

TITLE

Vapor Compression Utilizing Ionic Liquid as Compressor Lubricant

5 This application claims the benefit of U.S. Provisional Application No. 60/809,622, filed May 31, 2006, which is incorporated in its entirety as a part hereof for all purposes.

10 Technical Field

 This invention relates to the operation or execution of a vapor compression cycle using a compressor, wherein at least one ionic liquid is provided as a lubricant for the compressor.

15

Background

 Many refrigeration, heating and air-cooling and air-heating systems operate using a vapor compression cycle. This cycle consists of compressing refrigerant gas from a low pressure to a high pressure. The high-pressure gas
20 exits a device used for increasing the pressure of the refrigerant, typically a compressor, and is condensed to a liquid, typically by a heat exchanger called a condenser, where heat is removed from the refrigerant. The high-pressure liquid refrigerant is lowered in pressure through an expansion device (valve or capillary tube) and the refrigerant liquid expands to a gas through a second heat exchanger
25 called an evaporator where heat is absorbed by the refrigerant. This process of removing and absorbing heat produces a heating and cooling affect, which can be used for refrigeration, air-conditioning, and heating.

 While all of these steps are important, the compression step is a crucial
30 part of the cycle (Fluorocarbon Refrigerants Handbook, R. C. Downing, Prentice-Hall, Inc. 1988). When a compressor is used to mechanically increase the pressure of a refrigerant, a lubricant (*i.e.* oil) is needed in the compressor to lubricate the compressor bearings and other moving parts, and the properties of

the oil must be suitable for this purpose. Often the oil leaves the compressor by slipping (i.e. leaking) past the piston rings in reciprocating compressors, by entrainment with the refrigerant, and by excessive foaming as refrigerant is released from solution in the oil. A small amount of oil circulating with the refrigerant may not cause any significant problems; however, if the oil accumulates in the system over time, the oil level in the compressor can become critically low and the compressor bearings and other moving parts can overheat and fail. Many sealed (hermetic) compressors are used for refrigerators, window air-conditioners, residential heat pumps, and commercial air-handlers. These compressors are precharged at the factory with the correct amount of oil, and a drop in oil level can reduce the life expectancy for these machines.

Problems related to compressor oil migration in a vapor compression system are related to the fact that oil may accumulate in the evaporator and cause a decrease in the cooling capacity of the system. Designers of refrigeration and air-conditioning systems often install oil separators in the discharge line of the compressor to trap oil and return it to the compressor crankcase. The oil is pushed along in the condenser by the liquid refrigerant and in the evaporator and suction line back to the compressor by the velocity of the refrigerant gas. The refrigerant pipes can also be designed to allow the oil to flow downhill back to the compressor using gravity. When oil leaves the compressor, however, it may or may not be soluble in the refrigerant, and the more soluble the refrigerant is in the oil, the better oil return to the compressor will be. The refrigerant dissolves in the oil, and reduces the viscosity of the oil that assists with the oil moving through the piping back to the compressor. Although these measures can be helpful, they are not 100% effective and may postpone the problem rather than solve it completely. For example, when there is a drop in the cooling capacity of the system, high temperatures in the compressor can lead to thermal breakdown and copper plating on valve surfaces, thus interfering with the proper operation of the compressor. Sludging and acid formation may limit the life of the system.

Several classes of commercial oils exist. Historically, the most common lubricants were natural or mineral-based oils (MO). Initially, when most refrigerants were based on chlorofluorocarbons (CFCs), the chlorine content in

the refrigerant [*i.e.* fluorotrichloromethane (CFC-11) and difluorodichloromethane (CFC-12)] provided excellent solubility with mineral oil. With the development of the hydrofluorocarbon (HFC) refrigerants to replace CFCs, mineral oil had little to no solubility with the new refrigerants [*e.g.* 1,1,1,2-tetrafluoroethane (HFC-134a)]. Therefore, new synthetic based lubricants such as polyalkene glycol-based oil (PAG) and polyolester-based oil (PO) were developed. Although the new synthetic oils had better solubility with the HFC refrigerants, the solubility was still not as good as that of CFCs and mineral-based oils. Furthermore the PAG and PO oils are not as effective as lubricants, and additive packages must be mixed with the synthetic oils to improve their lubricating performance. Refrigerant gases other than HFCs, such as non-fluorinated hydrocarbons and carbon dioxide (CO₂), have been proposed. Non-fluorinated hydrocarbons have excellent solubility in MOs, however no suitable lubricant has been found for CO₂ refrigeration and air-conditioning compressors.

15

Ionic liquids have been described as potential lubricants. Wang *et al* [WEAR (2004) 256:44-48] and Ye *et al* [Chem. Comm. (2001) 21:2244-2245] describe the tribological properties and predicted lubricant functionality of alkyylimidazolium hexafluorophosphate and alkyylimidazolium tetrafluoroborate, respectively. WO 05/35702 discloses the heat resistance and friction characteristics of lubricating oils containing ionic liquids as base oils.

20

A need nevertheless remains for lubricants that are well suited for use in conjunction with the current and new refrigerants that are used in a vapor compression system. A lubricant with high solubility for HFCs and CO₂, for example, would be desirable to replace, or be mixed with, traditional synthetic oils such as PAGs and POs to increase the operating lifetime of vapor compression systems. As a result, ionic liquids are disclosed herein, for use with various refrigerants in a vapor compression system, as type of compressor lubricant having a balance of properties that is superior to that of the current synthetic lubricants.

30

Summary

The present invention relates to the use of ionic liquids as lubricants in the compressor of a vapor compression system that provides cooling or heating.

5

In one embodiment, this invention provides an apparatus for temperature adjustment that includes

(a) a compressor that increases the pressure of the vapor of at least one refrigerant, wherein the compressor comprises moving parts that are lubricated by
10 at least one ionic liquid;

(b) a condenser that receives refrigerant vapor that is passed out of the compressor, and condenses the vapor under pressure to a liquid;

(c) a pressure reduction device that receives liquid refrigerant that is passed out of the condenser, and reduces the pressure of the liquid to form a
15 mixture of refrigerant in liquid and vapor form;

(d) an evaporator that receives the mixture of liquid and vapor refrigerant that is passed out of the pressure reduction device, and evaporates the remaining liquid in the mixture to form refrigerant vapor; and

(e) a conduit that returns to the compressor refrigerant vapor that is passed
20 out of the evaporator.

In another embodiment, this invention provides an apparatus for temperature adjustment that includes

(a) a compressor that increases the pressure of the vapor of at least one
25 refrigerant;

(b) a condenser that receives refrigerant vapor that is passed out of the compressor, and condenses the vapor under pressure to a liquid;

(c) a pressure reduction device that receives liquid refrigerant that is passed out of the condenser, and reduces the pressure of the liquid to form a
30 mixture of refrigerant in liquid and vapor form;

(d) an evaporator that receives the mixture of liquid and vapor refrigerant that is passed out of the pressure reduction device, and evaporates the remaining liquid in the mixture to form refrigerant vapor; and

(e) a conduit that returns to the compressor refrigerant vapor that is passed out of the evaporator;

wherein a refrigerant is admixed with at least one ionic liquid.

5 Either of these apparatus may adjust temperature by absorbing heat from, or transferring heat to, an object, medium or space. In such an apparatus, the condenser may for example, be located in proximity to an object, medium or space to be heated; or the evaporator may be located in proximity to an object, medium or space to be cooled.

10

In a further embodiment, this invention provides a process for adjusting the temperature of an object, medium or a space by

(a) providing a mechanical device having moving parts to increase the pressure of the vapor of at least one refrigerant, and providing at least one ionic
15 liquid to lubricate the moving parts of the device;

(b) condensing the refrigerant vapor under pressure to a liquid;

(c) reducing the pressure of the liquid refrigerant to form a mixture of refrigerant in liquid and vapor form;

(d) evaporating the liquid refrigerant to form refrigerant vapor; and

20 (e) repeating step (a) to increase the pressure of the refrigerant vapor formed in steps (c) and (d).

In yet another embodiment, this invention provides a process for adjusting the temperature of an object, medium or a space by

25 (a) increasing the pressure of the vapor of at least one refrigerant;

(b) condensing the refrigerant vapor under pressure to a liquid;

(c) reducing the pressure of the liquid refrigerant to form a mixture of refrigerant in liquid and vapor form;

(d) evaporating the liquid refrigerant to form refrigerant vapor;

30 (e) separating from the refrigerant vapor any ionic liquid present therein; and

(f) repeating step (a) to increase the pressure of the refrigerant vapor formed in steps (c) and (d).

In either of such processes, temperature may be adjusted by absorbing heat from, or transferring heat to, an object, medium or space. In such a process, the refrigerant vapor may in step (b) be condensed to a liquid in proximity to an object, medium or space to be heated; or the liquid refrigerant may in step (d) be
5 evaporated to form refrigerant vapor in proximity to an object, medium or space to be cooled.

In yet another embodiment, this invention provides a method for operating a mechanical compressor that has moving parts by providing an ionic liquid as the
10 lubricant for the moving parts.

Brief Description of the Drawings

FIG. 1. is a schematic diagram of a simple vapor compression cycle.

15 **FIG. 2.** is a schematic diagram of a simple vapor compressor.

FIG. 3 shows measured isothermal solubility data (in mole fraction) of the system HFC-32 + [bmim][PF₆] as a function of pressure. Filled circles (●) represent measured isothermal data at 10 °C, filled triangles (▲) represent measured isothermal data at 25 °C, filled squares (■) represent measured
20 isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 4 shows measured isothermal solubility data (in mole fraction) of the system HFC-125 + [bmim][PF₆] as a function of pressure. Filled circles (●) represent measured isothermal data at 10 °C, filled triangles (▲) represent
25 measured isothermal data at 25 °C, filled squares (■) represent measured isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 5 shows measured isothermal solubility data (in mole fraction) of the system HFC-134a + [bmim][PF₆] as a function of pressure. Filled circles (●)
30 represent measured isothermal data at 10 °C, filled triangles (▲) represent measured isothermal data at 25 °C, filled squares (■) represent measured isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 6 shows measured isothermal solubility data (in mole fraction) of the system HFC-143a + [bmim][PF₆] as a function of pressure. Filled circles (●) represent measured isothermal data at 10 °C, filled triangles (▲) represent measured isothermal data at 25 °C, filled squares (■) represent measured isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 7 shows measured isothermal solubility data (in mole fraction) of the system HFC-152a + [bmim][PF₆] as a function of pressure. Filled circles (●) represent measured isothermal data at 10 °C, filled triangles (▲) represent measured isothermal data at 25 °C, filled squares (■) represent measured isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 8 shows measured isothermal solubility data (in mole fraction) of the system HFC-32 + [bmim][BF₄] as a function of pressure. Filled circles (●) represent measured isothermal data at 10 °C, filled triangles (▲) represent measured isothermal data at 25 °C, filled squares (■) represent measured isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 9 shows measured isothermal solubility data (in mole fraction) of the system HFC-23 + [bmim][PF₆] as a function of pressure. Filled circles (●) represent measured isothermal data at 10 °C, filled triangles (▲) represent measured isothermal data at 25 °C, filled squares (■) represent measured isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 10 shows measured isothermal solubility data (in mole fraction) of the system HFC-23 + [emim][PF₆] as a function of pressure. Filled squares (■) represent measured isothermal data at 60 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 11 shows measured isothermal solubility data (in mole fraction) of the system HFC-32 + [dmpim][TMeM] as a function of pressure. Filled circles (●) represent measured isothermal data at 10 °C, filled triangles (▲) represent measured isothermal data at 25 °C, filled squares (■) represent measured

isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 12 shows measured isothermal solubility data (in mole fraction) of the system HFC-32 + [emim][BEI] as a function of pressure. Filled circles (●) represent measured isothermal data at 10 °C, filled triangles (▲) represent measured isothermal data at 25 °C, filled squares (■) represent measured isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 13 shows measured isothermal solubility data (in mole fraction) of the system HFC-32 + [emim][BMeI] as a function of pressure. Filled circles (●) represent measured isothermal data at 10 °C, filled triangles (▲) represent measured isothermal data at 25 °C, filled squares (■) represent measured isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 14 shows measured isothermal solubility data (in mole fraction) of the system HFC-32 + [pmpy][BMeI] as a function of pressure. Filled circles (●) represent measured isothermal data at 10 °C, filled triangles (▲) represent measured isothermal data at 25 °C, filled squares (■) represent measured isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 15 shows measured isothermal solubility data (in mole fraction) of the system HFC-32 + [bmpy][BMeI] as a function of pressure. Filled circles (●) represent measured isothermal data at 10 °C, filled triangles (▲) represent measured isothermal data at 25 °C, filled squares (■) represent measured isothermal data at 50 °C, and filled diamonds (◆) represent measured isothermal data at 75 °C. Solid lines represent data trends.

