



US009950978B2

(12) **United States Patent**
Martin(10) **Patent No.:** **US 9,950,978 B2**
(45) **Date of Patent:** **Apr. 24, 2018**(54) **METHOD OF FORMING ALKOXYLATED FLUOROALCOHOLS**(71) Applicant: **Tyco Fire & Security GmbH**,
Neuhausen am Rheinfall (CH)
(72) Inventor: **Thomas Joseph Martin**, Mansfield, TX
(US)
(73) Assignee: **TYCO FIRE & SECURITY GMBH**
(CH)

(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 0 days.

(21) Appl. No.: **14/613,222**(22) Filed: **Feb. 3, 2015**(65) **Prior Publication Data**

US 2015/0148565 A1 May 28, 2015

Related U.S. Application Data

(63) Continuation of application No. 12/325,329, filed on Dec. 1, 2008, now Pat. No. 8,946,486.

(60) Provisional application No. 60/991,982, filed on Dec. 3, 2007.

(51) **Int. Cl.****C07C 41/03** (2006.01)
C07C 231/12 (2006.01)
C07C 303/40 (2006.01)(52) **U.S. Cl.**CPC **C07C 41/03** (2013.01); **C07C 231/12** (2013.01); **C07C 303/40** (2013.01)(58) **Field of Classification Search**CPC **C07C 41/03**
See application file for complete search history.(56) **References Cited****U.S. PATENT DOCUMENTS**2,600,378 A 6/1952 Dickey et al.
2,723,999 A 11/1955 Cowan et al.
3,359,331 A 12/1967 Baker et al.
3,676,477 A 7/1972 Chay et al.
3,944,586 A 3/1976 Knowles
3,944,653 A 3/1976 Stewart et al.
3,948,668 A 4/1976 Hayek et al.
3,980,715 A 9/1976 Szur
4,169,104 A 9/1979 Burt et al.
4,240,956 A 12/1980 Weil et al.
4,287,118 A 9/1981 Muldrow, Jr.
4,408,043 A 10/1983 Seale et al.
4,456,697 A 6/1984 Yang
4,483,941 A 11/1984 Yang
4,490,561 A 12/1984 Yang et al.
4,594,200 A 6/1986 Penny
4,983,778 A 1/1991 Ploog
4,992,211 A 2/1991 Casciani
5,002,678 A 3/1991 Vanover et al.
5,026,923 A 6/1991 Kemp
5,041,576 A 8/1991 Wasfi
5,057,628 A 10/1991 Edwards et al.
5,210,323 A 5/1993 Wimmer et al.5,290,912 A 3/1994 Watabe et al.
5,567,857 A 10/1996 Huang et al.
5,608,116 A * 3/1997 Halling B01F 17/0035
568/614
5,852,148 A 12/1998 Behr et al.
5,910,599 A 6/1999 Tanaka et al.
6,028,230 A 2/2000 Le-Khac et al.
6,048,952 A 4/2000 Behr et al.
6,201,101 B1 3/2001 Tozzola et al.
6,340,779 B1 1/2002 Enna et al.
6,365,541 B1 4/2002 Wimmer et al.
6,365,769 B1 4/2002 Behr et al.
6,395,083 B2 5/2002 Enna et al.
6,455,459 B1 9/2002 Bedard et al.
6,734,327 B2 5/2004 Bedard et al.
8,058,480 B2 11/2011 Moloy
8,058,481 B2 * 11/2011 Moloy C07C 41/03
560/239
8,067,329 B2 * 11/2011 Moloy B01J 31/0239
502/164
8,946,486 B2 * 2/2015 Martin C07C 41/03
568/615
2004/0266981 A1 12/2004 Yokoyama et al.
2006/0069220 A1 * 3/2006 Meurs C07C 41/03
526/160
2009/0143621 A1 * 6/2009 Martin C07C 41/03
564/96**FOREIGN PATENT DOCUMENTS**DE 111522 A 2/1975
GB 1 410 328 10/1975
JP 2005-46749 2/2005

(Continued)

OTHER PUBLICATIONS

Briggs (Kirk-Othmer Encyclopedia of Chemical Technology, vol. 4, Jul. 2001, pp. 241-294).*

"Ethoxylation of PFPE," experimental data, submitted with Opposition filed in European Application No. 08 856 599.9, 1 page (submitted to EPO on Jan. 5, 2015).

