

US005710345A

United States Patent [19]

Navarrini et al.

[11] **Patent Number:** 5,710,345[45] **Date of Patent:** Jan. 20, 1998

[54] **FLOORINATED POLYMERS AND
COPOLYMERS CONTAINING CYCLIC
STRUCTURES**

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[21] **Appl. No.:** 749,504

[22] **Filed:** Nov. 15, 1996

Related U.S. Application Data

[62] **Division of Ser. No. 441,197, May 15, 1995, Pat. No. 5,589,557.**

[30] Foreign Application Priority Data

May 19, 1994 [IT] Italy MI94A1011

[51] **Int. Cl.⁶** C07C 43/03

[52] **U.S. Cl.** 568/596; 562/849

[58] **Field of Search** 562/849; 568/596

[56] References Cited**U.S. PATENT DOCUMENTS**

3,418,302	12/1968	Darby	260/87.5
4,864,006	9/1989	Gianetti et al.	526/209
4,908,461	3/1990	Hung	549/455
4,910,276	3/1990	Nakamura et al.	526/247
5,182,342	1/1993	Feiring et al.	526/206
5,225,576	7/1993	Navarrini et al.	549/449
5,235,074	8/1993	Navarrini et al.	549/449
5,245,054	9/1993	Navarrini et al.	549/455
5,247,035	9/1993	Besecke et al.	526/247
5,260,492	11/1993	Feirung et al.	568/685
5,264,508	11/1993	Ishibe et al.	526/247
5,296,617	3/1994	Navarrini et al.	549/455

5,343,852	9/1994	Ishibe et al.	526/247
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FOREIGN PATENT DOCUMENTS

0247379	12/1989	European Pat. Off.	
0633257	1/1995	European Pat. Off.	
1106344	3/1968	United Kingdom	526/247

OTHER PUBLICATIONS

Inorganic Chemistry, vol. 7, No. 3, 1968, pp. 624-626,
"Some Reactions of Bis(Fluoroxo)Difluoromethane,
CF₂(OF)₂," Hohorst et al.

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[57] ABSTRACT

Polymers and copolymers of one or more fluorinated dienes of structure CFX¹=CX²—O—CX³X⁴—O—CX²=CX¹F, wherein X¹ and X² are F, Cl or H, and X³ and X⁴ are F or CF₃, wherein said dienes essentially form cyclic repetitive units, the comonomers used for preparing copolymers being ethylene unsaturated fluorinated compounds.

Processes for preparing the fluorinated dienes having said structure which comprise the reaction in solution between one halogenated olefin and one hypofluorite CX³X⁴(OF)₂, wherein X³ and X⁴ are F or CF₃, and the dehalogenation or dehydrohalogenation of the linear adduct between one hypofluorite molecule and two olefin molecules.

The polymers and copolymers of the invention are particularly suitable for preparing coatings for applications at high temperatures.

13 Claims, No Drawings

FLOORINATED POLYMERS AND COPOLYMERS CONTAINING CYCLIC STRUCTURES

This application is a division of application Ser. No. 08/441,197 filed May 15, 1995 which application is now: U.S. Pat. No. 5,589,557.

The present invention relates to fluorinated polymers and copolymers containing cyclic structures in the main chain. Specifically the present invention relates to fluorinated polymers and copolymers containing cyclic structures obtained by cyclopolymerization of dienes.

The polymers of the invention do not show gelling phenomena during the process for preparing them. This is important, as a matter of fact the possible formation of gels would lead to polymers having mechanical, processability and solubility characteristics unsuitable to some applications. On the other hand, the separation of gels from polymeric crude products is difficult and not always possible.

The invention relates therefore in particular to fluorinated polymers and copolymers, containing said cyclic structures, thermoprocessable and having improved thermostability combined with high solubility and absence of gels.

Said polymers and copolymers can be used in particular in preparing coatings for applications at high temperatures, where it is necessary to have high thermal stability and also high solubility to obtain solutions having high concentration of polymer.

It is known in the art that polymers containing cyclic structures in the main chain can be obtained by radical polymerization of non conjugated dienes. In particular the cyclopolymerization of fluorinated non conjugated dienes is described in U.S. Pat. Nos. 3,418,302, 4,910,276 and 5,260,492.

In U.S. Pat. No. 3,418,302 it is stressed in general that the formation of not crosslinked saturated linear polymers from compounds containing two double bonds is extremely difficult and in particular how it is possible in the case of perfluorodimethylenebisvinylether with formation of cyclic structures in the main chain only on condition that the monomer is very diluted in the polymerization step, its concentration having to be lower than 12% by weight of the total monomers and diluents. In fact if the concentration of the dienic monomer is higher, gel phenomena occur which lead to the above mentioned drawbacks.

To avoid the gel formation it was suggested in the art to resort to the use of monomers having particular molecular structures.

For instance, in U.S. Pat. No. 4,910,276 it is described the cyclopolymerization of divinyl fluorinated monomers also at high concentration of the same during polymerization, but a necessary condition for its occurrence is that the monomers have two vinylic groups with different reactivity, in practice said groups must have different structure. For the homopolymers according to this invention the thermostability data show a maximum temperature of 475° C. in relation to a weight loss of 10% because of thermal effect, while as to solubility the highest value is 7%.

Likewise U.S. Pat. No. 5,260,492 describes the cyclopolymerization of non conjugated fluorinated dienes, in particular of the ω -alkenyl-vinylethers type, but also in this case monomers having unsaturated groups with different reactivity are used.

Also polymers and copolymers obtained according to the last patents do not show in general an high thermal stability. In the case of homopolymers there is indeed a thermostability just slightly higher than that of the polymers according

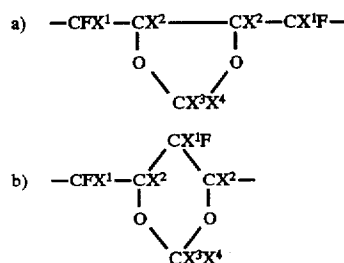
to U.S. Pat. No. 4,910,276: the maximum attainable temperature before loosing 10% by weight because of thermal effect is about 485° C.

Moreover, in the preparation of polymers according to said patents one has, to a different extent, the rearrangement of the vinylether group with formation of acylfluoride —COF, which, as known in the art, leads to a reduction of the thermal stability of the polymers themselves. If it is desired to reduce the content of —COF of said polymers, one could resort to a subsequent fluorination process, according to what known in the art, such resort, however, would make more complex the preparation of the polymers themselves.

It has now been surprisingly and unexpectedly found that it is possible to obtain by diene cyclopolymerization linear fluorinated polymers thermally more stable than those of the prior art, the content in fluorine being same, free from gels and endowed with high solubility, if a particular class of non conjugated fluorinated dienes is used. Such dienes, contrary to what reported in U.S. Pat. No. 4,910,276, cyclopolymerize even though they have two vinylic groups with the same reactivity. Moreover, unexpectedly, with the monomers of the invention, rearrangement phenomena and formation of —COF groups do not occur.

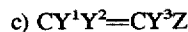
According to the present invention it has surprisingly and unexpectedly found that fluorinated polymers and copolymers with cyclic structures in main chain having said characteristics, can be obtained by using, as diene, bisvinylloxymethanes having the structure $\text{CFX}^1=\text{CX}^2-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CX}^2=\text{CX}^1\text{F}$, wherein X^1 and X^2 , equal to or different from each other, are F, Cl or H; X^3 and X^4 , equal to or different from each other, are F or CF_3 .

Object of the present invention are therefore polymers and copolymers derived from one or more of said bisvinylloxymethanes forming essentially only repetitive cyclic units having the structures a) and b):

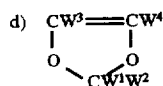


wherein X^1 , X^2 , X^3 and X^4 have the meaning indicated above.