FIG. 16 shows measured isothermal solubility data at 25 °C of the systems HFC-32 + eight different ionic liquids as a function of pressure for comparison. Open diamonds (◇) represent measured isothermal data for HFC-32 + 1-ethyl-3-methylimidazolium bis(pentafluoroethylsulfonyl)imide, open circles (○) represent measured isothermal data for HFC-32 + 1-propyl-2,3-dimethylimidazolium tris(trifluoromethylsulfonyl)methide at 25 °C, open squares (□) represent measured isothermal data for HFC-32 + 1-propyl-2,3-

dimethylimidazolium bis(trifluoromethylsulfonyl)imide at 25 °C, closed diamonds (◆) represent measured isothermal data for HFC-32 + 3-methyl-1-propylpyridinium bis(trifluoromethylsulfonyl)imide, open triangles (Δ) represent measured isothermal data for HFC-32 + 1-butyl-3-methylimidazolium hexafluorophosphate at 25 °C, filled circles (●) represent measured isothermal data for HFC-32 + 1-butyl-3-methylimidazolium tetrafluoroborate at 25 °C, filled squares (■) represent measured isothermal data for HFC-32 + 1,3-dioctylimidazolium iodide at 25 °C, and filled triangles (▲) represent measured isothermal data for HFC-32 + 1-octyl-3-methylimidazolium iodide at 25 °C. Solid lines represent data trends.

FIG. 17 shows measured isothermal solubility data (in mole fraction) at 10 °C of the systems HFC-32, HFC-152a, HFC-134a, HFC-125, and HFC-143a + [bmim][PF₆] in terms of absolute pressure divided by the gas saturation pressure at 10 °C shown by ratio (P/P_0). Open diamonds (◇) represent measured isothermal data for HFC-32 at 10 °C with $P_0 = 11.069$ bar, open cross hatch (X) represents measured isothermal data for HFC-152a at 10 °C with $P_0 = 3.7277$ bar, filled circles (●) represent measured isothermal data for HFC-134a at 10 °C with $P_0 = 4.1461$ bar, open squares (□) represent measured isothermal data for HFC-125 at 10 °C with $P_0 = 9.0875$ bar, filled circles (●) represent measured isothermal data for HFC-143a at 10 °C with $P_0 = 8.3628$ bar. Solid lines represent data trend and dashed line represents Raoult's Law.

FIG. 18 shows a schematic diagram of the gravimetric microbalance used for measuring gas absorption in the ionic liquids. In the diagram j_1 , j_2 , and j_3 refer to the counter-weight, hook and chain, respectively; i_1 , i_2 and i_3 refer to the sample container, wire and chain, respectively, W_g refers to the force due to gravity; and B refers to the force due to buoyancy.

Detailed Description

In the description of this invention, the following definitional structure is provided for certain terminology as employed in various locations in the specification:

An "ionic liquid" is an organic salt that is fluid at about 100°C or below, as more particularly described in *Science* (2003) 302:792-793. A "fluorinated ionic liquid" is an ionic liquid having at least one fluorine on either the cation or the anion. A "fluorinated cation" or "fluorinated anion" is a cation or anion, respectively, comprising at least one fluorine.

A "halogen" is bromine, iodine, chlorine or fluorine.

A "heteroaryl" is an alkyl group having a heteroatom.

A "heteroatom" is an atom other than carbon in the structure of an alkanyl, alkenyl, cyclic or aromatic compound.

A "hydrofluorocarbon" is a compound comprising fluorine, carbon, and at least one hydrogen atom. A hydrofluorocarbon compound (HFC) includes a hydrochlorofluorocarbon (HCFC), wherein HFC and HCFC are common terms used to define refrigerants [see, for example, Ralph C. Downing, *Fluorocarbon Refrigerants Handbook*, Prentice-Hall, Inc., Englewood Cliffs, NJ (1988)]. Hydrofluorocarbon compounds also include compounds selected from the group consisting of hydrofluoroethers, hydrofluoroketones, hydrofluoroaromatics and hydrofluoroolefins; representative examples of which include methyl nonafluoroisobutyl ether, methyl nonafluorobutyl ether, ethyl nonafluoroisobutyl ether, ethyl nonafluorobutyl ether, and 3-ethoxy-1,1,1,2,3,4,4,5,5,6,6,6-dodecafluoro-2-trifluoromethylhexane. Hydrofluorocarbon compounds also include compounds wherein one or more substituents is bromine, chlorine or iodine.

“Optionally substituted with at least one member selected from the group consisting of”, when referring to an alkane, alkene, alkoxy, fluoroalkoxy, perfluoroalkoxy, fluoroalkyl, perfluoroalkyl, aryl or heteroaryl radical or moiety, means that one or more hydrogens on a carbon chain of the radical or moiety may
5 be independently substituted with one or more of the members of a recited group of substituents. For example, a substituted $-C_2H_5$ radical or moiety may, without limitation, be $-CF_2CF_3$, $-CH_2CH_2OH$ or $-CF_2CF_2I$ where the group or substituents consist of F, I and OH.

10 A “refrigerant” is a fluidic substance such as a fluorocarbon (FC), hydrofluorocarbon (HFC), chlorofluorocarbon (CFC), hydrochlorofluorocarbon (HCFC), or carbon dioxide that may be used as a thermal energy transfer vehicle. A refrigerant, when it changes phase from liquid to vapor (evaporates), removes heat from the surroundings; and when it changes phase from vapor to liquid
15 (condenses), adds heat to the surroundings. Although the term refrigerant may typically carry the connotation of a substance used only for cooling, the term is used herein in the generic sense of a thermal energy transfer vehicle or substance that is applicable for use in a system or apparatus that may be used for heating or cooling by the absorption or rejection of heat in a selected location. For the
20 present invention, the term “refrigerant” can be used to indicate one fluidic substance as described above, or may indicate a blend or mixture of two or more such fluidic substances.

A “vacuum” is a pressure less than 1 bar but greater than 10^{-4} bar for
25 practical use in extractive distillation equipment.

The present invention relates to the use of ionic liquids as lubricants in the operation of a compressor in a vapor compression system that provides cooling or
30 heating. This invention thus also relates to an apparatus for adjusting temperature that operates or executes a vapor compression cycle. The invention also provides a process for temperature adjustment, either cooling or heating, utilizing a vapor compression system in which an ionic liquid is used as a lubricant.

Vapor compression cycles for cooling or heating are known in the art from sources such as *Application Guide for Absorption Cooling/Refrigeration Using Recovered Heat* [Dorgan Cb, *et al.* (American Society of Heating, Refrigeration and Air Conditioning Engineers, Inc., 1995, Atlanta, GA, Chapter 5)]. A schematic diagram for a simple vapor compression system is shown in Figure 1. The system is composed of condenser and evaporator units with an expansion valve and a vapor compressor. Figure 2 provides a schematic for a simple vapor compressor.

In Fig. 1, the pressure of the vapor of one or more refrigerants is increased mechanically by the operation of a compressor. The compressor has moving parts that are lubricated by a lubricant. In this invention, one or more ionic liquids serve as the lubricant. The refrigerant having increased pressure is passed through a conduit to a condenser where it is condensed to liquid form. The action of condensing the refrigerant generates heat, which is transferred or rejected to the surroundings, which may be an object, medium or space to be heated. The condensed refrigerant in liquid form is passed out of the condenser and is received into a pressure reduction device such as an expansion valve, where some of the liquid is converted to refrigerant vapor, creating a mixture of refrigerant in liquid and vapor form. From there, the mixture flows out of the pressure reduction device and is received by an evaporator where the liquid refrigerant is evaporated to vapor form. As heat is absorbed by, and flows into, the evaporator by virtue of the transformation of the refrigerant from liquid to vapor form, the surroundings (such as an object, medium or space) lose heat and are cooled. The refrigerant, now all or essentially all in vapor form, is returned to the compressor where the same cycle commences again. Operation or execution of a vapor compression cycle in this manner can thus be used for temperature adjustment as, by the absorption or expulsion heat an object (for example a conduit or a container), a medium (for example a fluid such as air or water) or a space can be heated or cooled as desired.

In one embodiment, therefore, this invention provides an apparatus for temperature adjustment, to operate or execute a vapor compression cycle as described above, that includes (a) a compressor that increases the pressure of the

vapor of at least one refrigerant, wherein the compressor comprises moving parts that are lubricated by at least one ionic liquid; (b) a condenser that receives refrigerant vapor that is passed out of the compressor, and condenses the vapor under pressure to a liquid; (c) a pressure reduction device that receives liquid
5 refrigerant that is passed out of the condenser, and reduces the pressure of the liquid to form a mixture of refrigerant in liquid and vapor form; (d) an evaporator that receives the mixture of liquid and vapor refrigerant that is passed out of the pressure reduction device, and evaporates the remaining liquid in the mixture to form refrigerant vapor; and (e) a conduit that returns to the compressor
10 refrigerant vapor that is passed out of the evaporator.

In another embodiment, this invention provides an apparatus for temperature adjustment that includes (a) a compressor that increases the pressure of the vapor of at least one refrigerant; (b) a condenser that receives refrigerant
15 vapor that is passed out of the compressor, and condenses the vapor under pressure to a liquid; (c) a pressure reduction device that receives liquid refrigerant that is passed out of the condenser, and reduces the pressure of the liquid to form a mixture of refrigerant in liquid and vapor form; (d) an evaporator that receives the mixture of liquid and vapor refrigerant that is passed out of the pressure
20 reduction device, and evaporates the remaining liquid in the mixture to form refrigerant vapor; and (e) a conduit that returns to the compressor refrigerant vapor that is passed out of the evaporator; wherein a refrigerant is admixed with at least one ionic liquid. In a vapor compression system where an ionic liquid is used as a lubricant for a mechanical compressor, the refrigerant as is circulated
25 through the vapor compression cycle may contain some amount of an ionic liquid lubricant where the lubricant has leaked out of its ordinary location and into other parts of the system where it is conceptually not intended to be.

Either of such apparatus may adjust temperature by absorbing heat from,
30 or transferring heat to, an object, medium or space. In such an apparatus, the condenser may for example, be located in proximity to an object, medium or space to be heated; or the evaporator may be located in proximity to an object, medium or space to be cooled.