Docks, "Boric Acid Esters," *Kirk-Othmer Encyclopedia of Chemical Technology*, John Wiley & Sons, Inc., 10 pages (Dec. 4, 2000).
DuPont™ Zonyl® FSH Fluorosurfactant, Product Sheet, 2 pages (2001).

DuPont™ Zonyl® FSN-100 Fluorosurfactant, Product Sheet, 2 pages (2005).

DuPont™ Zonyl® FSO Fluorosurfactant, Product Sheet, 2 pages (2005).

Extended European Search Report, European Application No. 08 856 599.9, 5 pages (dated Oct. 7, 2011).

Golden et al., "Disproportionation of Alkoxyborohydrides: A ¹¹B NMR Study of the Reaction between Sodium Borohydride and Fluorinated Alcohols and Phenols. The Preparation of Tris(fluoroalkoxy)- and Tris(fluorophenoxy)borohydrides," *Inorganic Chemistry*, 31:1533-1535 (1992).

(Continued)

Primary Examiner — Rosalynd A Keys(74) *Attorney, Agent, or Firm* — Perkins Coie LLP(57) **ABSTRACT**

A method of forming an alkoxyated fluoroalcohol is accomplished by providing a boron compound having or providing at least one boron-oxygen bond and an iodine source. The boron compound and iodine source are combined with reactants of a fluoroalcohol and an alkylene oxide in the presence of a base. The reactants are allowed to react to form an alkoxyated fluoroalcohol reaction product.

9 Claims, No Drawings

(56)

References Cited

FOREIGN PATENT DOCUMENTS

SU	570590 A	9/1977
WO	WO 95/35272 A1	12/1995
WO	WO 96/28407 A2	9/1996
WO	WO 98/19743 A1	5/1998
WO	WO 98/23553 A1	6/1998
WO	WO 03/044074 A1	5/2003
WO	WO 2006/034997 A1	4/2006

OTHER PUBLICATIONS

International Search Report and Written Opinion, International Application No. PCT/US08/85217 (published under WO 2009/073641), 6 pages (dated Feb. 4, 2009).

Landesman et al., "Fluoroalkyl Borate Esters," *Inorganic Chemistry*, 3(6):896-898 (Jun. 1964).

Opposition Against European Patent 2215042 B1 (08856599.9)—Statement of Grounds, Facts and Arguments, European Application No. 08 856 599.9, 25 pages (Jan. 5, 2015).

Summons to Oral Proceedings, European Application No. 08 856 599.9, 18 pages (Jun. 9, 2016).

U.S. Appl. No. 60/991,982, filed Dec. 3, 2007, Parent U.S. Appl. No. 14/613,222.

Young et al., "Perhaloalkyl Hypochlorites. V. Perfluoroalkyl Borate Esters from Reactions with Boron Trichloride," *Inorganic Chemistry*, 10(12):2810-2812 (1971).

* cited by examiner

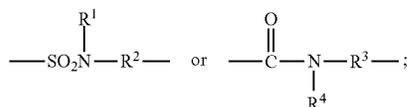
3

illustrating various embodiments of the invention and should not be construed as a limitation to the scope and applicability of the invention. While the methods and compositions of the present invention are described herein as utilizing or comprising certain materials, it should be understood that the method or composition could optionally comprise two or more chemically different materials. In addition, the composition can also comprise some components other than the ones cited. In the summary and detailed description, each numerical value should be read once as modified by the term "about" (unless already expressly so modified), and then read again as not so modified unless otherwise indicated in context. Also, in the summary and description, it should be understood that a concentration, value or amount listed or described as being useful, suitable, or the like, is intended that any and every concentration, value or amount within the range, including the end points, is to be considered as having been stated. For example, "a range of from 1 to 10" is to be read as indicating each and every possible number along the continuum between about 1 and about 10. Thus, even if specific data points within the range, or even no data points within the range, are explicitly identified or refer to only a few specific, it is to be understood that the inventor appreciates and understands that any and all data points within the range are to be considered to have been specified, and that the inventor is in possession of the entire range and all points within the range.

