2 kPa (psi)	MI	HLM/MI	Density (g/ml)
21	tri-n-butylphosphate	23,000 ^b	138 (20)	0.72	62	0.9675
22	triethylphosphite	49,400 ^c	138 (20)	0.49	69	0.9633

^b Mw/Mn=12.2, flexural modulus 1673 MPa, RDI=1.59, ESCR 49 hrs.

^c Mw/Mn=11.7, flexural modulus 1698 MPa, RDI=1.62, ESCR 80 hrs.

Example V

In order to evaluate the effect of the levels of the three reactants upon the resulting catalysts a series of catalysts were prepared using different amounts of vanadium, phosphorus, and aluminum compounds.

A first catalyst was prepared without the use of any phosphorus compound. VCl_4 and EADC were reacted at a 1/1 molar ratio in CH_2Cl_2 to obtain 1.82 g of solid from 6.30 mmoles of VCl_4 implying 17.6 weight percent V in the solid.

In another preparation 12.6 mmoles of triethylphosphite were combined with 6.3 mmoles of VCl_4 in CH_2Cl_2 . It was noted that 12.6 mmoles of EADC was needed to completely precipitate the vanadium. The reaction produced 1.38 g of catalyst with a calculated weight percent vanadium of 23.1.

In yet another preparation the P/V ratio and the Al/V ratio was raised to 3. Precipitation was not complete. However, 0.07 g of particulate catalyst was recovered.

Moving in the other direction, the amounts of triethylphosphite and EADC were kept at 6.3 mmoles while the VCl_4 was raised to 10.4 mmoles, and in another preparation to 20.8 mmoles. This results in P/V ratios of 0.6 and 0.3. The former yielded 2.43 g of particulate solid with a calculated weight percent vanadium of 21.8 and the later 4.27 g of particulate solid with a calculated 24.8 weight percent V.

The effect of these catalysts in polymerizing ethylene is summarized in Table IV.

TABLE IV
Varying the P/V ratio

Run No.	P/V	Productivity (g/g)	MI ^a	HLM/MI	Mw/Mn	RDI
23	0	3,750	0.22	111		
24	0.30	28,000	0.08	122		
25	0.30	34,900 ^b	0.05	136	16.0	1.79
26	0.60	30,500	0.22	57		
16	1.0	70,200	1.4	49	9.4	1.56
27	2.0	28,200	1.4	30	7.0	1.48
28	3.0	49,300	0.95	30	6.4	1.42

^a All polymerizations were carried out at 100°C in 2 l of isobutane with ethylene pressure of 1586 kPa (230 psi) and 414 kPa (60 psi) of hydrogen measured as the pressure drop from a 0.325 l vessel. As usual 0.5 mmole of triethylaluminum and 3 ml of CH_2Cl_2 were charged to the reactor.

^b The amount of triethylaluminum was increased to 1.5 mmole.

The catalyst prepared with the phosphite is clearly not as active as the inventive catalysts. Variation in the ratio of phosphorus to vanadium gives some control over the molecular weight distribution. Values of the HLM/MI ratio, heterogeneity index (Mw/Mn) and (RDI) rheological distribution index all decline as the P/V ratio increases. This indicates a narrowing of the molecular weight distribution as the P/V ratio increases. Also as the P/V ratio increases the amount of aluminum required to give complete precipitation of the vanadium increases.

Example VI

The catalyst employed in Run 4, i.e., one produced from VCl_4 , EADC, and triethylphosphite, was employed in a series of polymerizations in which ethylene was copolymerized with 1-butene. For comparison purposes a couple of copolymerizations were also conducted using a high activity titanium catalyst. The results are summarized in Table V.