An apparatus of this invention may be deployed for use in, or fabricated or operated as, a refrigerator, a freezer, an ice machine, an air conditioner, an industrial cooling system, a heater or heat pump. Each of these instruments may be situated in a residential, commercial or industrial setting, or may be
5 incorporated into a mobilized device such as a car, truck, bus, train, airplane, or other device for transportation, or may be incorporated into a piece of equipment such as a medical instrument.

In another embodiment, this invention provides a process for adjusting the
10 temperature of an object, medium or a space by (a) providing a mechanical device to increase the pressure of the vapor of at least one refrigerant where the device has moving parts, and providing at least one ionic liquid to lubricate the moving parts of the device; (b) condensing the refrigerant vapor under pressure to a liquid; (c) reducing the pressure of the liquid refrigerant to form a mixture of
15 refrigerant in liquid and vapor form; (d) evaporating the liquid refrigerant to form refrigerant vapor; (e) repeating step (a) to increase the pressure of the refrigerant vapor formed in steps (c) and (d).

In another embodiment, this invention provides a process for adjusting the
20 temperature of an object, medium or a space by (a) passing the vapor of at least one refrigerant through a compressor that has moving parts to increase the pressure of the refrigerant vapor, and providing at least one ionic liquid to lubricate the moving parts of the compressor; (b) passing the refrigerant vapor out of the compressor and into a condenser to condense the refrigerant vapor
25 under pressure to a liquid; (c) passing the refrigerant in liquid form out of the condenser to a pressure reduction device to reduce the pressure of the liquid refrigerant to form a mixture of refrigerant in liquid and vapor form; (d) passing the mixture to an evaporator to evaporate the liquid refrigerant to form refrigerant vapor; (e) passing the refrigerant vapor throughout a conduit to the compressor
30 to repeat step (a) to increase the pressure of the refrigerant vapor formed in steps (c) and (d).

In yet another embodiment, this invention provides a process for adjusting the temperature of an object, medium or a space by (a) increasing the pressure of

the vapor of at least one refrigerant; (b) condensing the refrigerant vapor under pressure to a liquid; (c) reducing the pressure of the liquid refrigerant to form a mixture of refrigerant in liquid and vapor form; (d) evaporating the liquid refrigerant to form refrigerant vapor; (e) separating from the refrigerant vapor any 5 ionic liquid present therein; and (f) repeating step (a) to increase the pressure of the refrigerant vapor formed in steps (c) and (d). In a vapor compression system where an ionic liquid is used as a lubricant for a mechanical compressor, the refrigerant as is circulated through the vapor-compression cycle may contain some amount of an ionic liquid lubricant where the lubricant has leaked out of its 10 ordinary location and into other parts of the system where it is conceptually not intended to be. When refrigerant that is thus admixed with an ionic liquid is returned to the compressor for re-pressurizing, the ambient temperature will be at a level at which the refrigerant will be in vapor form but an ionic liquid will be in liquid form. The return conduit may thus be designed to permit the ionic liquid, 15 as a liquid, to be separated from the refrigerant by being drained out through an opening into the lubricant sump while the refrigerant vapor, as a vapor, passes alone on into the compressor for re-pressurization.

In any of such processes, temperature may be adjusted by absorbing heat 20 from, or transferring heat to, an object, medium or space. In such a process, the refrigerant vapor may in step (b) be condensed to a liquid in proximity to an object, medium or space to be heated; or the liquid refrigerant may in step (d) be evaporated to form refrigerant vapor in proximity to an object, medium or space to be cooled.

25

In one embodiment of the invention, a refrigerant suitable for use in a vapor compression system hereof may be selected from one or more members of the group consisting of CHClF_2 (R-22), CHF_3 (R-23), CH_2F_2 (R-32), CH_3F (R-41), CHF_2CF_3 (R-125), CH_2FCF_3 (R-134a), $\text{CHF}_2\text{OCHF}_2$ (E-134), CH_3CClF_2 (R-142b), CH_3CF_3 (R-143a), CH_3CHF_2 (R-152a), $\text{CH}_3\text{CH}_2\text{F}$ (R-161), CH_3OCH_3 (E170), $\text{CF}_3\text{CF}_2\text{CF}_3$ (R-218), CF_3CHF_2 (R-227ea), $\text{CF}_3\text{CH}_2\text{CF}_3$ (R-236fa), $\text{CH}_2\text{FCF}_2\text{CHF}_2$ (R-245ca), $\text{CHF}_2\text{CH}_2\text{CF}_3$ (R-245fa), $\text{CH}_3\text{CH}_2\text{CH}_3$ (R-290), $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_3$ (R-600), $\text{CH}(\text{CH}_3)_2\text{CH}_3$ (R-600a), $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3$ (R-601), $(\text{CH}_3)_2\text{CHCH}_2\text{CH}_3$ (R-601a), $\text{CH}_3\text{CH}_2\text{OCH}_2\text{CH}_3$ (R-610), NH_3 , CO_2 , and

$\text{CH}_3\text{CH}=\text{CH}_2$, wherein the common name is given in parentheses following the chemical formula. In another embodiment of the invention, the refrigerant useful for the invention can be a blend selected from the group consisting of R-125/R-143a/R-134a (44.0/52.0/4.0) (R-404A), R-32/R-125/R-134a (20.0/40.0/40.0) (R-407A), R-32/R-125/R-134a (10.0/70.0/20.0) (R-407B), R-32/R-125/R-134a (23.0/25.0/52.0) (R-407C), R-32/R-125/R-134a (15.0/15.0/70.0) (R-407D), R-32/R-125/R-134a (25.0/15.0/60.0) (R-407E), R-32/R-125 (50.0/50.0) (R-410A), R-32/R-125 (45.0/55) (R-410B), R-218/R-134a/R-600a (9.0/88.0/3.0) (R-413A), R-125/R-134a/R-600 (46.6/50.0/3.4) (417A), R-125/R-134a/E170 (77.0/19.0/4.0) (R-419A), R-134a/R-142b (88.0/12.0) (R-420A), R-134a/R-142b (80.6/19.4), R-125/R-134a (58.0/42.0) (R-421A), R-125/R-134a (85.0/15.0) (R-421B), R-125/R-134a/R-600a (85.1/11.5/3.4) (R-422A), R-125/R-134a/R-600a (55.0/42.0/3.0) (R-422B), R-125/R-134a/R-600a (82.0/15.0/3.0) (R-422C), R-125/R-134a/R-600a (65.1/31.5/3.4) (R-422D), R-134a/R-227ea (52.5/47.5) (R423A), R-125/R-134a/R-600a/R-600/R-601a (50.5/47.0/0.9/1.0/0.6) (R-424A), R-32/R-134a/R-227ea (18.5/69.5/12.0) (R-425A), R-125/R-134a/R-600/R-601a (5.1/93.0/1.3/0.6) (R-426A), R-32/R-125/R-143a/R-134a (15.0/25.0/10.0/50.0) (R-427A), R-32/R-125/R-143a/R-134a (2.0/41.0/50.0/7.0), R-32/R-125/R-143a/R-134a (10.0/33.0/36.0/21.0), R-125/R-143a/R-290/R-600a (77.5/20.0/0.6/1.9) (R-428A), and R-125/R-143a (50.0/50.0) (R-507A), wherein the weight percent of the individual components relative to the other refrigerant components in the blend and the common name are given in parentheses following each blend.

In another embodiment of the invention, the refrigerant can be selected from one or more members of the group consisting of R-22, R-32, R-125, R-134a, R-404A, R-410A, R-413A, R-422A, R-422D, R-423A, R-426A, R-427A and R-507A.

In another embodiment of the invention, a hydrofluorocarbon for use as a refrigerant herein is selected from one or more members of the group consisting of trifluoromethane (HFC-23), difluoromethane (HFC-32), pentafluoroethane (HFC-125), 1,1,2,2-tetrafluoroethane (HFC-134), 1,1,1,2-tetrafluoroethane (HFC-134a), 1,1,1-trifluoroethane (HFC-143a), 1,1-difluoroethane (HFC-152a), fluoroethane (HFC-161), R-404A, R-407C, and R-410A.

In another embodiment, a hydrofluorocarbon is selected from one or more members of the group consisting of pentafluoroethane (HFC-125), 1,1,2,2-tetrafluoroethane (HFC-134), 1,1,1,2-tetrafluoroethane (HFC-134a), R-404A, R-407C, and R-410A.

Mixtures of refrigerants or blends may also be used.

In another embodiment, the refrigerant can be at least one fluoroolefin selected from the group consisting of:

- (i) fluoroolefins of the formula E - or Z - $R^1CH=CHR^2$, wherein R^1 and R^2 are, independently, C_1 to C_6 perfluoroalkyl groups, and wherein the total number of carbons in the compound is at least 5;
- (ii) cyclic fluoroolefins of the formula $cyclo-[CX=CY(CZW)_n]$, wherein X, Y, Z, and W, independently, are H or F, and n is an integer from 2 to 5; and
- (iii) fluoroolefins selected from the group consisting of:

2,3,3-trifluoro-1-propene ($CHF_2CF=CH_2$);	1,1,2-trifluoro-1-propene ($CH_3CF=CF_2$);
1,2,3-trifluoro-1-propene ($CH_2FCF=CF_2$);	1,1,3-trifluoro-1-propene ($CH_2FCH=CF_2$);
1,3,3-trifluoro-1-propene ($CHF_2CH=CHF$);	1,1,1,2,3,4,4,4-octafluoro-2-butene ($CF_3CF=CFCF_3$);
1,1,2,3,3,4,4,4-octafluoro-1-butene ($CF_3CF_2CF=CF_2$);	1,1,1,2,4,4,4-heptafluoro-2-butene ($CF_3CF=CHCF_3$);
1,2,3,3,4,4,4-heptafluoro-1-butene ($CHF=CFCF_2CF_3$);	1,1,1,2,3,4,4-heptafluoro-2-butene ($CHF_2CF=CFCF_3$);
1,3,3,3-tetrafluoro-2-(trifluoromethyl)-1-propene ($((CF_3)_2C=CHF)$);	1,1,3,3,4,4,4-heptafluoro-1-butene ($CF_2=CHCF_2CF_3$);
1,1,2,3,4,4,4-heptafluoro-1-butene ($CF_2=CFCHF_2CF_3$);	1,1,2,3,3,4,4-heptafluoro-1-butene ($CF_2=CFCF_2CHF_2$);
2,3,3,4,4,4-hexafluoro-1-butene ($CF_3CF_2CF=CH_2$);	1,3,3,4,4,4-hexafluoro-1-butene ($CHF=CHCF_2CF_3$);
1,2,3,4,4,4-hexafluoro-1-butene ($CHF=CFCHF_2CF_3$);	1,2,3,3,4,4-hexafluoro-1-butene ($CHF=CFCF_2CHF_2$);
1,1,2,3,4,4-hexafluoro-2-butene ($CHF_2CF=CFCHF_2$);	1,1,1,2,3,4-hexafluoro-2-butene ($CH_2FCF=CFCF_3$);
1,1,1,2,4,4-hexafluoro-2-butene	

