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- (54) **LABILE BROMINE FIRE SUPPRESSANTS**
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- (58) **Field of Classification Search** **252/2, 252/3, 4, 5, 8**
See application file for complete search history.

- (56) **References Cited**

U.S. PATENT DOCUMENTS

955,316 A * 4/1910 Breslauer 169/47
2,732,410 A * 1/1956 Farlow et al. 570/150
2,791,490 A * 5/1957 Willcox 423/613
2,918,125 A * 12/1959 Sweetman 166/297
3,076,507 A * 2/1963 Sweetman 166/297
3,266,674 A * 8/1966 Martin 222/146.5
3,372,208 A * 3/1968 Waters et al. 528/108
3,479,286 A * 11/1969 Gambaretto et al. 252/8
3,637,022 A * 1/1972 Kelly et al. 169/44
3,657,030 A * 4/1972 Porter 438/703
3,663,283 A * 5/1972 Hebert et al. 106/401
3,812,239 A * 5/1974 Swaney 423/440
3,903,037 A * 9/1975 Deets 524/228
3,959,741 A * 5/1976 Meinzer 372/89
3,985,689 A * 10/1976 Ferguson 521/77
4,022,872 A * 5/1977 Carson et al. 423/297
4,043,950 A * 8/1977 Wilmsen et al. 428/308.4
4,048,290 A * 9/1977 Lee 423/336
4,125,161 A * 11/1978 Chammas 166/297
4,129,513 A * 12/1978 Stangroom 252/78.1
4,241,042 A * 12/1980 Matijevec et al. 423/610
4,250,960 A * 2/1981 Chammas 166/55
4,292,290 A * 9/1981 Tunison, III 423/336
4,397,977 A * 8/1983 Sandler 524/288
4,406,797 A * 9/1983 Altman et al. 252/5
4,415,029 A * 11/1983 Pratt et al. 166/212
4,428,430 A * 1/1984 Terrell et al. 166/298
4,481,119 A * 11/1984 Rhein et al. 252/2
4,494,601 A * 1/1985 Pratt et al. 166/55
4,619,318 A * 10/1986 Terrell et al. 166/55
4,819,728 A * 4/1989 Lafitte 166/298
4,822,513 A * 4/1989 Corby 510/234
4,889,187 A * 12/1989 Terrell et al. 166/298
4,949,789 A * 8/1990 Lafitte 166/298

4,971,146 A * 11/1990 Terrell 166/55
5,017,525 A * 5/1991 Birchall et al. 501/87
5,055,208 A * 10/1991 Stewart et al. 252/8
5,080,177 A * 1/1992 Robin et al. 169/46
5,082,575 A * 1/1992 Yamaguchi 252/2
5,084,190 A * 1/1992 Fernandez 252/8
5,093,013 A * 3/1992 Sprague 252/8
5,120,689 A * 6/1992 Birchall et al. 501/99
5,135,054 A * 8/1992 Nimitz et al. 169/46
5,165,916 A * 11/1992 Cheng et al. 423/346
5,192,714 A * 3/1993 Suguro et al. 438/631
5,219,474 A * 6/1993 Song et al. 252/8
5,235,000 A * 8/1993 McKinnie 525/355
5,236,611 A * 8/1993 Shiflett 252/67
5,287,920 A * 2/1994 Terrell 166/55
5,320,174 A * 6/1994 Terrell et al. 166/297
5,444,102 A * 8/1995 Nimitz et al. 521/131
5,491,028 A * 2/1996 Sarin et al. 428/408
5,520,826 A * 5/1996 Reed et al. 252/5
5,562,764 A * 10/1996 Gonzalez 106/437
5,573,744 A * 11/1996 Gebben et al. 423/613
RE40,651 E * 3/2009 Huntington et al. 252/2
2006/0273223 A1 * 12/2006 Haaland et al. 244/129.2
2007/0119602 A1 * 5/2007 Haaland et al. 169/14
2007/0119603 A1 * 5/2007 Haaland et al. 169/54
2008/0115950 A1 * 5/2008 Haaland et al. 169/62