In forming alkoxyated fluoroalcohols in accordance with the invention, the fluoroalcohol may be a fluoroalcohol having the generalized structure according to Equations (1) or (2) below:



where $\text{F}(\text{CF}_2)_m$ denotes a perfluorinated alkyl group, where m is usually, but not necessarily, 2-20, and more typically 4-14. In certain embodiments m may be from 2 to 8, and in still other embodiments m may be from 2 to 6. The alkyl group $\text{F}(\text{CF}_2)_m$ is typically linear but may be branched, as well. The linking group A may be a connecting group of a $\text{—}(\text{CH}_2)_n\text{—}$, which may be branched or linear,



where n is from 1 to 6 or more, R^1 and R^4 are each independently selected from one of hydrogen, a halogen or an alkyl group containing from 1-30 carbon atoms, and R^2 and R^3 are each independently selected from an alkylene group containing from 2 to 30 carbon atoms. Typically, the linking group A is methylene or ethylene. In certain embodiments, the fluoroalcohol may include one or more hydrogens substituted for fluorine, such as the group $\text{H}(\text{CF}_2)_m$ for example. The fluoroalcohols may have molecular weights of from about 200 to about 2000 g/mol, more typically from about 300 to about 500 g/mol. Non-limiting examples of suitable fluoroalcohols are those described in U.S. Pat. Nos. 4,490,561; 5,590,561 and 5,608,116, which are each incorporated herein by reference in their entireties for all purposes.

The alkoxyating reaction is carried out using a catalyst system that includes a boron compound. Most alkoxyating

4

agents are not appreciably soluble in the fluoroalcohols. The boron compound may complex with the alkoxyating agent facilitating incorporation and reaction of the alkoxyating with the fluoroalcohol. The catalyst or catalytic properties of the boron compound may be formed or provided in-situ, with the same or similar functionality as borohydride or boron trifluoride catalysts. As used herein, "in-situ" refers to the formation of the catalyst in the presence of the fluoroalcohol or at least one of the reactants. The boron compound is one that has or provides at least one boron-oxygen bond. In particular, the boron compound may have or provide one (1) to four (4) boron-oxygen bonds per boron atom. To produce a boron-based catalyst, the boron compound may be added to the fluoroalcohol or a stand-in alcohol to provide or create a catalyst species capable of carrying out the alkoxyating reactions. The boron compound may have, but is not limited to, the general structure shown in Equation (3) below:



where R^5 and R^6 are each independently H, an alkyl group (e.g. CH_3 , CH_3CH_2 , etc.), an aryl, an alkylaryl, an alkyl halide, a halogen (e.g. F, Cl, Br, I), or —OR^8 , and wherein R^7 and R^8 are each independently H, an alkyl group, a carbonyl, a carboxyl, an aryl or an alkylaryl. While the boron compound of Equation 3 is shown as having three substituents, the boron compound also has an empty orbital that can complex with a fourth molecule, which is typically neutral or uncharged. This fourth molecule may include an alcohol, such as methanol, ethanol, etc., or an ether, such as diethyl ether, tetrahydrofuran, etc.

The boron compound may be an oxide of boron, an alkyl borate, a boric acid, a boric acid anhydride, a boronic acid, a borinic acid or salts or esters of said acids or combinations of these. Examples of the boron compound may include boric acid (H_3BO_3), meta boric acid (HBO_2), tetra boric acid ($\text{H}_2\text{B}_4\text{O}_7$), boron oxide (B_2O_3), trimethyl borate, triethyl borate, triisopropyl borate, tripropyl borate, tributyl borate, and tetraborate salts, such as sodium tetraborate (borax), potassium tetraborate, etc. Generally, any oxide, acid, salt or ester of boron may be useful in the present invention. Further examples of suitable boron compounds are described in U.S. Patent Application Publication No. 2006/0069220A1, which is herein incorporated by reference for all purposes. Examples of boric acid esters are described in Boric Acid Esters, Kirk-Othmer Encyclopedia of Chemical Technology, pp. 1-10, John Wiley & Sons, Inc., which is herein incorporated by reference. In certain instances, the boron compound may provide the boron-oxygen bond in-situ.

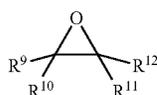
The boron compound may be used in an amount of from about 0.01 mole % to about 20 mole % by total moles of fluoroalcohol, more particularly from about 0.05 mole % to about 5 mole % by total moles of fluoroalcohol.