	(CHF ₂ CH=CFCF ₃);	1,1,1,3,4,4-hexafluoro-2-butene
	(CF ₃ CH=CFCHF ₂);	1,1,2,3,3,4-hexafluoro-1-butene
	(CF ₂ =CFCF ₂ CH ₂ F);	1,1,2,3,4,4-hexafluoro-1-butene
	(CF ₂ =CFCHFCHF ₂);	3,3,3-trifluoro-2-(trifluoromethyl)-1-propene
5	(CH ₂ =C(CF ₃) ₂);	1,1,1,2,4-pentafluoro-2-butene
	(CH ₂ FCH=CFCF ₃);	1,1,1,3,4-pentafluoro-2-butene
	(CF ₃ CH=CFCH ₂ F);	3,3,4,4,4-pentafluoro-1-butene
	(CF ₃ CF ₂ CH=CH ₂);	1,1,1,4,4-pentafluoro-2-butene
	(CHF ₂ CH=CHCF ₃);	1,1,1,2,3-pentafluoro-2-butene
10	(CH ₃ CF=CFCF ₃);	2,3,3,4,4-pentafluoro-1-butene
	(CH ₂ =CFCF ₂ CHF ₂);	1,1,2,4,4-pentafluoro-2-butene
	(CHF ₂ CF=CHCHF ₂);	1,1,2,3,3-pentafluoro-1-butene
	(CH ₃ CF ₂ CF=CF ₂);	1,1,2,3,4-pentafluoro-2-butene
	(CH ₂ FCF=CFCF ₂);	1,1,3,3,3-pentafluoro-2-methyl-1-propene
15	(CF ₂ =C(CF ₃)(CH ₃));	2-(difluoromethyl)-3,3,3-trifluoro-1-propene
	(CH ₂ =C(CHF ₂)(CF ₃));	2,3,4,4,4-pentafluoro-1-butene
	(CH ₂ =CFCHF ₂ CF ₃);	1,2,4,4,4-pentafluoro-1-butene
	(CHF=CFCH ₂ CF ₃);	1,3,4,4,4-pentafluoro-1-butene
	(CHF=CHCHF ₂ CF ₃);	1,3,3,4,4-pentafluoro-1-butene
20	(CHF=CHCF ₂ CHF ₂);	1,2,3,4,4-pentafluoro-1-butene
	(CHF=CFCHFCHF ₂);	3,3,4,4-tetrafluoro-1-butene
	(CH ₂ =CHCF ₂ CHF ₂);	1,1-difluoro-2-(difluoromethyl)-1-propene
	(CF ₂ =C(CHF ₂)(CH ₃));	1,3,3,3-tetrafluoro-2-methyl-1-propene
	(CHF=C(CF ₃)(CH ₃));	3,3-difluoro-2-(difluoromethyl)-1-propene
25	(CH ₂ =C(CHF ₂) ₂);	1,1,1,2-tetrafluoro-2-butene
	(CF ₃ CF=CHCH ₃);	1,1,1,3-tetrafluoro-2-butene
	(CH ₃ CF=CHCF ₃);	1,1,1,2,3,4,4,5,5,5-decafluoro-2-pentene
	(CF ₃ CF=CFCF ₂ CF ₃);	1,1,2,3,3,4,4,5,5,5-decafluoro-1-pentene
	(CF ₂ =CFCF ₂ CF ₂ CF ₃);	1,1,1,4,4,4-hexafluoro-2-(trifluoromethyl)-2-butene
	((CF ₃) ₂ C=CHCF ₃);	
30	(CF ₃ CF=CHCF ₂ CF ₃);	1,1,1,2,4,4,5,5,5-nonafluoro-2-pentene
	(CF ₃ CH=CFCF ₂ CF ₃);	1,1,1,3,4,4,5,5,5-nonafluoro-2-pentene
	(CHF=CFCF ₂ CF ₂ CF ₃);	1,2,3,3,4,4,5,5,5-nonafluoro-1-pentene
	(CF ₂ =CHCF ₂ CF ₂ CF ₃);	1,1,3,3,4,4,5,5,5-nonafluoro-1-pentene
	(CF ₂ =CFCF ₂ CF ₂ CHF ₂);	1,1,2,3,3,4,4,5,5-nonafluoro-1-pentene
35	(CHF ₂ CF=CFCF ₂ CF ₃);	1,1,2,3,4,4,5,5,5-nonafluoro-2-pentene
	(CF ₃ CF=CFCF ₂ CHF ₂);	1,1,1,2,3,4,4,5,5-nonafluoro-2-pentene
	(CF ₃ CF=CFCHF ₂ CF ₃);	1,1,1,2,3,4,5,5,5-nonafluoro-2-pentene
		1,2,3,4,4,4-hexafluoro-3-(trifluoromethyl)-1-butene

- (CHF=CFCF(CF₃)₂); 1,1,2,4,4,4-hexafluoro-3-(trifluoromethyl)-1-butene (CF₂=CFCH(CF₃)₂); 1,1,1,4,4,4-hexafluoro-2-(trifluoromethyl)-2-butene (CF₃CH=C(CF₃)₂); 1,1,3,4,4,4-hexafluoro-3-(trifluoromethyl)-1-butene (CF₂=CHCF(CF₃)₂);
- 5 2,3,3,4,4,5,5,5-octafluoro-1-pentene (CH₂=CFCF₂CF₂CF₃); 1,2,3,3,4,4,5,5-octafluoro-1-pentene (CHF=CFCF₂CF₂CHF₂); 3,3,4,4,4-pentafluoro-2-(trifluoromethyl)-1-butene (CH₂=C(CF₃)CF₂CF₃); 1,1,4,4,4-pentafluoro-3-(trifluoromethyl)-1-butene (CF₂=CHCH(CF₃)₂); 1,3,4,4,4-pentafluoro-3-(trifluoromethyl)-1-butene (CHF=CHCF(CF₃)₂); 1,1,4,4,4-pentafluoro-2-(trifluoromethyl)-1-butene (CF₂=C(CF₃)CH₂CF₃); 3,4,4,4-tetrafluoro-3-(trifluoromethyl)-1-butene ((CF₃)₂CFCH=CH₂); 3,3,4,4,5,5,5-heptafluoro-1-pentene (CF₃CF₂CF₂CH=CH₂); 2,3,3,4,4,5,5-heptafluoro-1-pentene (CH₂=CFCF₂CF₂CHF₂); 1,1,3,3,5,5,5-heptafluoro-1-butene (CF₂=CHCF₂CH₂CF₃); 1,1,1,2,4,4,4-heptafluoro-3-methyl-2-butene (CF₃CF=C(CF₃)(CH₃)); 2,4,4,4-tetrafluoro-3-(trifluoromethyl)-1-butene (CH₂=CFCH(CF₃)₂); 1,4,4,4-tetrafluoro-3-(trifluoromethyl)-1-butene (CHF=CHCH(CF₃)₂);
- 20 1,1,1,4-tetrafluoro-2-(trifluoromethyl)-2-butene (CH₂FCH=C(CF₃)₂); 1,1,1,3-tetrafluoro-2-(trifluoromethyl)-2-butene (CH₃CF=C(CF₃)₂); 1,1,1-trifluoro-2-(trifluoromethyl)-2-butene ((CF₃)₂C=CHCH₃); 3,4,4,5,5,5-hexafluoro-2-pentene (CF₃CF₂CF=CHCH₃); 1,1,1,4,4,4-hexafluoro-2-methyl-2-butene (CF₃C(CH₃)=CHCF₃); 3,3,4,5,5,5-hexafluoro-1-pentene (CH₂=CHCF₂CHFCF₃); 4,4,4-trifluoro-3-(trifluoromethyl)-1-butene (CH₂=C(CF₃)CH₂CF₃); 1,1,2,3,3,4,4,5,5,6,6,6-dodecafluoro-1-hexene (CF₃(CF₂)₃CF=CF₂); 1,1,1,2,2,3,4,5,5,6,6,6-dodecafluoro-3-hexene (CF₃CF₂CF=CFCF₂CF₃); 1,1,1,4,4,4-hexafluoro-2,3-bis(trifluoromethyl)-2-butene ((CF₃)₂C=C(CF₃)₂); 1,1,1,2,3,4,5,5,5-nonafluoro-4-(trifluoromethyl)-2-pentene ((CF₃)₂CFCF=CFCF₃); 1,1,1,4,4,5,5,5-octafluoro-2-(trifluoromethyl)-2-pentene ((CF₃)₂C=CHC₂F₅); 1,1,1,3,4,5,5,5-octafluoro-4-(trifluoromethyl)-2-pentene ((CF₃)₂CFCF=CHCF₃); 3,3,4,4,5,5,6,6,6-nonafluoro-1-hexene (CF₃CF₂CF₂CF₂CH=CH₂); 4,4,4-trifluoro-3,3-bis(trifluoromethyl)-1-butene (CH₂=CHC(CF₃)₃); 1,1,1,4,4,4-hexafluoro-2-(trifluoromethyl)-3-methyl-2-butene
- 35

5 $((CF_3)_2C=C(CH_3)(CF_3))$; 2,3,3,5,5,5-hexafluoro-4-(trifluoromethyl)-1-pentene $(CH_2=CFCF_2CH(CF_3)_2)$;
 1,1,1,2,4,4,5,5,5-nonafluoro-3-methyl-2-pentene
 $(CF_3CF=C(CH_3)CF_2CF_3)$; 1,1,1,5,5,5-hexafluoro-4-(trifluoromethyl)-2-pentene $(CF_3CH=CHCH(CF_3)_2)$;
 3,4,4,5,5,6,6,6-octafluoro-2-hexene $(CF_3CF_2CF_2CF=CHCH_3)$;
 3,3,4,4,5,5,6,6-octafluoro-1-hexene $(CH_2=CHCF_2CF_2CF_2CHF_2)$;
 1,1,1,4,4-pentafluoro-2-(trifluoromethyl)-2-pentene
 10 $((CF_3)_2C=CHCF_2CH_3)$; 4,4,5,5,5-pentafluoro-2-(trifluoromethyl)-1-pentene $(CH_2=C(CF_3)CH_2C_2F_5)$; 3,3,4,4,5,5,5-heptafluoro-2-methyl-1-pentene $(CF_3CF_2CF_2C(CH_3)=CH_2)$; 4,4,5,5,6,6,6-heptafluoro-2-hexene $(CF_3CF_2CF_2CH=CHCH_3)$; 4,4,5,5,6,6,6-heptafluoro-1-hexene $(CH_2=CHCH_2CF_2C_2F_5)$; 1,1,1,2,2,3,4-heptafluoro-3-hexene $(CF_3CF_2CF=CFC_2H_5)$; 4,5,5,5-tetrafluoro-4-(trifluoromethyl)-1-pentene $(CH_2=CHCH_2CF(CF_3)_2)$; 1,1,1,2,5,5,5-heptafluoro-4-methyl-2-pentene $(CF_3CF=CHCH(CF_3)(CH_3))$;
 15 1,1,1,3-tetrafluoro-2-(trifluoromethyl)-2-pentene
 $((CF_3)_2C=CFC_2H_5)$; 1,1,1,2,3,4,4,5,5,6,6,7,7,7-tetradecafluoro-2-heptene $(CF_3CF=CFCF_2CF_2C_2F_5)$; 1,1,1,2,2,3,4,5,5,6,6,7,7,7-tetradecafluoro-3-heptene $(CF_3CF_2CF=CFCF_2C_2F_5)$;
 20 1,1,1,3,4,4,5,5,6,6,7,7,7-tridecafluoro-2-heptene
 $(CF_3CH=CFCF_2CF_2C_2F_5)$; 1,1,1,2,4,4,5,5,6,6,7,7,7-tridecafluoro-2-heptene $(CF_3CF=CHCF_2CF_2C_2F_5)$; 1,1,1,2,2,4,5,5,6,6,7,7,7-tridecafluoro-3-heptene $(CF_3CF_2CH=CFCF_2C_2F_5)$;
 25 1,1,1,2,2,3,5,5,6,6,7,7,7-tridecafluoro-3-heptene
 $(CF_3CF_2CF=CHCF_2C_2F_5)$; $CF_2=CFOCF_2CF_3$ (PEVE) and $CF_2=CFOCF_3$ (PMVE).