In the case of copolymers, at least one of the other comonomers is a compound with an ethylene unsaturation. Utilizable comonomers of this type are olefins, optionally containing also chlorine and/or fluorine atoms, alkylvinylethers and dioxolic compounds. More precisely olefins have the structure of the type



wherein Y^1 , Y^2 , Y^3 , equal to or different from each other, are F, Cl or H; Z is F, Cl, H or a perfluoroalkyl radical R' containing from one to five carbon atoms, with the proviso that the olefin contains 2 chlorine atoms at most and preferably no more than one chlorine atom; alkylvinylethers have the same structure c) wherein Z is $\text{O---R}'$ with R' equal to a perfluoroalkyl radical having from one to five carbon atoms, with the same limitations relating to the content of chlorine atoms. The dioxolic compounds have the structure:



wherein W^1 and W^2 , equal to or different from each other, represent F or CF_3 , W^3 represents F or H, and W^4 represents F, H, R' or $O-R'$ with R' equal to a perfluoroalkyl radical having from 1 to 5 carbon atoms. The preparation of said dioxolic compounds is described in U.S. Pat. Nos. 4,908, 461, 5,245,054, 5,296,617 when W^4 is different from R' and $O-R'$. When W^4 is R' or $O-R'$, for different dioxoles than the ones cited in said U.S. patents see the European patent application EP-A-633257.

Preferred polymers and copolymers are those derived from monomers with $X^2=X^1$, having the structure $CFX^1=CX^1-O-CX^3X^4-O-CX^1=CF$, wherein X^1 is F, Cl or H; X^3 and X^4 , equal to or different from each other, are F or CF_3 ; more preferred polymers and copolymers are those derived from monomers having the structure $CF_2=CF-O-CX^3X^4-O-CF=CF_2$, wherein X^3 and X^4 , equal to or different from each other, are F or CF_3 and even more preferred are those derived from perfluorobisvinylloxymethane $CF_2=CF-O-CF_2-O-CF=CF_2$.

As comonomers, one or more of the following compounds can be preferably used: tetrafluoroethylene, vinylidene fluoride, ethylene, chlorotrifluoroethylene, trifluoroethylene, perfluoromethylvinylether, perfluoropropylvinylether, perfluoropropene, perfluoro 2,2-dimethyl-1,3-dioxole, perfluoro 1,3-dioxole, 2,2,4-trifluoro-5-trifluoromethoxy-1,3-dioxole.

The obtained polymers show the combination of the aforesaid cyclic structures in main chain, the 5 atoms ring structures being prevalent, while both open structures containing double bonds in side chain and carbonylic groups deriving from the rearrangement of the perfluorovinylether group are substantially absent.

The polymers and copolymers according to the present invention, besides being chemically stable, have a surprising thermal stability, high transparency and in some cases, even though they have a high content in fluorine, are soluble in some common solvents, such as for instance acetone, diethyleneglycol dimethylether and N,N-dimethylformamide. The perfluorobisvinylloxymethane homopolymer and some copolymers according to this invention show on the other hand a good solubility in the fluorinated solvents, such as hydrochlorofluorocarbons, fluorohydrocarbons, perfluoroamines, perfluoroethers, similar solvents and their mixtures and high solubility in Fluorinert® FC 75 (perfluoro-2-butyltetrahydrofuran) and in perfluoropolyethers Fomblin® and Galden® having, optionally, one or more H in one or in both the ending groups (e.g. $-CF_2H$).

In particular, the perfluorobisvinylloxymethane homopolymer according to this invention, has a weight loss of 10% only at a temperature of about 530° C. and it shows an high solubility in the perfluoropolyethers Fomblin® and Galden® and in Fluorinert® FC 75.

These properties make such polymers and copolymers very suitable to be used also for certain applications as protective coating both of electric wires and particularly of metals, in applications of optical type, particularly in the manufacture of optical fibers, where the low refraction index and the high transparency of these products are used also in the area of wave lengths corresponding to the ultraviolet, and in the preparation of manufactured articles where high thermal stability is required.

The polymers and copolymers of the present invention can be prepared by radical polymerization in solution or in absence of solvent, in suspension or in mass.

General polymerization methodologies utilizable in a non aqueous medium are described for instance in U.S. Pat. Nos. 4,864,006 and 5,182,342, while methodologies utilizable in an aqueous medium are described in EP Patent 247379 and again in U.S. Pat. No. 5,182,342. Any substance capable of generating radicals under the chosen reaction conditions can be used as polymerization initiator, in particular perfluoropropionylperoxide, benzoylperoxide, azobisisobutyronitrile or percarbonates can be used. It is also possible to start the polymerization using systems of redox couples, for instance those described in Prog. Polym. Sci., vol. 8, page 61, 1982.

The fluorinated derivatives of bisvinylloxymethane having the structure $CFX^1=CX^2-O-CX^3X^4-O-CX^2=CF$, wherein X^1 and X^2 , equal to or different from each other, are F, Cl or H; X^3 and X^4 , equal to or different from each other, are F or CF_3 ; X^1 and X^2 , equal to or different from each other, are X^1 or X^2 , with the proviso that when X^1 is different from X^2 also X^1 is to be different from X^2 , are new as such and are obtainable according to an innovative process. Among these compounds, those wherein $X^1=X^1$ and $X^2=X^2$ cyclopolymerize without giving $-COF$ groups in the polymer; this occurs in particular in case of perfluorobisvinylloxymethane $CF_2=CF-O-CF_2-O-CF=CF_2$.

A further object of the present invention is therefore the process for preparing fluorinated derivatives of bisvinylloxymethane having the structure $CFX^1=CX^2-O-CX^3X^4-O-CX^2=CF$, wherein X^1 and X^2 equal to or different from each other are F, Cl or H, X^3 and X^4 equal to or different from each other are F or CF_3 , X^1 and X^2 equal to or different from each other are X^1 or X^2 , with the proviso that when X^1 is different from X^2 also X^1 is different from X^2 , which comprises:

- i) addition of an olefin of formula $CX^1X^5=CX^2X^6$, wherein X^1 , X^2 , X^5 and X^6 equal to or different from each other are F, Cl, H or Br, the Br atoms being 2 at most and in such a case bound to different carbon atoms, X^1 and X^5 not being both F, X^2 and X^6 not being both F, X^1 and X^2 being F only if X^5 and X^6 are different from F and not being both H, in a reactor where an hypofluorite of general formula $CX^3X^4(OF)_2$ wherein X^3 and X^4 equal to or different from each other are F or CF_3 , is essentially always present, dissolved in an inert solvent having a concentration comprised between 0.001M and 10M, at a temperature from -140° C. to +60° C.;
- ii) separation of the reaction product between two olefin molecules and one hypofluorite molecule from the reaction mixture obtained in i) by fractional distillation.
- iii) dehalogenation or dehydrohalogenation of the product obtained in ii), where the eliminated halogen atoms are Cl or Br.

The Applicant has surprisingly found experimental conditions such as to substantially and not predictably modify the reactivity of the hypofluorites having the general formula $CXY(OF)_2$ towards the olefins when one operates in solution, succeeding in obtaining with good yield linear reaction products between one molecule of hypofluorite and two molecules of olefin.

According to U.S. Pat. Nos. 5,225,576 and 5,235,074, such reactivity leads in fact essentially to the formation of a 1,3-dioxolane derivative and to the fluorination product of the olefin itself.

Linear addition products between hypofluorite and olefins in the presence of solvent are described in Inorganic Chemistry vol. 7, No. 3, 1968, page 624-6, where the hypofluorite

and the olefin are first condensed in stoichiometric ratio at -184°C . and by subsequent heating they are reacted in absence of an excess of hypofluorite, conditions under which the preferred product is not the linear product but the dioxolanic compound, as described in the aforesaid patents.

The experimental conditions which surprisingly allow to obtain the linear addition product as preferred product refer to the addition methodology of the reactants used and to the polarity of the reaction medium.

It has been surprisingly found as a matter of fact that by adding the olefin to the hypofluorite dissolved in a solvent, provided that it does not react with the hypofluorite, with a concentration from 0.001M to 10M, preferably from 0.1M to 4M, at a temperature from -140°C . to $+60^{\circ}\text{C}$. and preferably from -120°C . to 0°C ., the reaction proceeds with a clear increase of the formation of the aforesaid linear reaction product. The inert solvent can be an ordinary one with low polarity, for instance dichlorodifluoromethane, trichlorofluoromethane, Fluorinert® FC 75 or a perfluoropolyether Fomblin® and Galden®, but preferably the solvent used is a polar one. The polar solvent is preferably selected from hydrogenfluorocarbons, hydrogenchlorocarbons, fluorochlorocarbons, hydrogenchlorofluorocarbons, trifluoroacetic acid, trifluoroacetic anhydride, acetic nitrile, hydrofluoric acid, sulphur dioxide, trifluoromethanesulphonic acid, $\text{CF}_2\text{Cl}-\text{CFCl}-\text{SO}_2\text{F}$, mixtures of the same and mixtures of one or more of said solvents also in small percentage with an ordinary solvent having low polarity. Among chlorinated and fluorochlorinated solvents the ones containing also hydrogen in the molecule are preferred, because of their lower impact on the decrease in the atmosphere ozone layer.

