

TITLE

HFO-1234ZE, HFO-1225ZC AND HFO-1234YF CONTAINING COMPOSITIONS AND PROCESSES FOR PRODUCING AND USING THE COMPOSITIONS

5 This Application claims the benefit of PCT Application No.
PCT/US20/029690, filed on April 24, 2020. The disclosure of Application No.
PCT/US20/029690 is hereby incorporated by reference.

FIELD OF THE INVENTION

10 The present invention relates to tetrafluoropropene compositions and methods
for making and using the compositions. The invention particularly concerns
compositions comprising E-1,3,3,3-tetrafluoropropene (HFO-1234ze),
1,1,3,3,3-pentafluoropropene (HFO-1225zc), and 2,3,3,3-tetrafluoropropene
(HFO-1234yf), and uses thereof as well as a method for producing a product
comprising 1,3,3,3-tetrafluoropropene (HFO-1234ze), 1,1,3,3,3-pentafluoropropene
15 (HFO-1225zc), and 2,3,3,3-tetrafluoropropene (HFO-1234yf) from 1,1,1,3,3-
pentafluoropropane (HFC-245fa).

BACKGROUND OF THE INVENTION

20 The fluorocarbon industry has been working for the past few decades to find
replacement refrigerants for the ozone depleting chlorofluorocarbons (CFCs) and
hydrochlorofluorocarbons (HCFCs) being phased out as a result of the Montreal
Protocol. The solution for many applications has been the commercialization of
hydrofluorocarbon (HFC) compounds for use as refrigerants, solvents, fire
extinguishing agents, blowing agents and propellants. These new compounds, such
as HFC refrigerants, HFC-134a and HFC-125 being the most widely used at this
25 time, have zero ozone depletion potential and thus are not affected by the current
regulatory phase-out as a result of the Montreal Protocol.

30 In addition to ozone depleting concerns, global warming is another
environmental concern in many of these applications. Thus, there is a need for
compositions that meet both low ozone depletion standards as well as having low
global warming potentials. Certain hydrofluoroolefin compositions are believed to

meet both goals. Thus, there is also a need for economical manufacturing processes that provide these compositions.

HFO-1234ze ($\text{CF}_3\text{CH}=\text{CHF}$) and HFO-1234yf ($\text{CF}_3\text{CF}=\text{CH}_2$), both having zero ozone depletion and low global warming potential, have been identified as potential refrigerants. Particularly the E-isomer of HFO-1234ze (HFO-1234zeE) has proven to be useful for refrigeration applications. US Patent No 7,862,742 discloses compositions comprising HFO-1234ze and HFO-1234yf. U.S. Patent No. 9,302,962 discloses methods for making HFO-1234ze. The disclosures of U.S. Patent No. 7,862,742 and U.S. Patent No. 9,302,962 are hereby incorporated by reference in their entirety.

Catalytic dehydrofluorination of HFC-245fa in general produces a mixture of both the E-isomer as well as the Z-isomer of HFC-1234ze. Depending on the particular catalyst chosen, the amount of the Z-isomer can vary between 15% to 23%. Dehydrofluorination in the liquid phase using aqueous solutions of caustic or other strong bases also produces mixture of both isomers. Although the ratio of the two isomers can be shifted somewhat by temperature, about 13% to about 15% of the Z-isomer is typically formed. As the E-isomer is the most useful for refrigeration applications, after separation of the E-isomer from the Z-isomer, the Z-isomer is typically either isomerized to the E-isomer in a separate step or converted back to 245fa through addition of hydrogen fluoride. Both alternatives require additional steps which add cost.

There is a need in this art for art for compositions comprising HFO-1234zeE which can maintain the favorable properties of HFO-1234zeE. There is a further need for a process that can produce compositions of HFO-1234ze and HFO-1234yf that minimizes or eliminates the need for purification or separation steps for removing excess quantities of HFO-1234yf.

BRIEF DESCRIPTION OF THE INVENTION

Described is a fluoropropene composition comprising E-1,3,3,3-tetrafluoropropene, 1,1,3,3,3-pentafluoropropene, and 2,3,3,3-tetrafluoropropene. The total amount of 1,1,3,3,3-pentafluoropropene and

2,3,3,3-tetrafluoropropene in the composition is 1.0 wt.% or less based on the total weight of the fluoropropene composition.

In addition, the present disclosure includes a method of producing a mixture of 1,3,3,3-tetrafluoropropene and 2,3,3,3-tetrafluoropropene, comprising contacting
5 a mixture of 1,1,1,3,3-pentafluoropropane and Z-1,3,3,3-tetrafluoropropene in the gas phase with a catalyst comprising at least one catalyst selected from the group consisting of fluorinated Cr_2O_3 and Cr/Ni on fluorinated alumina, preferably in the presence of an oxygen containing gas, to form a mixture comprising Z-1,3,3,3-tetrafluoropropane, E-1,3,3,3-tetrafluoropropene, 1,1,3,3,3-pentafluoropropene,
10 2,3,3,3-tetrafluoropropene, hydrogen fluoride, and optionally unreacted 1,1,1,3,3-pentafluoropropane. One embodiment the inventive method produces a useful composition without the need for purification or separation steps including steps for removing excess quantities of 2,3,3,3-tetrafluoropropene (HFO-1234yf) or 1,1,3,3,3-pentafluoropropene.

15 Further still, the present disclosure includes fluoropropene compositions formed from the method of contacting a mixture of 1,1,1,3,3-pentafluoropropane and Z-1,3,3,3-tetrafluoropropene in the gas phase with a catalyst comprising at least one catalyst selected from the group consisting of fluorinated Cr_2O_3 and Cr/Ni on fluorinated alumina, optionally in the presence of an oxygen containing gas.

20 In one embodiment, the inventive process produces a composition comprising HFO-1234ze(E), HFO-1225zc and HFO-1234yf and the composition is useful as a refrigerant. In another embodiment, the composition comprises HFO-1225zc and a near azeotropic composition comprising HFO-1234ze(E) and HFO-1234yf. In a further embodiment, the composition comprises a three component near azeotropic
25 composition comprising HFO-1234ze(E), HFO-1234yf and HFO-1225zc.

One embodiment relates to any combination of the foregoing wherein the 2,3,3,3-tetrafluoropropene is present in the fluoropropene composition in an amount of 0.0001 to 0.9 wt.% based on the total weight of the fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the
30 2,3,3,3-tetrafluoropropene is present in the fluoropropene composition in an

amount of 0.001 to 0.8 wt.% based on the total weight of the fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the 2,3,3,3-tetrafluoropropene is present in the fluoropropene composition in an amount of 0.1 to 0.6 wt.% based on the total weight of the fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the 2,3,3,3-tetrafluoropropene is present in the fluoropropene composition in an amount of 0.3 to 0.4 wt.% based on the total weight of the fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the 2,3,3,3-tetrafluoropropene is present in an amount of 0.01 to 1.0 mol%.

One embodiment relates to any combination of the foregoing wherein the 2,3,3,3-tetrafluoropropene is present in an amount of 0.1 to 0.9 mol%.

One embodiment relates to any combination of the foregoing wherein the 2,3,3,3-tetrafluoropropene is present in an amount of 0.2 to 0.4 mol%.

One embodiment relates to any combination of the foregoing wherein the 2,3,3,3-tetrafluoropropene is present in an amount of 0.3 to 0.4 mol%.

Another embodiment relates to any combination of the foregoing wherein the 1,1,3,3,3-pentafluoropropene is present in the fluoropropene composition in an amount of 0.0001 to 0.9 wt.%, preferably in an amount of 0.001 to 0.8 wt.%, more preferably in an amount of 0.01 to 0.6 wt.%, and most preferably in an amount of 0.1 to 0.4 wt.% based on the total weight of the fluoropropene composition.

Another embodiment relates to any combination of the foregoing wherein the E-1,3,3,3-tetrafluoropropene is present in the fluoropropene composition in an amount of 50.0 wt.% or more, preferably in an amount of 75.0 wt.% or more, more preferably in an amount of 99.0 wt.% or more, even more preferably in an amount of 99.5 wt.% or more, and most preferably in an amount of 99.8 wt.% or more, based on the total weight of the fluoropropene composition.

Another embodiment relates to any combination of the foregoing the total amount of 2,3,3,3-tetrafluoropropene and 1,1,3,3,3-pentafluoropropene in the fluoropropene composition is 0.001 to 0.9 wt.%, preferably is 0.1 to 0.8 wt.%, and most preferably is 0.3 to 0.5 wt.% based on the total weight of the fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the fluoropropene composition additionally comprises R-134, preferably in an amount of 1.0 to 40.0 wt.%, more preferably in an amount of 30.0 to 40.0 wt.%, and most preferably in an amount of 35.0 to 40.0 wt.% based on the total weight of the fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the fluoropropene composition additionally comprises R-1336mzzE and/or R-227ea, preferably in an amount of 15.0 to 20.0 wt.% of R-1336mzzE and of 2.0 to 5.0 wt.% of R-227ea based on the total weight of the fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the fluoropropene composition additionally optionally comprises one or more of R-143a, R-152a, TFP (trifluoropropyne), R-1233xf, R-1233zd(E), R-1233zd(Z), R236fa, and at least one HFO-1234 isomer including at least one of HFO-1234zc, HFO-1234yc and HFO-1234ye.

One embodiment relates to any combination of the foregoing wherein the sum total of the amounts of R-143a, R-152a, TFP, R-1233xf, R-1233zd(E), and R-1233zd(Z) is between 0.001 mole percent and 2 mole percent, based on the total fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the fluoropropene composition includes R-1233zd(E) in an amount of 0.7 mole percent to 1.15 mole percent, based on the total fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the fluoropropene composition includes R-1233zd(Z) in an amount of 0.05 mole percent to 0.25 mole percent, based on the total fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the fluoropropene composition includes R-143a in an amount of 0.05 mole percent to 0.25 mole percent, based on the total fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the
 5 fluoropropene composition optionally comprises one or more of 1224yd, 1224zc, 1326mxz, 113, 32, 23, trifluoropropyne, 356mff, 1326mxz, HFC-245fa and HFC-245cb.

One embodiment relates to any combination of the foregoing wherein the sum
 total of the amounts 1224yd, 1224zc, 1326mxz, 113, 32, 23, trifluoropropyne,
 10 356mff, 1326mxz, HFC-245fa and HFC-245cb is between 0.001 mole percent and 2 mole percent, based on the total fluoropropene composition.

One embodiment relates to any combination of the foregoing wherein the composition is near-azeotropic.

A further embodiment of the invention relates to a method of producing a mixture
 15 of 1,3,3,3-tetrafluoropropene and 2,3,3,3-tetrafluoropropene, comprising:

(a) contacting a mixture of 1,1,1,3,3-pentafluoropropane and
 Z-1,3,3,3-tetrafluoropropene in the gas phase with a catalyst comprising at
 least one catalyst selected from the group consisting of fluorinated Cr_2O_3 and
 Cr/Ni on fluorinated alumina, optionally in the presence of an oxygen
 20 containing gas, to form a mixture comprising Z-1,3,3,3-tetrafluoropropene,
 E-1,3,3,3-tetrafluoropropene, 1,1,3,3,3-pentafluoropropene, 2,3,3,3-
 tetrafluoropropene, hydrogen fluoride, and optionally unreacted
 1,1,1,3,3-pentafluoropropane,

wherein the mixture comprises 0.00001 to 1.0 wt.% 2,3,3,3-tetrafluoropropene
 25 and greater than 0 and less than 1.0 wt.% 1,1,3,3,3-pentafluoropropene.