An iodine co-catalyst is also used in the catalyst system. The iodine co-catalyst is provided from an iodine source, which may be an alkali metal iodide, alkaline earth metal iodide or elemental iodine and combinations of these. Generally, all iodine salts may be useful as the iodine source. Examples of suitable iodine sources include lithium iodide, sodium iodide, potassium iodide, calcium iodide, ammonium iodide, elemental iodine or combinations of these. The iodine source is used in an amount of from about 0.05% to

5

about 10% by weight of the fluoroalcohol, more particularly from about 0.1% to about 2% by weight of the fluoroalcohol. If the iodine source is other than sodium iodide (NaI) or elemental iodine (I₂), higher amounts of the iodine source may be used. In certain instances, approximately twice as much of the other iodine sources may be used. Thus, for example, if sodium iodide or elemental iodine are typically used in an amount of from about 0.05% to about 5% by weight of the fluoroalcohol, other iodine sources may be used in amounts of from about 0.1% to 10% by weight of the fluoroalcohol for the alkoxylation of the fluoroalcohol.

An alkoxyating agent of an alkylene oxide used for the fluoroalcohol alkoxylation may have the general structure of Equation (4) below:



where R⁹, R¹⁰, R¹¹ and R¹² are each independently one of hydrogen, an alkyl group, an alkyl alcohol, an alkyl halide, an aryl, an alkylaryl or an allyl ether. Particularly useful are those alkylene oxides of ethylene oxide, propylene oxide, butylene oxide, glycidol, epichlorohydrin, styrene oxide, oxetane, tetrahydrofuran, 1,4-dioxane, allyl glycidyl ether, fluorinated cyclic ethers (e.g., fluorinated ethylene oxide, fluorinated propylene oxide, etc.) or mixtures of these. Ethylene oxide and propylene oxide are particularly useful in carrying out the alkoxylation reactions. Examples of suitable alkylene oxides for use in the alkoxylation reactions are described in U.S. Patent Application Publication No. 2006/0069220A1 and U.S. Pat. No. 5,608,116, which are each incorporated herein by reference.

The alkoxyating agent may be used in an amount of from about 1 mole % to about 100 mole % by total moles of the fluoroalcohol, more typically from about 2 mole % to about 20 mole %, and still more typically from about 3 mole % to about 12 mole % by total moles of the fluoroalcohol. The alkoxyating agent may be used in stoichiometric amounts to provide the desired degree of alkoxylation.

An alkali or base is also used in the reaction. While not wanting to be bound by any particular theory, it is believed that the base deprotonates the fluoroalcohol to allow it to react with the boron compound to form a borate ester in situ. Examples of suitable bases include metal hydroxides, such as sodium hydroxide (NaOH), sodium hydride (NaH), and those having the general structure R—O—Na, where R is an alkyl, such as sodium methylate, sodium ethoxide, etc. Other bases may include KOH, Ca(OH)₂, Mg(OH)₂, etc., however, NaOH has been shown to work particularly well in testing. The base may be used in an amount of 0.05% to about 5% by weight of the fluoroalcohol, more particularly from about 1% to about 5% by weight of the fluoroalcohol.

In certain instances the reaction can be carried out under an inert atmosphere (nitrogen, argon, etc.) or in air. The reaction can also be carried out under a reduced pressure or under vacuum or under reactant pressure (e.g., ethylene oxide vapor pressure).

Although not necessary, solvents may be combined with the fluoroalcohol in certain instances, such as for temperature control and to facilitate solubility of the catalyst system, reactants or reaction products. Examples of suitable solvents include, polar and non-polar aprotic solvents such as dimethyl sulfoxide, N-methyl pyrrolidone, N,N-dimethyl for-

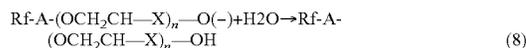
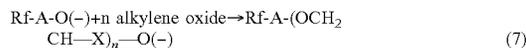
6

mamide, glyme, diglyme, etc., acetone, acetonitrile, hydrocarbons, glycol ether esters, esters, fluorocarbon solvents, etc. Water is generally excluded from alkoxylation reactions by a drying process due to its initiation of unwanted, homo-polymerization.

To carry out the alkoxyating reaction, the iodine source and any base may be initially added to the fluoroalcohol. When added to the fluoroalcohol, NaOH and other bases may form water. It may be desirable to remove such water from the fluoroalcohol prior to adding the boron compound. Water may compete against the fluoroalcohol and react with the boron compound to produce unwanted side products or sub-optimal catalytic species, or hamper the desired in situ catalyst formation. It may be desirable to therefore add the NaOH or other base prior to the salts and other components for this reason as well. The solution may then be heated to remove any water to dry the fluoroalcohol. The iodine source may then be added followed by addition of the boron compound. In certain instances, such as where boric acid is used in combination with NaOH or another base, water may also be formed from the reaction of boric acid and base. This water may also be removed through heating of the solution. After any drying, the alkoxyating agent may be added. This may be added gradually as the reaction is carried out. The reaction temperatures may be kept at from about ambient to 200° C., with typical reaction temperatures being from about 90° C. to about 150° C. Mixing and heating, if any, may be continued during the reaction.