FOREIGN PATENT DOCUMENTS

JP 56-9230 * 1/1981
SU 981338 * 6/1981
WO WO 93/15794 * 8/1993

OTHER PUBLICATIONS

Letter dated Aug. 18, 2006, from J. Sawtelle to T. Howard.*
A.N. Baratov, A Review of Investigations on the Chemical Inhibition of Flames, Problems in Combustion and Extinguishment, 1968, 29–51, TsNIPO MOOP, Moscow.*
W.A. Rosser, et al., The Effect of Metal Salts on Premixed Hydrocarbon—Air Flames, Combustion and Flame, Mar. 1963, 107–119, vol. 7, Butterworths, London.*
D.R. Miller, et al., Effects of Various Inhibitors on Hydrogen—Air Flame Speeds, Combustion and Flame, Mar. 1963, 137–142, vol. 7, Butterworths, London.*
Letter dated Aug. 15, 2006, from T. Howard to J. Sawtelle.*
Final Technical Report FR-4021 Halogen Replacement for Aviation Systems, Peter D. Haaland & John H Huntington, May 1994–Nov. 1994.*
W.A. Rosser, et al., The Quenching of Premixed Flames by Volatile Inhibitors, ICombustion and Flame, Mar. 1966, 287–294, vol. 10, Butterworths, london.*

(Continued)

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(57) **ABSTRACT**

A class of fire suppressant compounds which have labile bromine atoms bound to atoms other than carbon have been discovered to be more effective at suppressing fires than Halon 1211 and Halon 1301. Moreover, this class of fire suppressant compounds hydrolyze or oxidize rapidly in the troposphere and as a consequence thereof, they have minimal ozone depletion potential.

30 Claims, No Drawings

OTHER PUBLICATIONS

G. Lask & H. GG. Wagner, Influence of Additives on the Velocity of Laminar Flames, 8th Symposium (International) on Combustion, 1962, 432-438, Williams & Wilkens Co., Baltimore.*

The Montreal Protocol on Substances that Deplete the Ozone Layer, 2000, 1-47, UNEP Ozone Secretariat UN Environment Programme, Kenya.*

Chemical Abstracts No. 117:134014, "Preparation of a Fire Extinguishing Composition", Valukonis et al. (1991).*

Chemical Abstracts No. 98:129086. "Combustion inhibitor for hydrocarbon-air Mixture", Petrova et al. (1982).*

CRC Handbook of Chemistry and Physics, 63rd Edition, pp.B-84, B-97, B-106, B-125, B-127, B-143, B-157, B-159 and C-525, (1982).*

Hawley's Condensed Chemical Dictionary, 11u-Edition, pp. 164, 169, 170, 173, 329, 642, 913, and 1040, Van Nostrand Reinhold Company, N.Y. (1987).*

* cited by examiner

LABILE BROMINE FIRE SUPPRESSANTS

Matter enclosed in heavy brackets [] appears in the original patent but forms no part of this reissue specification; matter printed in italics indicates the additions made by reissue.

Notice: More than one reissue application has been filed for the reissue of U.S. Pat. No. 5,626,786. The reissue applications are application Ser. No. 10/893,705, application number not yet assigned, and application number not yet assigned (the present application), all of which are divisional reissues of U.S. Pat. No. 5,626,786.

GOVERNMENT RIGHTS

This invention was made with support by the U.S. Government. The Government may have certain rights in this invention.

BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention generally relates to prevention and extinguishment of fires combustible materials by utilizing a composition of matter group which is highly efficient and environmentally friendly. More particularly, the invention relates to prevention and extinguishment of fires of combustible materials by using a group of fire suppressants having labile bonds between bromine atoms and atoms other than carbon.

2. Description of the Prior Art

Fire suppression by halogenated alkanes is well-established in both the scientific literature and commercial practice as taught, for example, in R. G. Gann ed., *Halogenated Fire Suppressants*, ACS Symposium Series 16 (American Chemical Society, Washington, D.C.) 1975. The two most widely used halogenated suppressants are Halon 1301 (CF₃Br) and Halon 1211 (CF₂ClBr). These compounds are very stable, so they survive long enough in the troposphere to be gradually transported to the stratosphere, where they are photolyzed by solar ultraviolet radiation to produce free radicals that catalyze ozone depletion as taught, for example, in J. G. Anderson, D. W. Toohey, and W. F. Brune, *Science*, 251, 39 (1991). Production of these materials has therefore been internationally prohibited after January, 1994 by the Montreal Protocol on Substances that Deplete the Ozone Layer. The problem is therefore to find fire suppression materials and methods which are at least as effective as Halon 1301 and Halon 1211 but which do not deplete the ozone layer.