The olefin used in the process is selected depending on the desired intermediate or dienic compound. Preferred olefins are $\text{CFCI}=\text{CFCI}$, $\text{CHCl}=\text{CHCl}$, $\text{CHCl}=\text{CCl}_2$, $\text{CCl}_2=\text{CCl}_2$, $\text{CH}_2=\text{CF}_2$, $\text{CF}_2=\text{CF}_2$, $\text{CFH}=\text{CFCI}$, $\text{CFCI}=\text{CHCl}$, $\text{CH}_2=\text{CHCl}$, $\text{CH}_2=\text{CCl}_2$, $\text{CH}_2=\text{CFCl}$.

The process can be carried out both in a continuous way, introducing the olefin in a reactor where hypofluorite is always present at a concentration kept constant and in a discontinuous way by adding the olefin to an hypofluorite solution in a suitable solvent without restoring the concentration of the hypofluorite which progressively reacts.

The dehalogenation or dehydrohalogenation of the product obtained in the phase ii) of the process can be carried out according to one of the methods described in the art.

The process is preferably used for the preparation of bisvinylloxymethane derivatives having the structure $\text{CFX}^1=\text{CX}^1-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CX}^1=\text{CX}^1\text{F}$, wherein X^1 is F, Cl or H; X^3 and X^4 , equal to or different from each other, are F or CF_3 , using an olefin of formula $\text{CX}^1\text{X}^5=\text{CX}^1\text{X}^6$, where X^1 , X^5 and X^6 , equal to or different from each other, are F, Cl, Br or H, X^1 and X^5 not being both F, X^1 and X^6 not being both F, X^1 being Br only when X^5 and X^6 equal to each other are different from Br, X^1 being F only when X^5 and X^6 are different from F and not being both H.

The process is used even more preferably for the preparation of bisvinylloxymethane derivatives having the structure $\text{CF}_2=\text{CF}-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CF}=\text{CF}_2$, wherein X^3 and X^4 , equal to or different from each other, are F or CF_3 , using an olefin of formula $\text{CFX}^2=\text{CFX}^6$, wherein X^2 is Cl, Br or H, and X^6 is Cl or Br.

It has been found, moreover, that if the solvent used in the reaction between olefin and hypofluorite is of polar type, the addition sequence of the reactants can be inverted. A further object of the present invention is therefore a process for

preparing fluorinated derivatives of bisvinylloxymethane having the structure $\text{CFX}^1=\text{CX}^2-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CX}^2=\text{CX}^1\text{F}$, wherein X^1 and X^2 , equal to or different from each other, are F, Cl or H; X^3 and X^4 , equal to or different from each other, are F or CF_3 ; X^1 and X^2 , equal to or different from each other, are X^1 or X^2 , with the proviso that if X^1 is different from X^2 also X^1 is to be different from X^2 , comprising:

i) addition of hypofluorite having the general formula $\text{CX}^3\text{X}^4(\text{OF})_2$ wherein X^3 and X^4 , equal to or different from each other, are F or CF_3 , to an olefin of formula $\text{CX}^1\text{X}^5=\text{CX}^2\text{X}^6$, wherein X^1 , X^2 , X^5 and X^6 , equal to or different from each other, are F, Cl, H or Br; the Br atoms being 2 at most and in such a case bound to different carbon atoms; X^1 and X^5 not being both F; X^2 and X^6 not being both F; X^1 and X^2 being F only if X^5 and X^6 are different from F and not being both H; said olefin being dissolved in an inert solvent of polar type, at a temperature from -140°C . to $+60^{\circ}\text{C}$., preferably from -120°C . to 0°C .;

ii) separation of the reaction product of two olefin molecules with one hypofluorite molecule from the reaction mixture obtained in i) by fractional distillation.

iii) dehalogenation or dehydrohalogenation of the product obtained in ii), where the eliminated halogen atoms are Cl or Br.

Preferably the inert solvent of polar type is selected from hydrogenfluorocarbons, hydrogenchlorocarbons, fluorochlorocarbons, hydrogenchlorofluorocarbons, trifluoroacetic acid, trifluoroacetic anhydride, acetic nitrile, hydrofluoric acid, sulphur dioxide, trifluoromethanesulphonic acid, $\text{CF}_2\text{Cl}-\text{CFCl}-\text{SO}_2\text{F}$, mixtures of the same and mixtures of one or more of said solvents also in small percentage with an ordinary low polarity solvent.

Also for this process the preferred olefins are $\text{CFCI}=\text{CFCI}$, $\text{CHCl}=\text{CHCl}$, $\text{CHCl}=\text{CCl}_2$, $\text{CCl}_2=\text{CCl}_2$, $\text{CH}_2=\text{CF}_2$, $\text{CF}_2=\text{CF}_2$, $\text{CFH}=\text{CFCI}$, $\text{CFCI}=\text{CHCl}$, $\text{CH}_2=\text{CHCl}$, $\text{CH}_2=\text{CCl}_2$, $\text{CH}_2=\text{CFCl}$.

The dehalogenation or dehydrohalogenation of the product obtained in step ii) of the process can be likewise carried out according to one of the methods described in the art.

Like the previous process, also this process is preferably used for the preparation of bisvinylloxymethane derivatives having the structure $\text{CFX}^1=\text{CX}^1-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CX}^1=\text{CX}^1\text{F}$, wherein X^1 is F, Cl or H; X^3 and X^4 , equal to or different from each other, are F or CF_3 , by using an olefin of formula $\text{CX}^1\text{X}^5=\text{CX}^1\text{X}^6$ wherein X^1 , X^5 and X^6 equal to or different from each other are F, Cl, Br or H, X^1 and X^5 not being both F, X^1 and X^6 not being both F, X^1 being Br only when X^5 and X^6 equal each other are different from Br, X^1 being F only when X^5 and X^6 are different from F and not being both H.

Likewise also this process is even more preferably used for preparing bisvinylloxymethane derivatives having the structure $\text{CF}_2=\text{CF}-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CF}=\text{CF}_2$, wherein X^3 and X^4 , equal to or different from each other, are F or CF_3 , using an olefin of formula $\text{CFX}^5=\text{CFX}^6$, wherein X^5 is Cl, Br or H, and X^6 is Cl or Br.

The fluorinated derivatives of bisethoxymethane obtainable after steps i) and ii) of the processes described above, possibly as a mixture of more compounds, are a further object of the present invention. These products, besides being used as intermediates for the synthesis of monomers, are also used as fluorinated solvents with low impact on the atmospheric ozone layer. The products without chlorine are preferred for this use.

The following examples are to be considered for illustrative purposes, but not limitative of the scope of the present invention.

EXAMPLES 1-16

Bisethoxymethane derivatives from olefin and hypofluorite: addition of the hypofluorite to the olefin in solvents having different polarity.

EXAMPLE 1

80 g (600 mmoles) of 1,2-dichloro-1,2-difluoroethylene and a solvent mixture formed by 32 g of SO₂ and 17 g of CH₂Cl₂ are introduced into a 125 ml multi-neck glass cylindrical reactor, equipped with mechanical stirrer, thermocouple, suction inlet for the reacting gaseous mixture and inert gas head outlet.

The so loaded reactor is brought to the temperature of -80° C. using a cryostat, and then a mixture of CF₂(OF)₂ (1.5 NI/h), CO₂ (2.2 NI/h) and helium (6 NI/h) are continuously added under mechanical stirring for 3.5 hours with an overall supply of hypofluorite equal to 234 mmoles.

The reaction is practically instantaneous. When the addition of the mixture containing hypofluorite is over, the products are separated from the reaction crude product by fractional distillation with a plate column at atmospheric pressure. The fractions having boiling point in the range comprised between -1° and +1° with respect to the temperatures indicated hereinafter, are collected:

- a) at +4° C. 19.3 g (112 mmoles) of CF₂Cl—CF₂Cl
- b) at +22° C. 17.4 g (131 mmoles) of unreacted CFCI=CFCI
- c) at 47/49° C. mixture formed by 24.4 g (113 mmoles) of 4,5-dichloro-2,2,4,5-tetrafluoro-1,3-dioxolane and 4 g (22 mmoles) of CClF₂—CCl₂F
- d) at +76° C. 8.6 g (40 mmoles) CF₂Cl—CFCI—OC(O)F
- e) at +70.5° C./54 mm Hg 20.4 g (52 mmoles) of CF₂Cl—CFCI—OCF₂O—CFCI—CF₂Cl.