Another embodiment of the invention relates to a method of producing a mixture of
 a fluoropropene of formula $\text{CF}_3\text{CH}=\text{CHF}$ and a fluoropropene of formula
 $\text{CF}_3\text{CF}=\text{CH}_2$, comprising:

(a) contacting a mixture of 1,1,1,3,3-pentafluoropropane and
 30 Z-1,3,3,3-tetrafluoropropene in the gas phase with a catalyst comprising at

least one catalyst selected from the group consisting of fluorinated Cr_2O_3 or Cr/Ni on fluorinated alumina, in the presence of an oxygen containing gas, to form a mixture comprising Z-1,3,3,3-tetrafluoropropene, E-1,3,3,3-tetrafluoropropene, 2,3,3,3-tetrafluoropropene, hydrogen fluoride, and optionally unreacted 1,1,1,3,3-pentafluoropropane, wherein the mixture includes 0.00001% to 1.00% 2,3,3,3-tetrafluoropropene.

One embodiment of the invention relates to any combination of the foregoing wherein said mixture of 1,1,1,3,3-pentafluoropropane and 1,3,3,3-tetrafluoropropene comprises at least 7 wt.% Z-1,3,3,3-tetrafluoropropene, preferably at least 10 wt.% Z-1,3,3,3-tetrafluoropropene.

One embodiment of the invention relates to any combination of the foregoing wherein at least 94%, preferably at least 98% of the 1,1,1,3,3-pentafluoropropane is converted to E-1,3,3,3-tetrafluoropropene.

One embodiment of the invention relates to any combination of the foregoing wherein the method further comprises

(b) recovering Z-1,3,3,3-tetrafluoropropene, or a mixture of Z-1,3,3,3-tetrafluoropropene and 1,1,1,3,3-pentafluoropropane, and recycling Z-1,3,3,3-tetrafluoropropene, or a mixture of Z-1,3,3,3-tetrafluoropropene and 1,1,1,3,3-pentafluoropropane back to step (a).

One embodiment of the invention relates to any combination of the foregoing wherein said hydrogen fluoride produced in step (a) is separated and recovered.

One embodiment of the invention relates to any combination of the foregoing wherein said oxygen containing gas is oxygen or air.

One embodiment of the invention relates to any combination of the foregoing wherein said mixture of 1,1,1,3,3-pentafluoropropane and 1,3,3,3-tetrafluoropropene comprises 0.1 to 0.8 wt.% 2,3,3,3-tetrafluoropropene, preferably 0.2 to 0.6 wt.% 2,3,3,3-tetrafluoropropene, more preferably 0.3 to 0.4 wt.% 2,3,3,3-tetrafluoropropene.

A further embodiment of the invention relates to any combination of the foregoing wherein said mixture of 1,1,1,3,3-pentafluoropropane and Z-1,3,3,3-tetrafluoropropene comprises at least 7% by weight Z-1,3,3,3-tetrafluoropropene.

One embodiment of the invention relates to any combination of the foregoing
5 wherein said mixture of 1,1,1,3,3-pentafluoropropane and Z-1,3,3,3-tetrafluoropropene comprises at least 10% by weight Z-1,3,3,3-tetrafluoropropene.

One embodiment of the invention relates to any combination of the foregoing wherein at least 94% by weight of the 1,1,1,3,3-pentafluoropropane is converted to E-isomer of 1,3,3,3-tetrafluoropropene.

10 One embodiment of the invention relates to any combination of the foregoing wherein at least 98% by weight of the 1,1,1,3,3-pentafluoropropane is converted to E-isomer of 1,3,3,3-tetrafluoropropene.

One embodiment of the invention relates to any combination of the foregoing and further comprising recovering Z-1,3,3,3-tetrafluoropropene, or a mixture of Z-
15 1,3,3,3-tetrafluoropropene and 1,1,1,3,3-pentafluoropropane, and recycling Z-1,3,3,3-tetrafluoropropene, or a mixture of Z-1,3,3,3-tetrafluoropropene and 1,1,1,3,3-pentafluoropropane back to step (a).

One embodiment of the invention relates to any combination of the foregoing wherein said hydrogen fluoride produced in step (a) is separated and recovered.

20 One embodiment of the invention relates to any combination of the foregoing wherein said oxygen containing gas is oxygen, or air.

One embodiment of the invention relates to any combination of the foregoing wherein the mixture includes 0.1 to 0.5 mol% 2,3,3,3-tetrafluoropropene.

One embodiment of the invention relates to any combination of the foregoing
25 wherein the mixture includes 0.2 to 0.4 mol% 2,3,3,3-tetrafluoropropene.

One embodiment of the invention relates to any combination of the foregoing wherein the mixture includes 0.3 to 0.4 mol% 2,3,3,3-tetrafluoropropene.

Another embodiment of the invention relates to any combination of the foregoing methods and to a fluoropropene composition produced by these methods.

A further embodiment of the invention relates to any combination of the foregoing embodiments and comprising a refrigerant composition comprising Z-1,3,3,3-tetrafluoropropene, E-1,3,3,3-tetrafluoropropene, 1,1,3,3,3-pentafluoropropene, and 2,3,3,3-tetrafluoropropene and at least one member

5 selected from the following groups:

- (a) comprising one or more of R-143a, R-152a, TFP, R-1233xf, R-1233zd(E), R-1233zd(Z), 1224yd, 1224zc, 1326mxz, 113, 32, 23, 356mff, 1326mxz, HFC-245fa, HFC-245cb, 1234zc, 1234yc, 1234ye, 134a, 1225ye (Z and E), 114, 124, and 236fa,
- 10 (b) comprising one or more of R-143a, R-152a, TFP, R-1233xf, R-1233zd(E), R-1233zd(Z), 1224yd, 1224zc, 1326mxz, 113, 32, 23, 356mff, 1326mxz, HFC-245fa and HFC-245cb,
- (c) comprising one or more of HFC-1234ye, HFC-1243zf, HFC-32, HFC-125, HFC-134, HFC-134a, HFC-143a, HFC-152a, HFC-161, HFC-227ea, 15 HFC-236ea, HFC-236fa, HFC-245fa, HFC-365mfc, R1336mzz(E), propane, n-butane, isobutane, 2-methylbutane, n-pentane, cyclopentane, dimethylether, CF_3SCF_3 , CO_2 , and CF_3I ;
- (d) combinations thereof.

One embodiment of the invention relates the fluoropropene composition
 20 further comprising a lubricant selected from the group consisting of mineral oils, such as paraffins, naphthenes and aromatics, alkylaryls, such as linear and branched alkyl alkylbenzenes, synthetic paraffins, synthetic naphthenes, poly(alphaolefins), polyol esters, polyalkylene glycols, and polyvinyl ethers.

One embodiment of the invention relates to a process for treating a surface,
 25 comprising:

providing an article; and

contacting the article with a heat transfer media;

wherein the heat transfer media comprises the fluoropropene composition of any combination of the foregoing embodiments.

One embodiment of the invention relates to a process for treating a surface, comprising:

providing a surface; and

contacting the surface with a treatment composition;

5 wherein the surface includes a treatable material deposited thereon; and

wherein the treatment composition comprises the fluoropropene composition of any combination of the foregoing embodiments.

One embodiment of the invention relates to said process for treating a surface, wherein the treatment composition substantially dissolves the treatable material.

10 One embodiment of the invention relates to a process forming a composition comprising:

providing a solute;

contacting the solute with a solvent;

15 wherein the solvent comprises the fluoropropene composition of any combination of the foregoing embodiments.

One embodiment of the invention relates to a refrigeration system, comprising:

an evaporator; a condenser; a compressor; an expansion device; and a heat transfer medium;

20 wherein the heat transfer medium comprises the fluoropropene composition of any combination of the foregoing embodiments.

One embodiment of the invention relates to a chiller apparatus comprising the fluoropropene composition of any combination of the foregoing embodiments.

25 One embodiment of the invention relates to the fluoropropene composition of any combination of the foregoing embodiments for use in a chiller apparatus, preferably in a medium temperature chiller apparatus.

One embodiment of the invention relates to the use of the fluoropropene composition of any combination of the foregoing embodiments in a chiller apparatus, preferably in a medium temperature chiller apparatus.

One embodiment of the invention relates to a method of producing cooling,
5 the method comprising: evaporating the composition of any combination of the foregoing embodiments in the vicinity of a body to be cooled and thereafter condensing the composition.

One embodiment of the invention relates to a method for replacing a high GWP refrigerant in a refrigeration, air conditioning, or heat pump apparatus,
10 wherein the high GWP refrigerant is selected from the group consisting of R134a, R22, R123, R11, R245fa, R114, R236fa, R124, R12, R410A, R407C, R417A, R422A, R507A, R502, and R404A, the method comprising providing the composition of any combination of the foregoing embodiments to the refrigeration, air conditioning, or heat pump apparatus that uses, used or is designed to use the
15 high GWP refrigerant.

One embodiment of the invention relates to a refrigeration apparatus, air-conditioning apparatus, or heat pump apparatus containing the composition of any combination of the foregoing embodiments, wherein the refrigeration apparatus preferably is a stationary or mobile refrigeration apparatus and the air-conditioning
20 apparatus preferably is a mobile air-conditioning apparatus, more preferably an automobile air-conditioning apparatus.

One embodiment of the invention relates to a process for transferring heat, comprising:

providing an article;
25 contacting the article with a heat transfer media;

wherein the heat transfer media comprises the fluoropropene composition of any combination of the foregoing embodiments and including a near azeotropic composition produced by the inventive method.

One embodiment of the invention relates to a process for treating a surface,
30 comprising:

providing a surface;

contacting the surface with a treatment composition;

wherein the surface includes a treatable material deposited thereon; and wherein the treatment composition comprises the fluoropropene composition of any

5 combination of the foregoing embodiments.

One embodiment of the invention relates to any combination of the foregoing wherein the treatment composition substantially dissolves the treatable material.

One embodiment of the invention relates to a process for forming a composition comprising:

10 providing a solute; contacting the solute with a solvent;

wherein the solvent comprises the fluoropropene composition of any of the foregoing embodiments.

Another embodiment of the invention relates to a refrigeration system, comprising:

15 an evaporator; a condenser;

a compressor; an expansion device;

and a heat transfer media;

wherein the heat transfer media comprises the fluoropropene composition of any combination of the foregoing embodiments and including a near azeotropic

20 composition produced by the inventive method.

The foregoing general description and the following detailed description are exemplary and explanatory only and are not restrictive of the invention, as defined in the appended claims. The various embodiments of the invention can be used alone or in combinations with each other. Other features and advantages of the present invention will be apparent from the following more detailed description of the preferred embodiment, taken in conjunction with the accompanying drawings, which illustrate, by way of example, the principles of the invention.

25

DETAILED DESCRIPTION OF THE INVENTION

Described is a fluoropropene composition comprising
 E-1,3,3,3-tetrafluoropropene, 1,1,3,3,3-pentafluoropropene, and
 2,3,3,3-tetrafluoropropene. The total amount of 1,1,3,3,3-pentafluoropropene and
 5 2,3,3,3-tetrafluoropropene in the composition is 1.0 wt.% or less based on the total
 weight of the fluoropropene composition.

In addition, the present disclosure includes a method of producing a mixture of a
 fluoropropene of formula $\text{CF}_3\text{CH}=\text{CHF}$, a fluoropropene of formula $\text{CF}_2=\text{CHCF}_3$
 and a fluoropropene of formula $\text{CF}_3\text{CF}=\text{CH}_2$, comprising contacting a mixture of
 10 1,1,1,3,3-pentafluoropropane and Z-1,3,3,3-tetrafluoropropene in the gas phase
 with a catalyst comprising at least one catalyst selected from the group consisting
 of fluorinated Cr_2O_3 or Cr/Ni on fluoride alumina, optionally in the presence of an
 oxygen containing gas, to form a mixture comprising Z-1,3,3,3-tetrafluoropropene,
 E-1,3,3,3-tetrafluoropropene, 1,1,3,3,3-pentafluoropropene, 2,3,3,3-
 15 tetrafluoropropene, and, optionally, unreacted 1,1,1,3,3-pentafluoropropane.
 Without wishing to be bound by any theory or explanation, a higher contact
 temperature and/or increased length of catalyst contact time can cause formation of
 an increased amount of 1,1,3,3,3-pentafluoropropene.