TABLE V
Copolymerizations

Run No.	Productivity (g/g)	C_2H_4 kPa (psi)	H_2^a kPa (psi)	H_2^b kPa (psi)	MI	Density (g/ml)
29	69,400 ^c	1586 (230)	173 (25)		0.18	0.9254
30	112,000 ^c	1586 (230)	173 (25)		0.13	0.9253
31	28,800	1586 (230)	173 (25)		V. high	0.9223
32	40,400	1586 (230)		414 (60)	9.8	0.9291
33	22,000	1035 (150)		138 (20)	55	0.9162
34	10,500	690 (100)		138 (20)	630	0.9014

^a All polymerizations were run at 80°C in 2 l of isobutane with 200 g of butene-1 and 0.5 mmole of triethylaluminum and 3 ml of CH_2Cl_2 . These runs contained hydrogen measured as the pressure drop from a 2.4 l vessel.

^b Hydrogen measured as the pressure drop from a 0.325 l vessel.

^c These results were obtained with a titanium catalyst.

Example VII

As noted in Example I, it was not found possible to obtain a particulate solid from the reaction of VCl_4 , EADC, and tri-n-butylphosphate in a liquid consisting essentially of hydrocarbon. A series of tests were made to determine what other liquids would allow the recovery of a particulate solid.

The liquids tested were CH_2Br_2 , 1-chlorobutane, and tetrachloroethylene. The liquids were first washed with concentrated H_2SO_4 and then aqueous potassium bicarbonate and water. Subsequently the liquids were dried over magnesium sulfate and then over Davison H-151™ alumina.

Addition of 1.22 ml of triethylphosphate (7.18 mmoles) to 7.18 mmoles of VCl_4 in 25 ml of 1-chlorobutane gave a dark red solution. As 4.8 ml of EADC solution (7.2 mmoles) was added, a burgundy colored solid formed. Some of the solid adhered to the reactor walls but 1.86 g of particulate solid was recovered. A productivity of 20,500 g/g was noted for this catalyst.

Similar results were obtained when the solution above was heated to 60°C before the addition of the EADC. Productivity for the catalyst was 30,300 g/g.

When the same amounts of the same reactants were used in tetrachloroethylene, a considerable amount of oily red liquid separated in the first step. However, addition of the EADC and stirring for 1 hour still permitted the isolation of 7.57 g of particulate solid. The activity of the catalyst was, however, very low, e.g. 1780. The low activity is attributed to agglomeration of the catalyst.

In contrast to the results obtained using triethylphosphate, the alternative liquids were not as suitable when the phosphorus compound was triethylphosphite. Addition of 7.18 mmoles, 1.23 ml of triethylphosphite to 7.18 mmoles of VCl_4 in 20 ml of 1-chlorobutane immediately gave an oily solid. Even heating to 60°C did not dissolve the solid. Similar results were obtained using tetrachloroethylene in the place of the 1-chlorobutane. When CH_2Br_2 was used there was still gummy solid formation but it was not as severe and after EADC addition 1.90 g of a rose colored powder was isolated.

Example VIII

All the polymerizations tabulated thus far have used triethylaluminum and CH_2Cl_2 as the cocatalyst system. Numerous other cocatalyst systems have been tested. A summary of the results obtained using other cocatalysts and the same catalyst as used in Run 4 is shown in Table VI.

TABLE VI
Various cocatalyst systems

Run No.	Cocatalyst	mmoles	Activator	mmoles	Productivity (g/g)	MI	HLM/MI	Condition ^a
35	DEAC	0.5	CH ₂ Cl ₂	47	6,830	1.9	57	A
36	TMA	0.5	CH ₂ Cl ₂	47	19,000	8.0	45	A
37	TIBA	0.5	CH ₂ Cl ₂	47	40,700	13.2	44	A
38	TIHA	0.7	CH ₂ Cl ₂	47	5,370	1.8	66	A
39	TEA	0.5	CH ₂ Cl ₂	47	37,300	25		A
40	TEA	0.5	CFCI ₃	33	16,600	8.2	94	A
41	TEA	0.5	CF ₂ Cl CFCl ₂	25	40,400	23		A
42	TEA	0.5	None		10,200	0.28	56	B
43	TEA	0.5	CH ₃ CCl ₂ CH ₃	29	0			B
44	TEA	0.5	CH ₃ CCl ₃	30	4,030	0.59	38	B
45	TEA	0.5	CH ₂ BrCl	33	14,500	33		B
46	TEA	0.5	CH ₂ ClCH ₂ Cl	38	13,000	1.6	41	B
47	TEA	0.5	CF ₂ ClCCl ₃	47	1,700	0.8	223	B
48	TEA	0.5	CFCI ₂ CFCl ₂	47	12,000	0.65	36	B
49	TEA	0.5	CF ₂ ClCF ₂ Cl	47	37,900	0.09	50	B
50	TEA	0.5	CF ₂ Cl ₂	64	36,900	0.82	46	B
51	TEA	0.5	CH ₂ Br ₂	47	31,100	2.3	53	B