The linear addition product yield, defined as ratio between the obtained moles of CF₂Cl—CFCI—OCF₂O—CFCI—CF₂Cl and the used moles of CF₂(OF)₂, is 22%. The product has been characterized by F¹⁹-NMR spectroscopy (in p.p.m. related to CFCI₃=O: 2F type O—CF₂—O at -50.5/-52.0; 4F type Cl—CF₂—C at -70.3/-72.3; 2F type C—CFCI—O at -77.5), and electronic impact mass spectroscopy, where the main peaks and the relative intensities are: 151 (100%), 101 (23%), 85 (15%), 66 (12%), 47 (12%), 28 (33.5%).

EXAMPLES 2-11

In these examples some solvents and co-solvents are tested at different temperatures under experimental conditions comparable with the ones of example 1. The data relating to the yield in linear addition product formed by two molecules of olefin and one molecule of hypofluorite and under the experimental conditions adopted are reported in Table 1.

EXAMPLE 12

In this example the use of a different olefin, the trans-1,2-dichloroethylene, is tested.

174 g (1.79 moles) of trans-1,2-dichloroethylene and a solvent mixture formed by 118 g of SO₂ and 247 g of CFCI₃ are introduced into a 500 ml multi-neck glass cylindrical reactor, equipped with mechanical stirrer, thermocouple, suction inlet for the reacting gaseous mixture and inert gas head outlet.

The so loaded reactor is brought to the temperature of -76° C. using a cryostat, and then a mixture of CF₂(OF)₂ (2 NI/h), CO₂ (1.0 NI/h) and helium (6 NI/h) are continuously

added under mechanical stirring for 8.5 hours with an overall supply of hypofluorite equal to 760 mmoles.

When the addition of hypofluorite is over, the sulphur dioxide and part of CFCI₃ are separated from the reaction crude product by distillation. Then the reaction products are separated by fractional distillation with a plate column at reduced pressure. The fractions having boiling point in the range between -1° and +1° with respect to the temperature indicated hereinafter, are collected:

- a) at +20° C./70 mm Hg mixture of 50 g of CFCIH—CFCIH and 25 g of unreacted trans-1,2-dichloroethylene;
- b) at +29° C./58 mm Hg 46 g of 4,5-dichloro-2,2-difluoro-1,3-dioxolane;
- c) at +12° C./24 mm Hg 20 g of CFCIH—CCIH—OC(O)F
- d) at +64° C./3 mm Hg 71 g of CFCIH—CHCl—OCF₂O—CHCl—CFCIH.

The linear addition product yield, defined as ratio between the obtained moles of CFCIH—CHCl—OCF₂O—CHCl—CFCIH and the used moles of CF₂(OF)₂, is 30%. The product has been characterized by F¹⁹-NMR spectroscopy (in p.p.m. related to CFCI₃=O: 2F type O—CF₂—O at -58.0/-61.5; 2F type CHFCl—CHCl— at -141/-146), H¹-NMR spectroscopy (in p.p.m. related to TMS=0: complex multiplet at +5.6/6.9) and mass spectroscopy (main peaks: 209, 211, 213 for C₄H₃OCl; 179, 181, 183 for C₃H₂OF₃Cl₂; 131, 133, 135 for C₂H₂OFCl; 15, 117, 119 for CH₂FCl₂).

EXAMPLES 13 AND 14

In these examples the use of trans-1,2-dichloroethylene is tested, with the same procedure of example 12, under different conditions reported in Table 1 along with the linear addition product yield.

EXAMPLE 15

79 g (0.6 moles) of trichloroethylene and a solvent mixture formed by 39 g of SO₂ and 83 g of CFCI₃ are introduced into a 500 ml multi-neck glass cylindrical reactor, equipped with mechanical stirrer, thermocouple, suction inlet for the reacting gaseous mixture and inert gas head outlet.

The so loaded reactor is brought to the temperature of -74° C. using a cryostat, and then a mixture of CF₂(OF)₂ (1 NI/h), CO₂ (0.5 NI/h) and helium (3 NI/h) are continuously added under mechanical stirring for 6 hours.

When the addition of hypofluorite is over, solvents and most volatile reaction products CHFCl—CFCI₂ and 4,4,5-trichloro-2,2,4-trifluoro-1,3-dioxolane are stripped off the crude reaction mixture.

33 g of isomeric, essentially pure bisethereal addition products, having molecular weight 382.8, are left in the kler. The bisethereal products can be separated by gaschromatography. The characterization is reported in example 18.

The linear addition product yield, defined as ratio between the moles of obtained bisethereal products and the used moles of CF₂(OF)₂, is 32%.

EXAMPLE 16

100 g (0.6 moles) of tetrachloroethylene and a solvent mixture formed by 39 g of SO₂ and 124 g of CFCI₃ are introduced into a 500 ml multi-neck glass cylindrical reactor, equipped with mechanical stirrer, thermocouple, suction inlet for the reacting gaseous mixture and inert gas head outlet.

The so loaded reactor is brought to the temperature of -54° C. using a cryostat, and then a mixture of CF₂(OF)₂ (1

NI/h), CO₂ (0.5 NI/h) and helium (3 NI/h) are continuously added under mechanical stirring for 6 hours.

When the addition of hypofluorite is over, solvents and most volatile reaction products CFCl₂—CFCl₂ and 4,4,5,5-tetrachloro-2,2-difluoro-1,3-dioxolane are stripped off the crude reaction mixture.

32 g of the essentially pure bisethereal addition product CFCl₂—CCl₂—OCF₂O—CCl₂—CFCl₂ are left in the kier.

The linear addition product yield, defined as ratio between the moles of obtained bisethereal product and the used moles of CF₂(OF)₂, is 26%.

The product CFCl₂—CCl₂—OCF₂O—CCl₂—CFCl₂ is characterized by F¹⁹-NMR spectroscopy, where it shows in p.p.m., related to CFCl₃=O, 2F type OCF₂O at -50.7 and 2F type CFCl₂ at -67.5

The mass spectroscopy analysis shows the following main peaks and relative intensities: 185 (100%), 148 (15%), 101 (12%).

TABLE 1

Formation of linear addition product between hypofluorite and olefin: addition of hypofluorite to the olefin(*)					
Ex.	Solvent (g)	Cosolvent (g)	Temperature (°C.)	Linear addition product yield (% by moles)	
1	SO ₂	32 CH ₂ Cl ₂	17	-80	22.0
2	CH ₂ Cl ₂	8		-90	10.5
3	SO ₂	32		-80	18.6
4	SO ₂	19.2		-80	17.5
5	SO ₂	19.2		-90	8.5
6	SO ₂	19.2		-60	17.0
7	SO ₂	64 CH ₂ Cl ₂	34	-80	18.3
8	CF ₂ Cl—CFCl—SO ₂ F	127		-80	18.5
9	HF	8		-80	12.4
10	(CF ₃ C(O)) ₂ O	8		-70	12.0
11	CF ₃ SO ₃ H	8		-55	11.0
12	SO ₂	118 CFCl ₃	247	-76	30.0
13	SO ₂	118 CH ₂ Cl ₂	153	-70	43.0
14	—	CHCl ₂ —CF ₃	250	-70	36.0
15	SO ₂	39 CFCl ₃	83	-74	32.0
16	SO ₂	39 CFCl ₃	124	-54	26.0

(*)The hypofluorite is CF₂(OF)₂; the olefin is 1,2-dichloro-difluoroethylene in the examples from 1 to 11, trans 1,2-dichloroethylene in the examples from 12 to 14, trichloroethylene in example 15 and tetrachloroethylene in example 16.

EXAMPLES 17-22

Bisethoxymethane derivatives from olefin and hypofluorite: addition of olefin to hypofluorite.

EXAMPLE 17

2 ml of CFCl₃ are loaded in a 10 ml two necks glass flask equipped with magnetic stirrer and internal thermocouple.

The reactor is brought to -196° C. and evacuated. Subsequently by vacuum line transfer 1 mmole of CF₂(OF)₂ is condensed therein at -196° C. The so loaded reactor is brought to -100° C. and the solution of CF₂(OF)₂ in CFCl₃ is homogenized under magnetic stirring. 3 mmoles of CFCl=CFCl are slowly added by condensation to the so obtained 0.5 molar solution of CF₂(OF)₂ in CFCl₃ and maintained at -100° C. so that the reaction heat immediately developed is effectively vented by the cooling system.