Representative of the prior art directed to the use of fluorocarbons which have no chlorine or bromine is U.S. Pat. No. 5,236,611 (Shiflet). These fluorocarbons are slowly transported into the stratosphere, but the catalytic efficiency of fluorine is very much smaller than that of chlorine, bromine, or iodine.

Representative of the prior art directed to the use of hydrogenated halocarbons, which are less stable than Halon 1301 or Halon 1211 in the troposphere, are U.S. Pat. Nos. 5,084,190 (Fernandez), 5,135,054 (Nimitz et al.); 5,093,013 (Sprague); and 5,080,177 (Robin et al.). It is well known that Halons containing chlorine or bromine will suppress fires in smaller quantities than those which contain only fluorine. However, it is understood by people practiced in the art of combustion that the principal source of heat release in hydrocarbon combustion is oxidation of hydrogen atoms to form water vapor. Thus hydrogenated halocarbons are expected to act chemically both as fuels and as fire suppressants.

SUMMARY OF THE INVENTION

In accordance with the teachings of the present invention compositions of matter having labile bromine atoms bound to atoms other than carbon have been discovered to have improved fire suppressant properties and to be environmentally friendly. Compounds which release bromine atoms and are commonly used as brominating agents in organic synthesis such as phosphorus tribromide (PBr₃), thionyl bromide (SOBr₂), boron tribromide (BBr₃), and the like are very efficient at extinguishing fires. Moreover, they hydrolyze or oxidize rapidly in the troposphere and consequently they have no stratospheric ozone depletion potential.

Additionally included in this class of fire suppressant compounds of labile bromine atoms bound to atoms other than carbon are silicon tetrabromide (SiBr₄), titanium tetrabromide (TiBr₄), iodine bromide (TBr), phosphorous oxybromide (POBr₃), bromine trifluoride (BrF₃), bromine pentafluoride (BrF₅), N-bromosuccinimide (C₄H₄O₂NBr) (the bromine is bound to nitrogen, not carbon), nitrosyl bromide (NOBr), chlorine bromide (ClBr), and cuprous bromide (CuBr). These compounds are used as brominating agents in chemical synthesis as taught, for example, in the Merck Index, ninth edition, (Merck & Co., Rahway, N.J., 1976) or suffer thermal decomposition with liberation of bromine at temperatures less than 200 degrees centigrade.

Examples of non-labile bromine compositions are found in such high-melting, ionically bound salts as lithium bromide (LiBr, m.p. 547° C.), calcium bromide (CaBr₂, m.p. 730° C.), or chromous bromide (CrBr₂, m.p. 842° C.), and other bromine-containing compositions that are thermally and oxidatively stable according to criteria familiar to people practiced in the art of synthetic chemistry.

In order to extinguish fires with a composition having one or more compounds of the aforesaid class of fire suppressants, equipment for delivering the composition incorporates such factors as specific geometry, gas flow, and flame conditions. A method of delivery of a composition having one or more liquid compounds of the aforesaid class of fire suppressants may employ a nonflammable, pressurized gas to propel the composition through a nozzle to the flame. Another method of delivery of liquid compositions may employ a deflagrating solid, gas-generating cartridge, such as is found in automotive airbags, to propel a mist of liquid to the flame. A third method for delivery of liquid compositions may employ a pressurized liquid propellant such as liquid carbon dioxide or liquid argon to atomize and direct a mist of suppressant onto the flame. Other methods for propelling powders or slurries of solid materials of the aforesaid class may employ a deflagrating solid gas generating cartridge and a wider nozzle such as would be used for an ordinary shotgun cartridge. Other methods for propelling gaseous materials of the aforesaid class may employ mixtures with pressurized inert propellants to aid transport of suppressant to the flame.