TABLE VI (continued)

Run No.	Cocatalyst	mmoles	Activator	mmoles	Productivity (g/g)	MI	HLM/MI	Condition ^a
52	TEA	0.5	CH ₂ Cl ₂	1.6	34,400	0.59	49	B
53	TEA	0.5	CH ₂ Cl ₂	7.8	43,800	0.36	53	B
54	TEA	0.5	CH ₂ Cl ₂	47	47,800	0.35	41	B
55	TEA	0.5	CH ₂ Cl ₂	78	38,800	0.33	51	B
56	TEA	1.7	CH ₂ Cl ₂	1.6	16,000	0.73	44	B
57	TEA	1.7	CH ₂ Cl ₂	7.8	39,800	0.43	49	B
58	TEA	1.7	CH ₂ Cl ₂	47	61,900	0.05	96	B
59	TEA	1.7	CH ₂ Cl ₂	78	46,400	0.59	54	B
60	TEA	3.3	CH ₂ Cl ₂	7.8	26,500	0.51	43	B
61	TEA	3.3	CH ₂ Cl ₂	47	39,300	0.28	54	B
62	TEA	3.3	CH ₂ Cl ₂	78	65,500	0.08	52	B
63	TEA	3.3	CH ₃ CH ₂ CH ₂ CH ₂ Cl	29	17,300	0.94	84	B

^a All polymerizations were carried out at 100°C in 2 l of isobutane with an ethylene pressure of 1586 kPa (230 psi) and the following:

Condition A—138 kPa (20 psi) of hydrogen measured from a 2.4 l vessel.

Condition B—414 kPa (60 psi) of hydrogen measured from a 0.325 l vessel.

Runs 35—39 show that there is little difference in behavior, except in terms of productivity. The best results were obtained with triethylaluminum and with triisobutylaluminum. In regard to the halocarbon activators, Run 43 reveals that 2,2-dichloropropane was not suitable. This is attributed to the reactivity of the secondary chlorides. Likewise Runs 40, 44, and 47 demonstrate that compounds having three chlorines on one carbon do not appear to be particularly good activators.

Runs 52—62 demonstrate that the activator to cocatalyst ratio can be varied over a wide range with little change in catalyst productivity. Especially good results were obtained with a $\text{CH}_2\text{Cl}_2/\text{TEA}$ molar ratio near 20.

Example IX

Addition of 20 ml of the 25 weight percent EADC heptane solution (30 mmoles) to 5.15 ml of triethylphosphite (30 mmoles) in 10 ml of heptane gave two phases which were colorless. When a 1 ml portion of the lower phase was added to 6.30 mmoles of VCl_4 in 20 ml of CH_2Cl_2 , a violet powder was isolated that was an active catalyst. When an attempt was made to form the catalyst with a heptane solution of VCl_4 only a sticky oil was obtained.

Example X

In another preparation VCl_4 was added to 20 ml of CH_2Cl_2 , then 5.0 ml of the EADC heptane solution was added and then 1.26 ml of triethylphosphite was added. The process yielded 2.27 g of a purple solid. The activity of the solid in ethylene polymerization was tested using 0.0102 g of the solid, 1 mmole of triethylaluminum, and 2 mmole of $\text{CFCI}_2\text{—CFCI}_2$. The catalyst was active. The productivity was 9,100 g/g/h. When the same reactants were combined in the most preferred reaction order of phosphite to VCl_4 followed by EADC, a solid was obtained which with the same cocatalyst system gave a productivity of 43,500 g/g/h.