In certain embodiments, the use of an alcohol that is a non-fluoroalcohol may be used. In such instances, this "stand-in" alcohol is used during catalyst synthesis using the same procedures or steps as with the fluoroalcohol described previously. The fluoroalcohol is then added to the mixture, with the fluoroalcohol displacing the non-fluoroalcohol, such as through evaporation or transesterification. The alcohols are typically low molecular weight alcohols, that have low boiling points so that they can be readily removed, such as through heating and evaporation. When removed in this manner, the volatile alcohols carry away residual water, further drying the reaction mixture. Examples of suitable high volatility alcohols are methanol, ethanol, n-propanol, isopropyl alcohol, butanol, etc. The stand-in alcohol may be used in an amount of from greater than 0 to about 5 mole % or more based on boron. Typically, the stand-in alcohol may be used in an amount of from about 3 mole % to about 4 mole %.

The alkoxyating reaction can be illustrated by the reaction shown in Equations (5) to (8) below:



where Rf is a perfluoroalkyl, such as F(CF₂)_m, M is the boron catalyst species, which may or may not carry a formal charge, A is the linking group described from Equation 2, and X may be H or an alkyl group (e.g., —CH₃), an alkyl alcohol, an alkyl halide, an aryl, an alkylaryl or an allyl ether, depending on the alkylene oxide used for alkoxylation, and n is an integer, which may range from about 1 to 100 or more. With Equations (5)-(9), certain exchange reactions, for example of a proton or catalyst species, are omitted, but are recognized to occur between various alco-

hol or alcohol ethoxylate molecules. It should be noted that the linking group A may also be absent from the above equations such as when fluoroalcohols having the structure of Equation 1 are used. The alkoxyated fluoroalcohol may then have the general formula of Equation (9) below:



Such alkoxyated fluoroalcohols formed from the above reactions are non-ionic alkoxyated fluoroalcohol products. The alkoxyated fluoroalcohols may also be subsequently derivatized to make anionic fluoroalcohols, such as through the use of, for example, chlorosulfonic acid and phosphorous pentoxide to make ether sulfates and phosphates, respectively, or other products, which may be anionic.

Further processing may be also conducted on the alkoxyated fluoroalcohols, such as catalyst removal, such as by filtration, distillation and other separation techniques. Dilution of the reaction product with solvents, such as water, alcohols, glycols, glycol ethers, etc., and mixtures of the same, may also be used if desired for final products.

The alkoxyated fluoroalcohols prepared in accordance with the invention are useful in formulating paints and coatings, inks, waxes, polishes, anti-corrosion coatings, leveling agents, wetting agents, oil field chemicals, etc.

The method of invention avoids the use of borohydride and boron trifluoride catalysts, while providing the generally the same degree or better reactivity. The boron compounds and other components used in the method of the invention are readily available and generally easily handled and safe to transport and store. Furthermore, resulting alcohols from borate ester transesterification that are formed during the method of the invention may be readily driven off. This is beneficial in that the evaporating alcohols entrain water, which can react with the alkoxyating agent (e.g. ethylene oxide) to produce polyalkylene oxide homopolymers. Such polymers provide no useful functionality and give a yield loss and stoichiometric imbalance for the active product. Furthermore, except where NaH is used as the base, no hydrogen gas is evolved, which reduces flammability.

The following examples serve to further illustrate the invention.

EXAMPLES

Example 1

A 139.3 g quantity of a fluoroalcohol having an average molecular weight of about 445 g/mol and the general formula, $\text{F}(\text{CF}_2)_m-\text{CH}_2\text{CH}_2-\text{OH}$, was charged to a 4-neck round bottom flask used as the reactor equipped with a mechanical stirrer, septa and a dry ice condenser. To the reactor was added 0.5 g NaOH, which was then heated to 120° C. under a slow nitrogen purge to dissolve the NaOH and dry the fluoroalcohol. Then, 0.6 g iodine and 0.7 g sodium iodide were added as the reactor cooled. At about 50° C., 1.48 ml of trimethyl borate was added, and the reactor was again heated slightly, being held at 60° C. (max.) for 45 min, and then heated again to the reaction temperature of between about 140 and 155° C. Ethylene oxide as added over the course of several hours, with a total of 123.5 g being reacted. The product readily dissolved in water to give a clear, transparent solution and had a surface tension of 22.7 dynes/cm (0.0227 N/m) at 0.01% w/w. This is contrasted with the starting fluoroalcohol, which had no solubility in water.