The primary advantage of the use of the class of fire suppressants of this discovery is to extinguish fires more efficiently with smaller volumes and masses of extinguishant than existing fire suppressants, without depleting the stratospheric ozone layer. PBr₃, POBr₃, SOBr₂, BBr₃, and the like react rapidly with water vapor or liquid to produce mild acids which precipitate with normal rain and are ultimately neutralized in soils. The short lifetimes of these materials also reduce toxicity of the suppressants since their simple acid decomposition products pose no chronic risk to pH buffered, living organisms.

These and other advantages and attainments of the present invention will become apparent to those skilled in the art

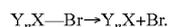
upon a reading of the detailed description wherein there are described several embodiments of the invention.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

In accordance with the teachings of the present invention a class of fire suppressants which have labile bromine atoms bound to atoms other than carbon are shown in Table I. These compounds are more effective at suppressing fire than Halon 1301 or Halon 1211 and have no ozone depletion potential.

The effectiveness of the class of suppressants described herein is a result of the relative ease with which bromine atoms are liberated in a flame environment. Halons 1301 and 1211 also liberate bromine atoms in a flame; however, the strength of the C—Br bond in these materials requires higher temperatures or longer interaction times than the compositions of matter described herein. The stability of the Halons against oxidation or hydrolysis in the troposphere is one indication of the stability of the C—Br bond. The compositions of matter described herein are not stable in the presence of water, oxygen, or heat, liberating bromine atoms under these conditions and thereby providing a catalyst for flame suppression.

Another indication of the stability of the C—Br bond is the measurement of its bond dissociation energy of 68 kilocalories per mole as taught by A. H. Sehon and M. Szwarc (Proceedings of the Royal Society (London), page 110, [1951]). This energy is larger than the bond energy of typical P—Br bonds (63 kcal/mole), I—Br bonds (43 kcal/mole), or S—Br bonds (52 kcal/mole) as taught by Streitwieser and Heathcock, Introduction to Organic Chemistry, (Macmillan:NY), 1976 and Cotton and Wilkinson, Advanced Inorganic Chemistry, (3rd ed. Wiley:NY), 1972. As is known to people practiced in the art of chemistry the interpretation of bond dissociation energies involves approximations based on the nature of the full molecular fragments (XY_n) from which bromine is liberated according to the reaction:



Thus lower average bond energies indicate the possibility of labile bonds in a group of materials, but further experimentation with specific materials is required to establish the lability of the bond compared with Halons.

TABLE I

Composition	Phase	% Br	Comments
PBr ₃	liquid	88	brominating agent
POBr ₃	liquid	83	brominating agent
SOBr ₂	liquid	77	brominating agent
BrF ₃	liquid	58	reactive solvent
BrF ₅	liquid	46	reactive solvent
PBr ₅	solid	92	brominating agent
TiBr ₄	solid	87	reacts with water
SiBr ₄	liquid	92	reacts with water
I ₂	solid	39	decomposes at 116° C.
CuBr	solid	29	brominating agent
NOBr	gas	73	brominating agent
BrF	gas	81	boils at -20° C.
C ₄ H ₄ O ₂ NBr	solid	44.9	decomposes at 170° C.
BBr ₃	liquid	96	boils at 90° C.
BrCl	gas	70	decomposes at 10° C.

In one embodiment of the invention liquid SOBr₂ is introduced as an air-pressurized mist into a 500,000 Btu/hr fire

resulting from kerosene flowing at a rate of 4 grams per second through a nozzle with cross-flowing compressed air to atomize the liquid into a fine mist. The fire is contained in a flame holder whose volume is approximately 8 liters and is further blown by an atmospheric cross-wind of 40 miles per hour. The fire is reproducibly and irreversibly extinguished with less than one gram of SOBr₂ in less than 0.2 seconds as confirmed by videotape records of the experiments. The same fire is not reproducibly suppressed with aliquots of 25 grams of CF₃Br added to the same location.

In another embodiment of the invention 0.2 cubic centimeters of PBr₃ is mixed with 0.7 cubic centimeters of liquid carbon dioxide. The liquid CO₂ propels the PBr₃ through a valve and into the flame zone, generated as herein above described, as it is opened, irreversibly and completely extinguishing the flame in the presence of flowing fuel, air, and hot surfaces.