Example XI

Another series of catalysts were prepared using aluminum compounds other than those mentioned in the preceding examples.

In one preparation ethylaluminum sesquichloride (EASC) was added to an equimolar mixture of 6.3 mmoles of VCl_4 and tri-n-butylphosphite in CH_2Cl_2 . A total of 3.15 mmoles of EASC was needed to give a precipitate.

In another run, the EASC was replaced by diethylaluminum ethoxide (DEAEO). Even at an Al/V ratio of 1 no solid was formed. After 24 hours, a thick gel was obtained from which 1.84 g of brown particulate solid was isolated.

In contrast to the difficulty of obtaining particulate solid using tri-n-butylphosphite and DEAEO, when DEAEO was used as a precipitating agent for a CH_2Cl_2 mixture of VCl_4 and triethylphosphite, a precipitate was obtained at an Al/V ratio of only 0.5. While some of the solid was sticky, 1.12 g of particulate solid was isolated and used as catalyst.

The DEAEO was also an effective precipitating agent for a CH_2Cl_2 solution of VCl_4 and triphenylphosphite. An Al/V ratio of 0.5 permitted most of the product to be isolated as a powder. When EADC was used on the CH_2Cl_2 solution of VCl_4 and triphenylphosphite, in contrast a sticky material was obtained from which no powder could be isolated.

The above described particulate catalysts were evaluated in the polymerization of ethylene using the same conditions as employed in Example II. The results are summarized in Table VII.

TABLE VII
VCl₄ catalysts

Run No.	Al compound	Phosphorus compound	Productivity (g/g)	H ₂ kPa (psi)	MI	HLM/MI	Density (g/ml)
64	EASC	tri-n-butylphosphate	6,860	138 (20)	7.3	79	
65	DEAO	tri-n-butylphosphate	33,700	138 (20)	1.1	54	0.9619
66	DEAO	triethylphosphite	40,800	138 (20)	7.5	51	0.9678
67	DEAO	triphenylphosphite	11,250	138 (20)	27		

This demonstrates that catalysts can be prepared using aluminum compounds other than ethylaluminum dichloride.

Example XII

Another series of ethylene polymerizations were conducted using a catalyst prepared from VCl_4 , triethylphosphite, and EADC and a cocatalyst comprising triethylaluminum (TEA) and 1,2-difluorotetrachloroethane as activator. The results are summarized in Table VIII.

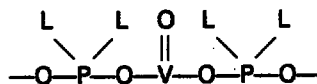
TABLE VIII
 $\text{CFCl}_2\text{CFCl}_2$ activator

Run No.	Activator mmoles	Cocatalyst mmoles	Productivity g/g
68	0.5	0.5	22,300
69	0.5	1.0	20,100
70	0.5	2.0	26,300
71	1.0	0.5	30,800
72	1.0	1.0	37,300
73	1.0	2.0	18,100
74	2.0	0.5	31,500
75	2.0	1.0	33,400
76	2.0	2.0	31,500
77	4.0	0.5	30,400
48	47	0.5	12,000

These results indicate that the best results in terms of activity are obtained when one uses at least 1 mmole of the activator and the molar ratio of activator to TEA is in the range of about 1/1 to about 8/1. A comparison of the results of Runs 68—77 with that of Run 48 indicates that the results will generally not be as desirable if one used very high molar ratios of 1,2-difluorotetrachloroethane to TEA. This is somewhat in contrast to the results obtained using dichloromethane as an activator. See Runs 52—62 of Table VI wherein it is shown that high productivities are obtained even at activator to cocatalyst molar ratios as high as 160/1. The best in Runs 52—62 are those using at least about 8 mmoles of activator and an activator/TEA molar ratio in the range of about 16/1 to about 28/1.