The weight percentages of the reaction products, determined by gaschromatography and by NMR, are: CF₂Cl—CF₂Cl 40.3%; 4,5-dichloro-2,2,4,5-tetrafluoro-1,3-dioxolane 50.7%; CF₂Cl—CFCl—OCF₂O—CFCl—CF₂Cl 9%. The linear addition product, defined as for example 1, is 9%.

EXAMPLES 18 AND 19

Under experimental conditions comparable with those of example 17, some solvents and CF₂(OF)₂ concentrations are tested at different temperatures. The data relating to the linear adduct yield and to the adopted experimental conditions are reported in Table 2.

EXAMPLE 20

20 ml of CHCl₂—CF₃ (d=1.5 g/l) are loaded into a 125 ml multi-neck glass cylindrical reactor, equipped with mechanical stirrer, thermocouple, suction inlet for the reacting gaseous mixture, inlet on the reactor top for the liquid reacting olefin, inert gas head outlet. The so loaded reactor is brought to -97° C. by means of a cryostat, and then a gaseous mixture of CF₂(OF)₂ (0.50 NI/h), CO₂ (0.25 NI/h) and He (1.5 NI/h) is continuously introduced under mechanical stirring until a 1.5M concentration of CF₂(OF)₂ in solution is achieved.

45

At this point CHCl=CCl₂ in admixture with hypofluorite is continuously added so as to keep constant the concentration of the latter in the solution during the addition. After 4 h and 30', the addition of the gaseous mixture containing CF₂(OF)₂ is interrupted and the addition of CHCl=CCl₂ is continued up to the complete conversion of CF₂(OF)₂ dissolved in the solution.

At the end of the reaction, the reaction crude product is stripped from the solvent and from the most volatile reaction by-products: CHFCI—CFCl₂ and 4,4,5-trichloro-2,2,4-trifluoro-1,3-dioxolane. 34.5 g of a mixture of bisethereal addition products formed by:

A) 71.5% of CCl₂F—CHCl—OCF₂O—CHCl—CCl₂F

B) 27.7% of CCl₂F—CHCl—OCF₂O—CCl₂—CHClF

C) 0.8% of CHClF—CCl₂—OCF₂O—CCl₂—CHClF

are obtained as residue.

The products are separated by preparative gaschromatography. The linear addition product yield, defined as ratio between the sum of the moles of products A, B and C and the moles of CF₂(OF)₂ converted, is 69%.

The products are characterized by F¹⁹-NMR spectroscopy where they show, in p.p.m., related to CFCl₃=O:

60

65

product A: 2F type CFCl_2 at -66.5 ; 2F type OCF_2O at $-59.3/-61.5$;

product B: 1F type CFCl_2 at -66.2 ; 1F type CFHCl at -138.3 ; 2F type OCF_2O at $-55.0/-56.3$;

product C: 2F type CFHCl at -138.4 ; 2F type OCF_2O at -50.6 .

The mass spectroscopy analysis shows the following main peaks and relative intensities: 151 (100%), 131 (23.5%), 114 (33%), 101 (20%), 79 (14%).

EXAMPLE 21

54 ml of $\text{CH}_2\text{F}-\text{CF}_3$ ($d=1.2$ g/l) are loaded into a 125 ml multi-neck glass cylindrical reactor, equipped with mechanical stirrer, thermocouple, suction inlet for the reacting gaseous mixture $\text{CF}_2(\text{OF})_2/\text{CO}_2$ /diluent gas, suction inlet for the reacting olefin, inert gas head outlet. The so loaded reactor is brought to -90°C . by means of a cryostat, and then a gaseous mixture of $\text{CF}_2(\text{OF})_2$ (1.0 NI/h), CO_2 (0.5 NI/h) and He (4 NI/h) is continuously introduced under mechanical stirring until a 0.2M concentration of $\text{CF}_2(\text{OF})_2$ in solution is achieved.

At this point $\text{CClF}=\text{CHF}$ (2.0 NI/h) is continuously added so as to keep constant the concentration of the hypofluorite in the solution during the addition. After 1 h and 40', the addition of the gaseous mixture containing $\text{CF}_2(\text{OF})_2$ is interrupted and the addition of $\text{CClF}=\text{CHF}$ is continued up to the complete conversion of $\text{CF}_2(\text{OF})_2$ dissolved in the solution.

At the end of the reaction, the reaction crude product is stripped from the solvent and from the most volatile reaction by-products: $\text{CHF}_2-\text{CF}_2\text{Cl}$ and 4-chloro-2,2,4,5-tetrafluoro-1,3-dioxolane. 16.5 g of a mixture of bisethereal addition products formed by:

A) 74% of $\text{CClF}_2-\text{CHF}-\text{OCF}_2\text{O}-\text{CHF}-\text{CClF}_2$

B) 23% of $\text{CClF}_2-\text{CHF}-\text{OCF}_2\text{O}-\text{CClF}-\text{CHF}_2$

C) 3% of $\text{CHF}_2-\text{CClF}-\text{OCF}_2\text{O}-\text{CClF}-\text{CHF}_2$ are obtained as residue.

The products are separated by preparative gaschromatography. The linear addition product yield, defined as ratio between the sum of the moles of products A, B and C and the moles of $\text{CF}_2(\text{OF})_2$ converted, is 61%. The products are characterized by F^{19} -NMR spectroscopy and H^1 -NMR spectroscopy.

By F^{19} -NMR spectroscopy the products show, in p.p.m., related to $\text{CFCl}_3=\text{O}$:

product A: 4F type CF_2Cl at -71.5 ; 2F type CHF at -140.3 ; 2F type OCF_2O at $-58.5/-60.0$;

product B: 2F type CF_2Cl at -71.5 ; 1F type CHF at -140.3 ; 1F type CFCl at -81.2 ; 2F type CHF_2 at $-131/-134$; 2F type OCF_2O at $-54.0/-56.6$;

product C: 2F type CFCl at -81.2 ; 4F type CHF_2 at $-131/-134$; 2F type OCF_2O at $-54.0/56.6$.

By H^1 -NMR spectroscopy the products show, in p.p.m., related to $\text{TMS}=\text{O}$:

product A: 2H type CHF at 5.8-6.0

product B: 1H type CHF at 5.8-6.0; 1H type CF_2H at 5.7-5.9-6.1

product C: 2H type CF_2H at 5.7-5.9-6.1

The electronic impact mass spectroscopy analysis shows the following main peaks and relative intensities: 185 (7%), 183 (20%), 119 (38%), 117 (100%), 67 (15%).

EXAMPLE 22

160 ml of $\text{CH}_2\text{F}-\text{CF}_3$ ($d=1.2$ g/l) are loaded into a 250 ml multi-neck glass cylindrical reactor, equipped with

mechanical stirrer, thermocouple, suction inlet for the reacting gaseous mixture $\text{CF}_2(\text{OF})_2/\text{CO}_2$ /diluent gas, suction inlet for the reacting olefin, inert gas head outlet. The so loaded reactor is brought to -95°C . by means of a cryostat, and then a gaseous mixture of $\text{CF}_2(\text{OF})_2$ (1.0 NI/h), CO_2 (0.5 NI/h) and He (4 NI/h) is continuously introduced under mechanical stirring until a 0.05M concentration of $\text{CF}_2(\text{OF})_2$ in solution is achieved.

At this point vinylidene fluoride $\text{CH}_2=\text{CF}_2$ (2.0 NI/h) is continuously added so as to keep constant the concentration of the hypofluorite in the solution during the addition. After 2 h and 40', the addition of the gaseous mixture containing $\text{CF}_2(\text{OF})_2$ is interrupted and the addition of $\text{CH}_2=\text{CF}_2$ is continued up to the complete conversion of $\text{CF}_2(\text{OF})_2$ dissolved in the solution.

At the end of the reaction, 21 g of $\text{CF}_3\text{CH}_2\text{OCF}_2\text{OCH}_2\text{CF}_3$ (b.p. $+51^\circ\text{C}/200$ mm Hg) are isolated by fractional distillation.

The linear addition product yield, defined as ratio between the moles of $\text{CF}_3\text{CH}_2\text{OCF}_2\text{OCH}_2\text{CF}_3$ and the moles of $\text{CF}_2(\text{OF})_2$ converted, is 67%. The product has $T_g=-127.9^\circ\text{C}$. The product is further characterized by F^{19} -NMR spectroscopy and mass spectroscopy.