Extinguishment of a similar fire, with a hydrocarbon fuel burn rate of 12 grams per second, by Halon 1301 is taught by Alvarez in chapter 3 of Gann (ibid.) to require between 90 and 130 grams per second of CF₃Br for suppression. Another example of a gasoline fire with similar heat output is taught by Ford in chapter 1 of Gann (ibid.) to require between 500 and 1500 grams of Halon 1301 for suppression. Another fire, in which 10 grams per second of jet fuel are burned in fast-flowing air at the Air Force Flight Dynamics Laboratory Engine Nacelle test facility (Wright-Patterson AFB, OH) requires between one and three kilograms of Halon 1301 for reproducible suppression. In each of these examples the quantity of Halon 1301 required to suppress a similar fire is between 100 and 1000 times greater than that required of the compositions of matter described herein above, of which SOBr₂ and PBr₃ are specific embodiments.

The labile bromine atoms and high proportion of bromine in the composition of matter listed in Table I provide a more efficient fire suppression formulation than the Halons, which typically have less bromine by weight (Halon 1301 and 1211 are 54% and 48% Br, respectively) and lesser proclivity for liberating bromine atoms when thermally or chemically activated in a combustion environment.

Methods for dispersing gas, liquid, or solid suppressants require designs based upon such factors as specific geometry of the locus of the fire, flow properties of the fire suppressants, and flame conditions of the fire. For example, fine mists of liquid are transported by fluid-dynamical drag forces along flow streamlines in the nacelle of an aircraft engine. The mists vaporize in hot zones, liberating bromine atoms by pyrolysis in precisely the regions where the heat released by combustion is most intense. Inasmuch as the drag coefficient is inversely proportional to the droplet diameter, as is known to people practiced in the art of fluid dynamics, there is a range of aerosol size distributions which most effectively deliver specific suppressants to specific fires. Another such factor for a gaseous composition is the mixing of suppressant flow with turbulent flames in a well-ventilated fire, which is affected by the delivery pressure, the nozzle contour and orientation, the mass-flow rate of the suppressant, and the fluid dynamical properties of the fire. Dispersing methods designed for suppressing fire in the nacelle of a jet engine differ from dispersing methods designed for suppressing fire in the engine compartment of a motor vehicle, the flu of a chimney, or the gas-handling manifold of a semiconductor processing clean-room.

Methods for preventing and extinguishing fires of jet fuel using a composition of matter class which is highly efficient and environmentally friendly is also disclosed by the present inventors in Final Technical Report FR-4021 (US Air Force Phase I SBIR Contract F33615-94-C-5005, November 1994).

Although preferred embodiments of the invention have been described, it will be understood that within the scope of this invention various changes may be made in the amount of fire suppressant, the composition of a fire suppressant mixture, and the method for dispersing fire suppressants which is generally stated consist of a class of fire suppressants and methods of dispersing such fire suppressants capable of carrying out the objects set forth as disclosed in the appended claims.

What is claimed is:

1. A fire suppressant composition consisting essentially of at least one brominated, non-carbon compound selected from the group consisting of PBr_3 , $POBr_3$, $SOBr_2$, BrF_3 , BrF_5 , PBr_5 , $TiBr_4$, $SiBr_4$, IBr , $CuBr$, $NOBr$, BrF , and BBr_3 , which is combined with a propellant such that the ozone depletion potential of the composition is less than 0.1.]

2. The fire suppressant composition of claim 1 wherein ozone depletion potential is defined on a scale where the ozone depletion caused by CFI_3 is 1.0.]

3. A fire suppressant composition consisting of at least one labile brominated compound selected from the group consisting of PBr_3 , $POBr_3$, $SOBr_2$, BrF_3 , BrF_5 , $TiBr_4$, $SiBr_4$, IBr , $CuBr$, $NOBr$, BrF , BBr_3 , and $BrCl$, which is combined with a propellant selected from the group consisting of CO_2 , N_2 , compressed air, and HCFC-123 (CF_3CCl_2H).]