Claims

1. A process for preparing a catalyst component comprising reacting in a liquid components (A) and (B), component (A) being comprised of (A1) at least one vanadium halide having a V—X linkage where X is halogen and at least one organophosphorus compound having a P—L linkage where L is selected from alkyl, alkoxy, aryl, aryloxy, cycloalkyl, cycloalkyloxy, cycloalkylaryl, alkylamino and haloalkyloxy radicals, or (A2) at least one compound having functionality of the formula



wherein each L is individually as defined hereinbefore, and (B) being at least one organoaluminum compound having the formula $\text{AlR}'_n\text{X}_{3-n}$ wherein X is halogen, each R' is individually selected from alkyl, aryl, cycloalkyl and alkoxy radicals, and n is a number from 1 to 3, and recovering the reaction product from said liquid, characterized in that said reaction is carried out in a liquid consisting substantially of nonaromatic halohydrocarbon.



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This demonstrates that catalysts can be prepared using aluminum compounds other than ethylaluminum dichloride.

Example XII

- 5 Another series of ethylene polymerizations were conducted using a catalyst prepared from VCl_4 , triethylphosphite, and EADC and a cocatalyst comprising triethylaluminum (TEA) and 1,2-difluorotetrachloroethane as activator. The results are summarized in Table VIII.

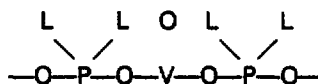
10 TABLE VIII
CFCI₂CFCI₂ activator

	Run No.	Activator mmoles	Cocatalyst mmoles	Productivity g/g
15	68	0.5	0.5	22,300
	69	0.5	1.0	20,100
20	70	0.5	2.0	26,300
	71	1.0	0.5	30,800
	72	1.0	1.0	37,300
25	73	1.0	2.0	18,100
	74	2.0	0.5	31,500
	75	2.0	1.0	33,400
30	76	2.0	2.0	31,500
	77	4.0	0.5	30,400
35	48	47	0.5	12,000

- These results indicate that the best results in terms of activity are obtained when one uses at least 1 mmole of the activator and the molar ratio of activator to TEA is in the range of about 1/1 to about 8/1. A comparison of the results of Runs 68—77 with that of Run 48 indicates that the results will generally not be as desirable if one used very high molar ratios of 1,2-difluorotetrachloroethane to TEA. This is somewhat in contrast to the results obtained using dichloromethane as an activator. See Runs 52—62 of Table VI wherein it is shown that high productivities are obtained even at activator to cocatalyst molar ratios as high as 160/1. The best in Runs 52—62 are those using at least about 8 mmoles of activator and an activator/TEA molar ratio in the range of about 16/1 to about 28/1.

Claims

1. A process for preparing a catalyst component comprising reacting in a liquid components (A) and (B), component (A) being comprised of (A1) at least one vanadium halide having a V—X linkage where X is halogen and at least one organophosphorus compound having a P—L linkage where L is selected from alkyl, alkoxy, aryl, aryloxy, cycloalkyl, cycloalkyloxy, cycloalkylaryl, alkylamino and haloalkyloxy radicals, or (A2) at least one compound having functionality of the formula



- wherein each L is individually as defined hereinbefore, and (B) being at least one organoaluminum compound having the formula $\text{AlR}'_n\text{X}_{3-n}$ wherein X is halogen, each R' is individually selected from alkyl, aryl, cycloalkyl and alkoxy radicals, and n is a number from 1 to 3, and recovering the reaction product from said liquid, characterized in that said reaction is carried out in a liquid consisting substantially of nonaromatic halohydrocarbon.



2. The process of claim 1 characterized in that said vanadium halide is reacted with said organophosphorus compound and the resulting product is then reacted with said organoaluminum compound.

3. The process of claims 1 or 2 characterized in that said halohydrocarbon liquid is selected from 1-chlorobutane, tetrachloroethylene, dibromomethane and dichloromethane.

4. The process of claim 3, characterized in that said halohydrocarbon liquid is dichloromethane.

5. The process of any of the preceding claims characterized in that said vanadium compound is selected from VCl_4 and VOCl_3 .