Example 2

A 141.3 g quantity of a fluoroalcohol having no solubility in water with the same general formula as that of Example

1 and having an average molecular weight of about 364 g/mol was charged to a reactor as described in Example 1. To the reactor was added 0.7 g NaOH, which was then heated to 125° C. under a nitrogen purge to dissolve the NaOH and dry the fluoroalcohol. Then, 0.7 g iodine and 0.8 g sodium iodide were added as the reactor cooled. At about 50° C., 3.9 ml of triisopropyl borate was added, and the reactor was again heated to 100° C. for 25 min, and then further heated to the reaction temperature of about 140° C. Ethylene oxide was added over the course of several hours between about 130 and 150° C., with a total of 148.7 g being reacted. The final product dissolved in DI water to give a clear, transparent, slightly yellow solution at 1% w/w with a cloud point of 78° C. Upon further dilution with DI water to 0.01% w/w, the solution exhibited a surface tension of 24.6 dynes/cm (0.0246 N/m).

Example 3

The procedures of Example 2 were repeated except that boric acid was used in place of triisopropyl borate. After charging 141.4 g of fluoroalcohol to the reactor, 0.8 g NaOH and 2.6 g boric acid were also added and heated to 125° C. under a nitrogen purge to dissolve the solids and dry the fluoroalcohol. Then, 0.7 g iodine and 0.8 g sodium iodide were added as the reactor cooled. The reactor was heated again to the reaction temperature of about 140° C. Ethylene oxide was added over the course of several hours between about 130 and 150° C., with a total of 55.6 g being consumed. The final product dispersed in DI water to give a hazy, slightly yellow dispersion at 1% w/w. Upon further dilution with DI water to 0.01% w/w, the solution exhibited a surface tension of 17.8 dynes/cm (0.0178 N/m).

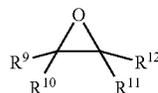
While the invention has been shown in some of its forms, it should be apparent to those skilled in the art that it is not so limited, but is susceptible to various changes and modifications without departing from the scope of the invention. Accordingly, it is appropriate that the appended claims should be construed broadly and in a manner to encompass such changes and modifications consistent with the scope of the invention.

I claim:

1. A process of forming alkoxyated fluoroalcohols, wherein the process comprises the steps of:

reacting

- (A) a boron and iodine based catalyst system, wherein boron is provided by at least one boron compound selected from the group consisting of an oxide of boron, an alkyl borate, a boric acid, a boronic acid anhydride, a boronic acid, salts of said acids, esters of said acids, and combinations thereof, and iodine is provided by at least one iodine source selected from the group consisting of an alkali metal iodide, alkaline earth metal iodide, elemental iodine and combinations thereof;
- (B) one or more fluoroalcohols of the formula $\text{F}(\text{CF}_2)_m-\text{CH}_2\text{CH}_2-\text{OH}$, where m is 2-20 and n is 1-6; and
- (C) a 1,2 alkylene oxide of the formula



wherein R^9 , R^{10} , R^{11} , and R^{12} are each independently hydrogen or alkyl, at a temperature from about ambient to about 200° C.

and a pressure from ambient atmospheric pressure to reactant pressure.

2. The process of claim 1, wherein the catalyst system comprises an alkyl borate.

3. The process of claim 1, wherein said boron compound is selected from the group consisting of trimethyl borate, triethyl borate, triisopropyl borate, tripropyl borate, tributyl borate, sodium tetraborate, potassium tetraborate and combinations thereof.

4. The process of claim 1, wherein the 1,2 alkylene oxide is used in an amount of from about 1 mole % to about 100 mole % by total moles of the fluoroalcohol.

5. The process of claim 1, wherein said process is carried out in the presence of an aprotic solvent.

6. The process of claim 1, wherein the iodine source is added in an amount from about 0.05% to 10% by weight of the fluoroalcohol.

7. The process of claim 1, wherein the boron source is added in an amount from about 0.01 mole % to 20 mole % by moles of the fluoroalcohol.

8. The process of claim 1, wherein the alkylene oxide comprises one or more alkylene oxides of ethylene oxide, propylene oxide or butylene oxide.

9. The process of claim 1, wherein the alkylene oxide is ethylene oxide.

* * * * *