By F^{19} -NMR spectroscopy the product shows 2F type OCF_2O at -64.0 p.p.m. and 6F type CF_3-CH_2 at -73.9 p.p.m., related to $\text{CFCl}_3=\text{O}$.

The electronic impact mass spectroscopy analysis shows the following main peaks and relative intensities: 179 (33.1%), 163 (28.2%), 149 (100%), 113 (17.7%), 83 (83.1%).

The linear adduct yield and the experimental conditions adopted in examples 17-22 are summarized in Table 2.

TABLE 2

Formation of linear addition product between hypofluorite and olefin: addition of olefin to hypofluorite^(*)

Ex. Solvent (g)	$\text{CF}_2(\text{OF})_2$ (moles/l)	Temperature ($^\circ\text{C}$.)	Linear addition product yield (% by moles)
17 CFCl_3	0.5	-100	9
18 CF_3-CFH_2	2	-95	19.2
19 $\text{CFCl}_2-\text{CFCl}-\text{SOF}$	2	-90	26.1
20 $\text{CHCl}_2-\text{CF}_3$	1.5	-97	69
21 $\text{CH}_2\text{F}-\text{CF}_3$	0.2	-90	61
22 $\text{CH}_2\text{F}-\text{CF}_3$	0.05	-95	67

^(*)The hypofluorite is $\text{CF}_2(\text{OF})_2$; the olefin is 1,2-dichloro-difluoroethylene in the examples from 17 to 19, trichloroethylene in example 20, 1-chloro-1,2-difluoroethylene in example 21 and vinylidene fluoride in example 22.

EXAMPLE 23

Preparation of perfluorobisvinylloxymethane (dehalogenation process).

500 ml of dimethylformamide, 90 g of Zn in powder previously activated by washing with 3N HCl and 100 mg of I_2 are loaded into a 1 l three necks flask, equipped with mechanical stirrer, thermometer, dripping funnel, distillation column with water cooler and collecting trap kept at -78°C . and connected to the vacuum of a mechanical pump. The internal temperature is brought to 80°C ., the vacuum is adjusted at 160 mm Hg and 88.4 g (229 mmoles) of $\text{CF}_2\text{Cl}-\text{CFCl}-\text{OCF}_2\text{O}-\text{CFCl}-\text{CF}_2\text{Cl}$ are added in drops.

Once the addition is over, the reaction mixture is kept under the conditions indicated above for 30 minutes, and then the vacuum is gradually increased up to 0.5 mm Hg.

After about 20 minutes the collecting flask is disconnected; it contains 51.3 g of a mixture consisting for 89% by weight of perfluorobisvinylloxymethane $\text{CF}_2=\text{CF}-\text{O}-\text{CF}_2\text{O}-\text{CF}=\text{CF}_2$ (45.6 g equal to 187 mmoles) and for 11% by weight of the compound of hemidehalogenation $\text{CF}_2=\text{CF}-\text{O}-\text{CF}_2-\text{O}-\text{CFCl}-\text{CF}_2\text{Cl}$ (5.6 g equal to 18 mmoles). After purification by distillation at atmospheric pressure with a plate column, 42 g (172 mmoles) of perfluorobisvinylloxymethane having boiling point at atmospheric pressure of 50° C. are isolated. The yield is 75%.

The product was characterized by F^{19} -NMR spectroscopy (in p.p.m., related to $\text{CFCl}_3=\text{O}$: 2F type $\text{CF}_2=\text{C}-$ at -114.0/-121.1; 1F type $\text{C}=\text{CF}-\text{O}$ at -136.1; 2F type $\text{O}-\text{CF}_2-\text{O}$ at -60.4); IR spectrum (absorption bands at 1840, 1338, 1298, 1248, 1192 cm^{-1}) and mass spectrum (electronic impact): 163 ($\text{M}^+-\text{C}_2\text{F}_3$); 147 ($\text{M}^+-\text{C}_2\text{F}_3\text{O}$); 135 ($\text{C}_2\text{F}_5\text{O}$); 119 (C_2F_5).

EXAMPLE 24

Preparation of bis-(2-fluorovinyl)-methane $\text{CFH}=\text{CH}-\text{OCF}_2\text{O}-\text{CH}=\text{CHF}$ (dehalogenation process).

50 ml of dimethylformamide, 21 g of Zn in powder previously activated by washing with 3N HCl and 10 mg of I_2 are loaded into a 150 ml three necks flask, equipped with mechanical stirrer, thermometer, dripping funnel, distillation column with water cooler and collecting trap kept at -78° C. and connected to the vacuum of a mechanical pump. The internal temperature is brought to 120° C., the vacuum is adjusted at 350 mm Hg and 17.7 g (56.4 mmoles) of $\text{CFHCl}-\text{CHCl}-\text{OCF}_2\text{O}-\text{CHCl}-\text{CFHCl}$ are added in drops.

Once the addition is over, the reaction mixture is kept under the conditions indicated above for 15 hours, and then the vacuum is gradually increased up to 200 mm Hg, after about 15 minutes the collecting flask containing 18.3 g of a mixture consisting of $\text{CFH}=\text{CH}-\text{OCF}_2\text{O}-\text{CH}=\text{CHF}$ and dimethylformamide is disconnected.

After purification of the mixture by washing twice with water and by distillation at reduced pressure with plates column, 4.6 g (26.7 mmoles) of 98% pure $\text{CFH}=\text{CH}-\text{OCF}_2\text{O}-\text{CH}=\text{CHF}$ having boiling point of 42.0° C. at 200 mm Hg are isolated. The yield is 47%.

The product was characterized by F^{19} -NMR spectroscopy where it shows in p.p.m., related to $\text{CFCl}_3=\text{O}$: 2F type $\text{O}-\text{CF}_2-\text{O}$ at -62.9; -63.3; -63.7; 1F type trans $\text{F}-\text{CH}=\text{CH}-\text{O}$ at -153.5; 1F type cis $\text{F}-\text{CH}=\text{CH}-\text{O}$ at -168.5; mass spectrum with electronic impact, wherein the main peaks and the relative intensities are: 45 (73%), 78(17%), 111 (100%), 172 (1%); and IR spectrum wherein the following main peaks are observed: 3110, 1719, 1693, 1374, 1301, 1202, 1106 and 1028 cm^{-1} .

EXAMPLE 25

Preparation of perfluorobisvinylloxymethane (dehydrohalogenation process)

3 g of bisetheral addition products mixture prepared in example 21 and 100 mg of tetrabutylammonium chloride, as phase transfer catalyst, are loaded into a 100 ml three necks flask, equipped with mechanical stirrer, thermometer, dripping funnel, distillation column with water cooler and collecting trap kept at -78° C. and connected to the vacuum of a mechanical pump. The internal temperature is brought to 60° C., the vacuum is adjusted at 200 mm Hg and 3 ml of a 40% KOH water solution are added in drops.

Once the addition is over, the reaction mixture is kept under the conditions indicated above for 30', and then the vacuum is gradually increased up to 100 mm Hg. After about 30' the collecting flask is disconnected; it contains 2 g of a mixture essentially consisting of perfluorobisvinylloxymethane, unreacted or only partially reacted starting product and water traces. From the crude reaction mixture, after dehydration with solid KOH, 1.3 g of perfluorobisvinylloxymethane and 0.5 g of not dehydrohalogenated products are isolated by distillation. The yield of the perfluorobisvinylloxymethane is 66%.

The product was characterized by F^{19} -NMR spectroscopy, IR spectroscopy and mass spectroscopy as in example 23.

EXAMPLE 26

Polymerization of perfluorobisvinylloxymethane without use of solvent.

The polymerization initiator consisting of 32 microliters of 6.5% by weight perfluoropropionylperoxide in $\text{CCl}_2\text{F}-\text{CF}_2\text{Cl}$ and 6.4 mmoles of perfluorobisvinylloxymethane are loaded into a 31.2 ml glass reactor for polymerization, equipped with magnetic stirring and opening for the loading and unloading of the reactants.

The so loaded reactor is cooled to -196° C., evacuated, brought to room temperature and cooled again twice; at the end of this operations sequence the reactor is kept at 30° C. for 1 hour under magnetic stirring. The reaction crude product so obtained looks like a viscous, transparent, colourless and homogeneous solution.