Certain dehydrofluorination reactions are well known in the art. The
 20 dehydrofluorination of HFC-245fa has been particularly studied. Both gas phase
 and liquid phases processes are known. 1,3,3,3-tetrafluoropropene (HFO-1234ze)
 exists as both a Z-isomer and an E-isomer about the double bond. Both gas phase
 and liquid phase processes are known to produce a mixture of both the Z- and E-
 isomers, with the E-isomer predominating. The selectivity for the production of the
 25 Z-isomer can vary from about 10% to about 23%, depending on the temperature,
 and choice of catalyst. The boiling point of the E-isomer at 1 atm is about -19°C ,
 while the boiling point of the Z-isomer is about 9°C . For many uses, the E-isomer is
 preferred. So as to minimize yield losses in the form of the generally unwanted Z-
 isomer, it becomes necessary to either add an isomerization step to isomerize the Z-
 30 isomer to the E-isomer or add a fluorination step to convert HFO-1234ze(Z) back to
 HFC-245fa.

The dehydrofluorination reaction according to embodiments of the present disclosure may result in compositions of HFO-1234ze(E), HFO-1225zc and HFO-1234yf that minimizes or eliminates the need for purification or separation steps for removing excess quantities of HFO-1234yf or HFO-1225zc. In some cases, the composition may be azeotropic or near azeotropic or include an azeotropic or near azeotropic composition. By azeotropic compositions it is meant a constant-boiling mixture of two or more substances that behave as a single substance. One manner to characterize an azeotropic composition is that the vapor produced by partial evaporation or distillation of a liquid has the same composition as the liquid from which it is evaporated or distilled (i.e., the mixture distills/refluxes without compositional change). Constant-boiling compositions are characterized as azeotropic because they exhibit either a maximum or minimum boiling point, as compared with that of the non-azeotropic mixture of the same compounds. An azeotropic composition will not fractionate within a refrigeration or air conditioning system during operation. Additionally, an azeotropic composition will not fractionate upon leakage from a refrigeration or air conditioning system. In the situation where one component of a mixture is flammable, fractionation during leakage could lead to a flammable composition either within the system or outside of the system.

By a near-azeotropic composition it is meant to refer to a substantially constant boiling liquid admixture of two or more compounds that behave essentially as a single substance. One manner to characterize a near-azeotropic composition is that the vapor produced by partial evaporation or distillation of a liquid has substantially the same composition as the liquid from which it was evaporated or distilled, that is, the admixture distills/refluxes without substantially compositional change. Another manner to characterize a near-azeotropic composition is that the bubble point vapor pressure and the dew point pressure of the composition at a particular temperature are substantially the same. Herein, a composition of the invention is near-azeotropic if, after 50 weight percent (50 wt.%) of the composition is removed, such as by evaporation or boiling off, the difference in vapor pressure, between the original composition and the composition remaining after 50 weight percent of the original composition has been removed, is less than about 10 percent (10%).

In accordance with one embodiment of the instant invention, the inventive compositions have a flammability rating of A2L as determined by ASHRAE Standard 34 and ASTM E681-09.

Many aspects and embodiments have been described above and are merely exemplary and not limiting. After reading this specification, skilled artisans appreciate that other aspects and embodiments are possible without departing from the scope of the invention.

Other features and benefits of any one or more of the embodiments will be apparent from the following detailed description, and from the claims.

Certain dehydrofluorinations are known in the art, and are preferably conducted in the vapor phase. The dehydrofluorination reaction may be conducted in any suitable reaction vessel or reactor, but it should preferably be constructed from materials which are resistant to the corrosive effects of hydrogen fluoride, such as nickel and its alloys, including Hastelloy, Monel, and Inconel, or vessels lined with fluoropolymers. These may be a single tube, or multiple tubes packed with a dehydrofluorination catalyst.

Useful catalysts for the process include chromium-based catalysts such as fluorinated chromium oxide, which catalyst may either be unsupported, or supported on a support such as activated carbon, graphite, fluoride graphite, or alumina fluoride. The chromium catalyst may either be used alone, or in the presence of a co-catalyst selected from nickel, cobalt, manganese or zinc salt. In one embodiment, a chromium catalyst is high surface area chromium oxide, or chromium/nickel on alumina fluoride (Cr/Ni/AlF_3), the preparation of which is reported in European Patent Application EP 0486333A1; the disclosure of which is hereby incorporated by reference. In another embodiment, the catalyst is fluorinated Guignet's green catalyst. Additional suitable catalysts include, but are not limited to, JM 62-2 (chrome catalyst available from Johnson Matthey), LV (chrome catalyst available from Chemours), JM-62-3 (chrome catalyst available from Johnson Matthey), and Newport Chrome (chrome catalyst available from Chemours). The chromium catalysts are preferably activated before use, typically by a procedure whereby the catalyst is heated to from 350°C to 400°C under a flow

of nitrogen for a period of time, after which the catalyst is heated under a flow of HF and nitrogen or air for an additional period of time.

In one embodiment, the Guignet's Green of the fluoride-activated Guignet's Green catalyst used in the present invention is made by reacting (fusing) boric acid with alkali metal dichromate at 500°C to 800°C, followed by hydrolysis of the reaction product, whereby said Guignet's Green contains boron, alkali metal, and water of hydration. The usual alkali metal dichromates are the Na and/or K dichromates. The reaction is typically followed by the steps of cooling the reaction product in air, crushing this solid to produce a powder, followed by hydrolysis, filtering, drying, milling and screening. The Guignet's Green is bluish green, but is known primarily as a green pigment, whereby the pigment is commonly referred to as Guignet's Green. When used as a catalyst, it is also referred to as Guignet's Green as disclosed in U.S. Pat. No. 3,413,363. In U.S. Pat. No. 6,034,289, Cr₂O₃ catalysts are disclosed as preferably being in the alpha form, and Guignet's Green is also disclosed as a commercially available green pigment having the composition: Cr₂O₃ 79–83 %, H₂O 16–18 %, B₂O₅ 1.5 to 2.7 % (sentence bridging cols. 2 and 3) that can be converted to the alpha form (col. 3, I. 3). U.S. Pat. No. 7,985,884 acknowledges the presence of alkali metal in the Guignet's Green in the composition of Guignet's Green disclosed in Example 1: 54.5% Cr, 1.43% B, 3,400 ppm Na, and 120 ppm K. The disclosure of the foregoing patents and patent applications is hereby incorporated by reference.

The physical shape of the catalyst is not critical and may, for example, include pellets, extrudates, powders, or granules. The fluoride activation of the catalyst is preferably carried out on the final shape of the catalyst.

In one embodiment, the instant invention relates to feeding a mixture of HFC-245fa and at least about 10% by weight of the Z-isomer of HFO-1234ze to a dehydrofluorination reactor in the presence of an oxygen containing gas in order to suppress the formation of additional Z-isomer so that the HFC-245fa converted by dehydrofluorination produces substantially only E-HFO-1234ze, HFO-1225zc and HFO-1234yf. Feeding less than about 10% will result in some suppression of the formation of additional Z-1234ze. Feeding greater than about 10% by weight of Z-1234ze simply results in the presence of additional material which must be

separated and recycled. The amount of Z-1234ze which is necessary to suppress the further formation of Z-isomer product is dependent to some extent on conversion. At 70% conversion of 245fa, about 10–11% Z-isomer in the feed is required. At 80% conversion, about 13% Z-isomer in the feed is required

5 In one embodiment, the reaction vessel can be held at a temperature of between 200°C and 425°C. In another embodiment, the reaction vessel can be held at a temperature of between 250°C and 350°C. In yet another embodiment, the reaction vessel can be held at a temperature of between 275°C and 325°C or between 350°C to 410°C.

10 The reaction pressure can be subatmospheric, atmospheric, or superatmospheric. In one embodiment, the reaction is conducted at a pressure of from 14 psig to about 100 psig. In another embodiment, the reaction is conducted at a pressure of from 14 psig to about 60 psig. In yet another embodiment, the reaction is conducted at a pressure of from 40 psig to about 85 psig. In yet another
15 embodiment, the reaction is conducted at a pressure of from 50 psig to 75 psig. In general, increasing the pressure in the reactor above atmospheric pressure will act to increase the contact time of the reactants in the process. Longer contact times will necessarily increase the degree of conversion in a process, without having to increase temperature.

20 Depending on the temperature of the reactor, and the contact time, the product mixture from the reactor will contain varying amounts of unreacted HFC-245fa. In certain embodiment, E-1,3,3,3-tetrafluoropropene, 1,1,3,3,3-pentafluoropropene, and HFO-1234yf may be separated from the Z-1,3,3,3-tetrafluoropropene, hydrogen fluoride, and any unreacted HFC-245fa, which are then recycled back to
25 the reactor with additional HFC-245fa. Hydrogen fluoride may be removed by scrubbing, by passing the reactor effluent through a solution of aqueous caustic, or hydrogen fluoride may be removed by distillation. In particularly suitable embodiments, the composition formed from the process of the present disclosure includes both 1,3,3,3-tetrafluoropropene (HFO-1234ze(E)), HFO-1225zc and
30 2,3,3,3-tetrafluoropropene (HFO-1234yf), which are not separated.

 In one embodiment, the reactor feed is preheated in a vaporizer to a temperature of from about 30°C to about 100°C. In another embodiment, the

reactor feed is preheated in a vaporizer to a temperature of from about 30°C to about 80°C.

In some embodiments, an inert diluent gas is used as a carrier gas for the hydrochlorofluoropropane. In one embodiment, the carrier gas is selected from
5 nitrogen, argon, helium, or carbon dioxide.

In one embodiment, the product mixture includes (on a mol basis) between 0.01% to 1.00% HFO-1234yf, alternatively between 0.05% to 0.95% HFO-1234yf, alternatively between 0.10% to 0.90% HFO-1234yf, alternatively between 0.20% to 0.80% HFO-1234yf, alternatively between 0.01% to 0.20% HFO-1234yf,
10 alternatively between 0.10% to 0.30% HFO-1234yf, alternatively between 0.20% to 0.40% HFO-1234yf, alternatively between 0.30% to 0.50% HFO-1234yf , alternatively between 0.30% to 0.40% HFO-1234yf, alternatively between 0.40% to 0.60% HFC-1234yf, alternatively between 0.50% to 0.70% HFO-1234yf, alternatively between 0.60% to 0.80% HFO-1234yf, alternatively between 0.70% to
15 0.70% HFO-1234yf, alternatively between 0.80% to 1.00% HFO-1234yf. In another embodiment, the foregoing product mixtures further comprises (on a mole basis) HFO-1225zc wherein the HFO-1225zc is present in an amount equal to 10% of the HFO-1234yf.