6. The process of any of the preceding claims characterized in that said vanadium compound is VCl_4 .

7. The process of any of the preceding claims characterized in that said organophosphorus compound is selected from triethylphosphite, tributylphosphate, tributylphosphite, triethylphosphate, trioctylphosphate, trimethylphosphite, hexamethylphosphoramide, diethylphosphite, tri(2-chloro-ethyl)-phosphite, trimethylphosphate, tri(2-ethyl hexyl)phosphate, triisopropylphosphite, triphenylphosphine, and triphenylphosphine oxide.

8. The process of any of the preceding claims characterized in that said organophosphorus compound is tri-n-butylphosphate.

9. The process of any of the preceding claims characterized in that said organophosphorus compound is triethylphosphite.

10. The process of any of the preceding claims characterized in that the P/V ratio of the reactants is in the range of 0.2/1 to 4/1 and the Al/P ratio of the reactants is in the range of 0.5/1 to 2/1.

11. The process of any of the preceding claims characterized in that said organoaluminum compound is selected from ethylaluminum dichloride, diethylaluminum chloride, trimethylaluminum, triisobutylaluminum, ethylaluminum sesquichloride and diethylaluminum ethoxide.

12. The process of any of the preceding claims characterized in that said organoaluminum compound is ethylaluminum dichloride.

13. The process of any of the preceding claims characterized in that the recovered catalyst component is washed free of hydrocarbon-soluble components.

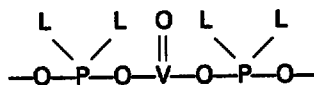
14. The use of the catalyst component as obtainable in any of claims 1 to 13 for the polymerization or copolymerization of α -olefins.

15. The embodiment of claim 14 characterized in that said polymerization is carried out in the presence of a cocatalyst system comprising at least one organoaluminum compound and at least one halogen-substituted alkane having from 1 to 10 carbon atoms as halohydrocarbon activator.

16. The embodiment of claim 15 characterized in that said activator is selected from chlorine-substituted alkanes having no more than two chlorine atoms on any primary carbon atom, no more than one chlorine atom on any secondary carbon atom, and no chlorine atom on any tertiary carbon atom.

Patentansprüche

1. Verfahren zur Herstellung einer Katalysatorkomponente, bei dem man in einer Flüssigkeit die Komponenten (A) und (B) umsetzt, wobei die Komponente (A) folgendes umfaßt: (A1) mindestens ein Vanadiumhalogenid mit einer V-X-Bindung, worin X Halogen bedeutet, und mindestens einer Organophosphorverbindung mit einer P-L-Bindung, worin L ausgewählt ist unter Alkyl-, Alkoxy-, Aryl-, Aryloxy-, Cycloalkyl-, Cycloalkyloxy-, Cycloalkylaryl-, Alkylamino- und Halogenalkyloxyresten oder (A2) mindestens eine Verbindung mit einer funktionellen Gruppe der Formel



worin die Reste L jeweils individuell der vorstehenden Definition entsprechen, und wobei (B) mindestens eine Organoaluminiumverbindung der Formel $\text{AlR}'_n\text{X}_{3-n}$ ist, worin X Halogen bedeutet, die Reste R' jeweils individuell unter Alkyl-, Aryl-, Cycloalkyl- und Alkoxyresten ausgewählt sind und n eine Zahl von 1 bis 3 ist, und das Reaktionsprodukt aus der Flüssigkeit gewinnt, dadurch gekennzeichnet, daß man die Umsetzung in einer Flüssigkeit durchführt, die im wesentlichen aus nicht-aromatischen Halogenkohlenwasserstoffen besteht.

2. Verfahren nach Anspruch 1, dadurch gekennzeichnet, daß das Vanadiumhalogenid mit der Organophosphorverbindung umgesetzt wird und das erhaltene Produkt anschließend mit der Organoaluminiumverbindung umgesetzt wird.