4. The fire suppressant composition of claim 3 wherein ozone depletion potential of the propellant equals 0.016.]

5. A fire suppressant composition consisting essentially of: at least one brominated, non-carbon compound in a liquid state selected from the group consisting of $TiBr_4$ and $SiBr_4$; and a propellant combined with the compound for propelling the composition such that sufficient bromine atoms are liberated from the composition to suppress a fire, wherein the composition has no ozone depletion potential.

6. The composition of claim 5, wherein the propellant is selected from the group consisting of CO_2 , N_2 , and compressed air.

7. The composition of claim 5 wherein said brominated compound is $TiBr_4$.

8. The composition of claim 5 wherein said brominated compound is $SiBr_4$.

9. The composition of claim 5 wherein said brominated compound is $TiBr_4$ and the propellant is nonflammable.

10. The composition of claim 5 wherein said brominated compound is $SiBr_4$ and the propellant is nonflammable.

11. The composition of claim 7 wherein said brominated compound is not stable in, and said sufficient bromine atoms are liberated in, the presence of oxygen.

12. The composition of claim 11 wherein said liberated bromine atoms are sufficient to catalytically suppress the fire.

13. The composition of claim 7 wherein said brominated compound is not stable in, and said sufficient bromine atoms are liberated in, the presence of heat.

14. The composition of claim 13 wherein said liberated bromine atoms are sufficient to catalytically suppress the fire.

15. The composition of claim 7 wherein said brominated compound is not stable in, and said sufficient bromine atoms are liberated in, the presence of water.

16. The composition of claim 15 wherein said liberated bromine atoms are sufficient to catalytically suppress the fire.

17. The composition of claim 8 wherein said brominated compound is not stable in, and said sufficient bromine atoms are liberated in, the presence of oxygen.

18. The composition of claim 17 wherein said liberated bromine atoms are sufficient to catalytically suppress the fire.

19. The composition of claim 8 wherein said brominated compound is not stable in, and said sufficient bromine atoms are liberated in, the presence of heat.

20. The composition of claim 19 wherein said liberated bromine atoms are sufficient to catalytically suppress the fire.

21. The composition of claim 8 wherein said brominated compound is not stable in, and said sufficient bromine atoms are liberated in, the presence of water.

22. The composition of claim 21 wherein said wherein said liberated bromine atoms are sufficient to catalytically suppress the fire.

23. The composition of claim 7 wherein the propellant is selected from the group consisting of a non-flammable, pressurized gas; a deflagrating, solid, gas-generating cartridge; and a pressurized liquid.

24. The composition of claim 8 wherein the propellant is selected from the group consisting of a non-flammable, pressurized gas; a deflagrating, solid, gas-generating cartridge; and a pressurized liquid.

25. A fire suppressant composition consisting essentially of $TiBr_4$ in a liquid state and a propellant combined with the $TiBr_4$ for propelling the composition to catalytically suppress a fire, wherein the composition has no ozone depletion potential.

26. The composition of claim 25, wherein the propellant is selected from the group consisting of CO_2 , N_2 , and compressed air.

27. The composition of claim 25 wherein the propellant is nonflammable.

28. The composition of claim 25 wherein the $TiBr_4$ is not stable in the presence of at least one of oxygen, heat, and water.

29. The composition of claim 25 wherein the propellant is selected from the group consisting of a non-flammable, pressurized gas; a deflagrating, solid, gas-generating cartridge; and a pressurized liquid.

30. A fire suppressant composition consisting essentially of $SiBr_4$ in a liquid state and a propellant combined with the $SiBr_4$ for propelling the composition to catalytically suppress a fire, wherein the composition has no ozone depletion potential.

31. The composition of claim 30, wherein the propellant is selected from the group consisting of CO_2 , N_2 , and compressed air.

32. The composition of claim 30 wherein the propellant is nonflammable.

33. The composition of claim 30 wherein the $SiBr_4$ is not stable in the presence of at least one of oxygen, heat, and water.

34. The composition of claim 30 wherein the propellant is selected from the group consisting of a non-flammable, pressurized gas; a deflagrating, solid, gas-generating cartridge; and a pressurized liquid.