In some embodiments, the fluoropropene composition additionally optionally
20 comprises one or more of R-143a, R-152a, TFP, R-1233xf, R-1233zd(E), or R-1233zd(Z). In some embodiments, the sum total of the amounts of R-143a, R-152a, TFP, R-1233xf, R-1233zd(E), and R-1233zd(Z) is between 0.00001 mole percent and 2 mole percent, based on the total fluoropropene composition. In one embodiment, the fluoropropene composition includes R-1233zd(E) in an amount of
25 0.7 mole percent to 1.15 mole percent, based on the total heat transfer media. In one embodiment, the fluoropropene composition includes R-1233zd(Z) in an amount of 0.05 mole percent to 0.25 mole percent, based on the total heat transfer media. . In one embodiment, the fluoropropene composition includes HFO-1234zeZ in an amount of 0.05 mole percent to 0.25 mole percent, based on the total fluoropropene
30 composition. In one embodiment, the fluoropropene composition includes R-143a in an amount of 0.05 mole percent to 0.25 mole percent, based on the total fluoropropene composition.

In other embodiments, the fluoropropene composition optionally comprises one or more of 1224yd, 1224zc, 1326mxz, 113, 32, 23, trifluoro propyne, 356mff, 1326mxz, HFC-245fa and HFC-245cb. The amount of the foregoing compounds can range from about 0.001 to about 1 mole%, about 0.001 to about 0.9 and, in some cases, about 0.001 to about 0.7 mole%.

In one particular embodiment, the sum total of the amounts 1224yd, 1224zc, 1326mxz, 113, 32, 23, trifluoro propyne, 356mff, 1326mxz, HFC-245fa and HFC-245cb is between 0.001 mole percent and 2 mole percent, based on the total fluoropropene composition. The amount of the foregoing compounds can range from about 0.001 to about 0.1 mole%, about 0.001 to about 0.09 and, in some cases, about 0.001 to about 0.07 mole%.

In another particular embodiment, the inventive composition can comprise greater than about 99wt% HFO-1234ze(E) and, for example, 99.5 to 99.99, 99.6 to 99.9 and in some cases about 99.7 to 99.99wt% HFO-1234ze(E) with the remainder comprising HFO-1225zc and HFO-1234yf. The inventive compositions can also contain at least one additional compound selected from the group consisting of HFC-134a, 245cb, 236fa, 1225ye isomers (e.g., E-1225ye and Z-1225ye), HFO-1234ze isomer (e.g., HFO-1234ze(Z)), HFC-245fa, HFC-124, HCFC-114, trifluoropropyne, HFC-152a and HFO-1234 isomers including at least one member selected from the group consisting of HFO-1234zc, HFO-1234yc and HFO-1234ye. The total combined amount of HFO-1225zc, HFO-1234yf and the additional compound(s) can range from greater than 0 to less than about 1 wt.%, and for example, greater than 0 to 0.3, greater than 0 to 0.1 and in some cases greater than 0 to 0.01 mol%. A specific Example of the foregoing composition is shown in Table A below

Table A

Components	
134a	3.2 ppm
1225zc	1.5 ppm
1234yf	47 ppm
245cb	Coelute with 1234yf
236fa	1.1 ppm
E-1234ze	99.98 wt.%
E-1225ye	0.6 ppm

Components	
1234 isomers*	6.1 ppm
245fa	20.5 ppm
124	4.6 ppm
Z-1234ze	87 ppm
114	14 ppm
trifluoropropyne	1 ppm
152a	0.5 ppm
Z-1225ye	4 ppm

* includes HFO-1234zc, HFO-1234yc and HFO-1234ye

The compositions of the present invention may be prepared by any convenient method to combine the desired amounts of the individual components. A preferred method is to weigh the desired component amounts and thereafter combine the components in an appropriate vessel. Agitation may be used, if desired.

An alternative means for making compositions of the present invention may be a method for making a refrigerant blend composition, wherein said refrigerant blend composition comprises a composition as disclosed herein, said method comprising (i) reclaiming a volume of one or more components of a refrigerant composition from at least one refrigerant container, (ii) removing impurities sufficiently to enable reuse of said one or more of the reclaimed components, (iii) and optionally, combining all or part of said reclaimed volume of components with at least one additional refrigerant composition or component.

A refrigerant container may be any container in which is stored a refrigerant blend composition that has been used in a refrigeration apparatus, air-conditioning apparatus or heat pump apparatus. Said refrigerant container may be the refrigeration apparatus, air-conditioning apparatus or heat pump apparatus in which the refrigerant blend was used. Additionally, the refrigerant container may be a storage container for collecting reclaimed refrigerant blend components, including but not limited to pressurized gas cylinders.

Residual refrigerant means any amount of refrigerant blend or refrigerant blend component that may be moved out of the refrigerant container by any method known for transferring refrigerant blends or refrigerant blend components.

Impurities may be any component that is in the refrigerant blend or refrigerant blend component due to its use in a refrigeration apparatus, air-conditioning apparatus or heat pump apparatus. Such impurities include but are not limited to refrigeration lubricants, being those described earlier herein, particulates including but not limited to metal, metal salt or elastomer particles, that may have come out of the refrigeration apparatus, air-conditioning apparatus or heat pump apparatus, and any other contaminants that may adversely effect the performance of the refrigerant blend composition.

Such impurities may be removed sufficiently to allow reuse of the refrigerant blend or refrigerant blend component without adversely effecting the performance or equipment within which the refrigerant blend or refrigerant blend component will be used.

It may be necessary to provide additional refrigerant blend or refrigerant blend component to the residual refrigerant blend or refrigerant blend component in order to produce a composition that meets the specifications required for a given product. For instance, if a refrigerant blend has 3 components in a particular weight percentage range, it may be necessary to add one or more of the components in a given amount in order to restore the composition to within the specification limits.

Compositions of the present invention have zero or low ozone depletion potential and low global warming potential (GWP). Additionally, the compositions of the present invention will have global warming potentials that are less than many hydrofluorocarbon refrigerants currently in use. One aspect of the present invention is to provide a refrigerant with a global warming potential of less than 1000, less than 500, less than 150, less than 100, or less than 50. Another aspect of the present invention is to reduce the net GWP of refrigerant mixtures by adding fluoroolefins to said mixtures.

The compositions of the present invention may be useful as low global warming potential (GWP) replacements for currently used refrigerants, including but not limited to R134a (or HFC-134a, 1,1,1,2-tetrafluoroethane), R22 (or HCFC-22, chlorodifluoromethane), R123 (or HFC-123, 2,2-dichloro-1,1,1-trifluoroethane), R11 (CFC-11, fluorotrichloromethane), R12 (CFC-12,

dichlorodifluoromethane), R245fa (or HFC-245fa, 1,1,1,3,3-pentafluoropropane), R114 (or CFC-114, 1,2-dichloro-1,1,2,2-tetrafluoroethane), R236fa (or HFC-236fa, 1,1,1,3,3,3-hexafluoropropane), R124 (or HCFC-124, 2-chloro-1,1,1,2-tetrafluoroethane), R407C (ASHRAE designation for a blend of 52 weight percent R134a, 25 weight percent R125 (pentafluoroethane), and 23 weight percent R32 (difluoromethane), R410A (ASHRAE designation for a blend of 50 weight percent R125 and 50 weight percent R32), R417A, (ASHRAE designation for a blend of 46.6 weight percent R125, 50.0 weight percent R134a, and 3.4 weight percent n-butane), R422A (ASHRAE designation for a blend of 85.1 weight percent R125, 11.5 weight percent R134a, and 3.4 weight percent isobutane), R404A, (ASHRAE designation for a blend of 44 weight percent R125, 52 weight percent R143a (1,1,1-trifluoroethane), and 4.0 weight percent R134a) and R507A (ASHRAE designation for a blend of 50 weight percent R125 and 50 weight percent R143a). Additionally, the compositions of the present invention may be useful as replacements for R12 (CFC-12, dichlorodifluoromethane) or R502 (ASHRAE designation for a blend of 51.2 weight percent CFC-115 (chloropentafluoroethane) and 48.8 weight percent HCFC-22).

The fluoropropene composition may be useful in various applications. In an embodiment, the fluoropropene composition may be used as a refrigerant. In some embodiments, the fluoropropene composition may be used as a replacement for older generation refrigerants (e.g., R404A, R502) to provide a more environmentally friendly composition. In some embodiments, the fluoropropene composition may be a hydrofluoroolefin composition. In an embodiment, the fluoropropene composition includes from 99 mole percent to 99.99 mole percent of 1,3,3,3-tetrafluoropropene (HFO-1234ze)(E) and from 0.0001 mole percent to 1.0 mole percent of 1,1,3,3,3-pentafluoropropene (HFO-1225zc) and 2,3,3,3-tetrafluoropropene (HFO-1234yf). In another embodiment, the fluoropropene composition is a near azeotropic composition that is substantially free of HFO-1234ze(Z). By substantially free, it is meant that the fluoropropene composition contains less than about 1000 ppm, less than about 500 ppm and typically less than about 100 ppm, HFO-1234ze(Z).

Often replacement refrigerants are most useful if capable of being used in the original refrigeration equipment designed for a different refrigerant. The compositions of the present invention may be useful as replacements for the above-mentioned refrigerants in original equipment. Additionally, the compositions of the present invention may be useful as replacements for the above-mentioned refrigerants in equipment designed to use the above-mentioned refrigerants.

In one embodiment, the foregoing inventive fluoropropene compositions can be blended with other fluorochemicals. This embodiment of the present invention relates to a refrigerant composition comprising the inventive composition (e.g., HFO-1234ze(E), HFO-1225zc and HFO-1234yf) and at least one compound selected from the group consisting of: HFC-1234ye, HFC-1243zf, HFC-32, HFC-125, HFC-134, HFC-134a, HFC-143a, HFC-152a, HFC-161, HFC-227ea, HFC-236ea, HFC-236fa, HFC-245fa, HFC-365mfc, R1336mzz(E), propane, n-butane, isobutane, 2-methylbutane, n-pentane, cyclopentane, dimethylether, CF₃SCF₃, CO₂, CF₃I and combinations thereof.

In one embodiment, the foregoing inventive fluoropropene compositions are combined with at least one additional refrigerant comprising a member selected from the group consisting of R32, R125, R134, R134a, 227ea, and R1336mzz(E). The amount of the at least one additional refrigerant can range from about 5 to about 95, about 50 to about 90 and in some cases about 60 to about 80wt.% of the refrigerant composition. In one particular embodiment, the inventive fluoropropene compositions can be employed as a source of HFO-1234ze for preparing R444, R446A/B, R447B, R448A, R450A, R456, R459A/B, R460A/B/C, R464A, 515A and 515B.

In some embodiments, the foregoing fluoropropene compositions may be used in a refrigeration system. One embodiment of a refrigeration system includes an evaporator, a condenser, a compressor, an expansion device, and a heat transfer medium. The heat transfer medium includes the fluoropropene composition. That is, the compositions of the present invention may further comprise a lubricant. The heat transfer medium or the compositions of the present invention can further comprise at least one lubricant including those suitable for use with refrigeration or air-conditioning apparatus. Among these lubricants are those conventionally used

in compression refrigeration apparatus utilizing chlorofluorocarbon refrigerants. Such lubricants and their properties are discussed in the 1990 ASHRAE Handbook, Refrigeration Systems and Applications, chapter 8, titled "Lubricants in Refrigeration Systems", pages 8.1 through 8.21, herein incorporated by reference.