3. Verfahren nach Anspruch 1 oder 2, dadurch gekennzeichnet, daß die Halogenkohlenwasserstoff-Flüssigkeit unter 1-Chlorbutan, Tetrachlorethylen, Dibrommethan und Dichlormethan ausgewählt wird.

4. Verfahren nach Anspruch 3, dadurch gekennzeichnet, daß es sich bei der Halogenkohlenwasserstoff-Flüssigkeit um Dichlormethan handelt.

5. Verfahren nach einem der vorstehenden Ansprüche, dadurch gekennzeichnet, daß die Vanadiumverbindung unter VCl_4 und VOCl_3 ausgewählt wird.



6. Verfahren nach einem der vorstehenden Ansprüche, dadurch gekennzeichnet, daß es sich bei der Vanadiumverbindung um VCl_4 handelt.

7. Verfahren nach einem der vorstehenden Ansprüche, dadurch gekennzeichnet, daß die Organophosphorverbindung ausgewählt wird unter Triethylphosphit, Tributylphosphat, Tributylphosphit, Triethylphosphat, Trioctylphosphat, Trimethylphosphit, Hexamethylphosphoramid, Diethylphosphit, Tri-(2-chlorethyl)-phosphit, Trimethylphosphat, Tri-(2-ethylhexyl)-phosphat, Triisopropylphosphit, Triphenylphosphin und Triphenylphosphinoxid.

8. Verfahren nach einem der vorstehenden Ansprüche, dadurch gekennzeichnet, daß es sich bei der Organophosphorverbindung um Tri-n-butylphosphat handelt.

9. Verfahren nach einem der vorstehenden Ansprüche, dadurch gekennzeichnet, daß es sich bei der Organophosphorverbindung um Triethylphosphit handelt.

10. Verfahren nach einem der vorstehenden Ansprüche, dadurch gekennzeichnet, daß das P/V-Verhältnis der Reaktanten im Bereich von 0,2/1 bis 4/1 und das Al/P-Verhältnis der Reaktanten im Bereich von 0,5/1 bis 2/1 liegt.

11. Verfahren nach einem der vorstehenden Ansprüche, dadurch gekennzeichnet, daß die Organoaluminiumverbindung ausgewählt ist unter Ethylaluminiumdichlorid, Diethylaluminiumchlorid, Trimethylaluminium, Triisobutylaluminium, Ethylaluminiumsesquichlorid und Diethylaluminiummethoxid.

12. Verfahren nach einem der vorstehenden Ansprüche, dadurch gekennzeichnet, daß es sich bei der Organoaluminiumverbindung um Ethylaluminiumdichlorid handelt.

13. Verfahren nach einem der vorstehenden Ansprüche, dadurch gekennzeichnet, daß die gewonnene Katalysatorkomponente von in Kohlenwasserstoffen löslichen Komponenten freigeschwenkt wird.

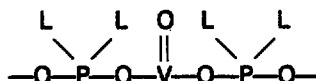
14. Verwendung der nach einem der Ansprüche 1 bis 13 erhältlichen Katalysatorkomponente für die Polymerisation oder Copolymerisation von α -Olefinen.

15. Ausführungsform nach Anspruch 14, dadurch gekennzeichnet, daß die Polymerisation in Gegenwart eines Kokatalysatorsystems durchgeführt wird, das mindestens eine Organoaluminiumverbindung und mindestens ein halogensubstituiertes Alkan mit 1 bis 10 Kohlenstoffatomen als Halogenkohlenwasserstoff-Aktivator enthält.

16. Ausführungsform nach Anspruch 15, dadurch gekennzeichnet, daß der Aktivator unter chloresubstituierten Alkanen mit einem Gehalt an nicht mehr als zwei Chloratomen an beliebigen primären Kohlenstoffatomen, nicht mehr als einem Chloratom an beliebigen sekundären Kohlenstoffatomen und ohne Chloratome an beliebigen tertiären Kohlenstoffatomen ausgewählt wird.