After distillation of the unreacted monomer and stripping under vacuum at the temperature of 120° C. for 3 hours, 340 mg of polymer are isolated.

The infrared analysis of the obtained polymer shows the absence of absorptions in the area of the carbonyl and in the area of the fluorinated double bond.

The F^{19} -NMR analysis is in accordance with the presence of the cyclic structures reported above, structure a) being 89% and structure b) being 11%. Neither unreacted vinyls nor acylfluorides appear.

The intrinsic viscosity of the polymer measured in Fluorinert® FC 75 is 66 ml/g. A Differential Scanning Calorimetry (DSC) test carried out on the polymers does not show any melting point, therefore the polymer is amorphous. The polymer T_g determined by DSC is 77.8° C. The thermogravimetric analysis (TGA) shows a weight loss of 2% at 488° C. and of 10% at 530° C.

EXAMPLE 27

Polymerization of perfluorobisvinylloxymethane in solution.

The polymerization initiator consisting of 50 microliters of 6.5% by weight perfluoropropionylperoxide in $\text{CCl}_2\text{F}-\text{CF}_2\text{Cl}$, 16.6 mmoles of $\text{CCl}_2\text{F}-\text{CF}_2\text{Cl}$ and 6.4 mmoles of perfluorobisvinylloxymethane are loaded into a 52.3 ml glass reactor for polymerization, equipped with magnetic stirring and opening for the loading and unloading of the reactants.

The so loaded reactor is cooled to -196° C., evacuated, brought to room temperature and cooled again twice; at the end of this operations sequence the reactor is kept at 30° C. for 2 hours under magnetic stirring. The reaction crude product so obtained looks like a viscous, transparent, colourless and homogeneous solution.

After distillation of the unreacted monomer and stripping under vacuum at the temperature of 120° C. for 3 hours, 450 mg of polymer are isolated.

15

The infrared analysis of the obtained polymer shows the absence of absorptions in the area of the carbonyl and of the fluorinated double bond.

The F^{19} -NMR analysis is in accordance with the presence of the cyclic structures reported above, structure a) being 89% and structure b) being 11%. Neither unreacted vinyls nor acylfluorides appear.

The intrinsic viscosity of the polymer measured in Fluorinert® FC 75 is 56 ml/g. The solubility of this homopolymeric sample in Fluorinert® FC 75 and in perfluoropolyethers Fomblin® and Galden® is higher than 20% by weight at room temperature. The 20% by weight solutions in these solvents are very viscous, but transparent and visually homogeneous; solutions even with concentration higher than 20% by weight are extremely viscous, but substantially homogeneous. The T_g of the polymer determined by DSC and the TGA are substantially the same as the T_g and the TGA of the polymer obtained in example 23.

EXAMPLE 28

Copolymer of perfluorobisvinylloxymethane with vinylidene fluoride.

3 ml of CCl_2F-CF_2Cl , the polymerization initiator consisting of 32 microliters of 6.5% by weight perfluoropropionylperoxide dissolved in CCl_2F-CF_2Cl , 2.20 mmoles of perfluorobisvinylloxymethane and 7.8 mmoles of vinylidene fluoride are loaded into a 52 ml glass reactor for polymerization, equipped with magnetic stirring and opening for the loading and unloading of the reactants.

The so loaded reactor is cooled to $-196^\circ C.$, evacuated, brought to room temperature and cooled again twice; at the end of this operation sequence the reactor is kept at $30^\circ C.$ for 16 hours under magnetic stirring. The reactor is brought to the temperature of the liquid nitrogen and connected to a vacuum system kept at the pressure of 10^{-3} mbar, it is then let to reach room temperature fractionating the vapours by traps cooled at $-80^\circ C.$, $-120^\circ C.$, and $-196^\circ C.$

The trap at $-80^\circ C.$ contains only the $CFCl_2-CF_2Cl$ used as solvent. The trap at $-120^\circ C.$ contains 6.90 mmoles of $CFCl_2-CF_2Cl$ and 0.26 mmoles of unreacted perfluorobisvinylloxymethane. The trap at $-196^\circ C.$ contains 4.5 mmoles of unreacted vinylidene fluoride.

After distillation of the solvent and of the unreacted monomers and stripping of the polymer under vacuum at the temperature of $120^\circ C.$ for 3 hours, 670 mg of polymer are isolated.

The weight balance determined by gaschromatography analysis of the content of the traps containing the unreacted monomers allows to calculate the amount of perfluorobisvinylloxymethane in the polymer, which results to be 37% by moles. The polymer results soluble in N,N-dimethylformamide.

The F^{19} -NMR analysis is in accordance with the presence of the cyclic structures reported above, structure a) being 91.4% and structure b) being 8.6%. Neither unreacted vinyls nor acylfluorides appear.

The polymer T_g determined by DSC is $17.2^\circ C.$ The DSC graph does not show any melting endotherm, therefore the polymer is amorphous. The TGA shows a weight loss of 2% at $412^\circ C.$ and of 10% at $450^\circ C.$

EXAMPLE 29

Copolymer of tetrafluoroethylene with perfluorobisvinylloxymethane.

8 ml of CCl_2F-CF_2Cl , the polymerization initiator consisting of 1.5 ml of 0.35% by weight perfluoropropionylp-

16

eroxide in CCl_2F-CF_2Cl , 0.33 mmoles of perfluorobisvinylloxymethane and 10 mmoles of tetrafluoroethylene are loaded into a 42 ml glass reactor for polymerization, equipped with magnetic stirring and opening for the loading and unloading of the reactants.

The so loaded reactor is cooled to $-196^\circ C.$, evacuated, brought to room temperature and cooled again twice; at the end of this operation sequence the reactor is kept at $40^\circ C.$ for 8 hours under magnetic stirring. The reaction crude product looks like a gelatinous mass. The reactor is brought to the temperature of the liquid nitrogen and connected to a vacuum system kept at the pressure of 10^{-3} mbar, it is then let to reach room temperature fractionating the vapours by traps cooled at $-80^\circ C.$, $-120^\circ C.$, and $-196^\circ C.$

The trap at $-80^\circ C.$ contains only the $CFCl_2-CF_2Cl$ used as solvent. The trap at $-120^\circ C.$ contains 4.90 mmoles of $CFCl_2-CF_2Cl$ and 0.16 mmoles of unreacted perfluorobisvinylloxymethane. The trap at $-196^\circ C.$ contains 0.12 mmoles of unreacted tetrafluoroethylene.

After distillation of the solvent and of the unreacted monomers and stripping of the polymer under vacuum at the temperature of $120^\circ C.$ for 3 hours, 1.030 g of polymer are isolated.

The weight balance made by gaschromatography analysis of the content of the traps containing the unreacted monomers allows to calculate the amount of perfluorobisvinylloxymethane in the polymer, which results to be 1.7% by moles.

The infrared analysis of the obtained polymer shows the absence of absorptions in the area of the carbonyl and of the fluorinated double bond.

The ΔH and the second melting point determined by DSC are respectively 9.8 cal/g and $314.5^\circ C.$

The TGA shows a weight loss of 2% at $502^\circ C.$ and of 10% at $540^\circ C.$ The polymer is hot moulded, at the temperature of $330^\circ C.$ and at the pressure of 122 atm in a transparent and tough film.

EXAMPLE 30

Copolymer of perfluorobisvinylloxymethane with perfluorodioxole.

20 microliters of 6.5% by weight perfluoropropionylperoxide dissolved in CCl_2F-CF_2Cl , 2.95 mmoles of perfluorobisvinylloxymethane and 0.6 mmoles of perfluoro-1,3-dioxole are loaded into a 42 ml glass reactor for polymerization, equipped with magnetic stirring and opening for the loading and unloading of the reactants.

The so loaded reactor is cooled to $-196^\circ C.$, evacuated, brought to room temperature and cooled again twice; at the end of this operation sequence the reactor is kept at $40^\circ C.$ for 8 hours under magnetic stirring. The reactor is brought to the temperature of the liquid nitrogen and connected to a vacuum system kept at the pressure of 10^{-3} mbar, it is then let to reach room temperature fractionating the vapours by traps cooled at $-90^\circ C.$, $-120^\circ C.$, and $-196^\circ C.$

The trap at $-90^\circ C.$ contains 1.92 mmoles of perfluorobisvinylloxymethane and $CFCl_2-CF_2Cl$ used as solvent. The trap at $-120^\circ C.$ contains 0.22 mmoles of perfluorodioxole and 0.01 mmoles of perfluorobisvinylloxymethane. The trap at $-196^\circ C.$ contains 0.1 mmoles of perfluorodioxole.