5 Lubricants of the present invention may comprise those commonly known as "mineral oils" in the field of compression refrigeration lubrication. Mineral oils comprise paraffins (i.e. straight-chain and branched-carbon-chain, saturated hydrocarbons), naphthenes (i.e. cyclic or ring structure saturated hydrocarbons, which may be paraffins) and aromatics (i.e. unsaturated, cyclic hydrocarbons containing one or more rings characterized by alternating double bonds).
 10 Lubricants of the present invention further comprise those commonly known as "synthetic oils" in the field of compression refrigeration lubrication. Synthetic oils comprise alkylaryls (i.e. linear and branched alkyl alkylbenzenes), synthetic paraffins and naphthenes, silicones, and poly-alpha-olefins. Representative
 15 conventional lubricants of the present invention are the commercially available BVM 100 N (paraffinic mineral oil sold by BVA Oils), naphthenic mineral oil commercially available under the trademark from Suniso[®] 3GS and Suniso[®] 5GS by Crompton Co., naphthenic mineral oil commercially available from Pennzoil under the trademark Sontex[®] 372LT, naphthenic mineral oil commercially
 20 available from Calumet Lubricants under the trademark Calumet[®] RO-30, linear alkylbenzenes commercially available from Shrieve Chemicals under the trademarks Zerol[®] 75, Zerol[®] 150 and Zerol[®] 500 and branched alkylbenzene, sold by Nippon Oil as HAB 22.

In one embodiment, the lubricant component can comprise those which have
 25 been designed for use with refrigerants and are miscible with the fluoropropene compositions of the present invention under compression refrigeration and air-conditioning apparatus' operating conditions. Such lubricants and their properties are discussed in "Synthetic Lubricants and High-Performance Fluids", R. L. Shubkin, editor, Marcel Dekker, 1993. Such lubricants include, but are not limited
 30 to, polyol esters (POEs) such as Castrol[®] 100 (Castrol, United Kingdom), polyalkylene glycols (PAGs) such as RL-488A from Dow (Dow Chemical, Midland, Michigan), and polyvinyl ethers (PVEs).

Lubricants of the present invention are selected by considering a given compressor's requirements and the environment to which the lubricant will be exposed. The amount of lubricant can range from about 1 to about 50, about 1 to about 20 and in some cases about 1 to about 3 weight percent of a refrigerant composition. In one particular embodiment, the foregoing refrigerant compositions are combined with a PAG lubricant for usage in an automotive A/C system having an internal combustion engine. In another particular embodiment, the foregoing refrigerant compositions are combined with a POE lubricant for usage in an automotive A/C system having an electric or hybrid electric drive train.

10 In one embodiment, a refrigerant composition comprises the inventive near azeotropic composition, at least one lubricant and at least one additive which can improve the refrigerant and air-conditioning system lifetime and compressor durability are desirable. In one aspect of the invention, the foregoing refrigerant compositions comprise at least one member selected from the group consisting of
15 acid scavengers, performance enhancers, and flame suppressants.

In another embodiment, the fluoropropene composition may be used in a process to transfer heat. The process may include providing an article and contacting the article with a heat transfer media including the fluoropropene composition. In some embodiments, the article may include electrical equipment
20 (e.g., circuit board, computer, display, semiconductor chip, or transformer), a heat transfer surface (e.g., heat sink), or article of clothing (e.g., a body suit).

In another embodiment, the fluoropropene composition may be used in a process for treating a surface. The process may include providing a surface having a treatable material deposited thereon and contacting the surface with a treatment
25 composition including the fluoropropene composition. In some embodiments, the treatment composition may substantially dissolve the treatable material.

In another embodiment, the fluoropropene composition may be used in a process for forming a composition. The process includes providing a solute and contacting the solute with a solvent including the fluoropropene composition. In
30 some embodiments, the fluoropropene composition may substantially dissolve the solute.

The present invention further relates to a chiller apparatus comprising the fluoropropene composition of the present invention.

The present invention further relates to the fluoropropene composition of the present invention for use in a chiller apparatus.

5 The chiller apparatus as referred to herein is preferably a low temperature chiller apparatus or a medium temperature chiller apparatus, more preferably a medium temperature chiller apparatus. Low temperature generally refers to an evaporator temperature of around or below about -17.8 °C (0°F), whereas medium temperature generally refers to evaporator temperature of around 0 °C to 10 °C or
10 below.

The present invention further relates to the use of the fluoropropene composition of the present invention in a chiller apparatus, preferably in a medium temperature chiller apparatus.

A chiller is a type of air conditioning/refrigeration apparatus. Chillers are
15 described for example in PCT application No. PCT/US2011/032072 (which corresponds to US 2013/025299; the disclosure of which is hereby incorporated by reference). The present disclosure is directed for example to a vapor compression chiller. Such vapor compression chillers may be either flooded evaporator chillers or direct expansion chillers. Both a flooded evaporator chiller and a direct
20 expansion chiller may be air-cooled or water-cooled. In the embodiment where chillers are water cooled, such chillers are generally associated with cooling towers for heat rejection from the system. In the embodiment where chillers are air-cooled, the chillers are equipped with refrigerant-to-air finned-tube condenser coils and fans to reject heat from the system. Air-cooled chiller systems are generally
25 less costly than equivalent-capacity water-cooled chiller systems including cooling tower and water pump. However, water-cooled systems can be more efficient under many operating conditions due to lower condensing temperatures.

Chillers, including both flooded evaporator and direct expansion chillers, may be coupled with an air handling and distribution system to provide comfort air
30 conditioning (cooling and dehumidifying the air) to large commercial buildings, including hotels, office buildings, hospitals, universities, etc. In another

embodiment, chillers, most likely air-cooled direct expansion chillers, have found additional utility in naval submarines and surface vessels.

To illustrate how chillers operate, reference is made to the Figures of PCT application No. PCT/US2011/032072. A water-cooled, flooded evaporator chiller is shown illustrated in Figure 1. In this chiller a first heat transfer medium, which is a warm liquid, which comprises water, and, in some embodiments, additives, such as a glycol (e.g., ethylene glycol or propylene glycol), enters the chiller from a cooling system, such as a building cooling system, shown entering at arrow 3, through a coil or tube bundle 9, in an evaporator 6, which has an inlet and an outlet.

The warm first heat transfer medium is delivered to the evaporator, where it is cooled by liquid refrigerant, which is shown in the lower portion of the evaporator. The liquid refrigerant evaporates at a temperature lower than the temperature of the warm first heat transfer medium which flows through coil 9. The cooled first heat transfer medium re-circulates back to the building cooling system, as shown by arrow 4, via a return portion of coil 9. The liquid refrigerant, shown in the lower portion of evaporator 6 in Figure 1, vaporizes and is drawn into a compressor 7, which increases the pressure and temperature of the refrigerant vapor. The compressor compresses this vapor so that it may be condensed in a condenser 5 at a higher pressure and temperature than the pressure and temperature of the refrigerant vapor when it comes out of the evaporator. A second heat transfer medium, which is a liquid in the case of a water-cooled chiller, enters the condenser via a coil or tube bundle 10 in condenser 5 from a cooling tower at arrow 1 in Figure 1. The second heat transfer medium is warmed in the process and returned via a return loop of coil 10 and arrow 2 to a cooling tower or to the environment. This second heat transfer medium cools the vapor in the condenser and causes the vapor to condense to liquid refrigerant, so that there is liquid refrigerant in the lower portion of the condenser as shown in Figure 1. The condensed liquid refrigerant in the condenser flows back to the evaporator through an expansion device 8, which may be an orifice, capillary tube or expansion valve. Expansion device 8 reduces the pressure of the liquid refrigerant, and converts the liquid refrigerant partially to vapor, that is to say that the liquid refrigerant flashes as pressure drops between the condenser and the evaporator. Flashing cools the refrigerant, i.e., both the liquid refrigerant and the refrigerant vapor to the saturation temperature at evaporator

pressure, so that both liquid refrigerant and refrigerant vapor are present in the evaporator.

It should be noted that for a single component refrigerant composition, the composition of the vapor refrigerant in the evaporator is the same as the composition of the liquid refrigerant in the evaporator. In this case, evaporation will occur at a constant temperature. However, if a refrigerant blend (or mixture) is used, as in the present invention, the liquid refrigerant and the refrigerant vapor in the evaporator (or in the condenser) may have different compositions. This may lead to inefficient systems and difficulties in servicing the equipment, thus a single component refrigerant is more desirable. An azeotrope or azeotrope-like composition will function essentially as a single component refrigerant in a chiller, such that the liquid composition and the vapor composition are essentially the same reducing any inefficiencies that might arise from the use of a non-azeotropic or non-azeotrope-like composition.

Chillers with cooling capacities above 700 kW generally employ flooded evaporators, where the refrigerant in the evaporator and the condenser surrounds a coil or tube bundle or other conduit for the heat transfer medium (i.e., the refrigerant is on the shell side). Flooded evaporators require larger charges of refrigerant, but permit closer approach temperatures and higher efficiencies.

Chillers with capacities below 700 kW commonly employ evaporators with refrigerant flowing inside the tubes and heat transfer medium in the evaporator and the condenser surrounding the tubes, i.e., the heat transfer medium is on the shell side. Such chillers are called direct-expansion (DX) chillers. One embodiment of a water-cooled direct expansion chiller is illustrated in Figure 2. In the chiller as illustrated in Figure 2, first liquid heat transfer medium, which is a warm liquid, such as warm water, enters an evaporator 6' at inlet 14. Mostly liquid refrigerant (with a small amount of refrigerant vapor) enters a coil or tube bundle 9' in the evaporator at arrow 3' and evaporates. As a result, first liquid heat transfer medium is cooled in the evaporator, and a cooled first liquid heat transfer medium exits the evaporator at outlet 16, and is sent to a body to be cooled, such as a building. In this embodiment of Figure 2, it is this cooled first liquid heat transfer medium that cools the building or other body to be cooled. The refrigerant vapor exits the

evaporator at arrow 4' and is sent to a compressor 7', where it is compressed and exits as high temperature, high pressure refrigerant vapor. This refrigerant vapor enters a condenser 5' through a condenser coil 10' or tube bundle at 1'. The refrigerant vapor is cooled by a second liquid heat transfer medium, such as water, in the condenser and becomes a liquid. The second liquid heat transfer medium enters the condenser through a condenser heat transfer medium inlet 20. The second liquid heat transfer medium extracts heat from the condensing refrigerant vapor, which becomes liquid refrigerant, and this warms the second liquid heat transfer medium in the condenser. The second liquid heat transfer medium exits through the condenser heat transfer medium outlet 18. The condensed refrigerant liquid exits the condenser through lower coil 10' as shown in Figure 2 and flows through an expansion device 12, which may be an orifice, capillary tube or expansion valve. Expansion device 12 reduces the pressure of the liquid refrigerant. A small amount of vapor, produced as a result of the expansion, enters the evaporator with liquid refrigerant through coil 9' and the cycle repeats.

Vapor-compression chillers may be identified by the type of compressor they employ. The present invention includes chillers utilizing centrifugal compressors as well as positive displacement compressors. In one embodiment, the compositions as disclosed herein are useful in chillers which utilizes a centrifugal compressor, herein referred to as a centrifugal chiller.

A centrifugal compressor uses rotating elements to accelerate the refrigerant radially, and typically includes an impeller and diffuser housed in a casing. Centrifugal compressors usually take fluid in at an impeller eye, or central inlet of a circulating impeller, and accelerate it radially outward. Some static pressure rise occurs in the impeller, but most of the pressure rise occurs in the diffuser section of the casing, where velocity is converted to static pressure. Each impeller-diffuser set is a stage of the compressor. Centrifugal compressors are built with from 1 to 12 or more stages, depending on the final pressure desired and the volume of refrigerant to be handled.

The pressure ratio, or compression ratio, of a compressor is the ratio of absolute discharge pressure to the absolute inlet pressure. Pressure delivered by a centrifugal compressor is practically constant over a relatively wide range of

capacities. The pressure a centrifugal compressor can develop depends on the tip speed of the impeller. Tip speed is the speed of the impeller measured at its outermost tip and is related to the diameter of the impeller and its revolutions per minute. The capacity of the centrifugal compressor is determined by the size of the passages through the impeller. This makes the size of the compressor more dependent on the pressure required than the capacity.