30 In yet another embodiment, the refrigerant can be a composition comprising the fluoroolefin as defined above with a flammable refrigerant as described in U.S. Provisional Application No. 60/732,581, which is incorporated in its entirety as a part hereof for all purposes. Flammable refrigerants comprise any compound, which may be demonstrated to propagate a flame under specified
 35 conditions of temperature, pressure and composition when mixed with air. Flammable refrigerants may be identified by testing under conditions specified by ASHRAE (American Society of Heating, Refrigerating and Air-Conditioning

Engineers, Inc.) Standard 34-2001, under ASTM (American Society of Testing and Materials) E681-01, with an electronic ignition source. Such tests of flammability are conducted with the refrigerant at 101 kPa (14.7 psia) and a specified temperature (typically 100 °C (212 °F), or room temperature, that being
5 about 23 °C (73 °F) at various concentrations in air in order to determine the lower flammability limit (LFL) and upper flammability limit (UFL) of the test compound in air. Flammable refrigerants include hydrofluorocarbons, such as difluoromethane (HFC-32), fluorolefins, such as 1,2,3,3-tetrafluoro-1-propene (HFC-1234ye), fluoroethers, such as $C_4F_9OC_2H_5$, hydrocarbon ethers, such as
10 dimethyl ether, hydrocarbons, such as propane, ammonia, and combinations thereof. One example of a refrigerant composition comprising a fluoroolefin refrigerant with a flammable refrigerant is a refrigerant composition comprising about 99.0 weight percent to about 63.0 weight percent C_3HF_5 (HFC-1225ye) and about 1.0 weight percent to about 37.0 weight percent HFC-32.

15

Refrigerants useful for the invention also include compositions comprising pentafluoroethane (R-125), 1,1,1,2-tetrafluoroethane (R-134a), and at least two hydrocarbons each having eight or fewer carbon atoms, as described in U.S. Provisional Application No. 60/876,406, which is incorporated in its entirety as a
20 part hereof for all purposes. In some embodiments, the hydrocarbon is C_4 to C_8 hydrocarbon, such as butanes, pentanes, hexanes, heptanes, octanes, C_4 to C_8 alkenes, C_4 to C_8 cycloalkanes, or mixtures thereof. In certain embodiments, the hydrocarbon components consist of n-butane (R-600) and n-pentane (R-601). In some embodiments, the pentafluoroethane is used at from about 13% to about
25 20% by weight of the composition, and in some embodiments, the 1,1,1,2-tetrafluoroethane is used at from about 70% to about 80% by weight of the composition. In some embodiments, the hydrocarbon component is used at from about 1% to about 6% by weight.

30 Refrigerants useful in this invention can be obtained commercially, or can be synthesized by the methods described in U.S. Provisional Application No. 60/732,581.

An ionic liquid useful as a lubricant herein can in principle be any ionic liquid that absorbs a refrigerant such as a hydrofluorocarbon or CO₂. Ideally, for maximum oil return to the compressor, the ionic liquid-based lubricant should have high solubility for the refrigerant and good friction/wear characteristics.

5

Ionic liquids are organic compounds that are liquid at room temperature (approximately 25°C). They differ from most salts in that they have very low melting points, they tend to be liquid over a wide temperature range, and have been shown to have high heat capacities. Ionic liquids have essentially no vapor pressure, and they can either be neutral, acidic or basic. The properties of an ionic liquid can be tailored by varying the cation and anion. A cation or anion of an ionic liquid useful for the present invention can in principle be any cation or anion such that the cation and anion together form an organic salt that is liquid at or below about 100°C.

15

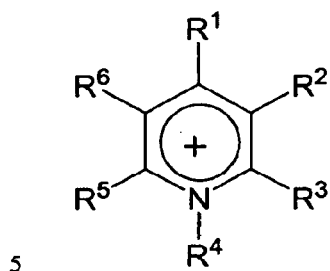
Many ionic liquids are formed by reacting a nitrogen-containing heterocyclic ring, preferably a heteroaromatic ring, with an alkylating agent (for example, an alkyl halide) to form a quaternary ammonium salt, and performing ion exchange or other suitable reactions with various Lewis acids or their conjugate bases to form the ionic liquid. Examples of suitable heteroaromatic rings include substituted pyridines, imidazole, substituted imidazole, pyrrole and substituted pyrroles. These rings can be alkylated with virtually any straight, branched or cyclic C₁₋₂₀ alkyl group, but preferably, the alkyl groups are C₁₋₁₆ groups, since groups larger than this may produce low melting solids rather than ionic liquids. Various triarylphosphines, thioethers and cyclic and non-cyclic quaternary ammonium salts may also been used for this purpose. Counterions that may be used include chloroaluminate, bromoaluminate, gallium chloride, tetrafluoroborate, tetrachloroborate, hexafluorophosphate, nitrate, trifluoromethane sulfonate, methylsulfonate, p-toluenesulfonate, hexafluoroantimonate, hexafluoroarsenate, tetrachloroaluminate, tetrabromoaluminate, perchlorate, hydroxide anion, copper dichloride anion, iron trichloride anion, zinc trichloride anion, as well as various lanthanum, potassium, lithium, nickel, cobalt, manganese, and other metal-containing anions.

30

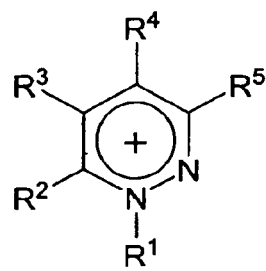
Ionic liquids may also be synthesized by salt metathesis, by an acid-base neutralization reaction or by quaternizing a selected nitrogen-containing compound; or they may be obtained commercially from several companies such
5 as Merck (Darmstadt, Germany) or BASF (Mount Olive, NJ).

Representative examples of ionic liquids useful herein included among those that are described in sources such as *J. Chem. Tech. Biotechnol.*, 68:351-356 (1997); *Chem. Ind.*, 68:249-263 (1996); *J. Phys. Condensed Matter*, 5: (supp
10 34B):B99-B106 (1993); *Chemical and Engineering News*, Mar. 30, 1998, 32-37; *J. Mater. Chem.*, 8:2627-2636 (1998); *Chem. Rev.*, 99:2071-2084 (1999); and WO 05/113,702 (and references therein cited). In one embodiment, a library, *i.e.* a combinatorial library, of ionic liquids may be prepared, for example, by preparing various alkyl derivatives of a quaternary ammonium cation, and varying
15 the associated anions. The acidity of the ionic liquids can be adjusted by varying the molar equivalents and type and combinations of Lewis acids.

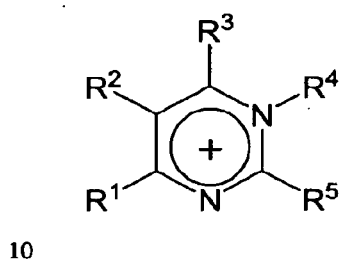
In another embodiment of the invention, ionic liquids suitable for use herein may have cations selected from the following Formulae:



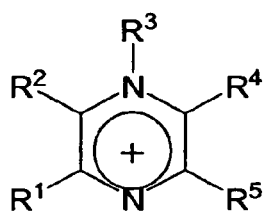
Pyridinium



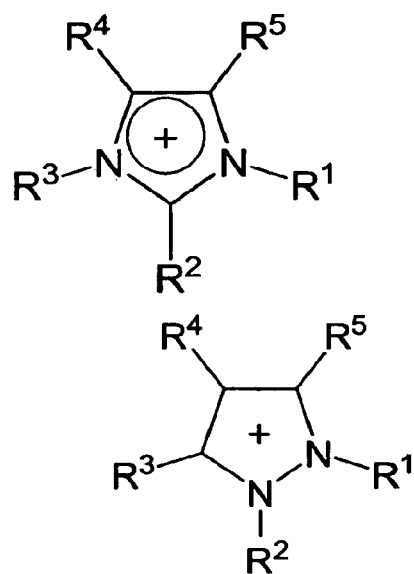
Pyridazinium



Pyrimidinium



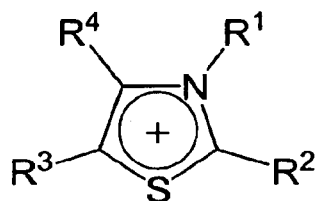
Pyrazinium



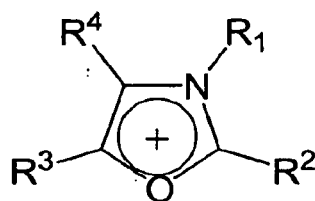
Imidazolium



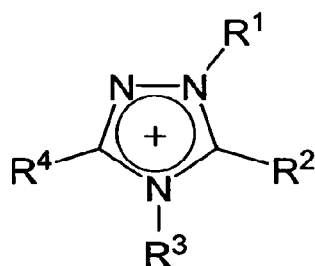
Pyrazolium



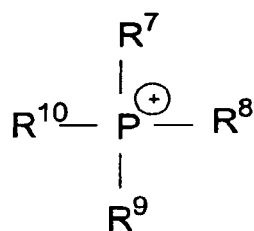
Thiazolium



Oxazolium

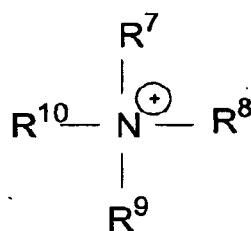


5 Triazolium



Phosphonium

and



Ammonium

wherein R^1 , R^2 , R^3 , R^4 , R^5 and R^6 are independently selected from the group consisting of:

- 10 (i) H
- (ii) halogen
- (iii) $-CH_3$, $-C_2H_5$, or C_3 to C_{25} straight-chain, branched or cyclic alkane or alkene, optionally substituted with at least one member selected from the group consisting of Cl, Br, F, I,
- 15 OH, NH_2 and SH;
- (iv) $-CH_3$, $-C_2H_5$, or C_3 to C_{25} straight-chain, branched or cyclic alkane or alkene comprising one to three heteroatoms selected from the group consisting of O, N, Si and S, and optionally substituted with at least one member selected
- 20 from the group consisting of Cl, Br, F, I, OH, NH_2 and SH;

- (v) C₆ to C₂₀ unsubstituted aryl, or C₃ to C₂₅ unsubstituted heteroaryl having one to three heteroatoms independently selected from the group consisting of O, N, Si and S; and
- (vi) C₆ to C₂₅ substituted aryl, or C₃ to C₂₅ substituted heteroaryl having one to three heteroatoms independently selected from the group consisting of O, N, Si and S; and wherein said substituted aryl or substituted heteroaryl has one to three substituents independently selected from the group consisting of:
- (1) -CH₃, -C₂H₅, or C₃ to C₂₅ straight-chain, branched or cyclic alkane or alkene, optionally substituted with at least one member selected from the group consisting of Cl, Br, F, I, OH, NH₂ and SH,
- (2) OH,
- (3) NH₂, and
- (4) SH;
- R⁷, R⁸, R⁹, and R¹⁰ are independently selected from the group consisting of:
- (vii) -CH₃, -C₂H₅, or C₃ to C₂₅ straight-chain, branched or cyclic alkane or alkene, optionally substituted with at least one member selected from the group consisting of Cl, Br, F, I, OH, NH₂ and SH;
- (viii) -CH₃, -C₂H₅, or C₃ to C₂₅ straight-chain, branched or cyclic alkane or alkene comprising one to three heteroatoms selected from the group consisting of O, N, Si and S, and optionally substituted with at least one member selected from the group consisting of Cl, Br, F, I, OH, NH₂ and SH;
- (ix) C₆ to C₂₅ unsubstituted aryl, or C₃ to C₂₅ unsubstituted heteroaryl having one to three heteroatoms independently selected from the group consisting of O, N, Si and S; and
- (x) C₆ to C₂₅ substituted aryl, or C₃ to C₂₅ substituted heteroaryl having one to three heteroatoms independently selected from the group consisting of O, N, Si and S; and wherein said substituted aryl or substituted heteroaryl has one to

three substituents independently selected from the group consisting of:

- (1) $-\text{CH}_3$, $-\text{C}_2\text{H}_5$, or C_3 to C_{25} straight-chain, branched or cyclic alkane or alkene, optionally substituted with at least one member selected from the group consisting of Cl, Br, F, I, OH, NH_2 and SH,
- (2) OH,
- (3) NH_2 , and
- (4) SH; and

wherein optionally at least two of R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , and R^{10} can together form a cyclic or bicyclic alkanyl or alkenyl group.