After distillation of the solvent and of the unreacted monomers and stripping of the polymer under vacuum at the temperature of $120^\circ C.$ for 2 hours, 280 mg of polymer are isolated.

The weight balance determined by gas chromatography analysis of the content of the traps containing the unreacted

monomers allows to calculate the amount of perfluorobisvinylmethane in the polymer, which results to be 78% by moles.

The polymer T_g determined by DSC is 94.5° C. The DSC graph does not show any melting endotherm, therefore the polymer is amorphous. The TGA shows a weight loss of 10% at 494° C.

EXAMPLE 31

Copolymer of bis(2-fluorovinyl)oxy-difluoromethane with vinylidene fluoride.

1 ml of 6.5% by weight perfluoropropionylperoxide dissolved in $\text{CCl}_2\text{F}-\text{CF}_2\text{Cl}$, 10 ml of $\text{CCl}_2\text{F}-\text{CF}_2\text{Cl}$ and 1.5 g of bis(2-fluorovinyl)oxy-difluoromethane (m.w.=172) are loaded into a 40 ml AISI-316 reactor for polymerization, equipped with magnetic stirring and opening for the loading and unloading of the reactants.

The so loaded reactor is cooled to -196° C., evacuated, brought to room temperature and cooled again twice; at the end of this operation sequence the reactor is kept at 30° C. and pressurized with vinylidene fluoride up to 20 atm. When the total pressure drops to 18 atm the reaction is blocked and gaseous reagents are removed at atmospheric pressure.

After distillation of the solvent and of the unreacted monomers and stripping of the polymer under vacuum at the temperature of 80° C. for 2 hours, 310 mg of polymer are isolated.

The polymer T_g determined by DSC is 39.9° C. The DSC graph does not show any melting endotherm, therefore the polymer is amorphous. The TGA shows a weight loss of 6% at 300° C.

We claim:

1. Fluorinated derivatives of bisvinylmethane of structure $\text{CFX}^1=\text{CX}^2-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CX}^2=\text{CX}^1\text{F}$, wherein X^1 and X^2 , equal to or different from each other, are F, Cl or H; X^3 and X^4 , equal to or different from each other, are F or CF_3 ; X^1 and X^2 , equal to or different from each other, are X^1 or X^2 , with the proviso that when X^1 is different from X^2 then X^1 is different from X^2 .

2. Fluorinated derivatives of bisvinylmethane of structure $\text{CFX}^1=\text{CX}^1-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CX}^1=\text{CX}^1\text{F}$ wherein X^1 is F, Cl or H; X^3 and X^4 , equal to or different from each other, are F or CF_3 .

3. Fluorinated derivatives of bisvinylmethane of structure $\text{CF}_2=\text{CF}-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CF}=\text{CF}_2$, wherein X^3 and X^4 , equal to or different from each other, are F or CF_3 .

4. Perfluorobisvinylmethane $\text{CF}_2=\text{CF}-\text{O}-\text{CF}_2-\text{O}-\text{CF}=\text{CF}_2$.

5. Process for the preparation of bisvinylmethane derivatives having the structure $\text{CFX}^1=\text{CX}^2-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CX}^2=\text{CX}^1\text{F}$, wherein X^1 and X^2 , equal to or different from each other, are F, Cl or H; X^3 and X^4 , equal to or different from each other, are F or CF_3 ; X^1 and X^2 , equal to or different from each other, are X^1 or X^2 , with the proviso that if X^1 is different from X^2 also X^1 is to be different from X^2 , comprising:

- addition of an olefin of formula $\text{CX}^1\text{X}^5=\text{CX}^2\text{X}^6$, wherein X^1 , X^2 , X^5 and X^6 equal to or different from each other are F, Cl, H or Br, the Br atoms being 2 at most and in such a case being bound to different carbon atoms, X^1 and X^5 not being both F, X^2 and X^6 not being both F, X^1 and X^2 being F only if X^5 and X^6 are different from F and not being both H, in a reactor where an hypofluorite of general formula $\text{CX}^3\text{X}^4(\text{OF})_2$ is essentially always present, where X^3 and X^4 equal to or different from each other, are F or CF_3 , dissolved in an inert solvent and having a concentration between 0.001M and 10M, at a temperature from -140° C. to +60° C.;

- separation of the reaction product of two olefin molecules and one hypofluorite molecule from the reaction mixture obtained in i) by fractional distillation

- dehalogenation or dehydrohalogenation of the product obtained in ii), wherein the eliminated halogen atoms are Cl or Br.

6. Process for preparing bisvinylmethane derivatives having the structure $\text{CFX}^1=\text{CX}^1-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CX}^1=\text{CX}^1\text{F}$, wherein X^1 is F, Cl or H; X^3 and X^4 , equal to or different from each other, are F or CF_3 , which comprises:

- addition of an olefin of formula $\text{CX}^1\text{X}^5=\text{CX}^1\text{X}^6$, wherein X^1 , X^5 and X^6 equal to or different from each other, are F, Cl, Br or H, X^1 and X^5 not being both F, X^1 and X^6 not being both F, X^1 being Br only if X^5 and X^6 equal to each other are different from Br, X^1 being F only if X^5 and X^6 are different from F and not both H, to an hypofluorite of general formula $\text{CX}^3\text{X}^4(\text{OF})_2$ wherein X^3 and X^4 equal to or different from each other, are F or CF_3 , dissolved in an inert solvent having a concentration between 0.001M and 10M, at a temperature from -140° C. to +60° C.;

- separation of the reaction product of two olefin molecules and one hypofluorite molecule from the reaction mixture obtained in i) by fractional distillation

- dehalogenation or dehydrohalogenation of the product obtained in ii), wherein the eliminated halogen atoms are Cl or Br.

7. Process for preparing bisvinylmethane derivatives having the structure $\text{CF}_2=\text{CF}-\text{O}-\text{CX}^3\text{X}^4-\text{O}-\text{CF}=\text{CF}_2$, wherein X^3 and X^4 , equal to or different from each other, are F or CF_3 , which comprises:

- addition of an olefin of formula $\text{CFX}^5=\text{CFX}^6$, wherein X^5 is Cl, Br or H, and X^6 is Cl or Br, to an hypofluorite of general formula $\text{CX}^3\text{X}^4(\text{OF})_2$ wherein X^3 and X^4 , equal to or different from each other, are F or CF_3 , dissolved in an inert solvent having a concentration between 0.001M and 10M, at a temperature from -140° C. to +60° C.;

- separation of the reaction product of two olefin molecules and one hypofluorite molecule from the reaction mixture obtained in i) by fractional distillation

- dehalogenation or dehydrohalogenation of the product obtained in ii), wherein the eliminated halogen atoms are Cl or Br.

8. Process according to anyone of the claims 5, 6 or 7 wherein the solution of the hypofluorite has a concentration from 0.1M to 4M.

9. Process according to anyone of the claims 5, 6 or 7 wherein the reaction temperature is from -120° C. to 0° C.

10. Process according to anyone of the claims 5, 6, 7, 8 or 9, characterized in that the solution of hypofluorite is prepared using an inert solvent of polar type.

11. Process according to claim 10 characterized in that the solvent is selected from hydrogenfluorocarbons, hydrogenchlorocarbons, fluorochlorocarbons, hydrogenfluorocarbons, trifluoroacetic acid, trifluoroacetic anhydride, acetic nitrile, hydrofluoric acid, sulphur dioxide, trifluoromethanesulphonic acid, $\text{CF}_2\text{Cl}-\text{CFCl}-\text{SO}_2\text{F}$, mixtures thereof and mixtures of one or more of the same with a solvent having low polarity.

12. Process according to claim 5 wherein the olefin is selected from: $\text{CFCl}=\text{CFCl}$, $\text{CHCl}=\text{CHCl}$, $\text{CHCl}=\text{CCl}_2$, $\text{CCl}_2=\text{CCl}_2$, $\text{CH}_2=\text{CF}_2$, $\text{CF}_2=\text{CF}_2$, $\text{CFH}=\text{CFCl}$, $\text{CFCl}=\text{CHCl}$, $\text{CH}_2=\text{CCl}_2$, $\text{CH}_2=\text{CFCl}$.

13. Fluorinated derivatives of bisethoxymethane obtained as intermediates after the steps i) and ii) of the processes of claims 5.

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