In another embodiment, the compositions as disclosed herein are useful in positive displacement chillers, which utilize positive displacement compressors, either reciprocating, screw, or scroll compressors. A chiller which utilizes a screw compressor will be hereinafter referred to as a screw chiller.

Positive displacement compressors draw vapor into a chamber, and the chamber decreases in volume to compress the vapor. After being compressed, the vapor is forced from the chamber by further decreasing the volume of the chamber to zero or nearly zero.

Reciprocating compressors use pistons driven by a crankshaft. They can be either stationary or portable, can be single or multi-staged, and can be driven by electric motors or internal combustion engines. Small reciprocating compressors from 5 to 30 hp are seen in automotive applications and are typically for intermittent duty. Larger reciprocating compressors up to 100 hp are found in large industrial applications. Discharge pressures can range from low pressure to very high pressure (>5000 psi or 35 MPa).

Screw compressors use two meshed rotating positive-displacement helical screws to force the gas into a smaller space. Screw compressors are usually for continuous operation in commercial and industrial application and may be either stationary or portable. Their application can be from 5 hp (3.7 kW) to over 500 hp (375 kW) and from low pressure to very high pressure (>1200 psi or 8.3 MPa).

Scroll compressors are similar to screw compressors and include two interleaved spiral-shaped scrolls to compress the gas. The output is more pulsed than that of a rotary screw compressor.

For chillers which use scroll compressors or reciprocating compressors, capacities below 150 kW, brazed-plate heat exchangers are commonly used for

evaporators instead of the shell-and-tube heat exchangers employed in larger chillers. Brazed-plate heat exchangers reduce system volume and refrigerant charge.

5 The present invention further relates to a process for producing cooling comprising evaporating the compositions of the present invention in the vicinity of a body to be cooled, and thereafter condensing said compositions.

The present invention further relates to a refrigeration, air-conditioning, or heat pump apparatus containing a composition of the present invention.

10 The present invention further relates to a mobile air-conditioning apparatus containing a composition of the present invention.

As used herein, mobile refrigeration apparatus or mobile air-conditioning apparatus refers to any refrigeration or air-conditioning apparatus incorporated into a transportation unit for the road, rail, sea or air. In addition, apparatus, which are meant to provide refrigeration or air-conditioning for a system independent of any
15 moving carrier, known as “intermodal” systems, are included in the present invention. Such intermodal systems include “containers” (combined sea/land transport) as well as “swap bodies” (combined road and rail transport). The present invention is particularly useful for road transport refrigerating or air-conditioning apparatus, such as automobile air-conditioning apparatus or refrigerated road
20 transport equipment.

In another embodiment, the present invention relates to blowing agent compositions comprising the fluoroolefin-containing compositions (e.g., near azeotropic containing compositions), as described herein for use in preparing foams. In other embodiments the invention provides foamable compositions, and
25 preferably polyurethane and polyisocyanate foam compositions, and method of preparing foams. In such foam embodiments, one or more of the present fluoroolefin-containing compositions are included as a blowing agent in foamable compositions, which composition preferably includes one or more additional components capable of reacting and foaming under the proper conditions to form a
30 foam or cellular structure. Any of the methods well known in the art, such as those described in “Polyurethanes Chemistry and Technology,” Volumes I and II,

Saunders and Frisch, 1962, John Wiley and Sons, New York, N.Y., which is incorporated herein by reference, may be used or adapted for use in accordance with the foam embodiments of the present invention.

The present invention further relates to a method of forming a foam
5 comprising: (a) adding to a foamable composition a fluoroolefin-containing composition of the present invention; and (b) reacting the foamable composition under conditions effective to form a foam.

Another embodiment of the present invention relates to the use of the fluoroolefin-containing compositions as described herein (e.g., compositions of
10 HFO-1234ze(E), HFO-1225zc and HFO-1234yf, which may be near-azeotropic), for use as propellants in sprayable compositions. Additionally, the present invention relates to a sprayable composition comprising the fluoroolefin-containing compositions as described herein. The active ingredient to be sprayed together with inert ingredients, solvents and other materials may also be present in a sprayable
15 composition. Preferably, the sprayable composition is an aerosol. Suitable active materials to be sprayed include, without limitations, cosmetic materials, such as deodorants, perfumes, hair sprays, cleaners, and polishing agents as well as medicinal materials such as anti-asthma and anti-halitosis medications.

The present invention further relates to a process for producing aerosol
20 products comprising the step of adding a fluoroolefin-containing composition as described herein to active ingredients in an aerosol container, wherein said composition functions as a propellant.

As used herein, the terms “comprises,” “comprising,” “includes,” “including,” “has,” “having” or any other variation thereof, are intended to cover a non-
25 exclusive inclusion. For example, a process, method, article, or apparatus that comprises a list of elements is not necessarily limited to only those elements but may include other elements not expressly listed or inherent to such process, method, article, or apparatus. Further, unless expressly stated to the contrary, “or” refers to an inclusive or and not to an exclusive or. For example, a condition A or B is satisfied by any one of the following: A is true (or present) and B is false (or not
30 present), A is false (or not present) and B is true (or present), and both A and B are true (or present).

The transitional phrase “consisting of” excludes any element, step, or ingredient not specified. If in the claim, such would close the claim to the inclusion of materials other than those recited except for impurities ordinarily associated therewith. When the phrase “consists of” appears in a clause of the body of a claim, rather than immediately following the preamble, it limits only the element set forth in that clause; other elements are not excluded from the claim as a whole. The transitional phrase “consisting essentially of” is used to define a composition, method that includes materials, steps, features, components, or elements, in addition to those literally disclosed provided that these additional included materials, steps, features, components, or elements do not materially affect the basic and novel characteristic(s) of the claimed invention, especially the mode of action to achieve the desired result of any of the processes of the present invention. The term ‘consisting essentially of’ occupies a middle ground between “comprising” and ‘consisting of’.

In the foregoing combinations of inventive embodiments, the compositions can comprise, consist essentially of or consist of HFO-1234ze(E), HFO-1225zc and HFO-1234yf.

Also, use of “a” or “an” are employed to describe elements and components described herein. This is done merely for convenience and to give a general sense of the scope of the invention. This description should be read to include one or at least one and the singular also includes the plural unless it is obvious that it is meant otherwise.

Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Although methods and materials similar or equivalent to those described herein can be used in the practice or testing of embodiments of the present invention, suitable methods and materials are described below. All publications, patent applications, patents, and other references mentioned herein are incorporated by reference in their entirety, unless a particular passage is cited. In case of conflict, the present specification, including definitions, will control. In addition, the materials, methods, and examples are illustrative only and not intended to be limiting.

EXAMPLES

The concepts described herein will be further described in the following examples, which do not limit the scope of the invention described in the claims.

Example 1

5 Table 1 shows the cooling performance of various refrigerant compositions comprising HFO-1234ze(E), HFO-1225zc and HFO-1234yf, by comparing cooling capacity (CAP_c) and energy efficiency (COP_c) to HFO-1234ze(E). The data are based on the following conditions.

10 Condenser temperature = 47.0°C
 Evaporator temperature = 7.0°C
 Subcool = 12.0 K
 Superheat = 3.0 K
 Compressor efficiency = 0.7

15 Average Heat Exchanger Temperature Set Points
 Superheat is included in refrigeration effect

 Cooling load = 3.517 kW

20 Compressor displacement = 0.00283168438736 (m³ / min)

Table 1

Composition (mass ratio)	CAP _c (kJ/m ³)	COP _c
R-1234zeE	2111.4	4.402
R-1234yf_R-1225zc_R-1234zeE W=_0.0_0.005_0.995	2112.4	4.402
R-1234yf_R-1225zc_R-1234zeE W=_0.00125_0.00375_0.995	2113.3	4.402
R-1234yf_R-1225zc_R-1234zeE W=_0.0025_0.0025_0.995	2114.2	4.402
R-1234yf_R-1225zc_R-1234zeE W=_0.00375_0.00125_0.995	2115.2	4.402
R-1234yf_R-1225zc_R-1234zeE W=_0.005_0.0_0.995	2116.1	4.402
R-515B	2100.4	4.393

Composition (mass ratio)	CAP _c (kJ/m ³)	COP _c
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0_0.00456_0.90644_0.089	2104.6	4.392
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00114_0.00342_0.90644_0.089	2103.7	4.392
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00228_0.00228_0.90644_0.089	2102.9	4.392
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00342_0.00114_0.90644_0.089	2102.1	4.392
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00456_0.0_0.90644_0.089	2101.3	4.393
R-515A	2096.1	4.389
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0_0.0044_0.8756_0.12	2100.1	4.389
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0011_0.0033_0.8756_0.12	2099.3	4.389
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0022_0.0022_0.8756_0.12	2098.5	4.389
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0033_0.0011_0.8756_0.12	2097.7	4.389
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0044_0.0_0.8756_0.12	2096.9	4.389
R-450A	2465.1	4.394
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.0_0.0029_0.5771_0.42	2467.7	4.394
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.00073_0.00218_0.5771_0.42	2467.2	4.394
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.00145_0.00145_0.5771_0.42	2466.8	4.394
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.00218_0.00073_0.5771_0.42	2466.3	4.394
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.0029_0.0_0.5771_0.42	2465.9	4.394
Refrigerant A (R-1234zeE_R-134 0.63 0.37)	2276.7	4.421
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.0_0.00315_0.62685_0.37	2279.3	4.421
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00079_0.00236_0.62685_0.37	2278.9	4.421

Composition (mass ratio)	CAP_c (kJ/m ³)	COP_c
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00157_0.00157_0.62685_0.37	2278.5	4.421
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00236_0.00079_0.62685_0.37	2278.1	4.421
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00315_0.0_0.62685_0.37	2277.7	4.421
Refrigerant B (R-1234zeE_R-1336mzzE_R-227ea W= 0.787 0.17_0.043)	1885.4	4.409
R-1225zc_R-1234yf_R-1234zeE_R-1336mzzE_R-227ea W=_0.0_0.00393_0.78307_0.17_0.043	1888.8	4.408
R-1225zc_R-1234yf_R-1234zeE_R-1336mzzE_R- 227ea_W=_0.00098_0.00295_0.78307_0.17_0.043	1888.1	4.408
R-1225zc_R-1234yf_R-1234zeE_R-1336mzzE_R- 227ea_W=_0.00197_0.00197_0.78307_0.17_0.043	1887.4	4.408
R-1225zc_R-1234yf_R-1234zeE_R-1336mzzE_R- 227ea_W=_0.00295_0.00098_0.78307_0.17_0.043	1886.7	4.408
R-1225zc_R-1234yf_R-1234zeE_R-1336mzzE_R-227ea W=_0.00393_0.0_0.78307_0.17_0.043	1886.1	4.408
R-448A	4718.7	4.214
R-1225zc_R-1234yf_R-1234zeE_R-125_R-134a_R-32 W=_0.0_0.20035_0.06965_0.26_0.21_0.26	4719	4.214
R-1225zc_R-1234yf_R-1234zeE_R-125_R-134a_R-32 W=_9e-05_0.20026_0.06965_0.26_0.21_0.26	4719	4.214
R-1225zc_R-1234yf_R-1234zeE_R-125_R-134a_R-32 W=_0.00017_0.20017_0.06965_0.26_0.21_0.26	4719	4.214
R-1225zc_R-1234yf_R-1234zeE_R-125_R-134a_R-32 W=_0.00026_0.20009_0.06965_0.26_0.21_0.26	4719	4.214
R-1225zc_R-1234yf_R-1234zeE_R-125_R-134a_R-32 W=_0.00035_0.2_0.06965_0.26_0.21_0.26	4718.9	4.214

Several compositions, and in particular the composition comprising HFO-1234ze(E), HFO-1225zc and HFO-1234yf, have higher capacity than pure HFO-1234ze(E) or others such as R-515B and R-450A. Energy efficiency (COP) for the present compositions in many cases is also similar to HFO-1234ze(E)

- 5 compositions indicating these could be replacement refrigerants for HFO-1234ze(E) or for others such as R-515B and R-450A in refrigeration and air-conditioning apparatus applications, and in stationary air-conditioning applications in particular.