In another embodiment, ionic liquids useful for the invention comprise fluorinated cations wherein at least one member selected from R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 and R^{10} comprises F^- .

In another embodiment, ionic liquids have anions selected from the group consisting of $[\text{CH}_3\text{CO}_2]^-$, $[\text{HSO}_4]^-$, $[\text{CH}_3\text{OSO}_3]^-$, $[\text{C}_2\text{H}_5\text{OSO}_3]^-$, $[\text{AlCl}_4]^-$, $[\text{CO}_3]^{2-}$, $[\text{HCO}_3]^-$, $[\text{NO}_2]^-$, $[\text{NO}_3]^-$, $[\text{SO}_4]^{2-}$, $[\text{PO}_4]^{3-}$, $[\text{HPO}_4]^{2-}$, $[\text{H}_2\text{PO}_4]^-$, $[\text{HSO}_3]^-$, $[\text{CuCl}_2]^-$, Cl^- , Br^- , I^- , SCN^- ; and preferably any fluorinated anion. Fluorinated anions of the invention include $[\text{BF}_4]^-$, $[\text{PF}_6]^-$, $[\text{SbF}_6]^-$, $[\text{CF}_3\text{SO}_3]^-$, $[\text{HCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{HFCCF}_2\text{SO}_3]^-$, $[\text{HCClFCF}_2\text{SO}_3]^-$, $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CF}_2\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{SO}_2)_3\text{C}]^-$, $[\text{CF}_3\text{CO}_2]^-$, $[\text{CF}_3\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CFHOCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{HCF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{ICF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[(\text{CF}_2\text{HCF}_2\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CFHCF}_2\text{SO}_2)_2\text{N}]^-$; and F^- . In another embodiment, ionic liquids comprise a cation selected from the group consisting of pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, triazolium, phosphonium, and ammonium as defined above; and an anion selected from the group consisting of $[\text{CH}_3\text{CO}_2]^-$, $[\text{HSO}_4]^-$, $[\text{CH}_3\text{OSO}_3]^-$, $[\text{C}_2\text{H}_5\text{OSO}_3]^-$, $[\text{AlCl}_4]^-$, $[\text{CO}_3]^{2-}$, $[\text{HCO}_3]^-$, $[\text{NO}_2]^-$, $[\text{NO}_3]^-$, $[\text{SO}_4]^{2-}$, $[\text{PO}_4]^{3-}$, $[\text{HPO}_4]^{2-}$, $[\text{H}_2\text{PO}_4]^-$, $[\text{HSO}_3]^-$, $[\text{CuCl}_2]^-$, Cl^- , Br^- , I^- , SCN^- ; and any fluorinated anion. In yet another embodiment, ionic liquids comprise a cation selected from the group consisting of pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, triazolium,

phosphonium, and ammonium as defined above; and an anion selected from the group consisting of $[\text{BF}_4]^-$, $[\text{PF}_6]^-$, $[\text{SbF}_6]^-$, $[\text{CF}_3\text{SO}_3]^-$, $[\text{HCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{HFCCF}_2\text{SO}_3]^-$, $[\text{HCCIFCF}_2\text{SO}_3]^-$, $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CF}_2\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{SO}_2)_3\text{C}]^-$, $[\text{CF}_3\text{CO}_2]^-$, $[\text{CF}_3\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCFHCF}_2\text{SO}_3]^-$,
 5 $[\text{CF}_3\text{CFHOCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{HCF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{ICF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[(\text{CF}_2\text{HCF}_2\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CFHCF}_2\text{SO}_2)_2\text{N}]^-$, and F^- .

In another embodiment, ionic liquids comprise a cation selected from the group consisting of pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, triazolium, phosphonium, and
 10 ammonium as defined above, wherein at least one member selected from R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , and R^{10} comprises F^- ; and an anion selected from the group consisting of $[\text{CH}_3\text{CO}_2]^-$, $[\text{HSO}_4]^-$, $[\text{CH}_3\text{OSO}_3]^-$, $[\text{C}_2\text{H}_5\text{OSO}_3]^-$, $[\text{AlCl}_4]^-$, $[\text{CO}_3]^{2-}$, $[\text{HCO}_3]^-$, $[\text{NO}_2]^-$, $[\text{NO}_3]^-$, $[\text{SO}_4]^{2-}$, $[\text{PO}_4]^{3-}$, $[\text{HPO}_4]^{2-}$, $[\text{H}_2\text{PO}_4]^-$, $[\text{HSO}_3]^-$,
 15 $[\text{CuCl}_2]^-$, Cl^- , Br^- , I^- , SCN^- ; and any fluorinated anion. In still another embodiment, ionic liquids comprise a cation selected from the group consisting of pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, triazolium, phosphonium, and ammonium as defined above, wherein at least one member selected from R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 ,
 20 R^9 , and R^{10} comprises F^- ; and an anion selected from the group consisting of $[\text{BF}_4]^-$, $[\text{PF}_6]^-$, $[\text{SbF}_6]^-$, $[\text{CF}_3\text{SO}_3]^-$, $[\text{HCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{HFCCF}_2\text{SO}_3]^-$, $[\text{HCCIFCF}_2\text{SO}_3]^-$, $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CF}_2\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{SO}_2)_3\text{C}]^-$, $[\text{CF}_3\text{CO}_2]^-$, $[\text{CF}_3\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CFHOCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{HCF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{ICF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$,
 25 $[(\text{CF}_2\text{HCF}_2\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CFHCF}_2\text{SO}_2)_2\text{N}]^-$, and F^- .

In another embodiment, an ionic liquid comprises 1-butyl-3-methylimidazolium, 1,2-dimethyl-3-propylimidazolium, 1-octyl-3-methylimidazolium, 1,3-dioctylimidazolium, 1-ethyl-3-methylimidazolium, 1-dodecyl-3-methylimidazolium, 1-heptyl-3-methylimidazolium, 3-methyl-1-propylpyridinium, 1-butyl-3-methylpyridinium, tetradecyl(trihexyl)phosphonium, or tributyl(tetradecyl)phosphonium as the cation and an anion selected from the group consisting of $[\text{CH}_3\text{CO}_2]^-$, $[\text{HSO}_4]^-$, $[\text{CH}_3\text{OSO}_3]^-$, $[\text{C}_2\text{H}_5\text{OSO}_3]^-$, $[\text{AlCl}_4]^-$, $[\text{CO}_3]^{2-}$, $[\text{HCO}_3]^-$, $[\text{NO}_2]^-$, $[\text{NO}_3]^-$, $[\text{SO}_4]^{2-}$, $[\text{PO}_4]^{3-}$, $[\text{HPO}_4]^{2-}$, $[\text{H}_2\text{PO}_4]^-$, $[\text{HSO}_3]^-$,

$[\text{CuCl}_2]^-$, Cl^- , Br^- , I^- , SCN^- , $[\text{BF}_4]^-$, $[\text{PF}_6]^-$, $[\text{SbF}_6]^-$, $[\text{CF}_3\text{SO}_3]^-$, $[\text{HCF}_2\text{CF}_2\text{SO}_3]^-$,
 $[\text{CF}_3\text{HFCCF}_2\text{SO}_3]^-$, $[\text{HCCIFCF}_2\text{SO}_3]^-$, $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CF}_2\text{SO}_2)_2\text{N}]^-$,
 $[(\text{CF}_3\text{SO}_2)_3\text{C}]^-$, $[\text{CF}_3\text{CO}_2]^-$, $[\text{CF}_3\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCFHCF}_2\text{SO}_3]^-$,
 $[\text{CF}_3\text{CFHOCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{HCF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{ICF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$,
 5 $[\text{CF}_3\text{CF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[(\text{CF}_2\text{HCF}_2\text{SO}_2)_2\text{N}]^-$, and $[(\text{CF}_3\text{CFHCF}_2\text{SO}_2)_2\text{N}]^-$.

In an even more specific embodiment, the at least one ionic liquid is selected from the group consisting of 1-butyl-3-methylimidazolium hexafluorophosphate $[\text{bmim}][\text{PF}_6]$, 1-butyl-3-methylimidazolium tetrafluoroborate $[\text{bmim}][\text{BF}_4]$, 1,2-dimethyl-3-propylimidazolium tris(trifluoromethylsulfonyl)methide $[\text{dmpim}][\text{TMeM}]$, 1-octyl-3-methylimidazolium iodide $[\text{omim}][\text{I}]$, 1,3-dioctylimidazolium iodide $[\text{doim}][\text{I}]$, 1-ethyl-3-methylimidazolium bis(pentafluoroethylsulfonyl)imide $[\text{emim}][\text{BEI}]$, 1,2-dimethyl-3-propylimidazolium bis(trifluoromethylsulfonyl)imide $[\text{dmpim}][\text{BMeI}]$, 3-methyl-1-propylpyridinium bis(trifluoromethylsulfonyl)imide $[\text{pmpy}][\text{BMeI}]$, 1-ethyl-3-methylimidazolium hexafluorophosphate $[\text{emim}][\text{PF}_6]$, 1-ethyl-3-methylimidazolium bis(trifluoroethylsulfonyl)imide $[\text{emim}][\text{BMeI}]$, 1-butyl-3-methylpyridinium bis(trifluoromethylsulfonyl)imide $[\text{bmpy}][\text{BMeI}]$, 1-ethyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate $[\text{emim}][\text{TFES}]$, 1-butyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate $[\text{bmim}][\text{TFES}]$, 1-dodecyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate $[\text{dmim}][\text{TFES}]$, 1-heptyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate $[\text{hmim}][\text{TFES}]$, 1-butyl-3-methylimidazolium acetate $[\text{bmim}][\text{Ac}]$, 1-butyl-3-methylimidazolium 2-(1,2,2,2-tetrafluoroethoxy)-1,1,2,2-tetrafluoroethanesulfonate $[\text{bmim}][\text{FS}]$, 1-butyl-3-methylimidazolium 1, 1, 2, 3, 3, 3-hexafluoropropanesulfonate $[\text{bmim}][\text{HFPS}]$, 1-butyl-3-methylimidazolium methyl sulfonate $[\text{bmim}][\text{MeSO}_4]$, 1-butyl-3-methylimidazolium thiocyanate $[\text{bmim}][\text{SCN}]$, 1-butyl-3-methylimidazolium 1, 1, 2-trifluoro-2-(perfluoroethoxy)ethanesulfonate $[\text{bmim}][\text{TPES}]$, 1-butyl-3-methylimidazolium 1, 1, 2-trifluoro-2-(trifluoromethoxy)ethanesulfonate $[\text{bmim}][\text{TTES}]$, 1-butyl-3-methylimidazolium 1,1,2-trifluoro-2-(trifluoromethoxy)ethanesulfonate $[\text{bmim}][\text{TTES}]$, 1-butyl-3-methylimidazolium 1,1,2-trifluoro-2-(perfluoroethoxy)ethanesulfonate $[\text{bmim}][\text{TPES}]$, 1-ethyl-3-methylimidazolium bis(pentafluoroethylsulfonyl)imide $[\text{emim}][\text{BEI}]$, 1-butyl-3-methylimidazolium 1,1,2,3,3-hexafluoropropanesulfonate

[bmim][HFPS], tetradecyl(trihexyl) phosphonium 1,1,2-trifluoro-2-(perfluoroethoxy) ethanesulfonate [6,6,6,14-P][TPES], tributyl(tetradecyl)phosphonium 1,1,2,3,3,3-hexafluoropropanesulfonate [4,4,4,14-P][HFPS].