Revendications

1. Procédé de préparation d'un constituant de catalyseur comprenant le fait de faire réagir dans un liquide les constituants (A) et (B), le constituant (A) étant constitué de (Al) au moins un halogénure de vanadium ayant une liaison V—X dans laquelle X est un atome d'halogène et au moins un composé organique du phosphore ayant une liaison P—L dans laquelle L est choisi parmi les radicaux alkyle, alcoxy, aryle, aryloxy, cycloalkyle, cycloalkyloxy, cycloalkylaryle, alkylamino et haloalkyloxy, ou (A2) au moins un composé ayant une fonctionnalité répondant à la formule:



45 dans laquelle chacun des L est individuellement tel que défini précédemment, et (B) étant au moins un composé organique de l'aluminium répondant à la formule AlR'_nX_{3-n} dans laquelle X est un atome d'halogène, chacun des R' est choisi individuellement parmi les radicaux alkyle, aryle, cycloalkyle et alcoxy, et n est un nombre de 1 à 3, et la récupération du produit de la réaction à partir de ce liquide, caractérisé en ce que cette réaction est effectuée dans un liquide essentiellement constitué d'un hydrocarbure halogéné non aromatique.

2. Procédé selon la revendication 1, caractérisé en ce que cet halogénure de vanadium est mis à réagir avec ce composé organique du phosphore et en ce que le produit obtenu est ensuite mis à réagir avec ce composé organique de l'aluminium.

3. Procédé selon les revendications 1 ou 2, caractérisé en ce que ce liquide halo-hydrocarboné est choisi parmi le 1-chlorobutane, le tétrachloréthylène, le dibromométhane et le dichlorométhane.

4. Procédé selon la revendication 3, caractérisé en ce que ce liquide halo-hydrocarboné est le dichlorométhane.

5. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que ce composé du vanadium est choisi parmi VCl_4 et $VOCl_3$.

6. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que ce composé de vanadium est VCl_4 .

7. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que ce composé organophosphoré est choisi parmi le phosphite de triéthyle, le phosphate de tributyle, le phosphite de tributyle, le phosphate de triéthyle, le phosphate de trioctyle, le phosphite de triméthyle,



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l'hexaméthylphosphoramide, le phosphite de diéthyle, le phosphite de tri(2-chloréthyle), le phosphate de triméthyle, le phosphate de tri(2-éthylhexyle), le phosphite de triisopropyle, la triphénylphosphine et l'oxyde de triphénylphosphine.

8. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que ce procédé organophosphoré est le phosphate de tri-n-butyle.

9. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que ce composé organophosphoré est le phosphite de triéthyle.

10. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que le rapport P/V est réactifs est dans l'intervalle de 0,2/1 à 4/1 et en ce que le rapport Al/P des réactifs est dans l'intervalle de 0,5/1 à 2/1.

11. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que ce composé organique de l'aluminium est choisi parmi le dichlorure d'éthylaluminium, le chlorure de diéthylaluminium, le triméthylaluminium, le triisobutylaluminium, le sesquichlorure d'éthylaluminium et l'éthylate de diéthylaluminium.

12. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que ce composé organique de l'aluminium est le dichlorure d'éthylaluminium.

13. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que le constituant de catalyseur récupéré est lavé jusqu'à absence de composés solubles dans les hydrocarbures.

14. Utilisation du constituant de catalyseur tel qu'il peut être obtenu dans l'une quelconque des revendications 1 à 13 pour la polymérisation ou la copolymérisation d' α -oléfines.

15. Mode de réalisation de la revendication 14, caractérisé en ce que cette polymérisation est effectuée en présence d'un système de cocatalyseurs comprenant au moins un composé organique de l'aluminium et au moins un alcane halogéno-substitué ayant 1 à 10 atomes de carbone comme activateur halohydrocarboné.

16. Mode de réalisation de la revendication 15, caractérisé en ce que cet activateur est choisi parmi des alcanes substitués par le chlore n'ayant pas plus de deux atomes de chlore sur tout atome de carbone primaire, pas plus d'un atome de chlore sur tout atome de carbone secondaire et pas d'atome de chlore sur tout atome de carbone tertiaire.