10 **Example 2**

Table 2 contains data on the impact of vapor leakage for various refrigerant compositions comprising HFO-1234ze(E), HFO-1225zc and HFO-1234yf, by comparing to HFO-1234ze(E).

- The following conditions were applied: A vessel is charged with an initial composition at a temperature of 25 °C, and the initial vapor pressure of the composition is measured. The composition is allowed to leak from the vessel, while the temperature is held constant, until 50 weight percent of the initial composition is removed, at which time the vapor pressure of the composition remaining in the vessel is measured. Delta P% gives the relative pressure difference in %.

Table 2

Composition (mass ratio)	delta P% 50% mass leak at 25°C
R-1234zeE	0.000
R-1234yf_R-1225zc_R-1234zeE W=_0.0_0.005_0.995	-0.023
R-1234yf_R-1225zc_R-1234zeE W=_0.00125_0.00375_0.995	-0.089
R-1234yf_R-1225zc_R-1234zeE W=_0.0025_0.0025_0.995	-0.158
R-1234yf_R-1225zc_R-1234zeE W=_0.00375_0.00125_0.995	-0.220

Composition (mass ratio)	delta P% 50% mass leak at 25°C
R-1234yf_R-1225zc_R-1234zeE W=_0.005_0.0_0.995	-0.285
R-515B	-0.001
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0_0.00456_0.90644_0.089	-0.255
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00114_0.00342_0.90644_0.089	-0.197
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00228_0.00228_0.90644_0.089	-0.139
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00342_0.00114_0.90644_0.089	-0.080
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00456_0.0_0.90644_0.089	-0.021
R-515A	-0.003
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0_0.0044_0.8756_0.12	-0.245
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0011_0.0033_0.8756_0.12	-0.190
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0022_0.0022_0.8756_0.12	-0.134
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0033_0.0011_0.8756_0.12	-0.078
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0044_0.0_0.8756_0.12	-0.022
R-450A	-1.86
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.0_0.0029_0.5771_0.42	-1.92
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.00073_0.00218_0.5771_0.42	-1.90
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.00145_0.00145_0.5771_0.42	-1.89
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.00218_0.00073_0.5771_0.42	-1.87
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.0029_0.0_0.5771_0.42	-1.86
Refrigerant A (R-1234ze E_R-134 0.63 0.37)	-0.084
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.0_0.00315_0.62685_0.37	-0.199

Composition (mass ratio)	delta P% 50% mass leak at 25°C
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00079_0.00236_0.62685_0.37	-0.175
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00157_0.00157_0.62685_0.37	-0.151
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00236_0.00079_0.62685_0.37	-0.127
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00315_0.0_0.62685_0.37	-0.103
Refrigerant B (R-1234zeE_R-1336mzzE_R-227ea W= 0.787 0.17_0.043)	-5.82
R-1225zc_R-1234yf_R-1234zeE_R-1336mzzE_R-227ea W=_0.0_0.00393_0.78307_0.17_0.043	-6.10
R-1225zc_R-1234yf_R-1234zeE_R-1336mzzE_R- 227ea_W=_0.00098_0.00295_0.78307_0.17_0.043	-6.04
R-1225zc_R-1234yf_R-1234zeE_R-1336mzzE_R- 227ea_W=_0.00197_0.00197_0.78307_0.17_0.043	-5.98
R-1225zc_R-1234yf_R-1234zeE_R-1336mzzE_R- 227ea_W=_0.00295_0.00098_0.78307_0.17_0.043	-5.91
R-1225zc_R-1234yf_R-1234zeE_R-1336mzzE_R-227ea W=_0.00393_0.0_0.78307_0.17_0.043	-5.85
R-448A	-14.4
R-1225zc_R-1234yf_R-1234zeE_R-125_R-134a_R-32 W=_0.0_0.20035_0.06965_0.26_0.21_0.26	-14.3
R-1225zc_R-1234yf_R-1234zeE_R-125_R-134a_R-32 W=_9e-05_0.20026_0.06965_0.26_0.21_0.26	-14.3
R-1225zc_R-1234yf_R-1234zeE_R-125_R-134a_R-32 W=_0.00017_0.20017_0.06965_0.26_0.21_0.26	-14.3
R-1225zc_R-1234yf_R-1234zeE_R-125_R-134a_R-32 W=_0.00026_0.20009_0.06965_0.26_0.21_0.26	-14.4
R-1225zc_R-1234yf_R-1234zeE_R-125_R-134a_R-32 W=_0.00035_0.2_0.06965_0.26_0.21_0.26	-14.4

The difference in vapor pressure between the original composition and the composition remaining after 50 weight percent is removed is less than about 10 percent for several compositions of the present invention. This indicates that

several of the compositions of the present invention would be azeotropic or near-azeotropic.

Example 3

Table 3 shows the cooling performance of an inventive refrigerant composition under typical medium temperature chiller apparatus conditions. The data are based on the following conditions.

Condenser temperature = 37.78 °C
 Evaporator temperature = 4.44 °C
 Superheat = 6.0 K
 Compressor efficiency = 0.85

Average Heat Exchanger Temperature Set Points
 100 % of superheat is included in refrigeration effect

Vapor Molar Quality Entering Evaporator: q_4
 cooling load = 1.00004194 kW
 compressor displacement = 0.1 (m³ / min)

Also included are GWP values for an inventive composition compared to HFO-1234ze(E).

Table 3

Composition (mass ratio)	CAP _c (kJ/m ³)	COP _c	GWP (AR 5)
R-1234zeE	1870	5.928	1
R-1234yf_R-1225zc_R-1234zeE W=_0.0_0.005_0.995	1874.1	5.926	1
R-1234yf_R-1225zc_R-1234zeE W=_0.00125_0.00375_0.995	1873.3	5.926	1
R-1234yf_R-1225zc_R-1234zeE W=_0.0025_0.0025_0.995	1872.5	5.927	1
R-1234yf_R-1225zc_R-1234zeE W=_0.00375_0.00125_0.995	1871.7	5.927	1
R-1234yf_R-1225zc_R-1234zeE W=_0.005_0.0_0.995	1870.9	5.927	1

As apparent, the composition comprising HFO-1234ze(E), HFO-1225zc and HFO-1234yf, has higher capacity than pure HFO-1234ze(E), but still maintains the very low GWP of HFO-1234ze(E). Energy efficiency (COP) is also comparable to HFO-1234ze(E) indicating the inventive compositions could be good replacement refrigerants for HFO-1234ze(E) in medium temperature chiller apparatus applications.

Example 4

According to ASHRAE Standard, heat of combustion is a measure for the influence of

compounds on flammability properties of materials. The results for various refrigerant compositions comprising HFO-1234ze(E), HFO-1225zc and HFO-1234yf, compared to HFO-1234ze(E) are given in Table 4.

Table 4

Composition (mass ratio)	Heat of Combustion (kJ/gm)
R-1234zeE	-10.2554
R-1234yf_R-1225zc_R-1234zeE W=_0.0_0.005_0.995	-10.2383
R-1234yf_R-1225zc_R-1234zeE W=_0.00125_0.00375_0.995	-10.2426
R-1234yf_R-1225zc_R-1234zeE W=_0.0025_0.0025_0.995	-10.2470
R-1234yf_R-1225zc_R-1234zeE W=_0.00375_0.00125_0.995	-10.2513
R-1234yf_R-1225zc_R-1234zeE W=_0.005_0.0_0.995	-10.2556
R-515B	
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.0_0.00456_0.90644_0.089	-9.6321
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00114_0.00342_0.90644_0.089	-9.6281
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00228_0.00228_0.90644_0.089	-9.6241

Composition (mass ratio)	Heat of Combustion (kJ/gm)
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00342_0.00114_0.90644_0.089	-9.6202
R-1225zc_R-1234yf_R-1234zeE_R-227ea W=_0.00456_0.0_0.90644_0.089	-9.6162
R-450A	
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.0_0.0029_0.5771_0.42	-8.6689
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.00073_0.00218_0.5771_0.42	-8.6664
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.00145_0.00145_0.5771_0.42	-8.6639
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.00218_0.00073_0.5771_0.42	-8.6614
R-1225zc_R-1234yf_R-1234zeE_R-134a W=_0.0029_0.0_0.5771_0.42	-8.6588
Refrigerant A (R-1234zeE_R-134 0.63 0.37)	
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.0_0.00315_0.62685_0.37	-8.9919
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00079_0.00236_0.62685_0.37	-8.9891
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00157_0.00157_0.62685_0.37	-8.9864
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00236_0.00079_0.62685_0.37	-8.9837
R-1225zc_R-1234yf_R-1234zeE_R-134 W=_0.00315_0.0_0.62685_0.37	-8.9809

The results indicate that the inventive compositions with 1225zc all are less flammable than the corresponding control. Therefore, compositions comprising HFO-1234ze(E), HFO-1225zc and HFO-1234yf are preferred in use.

Example 5

Table 5 discloses the reaction products of the dehydrofluorination of 245fa over various catalysts in the presence of Z-HFC-1234ze (in mol%).

Table 5

Catalyst	Others*	143a	152a	TFP	1234yf	1233xf
JM 62-2	0.15%	0.13%	0.00%	0.01%	0.35%	0.03%
LV	0.28%	0.14%	0.03%	0.02%	0.04%	0.00%
JM-62-3	0.28%	0.14%	0.02%	0.02%	0.24%	0.04%
Newport-Chrome	0.12%	0.13%	0.00%	0.00%	0.92%	0.00%
Catalyst	E-1233zd	Z-1233zd	Z-1234ze	E-1234ze	E+Z-1234ze	
JM 62-2	0.88%	0.13%	11.17%	87.13%	98.3%	
LV	1.03%	0.15%	10.9 %	87.4 %	98.3%	
JM-62-3	0.92%	0.14%	11 %	87.2 %	98.2%	
Newport-Chrome	0.92%	0.11%	10.5 %	87.3 %	97.8%	

*comprises 236fa, 1225zc, 1225ye (E and/or Z), and 1234zc

5

An inconel tube (½ inch OD) was filled with 10 cc (8 gm) of catalyst (see Table 5). After charging the reactor tube, the temperature of the catalyst bed was raised to 300 °C and purged with nitrogen (30 cc/min) for 200 minutes. Then the flow of nitrogen was reduced to 60 cc/min and HF was fed at 20 cc/min for 60 minutes. The temperature was increase to 325 °C for 300 minutes. The flow of nitrogen was then lowered to 30 cc/min and the flow of HF was raised to 30 cc/min for 30 minutes. The flow of nitrogen was then lowered to 12 cc/min and the flow of HF was raised to 48 cc/min for 60 minutes. The flow of nitrogen was then discontinued, and the flow of HF was raised to 48 cc/min for 30 minutes. The reactor temperature was then decreased to 250 °C for 30 minutes. Afterwards HF was turned off and the reactor was purged with 30 cc/min of nitrogen. The reactor temperature was then stabilized at 300 °C, the flow of nitrogen was turned off, and either CF₃CH₂CHF₂, or CF₃CH₂CHF₂ with 10.5-11% of Z-1234ze, was fed at 1.44 ml/hr. Contact time in the reactor was 45 seconds. The CF₃CH₂CHF₂ was vaporized at 50 °C. Part of the reactor effluent was passed through a series of valves and analyzed by GCMS. Amounts for Z-1234ze, 134a, 152b, TFP, 1234yf, 1233xf, E-1233zd, Z-1233zd and E + Z-1234ze are expressed as mole percent.

20

Results are summarized in Table 5. A grab sample was also taken for off-line GCMS analysis. It was surprisingly detected that the Others of Table 5 contained 236fa, 1225zc, 1225ye (E and Z), and 1234zc.

While any GCMS equipment, method and parameters, which are suitable for detecting the compounds that may be present in the inventive compositions, can be employed, one suitable method uses a single RTX-1 column.

Example 6

An inconel tube (1/2 inch {13 mm} OD) was filled with 5 cc (3.9 gm) of Cr₂O₃ catalyst (Louisville Cr) which was activated as described in Example 5. After activation, the flow of nitrogen was turned off, and the reactor temperature was set to 400°C. A flow of air (4 vol% O₂) and either CF₃CH₂CHF₂ (245fa alone), or CF₃CH₂CHF₂ with 13.3 mole % (corresponding to 11.5 wt%) of Z-1234ze, was fed at 0.67 ml/hr. Contact time in the reactor was 38 seconds. The CF₃CH₂CHF₂ was vaporized at 50°C. Part of the reactor effluent was passed through a series of valves and analyzed by GCMS. After 72 hours the flow of oxygen was stopped and the reactions were continued for another 72 hours in the absence of an oxygen containing gas. The catalysts started deactivating at a total time of about 120 hours for the reaction run with 245fa alone and at a total time of about 136 hours for the reaction run with 245fa containing Z-1234ze. Results are summarized in the following table with the amounts for Z-1234ze, 245fa and E-1234ze being expressed as mole percent:

Table 6

% Z-ze added	0	0	13.3	13.3
oxygen	yes	no	yes	no
Incoming composition	100/0	100/0	86.7/13.3	86.7/13.3
245fa conversion (%)	96.1	89.7	95.8	89.5
Z-ze in product (%)	22.9	21.1	22.7	20.8
% recovered 245fa	3.9	10.4	3.8	9.3
% E-ze	73.2	68.6	73.5	69.9
% yield E-ze	73.2	60.2	85.0	80.9
% selectivity E-ze	76.2	76.4	88.7	90.2

Note that not all the activities described above in the general description or the examples are required, that a portion of a specific activity may not be required, and that one or more further activities may be performed in addition to those described. Still further, the order in which activities are listed are not necessarily
5 the order in which they are performed.

Benefits, other advantages, and solutions to problems have been described above with regard to specific embodiments. However, the benefits, advantages, solutions to problems, and any feature(s) that may cause any benefit, advantage, or solution to occur or become more pronounced are not to be construed as a critical,
10 required, or essential feature of any or all the claims.

It is to be appreciated that certain features are, for clarity, described herein in the context of separate embodiments, may also be provided in combination in a single embodiment. Conversely, various features that are, for brevity, described in the context of a single embodiment, may also be provided separately or in any
15 subcombination. Further, reference to values stated in ranges include each and every value within that range.

While the invention has been described with reference to a preferred embodiment, it will be understood by those skilled in the art that various changes may be made, and equivalents may be substituted for elements thereof without
20 departing from the scope of the invention. In addition, many modifications may be made to adapt a particular situation or material to the teachings of the invention without departing from the essential scope thereof. Therefore, it is intended that the invention not be limited to the particular embodiment disclosed as the best mode contemplated for carrying out this invention, but that the invention will include all
25 embodiments falling within the scope of the appended claims.

Claims

1. A fluoropropene composition comprising E-1,3,3,3-tetrafluoropropene,
1,1,3,3,3-pentafluoropropene, and 2,3,3,3-tetrafluoropropene,
5 wherein the total amount of 1,1,3,3,3-pentafluoropropene and 2,3,3,3-
tetrafluoropropene is 1.0 wt.% or less, based on the total weight of the
fluoropropene composition.
2. The composition of claim 1, wherein the 2,3,3,3-tetrafluoropropene is present
in the fluoropropene composition in an amount of 0.0001 to 0.9 wt.%,
10 preferably in an amount of 0.001 to 0.8 wt.%, more preferably in an amount
of 0.1 to 0.6 wt.%, and most preferably in an amount of 0.3 to 0.4 wt.% based
on the total weight of the fluoropropene composition.
3. The composition of claim 1 or 2, wherein the 1,1,3,3,3-pentafluoropropene is
present in the fluoropropene composition in an amount of 0.0001 to 0.9 wt.%,
15 preferably in an amount of 0.001 to 0.8 wt.%, more preferably in an amount
of 0.01 to 0.6 wt.%, and most preferably in an amount of 0.1 to 0.4 wt.%
based on the total weight of the fluoropropene composition.
4. The composition of claim 1, 2 or 3, wherein the E-1,3,3,3-tetrafluoropropene
is present in the fluoropropene composition in an amount of 50.0 wt.% or
20 more, preferably in an amount of 75.0 wt.% or more, more preferably in an
amount of 99.0 wt.% or more, even more preferably in an amount of 99.5
wt.% or more, and most preferably in an amount of 99.8 wt.% or more, based
on the total weight of the fluoropropene composition.
5. The composition of claim 1, 2, 3 or 4, wherein the total amount of 2,3,3,3-
25 tetrafluoropropene and 1,1,3,3,3-pentafluoropropene in the fluoropropene
composition is 0.001 to 0.9 wt.%, preferably is 0.1 to 0.8 wt.%, and most
preferably is 0.3 to 0.5 wt.% based on the total weight of the fluoropropene
composition.
6. The composition of claim 1, 2, 3, 4 or 5, additionally comprising R-134,
30 preferably in an amount of 1.0 to 40.0 wt.%, more preferably in an amount of

30.0 to 40.0 wt.%, and most preferably in an amount of 35.0 to 40.0 wt.% based on the total weight of the fluoropropene composition.

7. The composition of claim 1, 2, 3, 4 or 5, additionally comprising R-1336mzzE and/or R-227ea, preferably in an amount of 15.0 to 20.0 wt.% of R-1336mzzE and of 2.0 to 5.0 wt.% of R-227ea based on the total weight of the fluoropropene composition.
8. The composition of any one of claims 1 to 7, wherein the composition is a near-azeotropic composition.
9. A method of producing a mixture of 1,3,3,3-tetrafluoropropene and 2,3,3,3-tetrafluoropropene, comprising:
 - (a) contacting a mixture of 1,1,1,3,3-pentafluoropropane and Z-1,3,3,3-tetrafluoropropene in the gas phase with a catalyst comprising at least one catalyst selected from the group consisting of fluorinated Cr_2O_3 and Cr/Ni on fluorinated alumina, optionally in the presence of an oxygen containing gas, to form a mixture comprising Z-1,3,3,3-tetrafluoropropene, E-1,3,3,3-tetrafluoropropene, 1,1,3,3,3-pentafluoropropene, 2,3,3,3-tetrafluoropropene, hydrogen fluoride, and optionally unreacted 1,1,1,3,3-pentafluoropropane, wherein the mixture comprises 0.00001 to 1.0 wt.% 2,3,3,3-tetrafluoropropene and greater than 0 and less than 1.0 wt.% 1,1,3,3,3-pentafluoropropene.
10. The method of claim 9, wherein said mixture of 1,1,1,3,3-pentafluoropropane and 1,3,3,3-tetrafluoropropene comprises at least 7 wt.% Z-1,3,3,3-tetrafluoropropene, preferably at least 10 wt.% Z-1,3,3,3-tetrafluoropropene.
11. The method of claim 9 or 10, wherein at least 94%, preferably at least 98% of the 1,1,1,3,3-pentafluoropropane is converted to E-1,3,3,3-tetrafluoropropene.
12. The method of claim 9, 10 or 11, further comprising
 - (b) recovering Z-1,3,3,3-tetrafluoropropene, or a mixture of Z-1,3,3,3-tetrafluoropropene and 1,1,1,3,3-pentafluoropropane, and recycling Z-1,3,3,3-tetrafluoropropene, or a mixture of Z-1,3,3,3-tetrafluoropropene and 1,1,1,3,3-pentafluoropropane back to step (a).

13. The method of any one of claims 9 to 12, wherein said hydrogen fluoride produced in step (a) is separated and recovered.
14. The method of any one of claims 9 to 13, wherein said oxygen containing gas is oxygen or air.
- 5 15. The method of any one of claims 9 to 14, wherein said mixture of 1,1,1,3,3-pentafluoropropane and 1,3,3,3-tetrafluoropropene comprises 0.1 to 0.8 wt.% 2,3,3,3-tetrafluoropropene, preferably 0.2 to 0.6 wt.% 2,3,3,3-tetrafluoropropene, more preferably 0.3 to 0.4 wt.% 2,3,3,3-tetrafluoropropene.
- 10 16. A fluoropropene composition obtained by the method of any one of claims 9 to 15.
17. The fluoropropene composition of any one of claims 1 to 8 or 16, further comprising a lubricant selected from the group consisting of mineral oils, such as paraffins, naphthenes and aromatics, alkylaryls, such as linear and

15 branched alkyl alkylbenzenes, synthetic paraffins, synthetic naphthenes, poly(alphaolefins), polyol esters, polyalkylene glycols, and polyvinyl ethers.
18. A process for transferring heat, comprising:

providing an article; and

contacting the article with a heat transfer media;

20 wherein the heat transfer media comprises the fluoropropene composition of any one of claims 1 to 8, 16 or 17.
19. A process for treating a surface, comprising:

providing a surface; and

contacting the surface with a treatment composition;

25 wherein the surface includes a treatable material deposited thereon; and

wherein the treatment composition comprises the fluoropropene composition of any one of claims 1 to 8, or 16.
20. The process of claim 19, wherein the treatment composition substantially dissolves the treatable material.

21. A process for forming a composition comprising:
- providing a solute;
 - contacting the solute with a solvent;
- wherein the solvent comprises the fluoropropene composition of any one of
5 claims 1 to 8, or 16.
22. A refrigeration system, comprising:
- an evaporator; a condenser; a compressor; an expansion device; and a heat transfer medium;
- wherein the heat transfer medium comprises the fluoropropene composition of
10 any one of claims 1 to 8, 16 or 17.
23. A chiller apparatus comprising the fluoropropene composition of any one of claims 1 to 8, 16 or 17.
24. The fluoropropene composition of any one of claims 1 to 8, 16 or 17 for use in a chiller apparatus, preferably in a medium temperature chiller apparatus.
- 15 25. Use of the fluoropropene composition of any one of claims 1 to 8, 16 or 17 in a chiller apparatus, preferably in a medium temperature chiller apparatus.
26. A method of producing cooling, the method comprising: evaporating the composition of any one of claims 1 to 8, 16 or 17 in the vicinity of a body to be cooled and thereafter condensing the composition.
- 20 27. A method for replacing a high GWP refrigerant in a refrigeration, air conditioning, or heat pump apparatus, wherein the high GWP refrigerant is selected from the group consisting of R134a, R22, R123, R11, R245fa, R114, R236fa, R124, R12, R410A, R407C, R417A, R422A, R507A, R502, and R404A, the method comprising providing the composition of any one of
25 claims 1 to 8, 16 or 17 to the refrigeration, air conditioning, or heat pump apparatus that uses, used or is designed to use the high GWP refrigerant.

28. A refrigeration apparatus, air-conditioning apparatus, or heat pump apparatus containing the composition of any one of claims 1 to 8, 16 or 17, wherein the refrigeration apparatus preferably is a stationary or mobile refrigeration apparatus and the air-conditioning apparatus preferably is a mobile air-conditioning apparatus, more preferably an automobile air-conditioning apparatus.
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