5

Mixtures of ionic liquids may also be useful as lubricants.

In another embodiment, vapor compression systems herein may utilize at least one refrigerant selected from the group consisting of CHClF_2 (R-22), CHF_3 (R-23), CH_2F_2 (R-32), CH_3F (R-41), CHF_2CF_3 (R-125), CH_2FCF_3 (R-134a), $\text{CHF}_2\text{OCHF}_2$ (E-134), CH_3CClF_2 (R-142b), CH_3CF_3 (R-143a), CH_3CHF_2 (R-152a), $\text{CH}_3\text{CH}_2\text{F}$ (R-161), CH_3OCH_3 (E170), $\text{CF}_3\text{CF}_2\text{CF}_3$ (R-218), $\text{CF}_3\text{CHFCF}_3$ (R-227ea), $\text{CF}_3\text{CH}_2\text{CF}_3$ (R-236fa), $\text{CH}_2\text{FCF}_2\text{CHF}_2$ (R-245ca), $\text{CHF}_2\text{CH}_2\text{CF}_3$ (R-245fa), $\text{CH}_3\text{CH}_2\text{CH}_3$ (R-290), $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_3$ (R-600), $\text{CH}(\text{CH}_3)_2\text{CH}_3$ (R-600a), $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3$ (R-601), $(\text{CH}_3)_2\text{CHCH}_2\text{CH}_3$ (R-601a), $\text{CH}_3\text{CH}_2\text{OCH}_2\text{CH}_3$ (R-610), NH_3 , CO_2 , and $\text{CH}_3\text{CH}=\text{CH}_2$; and at least one ionic liquid comprising pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, triazolium, phosphonium, or ammonium as the cation and an anion selected from the group consisting of $[\text{CH}_3\text{CO}_2]^-$, $[\text{HSO}_4]^-$, $[\text{CH}_3\text{OSO}_3]^-$, $[\text{C}_2\text{H}_5\text{OSO}_3]^-$, $[\text{AlCl}_4]^-$, $[\text{CO}_3]^{2-}$, $[\text{HCO}_3]^-$, $[\text{NO}_2]^-$, $[\text{NO}_3]^-$, $[\text{SO}_4]^{2-}$, $[\text{PO}_4]^{3-}$, $[\text{HPO}_4]^{2-}$, $[\text{H}_2\text{PO}_4]^-$, $[\text{HSO}_3]^-$, $[\text{CuCl}_2]^-$, Cl^- , Br^- , I^- , SCN^- , $[\text{BF}_4]^-$, $[\text{PF}_6]^-$, $[\text{SbF}_6]^-$, $[\text{CF}_3\text{SO}_3]^-$, $[\text{HCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{HFCCF}_2\text{SO}_3]^-$, $[\text{HCClFCF}_2\text{SO}_3]^-$, $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CF}_2\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{SO}_2)_3\text{C}]^-$, $[\text{CF}_3\text{CO}_2]^-$, $[\text{CF}_3\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CFHOCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{HCF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{ICF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[(\text{CF}_2\text{HCF}_2\text{SO}_2)_2\text{N}]^-$, and $[(\text{CF}_3\text{CFHCF}_2\text{SO}_2)_2\text{N}]^-$.

Vapor compression systems may also utilize at least one refrigerant blend selected from the group consisting of R-404A; R-407A; R-407B; R-407C; R-407D; R-407E; R-410A; R-410B; R-413A; 417A; R-419A; R-420A; 80.6 % R-134a and 19.4% R-142b (by weight); R-421A; R-421B; R-422A; R-422B; R-422C; R-422D; R-423A; R-424A; R-425A; R-426A; R-427A; 2.0% R-32, 41.0% R-125, 50.0% R-143a and 7.0% R-134a (by weight); 10.0% R-32, 33.0% R-125, 36.0% R-143a and 21.0% R-134a (by weight); R-428A; and R-507A; and at least

one ionic liquid comprising pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, triazolium, phosphonium, or ammonium as the cation and an anion selected from the group consisting of [CH₃CO₂]⁻, [HSO₄]⁻, [CH₃OSO₃]⁻, [C₂H₅OSO₃]⁻, [AlCl₄]⁻, [CO₃]²⁻, [HCO₃]⁻,
 5 [NO₂]⁻, [NO₃]⁻, [SO₄]²⁻, [PO₄]³⁻, [HPO₄]²⁻, [H₂PO₄]⁻, [HSO₃]⁻, [CuCl₂]⁻, Cl⁻, Br⁻, I⁻, SCN⁻, [BF₄]⁻, [PF₆]⁻, [SbF₆]⁻, [CF₃SO₃]⁻, [HCF₂CF₂SO₃]⁻, [CF₃HFCCF₂SO₃]⁻, [HCCIFCF₂SO₃]⁻, [(CF₃SO₂)₂N]⁻, [(CF₃CF₂SO₂)₂N]⁻, [(CF₃SO₂)₃C]⁻, [CF₃CO₂]⁻, [CF₃OCFHCf₂SO₃]⁻, [CF₃CF₂OCFHCf₂SO₃]⁻, [CF₃CFHOCF₂CF₂SO₃]⁻, [CF₂HCF₂OCF₂CF₂SO₃]⁻, [CF₂ICF₂OCF₂CF₂SO₃]⁻, [CF₃CF₂OCF₂CF₂SO₃]⁻,
 10 [(CF₂HCF₂SO₂)₂N]⁻, and [(CF₃CFHCF₂SO₂)₂N]⁻.

Vapor compression systems may also utilize at least one refrigerant selected from the group consisting of trifluoromethane (HFC-23), difluoromethane (HFC-32), pentafluoroethane (HFC-125), 1,1,2,2-
 15 tetrafluoroethane (HFC-134), 1,1,1,2-tetrafluoroethane (HFC-134a), 1,1,1-trifluoroethane (HFC-143a), 1,1-difluoroethane (HFC-152a), fluoroethane (HFC-161), R-404A, R-407C, R-410A, and CO₂; and at least one ionic liquid comprising pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, triazolium, phosphonium, or ammonium as
 20 the cation and an anion selected from the group consisting of [CH₃CO₂]⁻, [HSO₄]⁻, [CH₃OSO₃]⁻, [C₂H₅OSO₃]⁻, [AlCl₄]⁻, [CO₃]²⁻, [HCO₃]⁻, [NO₂]⁻, [NO₃]⁻, [SO₄]²⁻, [PO₄]³⁻, [HPO₄]²⁻, [H₂PO₄]⁻, [HSO₃]⁻, [CuCl₂]⁻, Cl⁻, Br⁻, I⁻, SCN⁻, [BF₄]⁻, [PF₆]⁻, [SbF₆]⁻, [CF₃SO₃]⁻, [HCF₂CF₂SO₃]⁻, [CF₃HFCCF₂SO₃]⁻, [HCCIFCF₂SO₃]⁻, [(CF₃SO₂)₂N]⁻, [(CF₃CF₂SO₂)₂N]⁻, [(CF₃SO₂)₃C]⁻, [CF₃CO₂]⁻,
 25 [CF₃OCFHCf₂SO₃]⁻, [CF₃CF₂OCFHCf₂SO₃]⁻, [CF₃CFHOCF₂CF₂SO₃]⁻, [CF₂HCF₂OCF₂CF₂SO₃]⁻, [CF₂ICF₂OCF₂CF₂SO₃]⁻, [CF₃CF₂OCF₂CF₂SO₃]⁻, [(CF₂HCF₂SO₂)₂N]⁻, and [(CF₃CFHCF₂SO₂)₂N]⁻.

Vapor compression systems may also utilize at least one fluoroolefin
 30 refrigerant selected from the group consisting of:

- (i) fluoroolefins of the formula *E*- or *Z*-R¹CH=CHR², wherein R¹ and R² are, independently, C₁ to C₆ perfluoroalkyl groups, and wherein the total number of carbons in the compound is at least 5;

(ii) cyclic fluoroolefins of the formula cyclo-[CX=CY(CZW)_n], wherein X, Y, Z, and W, independently, are H or F, and n is an integer from 2 to 5; and

(iii) fluoroolefins selected from the group consisting of:

- 5 2,3,3-trifluoro-1-propene (CHF₂CF=CH₂); 1,1,2-trifluoro-1-propene (CH₃CF=CF₂); 1,2,3-trifluoro-1-propene (CH₂FCF=CF₂); 1,1,3-trifluoro-1-propene (CH₂FCH=CF₂); 1,3,3-trifluoro-1-propene (CHF₂CH=CHF); 1,1,1,2,3,4,4,4-octafluoro-2-butene (CF₃CF=CFCF₃); 1,1,2,3,3,4,4,4-octafluoro-1-butene (CF₃CF₂CF=CF₂); 1,1,1,2,4,4,4-heptafluoro-2-butene (CF₃CF=CHCF₃);
- 10 1,2,3,3,4,4,4-heptafluoro-1-butene (CHF=CFCF₂CF₃); 1,1,1,2,3,4,4-heptafluoro-2-butene (CHF₂CF=CFCF₃); 1,3,3,3-tetrafluoro-2-(trifluoromethyl)-1-propene ((CF₃)₂C=CHF); 1,1,3,3,4,4,4-heptafluoro-1-butene (CF₂=CHCF₂CF₃); 1,1,2,3,4,4,4-heptafluoro-1-butene (CF₂=CFCHFCF₃); 1,1,2,3,3,4,4-heptafluoro-1-butene (CF₂=CFCF₂CHF₂); 2,3,3,4,4,4-hexafluoro-1-butene (CF₃CF₂CF=CH₂);
- 15 1,3,3,4,4,4-hexafluoro-1-butene (CHF=CHCF₂CF₃); 1,2,3,4,4,4-hexafluoro-1-butene (CHF=CFCHFCF₃); 1,2,3,3,4,4-hexafluoro-1-butene (CHF=CFCF₂CHF₂); 1,1,2,3,4,4-hexafluoro-2-butene (CHF₂CF=CFCHF₂); 1,1,1,2,3,4-hexafluoro-2-butene (CH₂FCF=CFCF₃); 1,1,1,2,4,4-hexafluoro-2-butene (CHF₂CH=CFCF₃); 1,1,1,3,4,4-hexafluoro-2-butene (CF₃CH=CFCHF₂); 1,1,2,3,3,4-hexafluoro-1-butene (CF₂=CFCF₂CH₂F); 1,1,2,3,4,4-hexafluoro-1-butene (CF₂=CFCHFCHF₂);
- 20 3,3,3-trifluoro-2-(trifluoromethyl)-1-propene (CH₂=C(CF₃)₂); 1,1,1,2,4-pentafluoro-2-butene (CH₂FCH=CFCF₃); 1,1,1,3,4-pentafluoro-2-butene (CF₃CH=CFCH₂F); 3,3,4,4,4-pentafluoro-1-butene (CF₃CF₂CH=CH₂); 1,1,1,4,4-pentafluoro-2-butene (CHF₂CH=CHCF₃); 1,1,1,2,3-pentafluoro-2-butene (CH₃CF=CFCF₃); 2,3,3,4,4-pentafluoro-1-butene (CH₂=CFCF₂CHF₂); 1,1,2,4,4-pentafluoro-2-butene (CHF₂CF=CHCHF₂); 1,1,2,3,3-pentafluoro-1-butene (CH₃CF₂CF=CF₂); 1,1,2,3,4-pentafluoro-2-butene (CH₂FCF=CFCHF₂); 1,1,3,3,3-pentafluoro-2-methyl-1-propene (CF₂=C(CF₃)(CH₃)); 2-(difluoromethyl)-3,3,3-trifluoro-1-propene (CH₂=C(CHF₂)(CF₃)); 2,3,4,4,4-pentafluoro-1-butene (CH₂=CFCHFCHFCF₃); 1,2,4,4,4-pentafluoro-1-butene (CHF=CFCH₂CF₃); 1,3,4,4,4-pentafluoro-1-butene (CHF=CHCHFCF₃); 1,3,3,4,4-pentafluoro-1-butene (CHF=CHCF₂CHF₂); 1,2,3,4,4-pentafluoro-1-butene (CHF=CFCHFCHF₂); 3,3,4,4-tetrafluoro-1-butene (CH₂=CHCF₂CHF₂); 1,1-difluoro-2-(difluoromethyl)-1-propene (CF₂=C(CHF₂)(CH₃)); 1,3,3,3-tetrafluoro-2-methyl-1-propene

- (CHF=C(CF₃)(CH₃)); 3,3-difluoro-2-(difluoromethyl)-1-propene
 (CH₂=C(CHF₂)₂); 1,1,1,2-tetrafluoro-2-butene (CF₃CF=CHCH₃); 1,1,1,3-
 tetrafluoro-2-butene (CH₃CF=CHCF₃); 1,1,1,2,3,4,4,5,5,5-decafluoro-2-pentene
 (CF₃CF=CFCF₂CF₃); 1,1,2,3,3,4,4,5,5,5-decafluoro-1-pentene
 5 (CF₂=CFCF₂CF₂CF₃); 1,1,1,4,4,4-hexafluoro-2-(trifluoromethyl)-2-butene
 ((CF₃)₂C=CHCF₃); 1,1,1,2,4,4,5,5,5-nonafluoro-2-pentene (CF₃CF=CHCF₂CF₃);
 1,1,1,3,4,4,5,5,5-nonafluoro-2-pentene (CF₃CH=CFCF₂CF₃); 1,2,3,3,4,4,5,5,5-
 nonafluoro-1-pentene (CHF=CFCF₂CF₂CF₃); 1,1,3,3,4,4,5,5,5-nonafluoro-1-
 pentene (CF₂=CHCF₂CF₂CF₃); 1,1,2,3,3,4,4,5,5,5-nonafluoro-1-pentene
 10 (CF₂=CFCF₂CF₂CHF₂); 1,1,2,3,4,4,5,5,5-nonafluoro-2-pentene
 (CHF₂CF=CFCF₂CF₃); 1,1,1,2,3,4,4,5,5,5-nonafluoro-2-pentene
 (CF₃CF=CFCF₂CHF₂); 1,1,1,2,3,4,5,5,5-nonafluoro-2-pentene
 (CF₃CF=CFCHFCF₃); 1,2,3,4,4,4-hexafluoro-3-(trifluoromethyl)-1-butene
 (CHF=CFCF(CF₃)₂); 1,1,2,4,4,4-hexafluoro-3-(trifluoromethyl)-1-butene
 15 (CF₂=CFCH(CF₃)₂); 1,1,1,4,4,4-hexafluoro-2-(trifluoromethyl)-2-butene
 (CF₃CH=C(CF₃)₂); 1,1,3,4,4,4-hexafluoro-3-(trifluoromethyl)-1-butene
 (CF₂=CHCF(CF₃)₂); 2,3,3,4,4,5,5,5-octafluoro-1-pentene (CH₂=CFCF₂CF₂CF₃);
 1,2,3,3,4,4,5,5,5-octafluoro-1-pentene (CHF=CFCF₂CF₂CHF₂); 3,3,4,4,4-
 pentafluoro-2-(trifluoromethyl)-1-butene (CH₂=C(CF₃)CF₂CF₃); 1,1,4,4,4-
 20 pentafluoro-3-(trifluoromethyl)-1-butene (CF₂=CHCH(CF₃)₂); 1,3,4,4,4-
 pentafluoro-3-(trifluoromethyl)-1-butene (CHF=CHCF(CF₃)₂); 1,1,4,4,4-
 pentafluoro-2-(trifluoromethyl)-1-butene (CF₂=C(CF₃)CH₂CF₃); 3,4,4,4-
 tetrafluoro-3-(trifluoromethyl)-1-butene ((CF₃)₂CFCH=CH₂); 3,3,4,4,5,5,5-
 heptafluoro-1-pentene (CF₃CF₂CF₂CH=CH₂); 2,3,3,4,4,5,5,5-heptafluoro-1-pentene
 25 (CH₂=CFCF₂CF₂CHF₂); 1,1,3,3,5,5,5-heptafluoro-1-butene
 (CF₂=CHCF₂CH₂CF₃); 1,1,1,2,4,4,4-heptafluoro-3-methyl-2-butene
 (CF₃CF=C(CF₃)(CH₃)); 2,4,4,4-tetrafluoro-3-(trifluoromethyl)-1-butene
 (CH₂=CFCH(CF₃)₂); 1,4,4,4-tetrafluoro-3-(trifluoromethyl)-1-butene
 (CHF=CHCH(CF₃)₂); 1,1,1,4-tetrafluoro-2-(trifluoromethyl)-2-butene
 30 (CH₂FCH=C(CF₃)₂); 1,1,1,3-tetrafluoro-2-(trifluoromethyl)-2-butene
 (CH₃CF=C(CF₃)₂); 1,1,1-trifluoro-2-(trifluoromethyl)-2-butene
 ((CF₃)₂C=CHCH₃); 3,4,4,5,5,5-hexafluoro-2-pentene (CF₃CF₂CF=CHCH₃);
 1,1,1,4,4,4-hexafluoro-2-methyl-2-butene (CF₃C(CH₃)=CHCF₃); 3,3,4,5,5,5-
 hexafluoro-1-pentene (CH₂=CHCF₂CHFCF₃); 4,4,4-trifluoro-3-(trifluoromethyl)-

- 1-butene ($\text{CH}_2=\text{C}(\text{CF}_3)\text{CH}_2\text{CF}_3$); 1,1,2,3,3,4,4,5,5,6,6,6-dodecafluoro-1-hexene
 $(\text{CF}_3(\text{CF}_2)_3\text{CF}=\text{CF}_2)$; 1,1,1,2,2,3,4,5,5,6,6,6-dodecafluoro-3-hexene
 $(\text{CF}_3\text{CF}_2\text{CF}=\text{CFCF}_2\text{CF}_3)$; 1,1,1,4,4,4-hexafluoro-2,3-bis(trifluoromethyl)-2-butene
 $((\text{CF}_3)_2\text{C}=\text{C}(\text{CF}_3)_2)$; 1,1,1,2,3,4,5,5,5-nonafluoro-4-(trifluoromethyl)-2-pentene
5 $((\text{CF}_3)_2\text{CFCF}=\text{CFCF}_3)$; 1,1,1,4,4,5,5,5-octafluoro-2-(trifluoromethyl)-2-pentene
 $((\text{CF}_3)_2\text{C}=\text{CHC}_2\text{F}_5)$; 1,1,1,3,4,5,5,5-octafluoro-4-(trifluoromethyl)-2-pentene
 $((\text{CF}_3)_2\text{CFCF}=\text{CHCF}_3)$; 3,3,4,4,5,5,6,6,6-nonafluoro-1-hexene
 $(\text{CF}_3\text{CF}_2\text{CF}_2\text{CF}_2\text{CH}=\text{CH}_2)$; 4,4,4-trifluoro-3,3-bis(trifluoromethyl)-1-butene
 $(\text{CH}_2=\text{CHC}(\text{CF}_3)_3)$; 1,1,1,4,4,4-hexafluoro-2-(trifluoromethyl)-3-methyl-2-butene
10 $((\text{CF}_3)_2\text{C}=\text{C}(\text{CH}_3)(\text{CF}_3))$; 2,3,3,5,5,5-hexafluoro-4-(trifluoromethyl)-1-pentene
 $(\text{CH}_2=\text{CFCF}_2\text{CH}(\text{CF}_3)_2)$; 1,1,1,2,4,4,5,5,5-nonafluoro-3-methyl-2-pentene
 $(\text{CF}_3\text{CF}=\text{C}(\text{CH}_3)\text{CF}_2\text{CF}_3)$; 1,1,1,5,5,5-hexafluoro-4-(trifluoromethyl)-2-pentene
 $(\text{CF}_3\text{CH}=\text{CHCH}(\text{CF}_3)_2)$; 3,4,4,5,5,6,6,6-octafluoro-2-hexene
 $(\text{CF}_3\text{CF}_2\text{CF}_2\text{CF}=\text{CHCH}_3)$; 3,3,4,4,5,5,6,6-octafluoro-1-hexene
15 $(\text{CH}_2=\text{CHCF}_2\text{CF}_2\text{CF}_2\text{CHF}_2)$; 1,1,1,4,4-pentafluoro-2-(trifluoromethyl)-2-pentene
 $((\text{CF}_3)_2\text{C}=\text{CHCF}_2\text{CH}_3)$; 4,4,5,5,5-pentafluoro-2-(trifluoromethyl)-1-pentene
 $(\text{CH}_2=\text{C}(\text{CF}_3)\text{CH}_2\text{C}_2\text{F}_5)$; 3,3,4,4,5,5,5-heptafluoro-2-methyl-1-pentene
 $(\text{CF}_3\text{CF}_2\text{CF}_2\text{C}(\text{CH}_3)=\text{CH}_2)$; 4,4,5,5,6,6,6-heptafluoro-2-hexene
 $(\text{CF}_3\text{CF}_2\text{CF}_2\text{CH}=\text{CHCH}_3)$; 4,4,5,5,6,6,6-heptafluoro-1-hexene
20 $(\text{CH}_2=\text{CHCH}_2\text{CF}_2\text{C}_2\text{F}_5)$; 1,1,1,2,2,3,4-heptafluoro-3-hexene
 $(\text{CF}_3\text{CF}_2\text{CF}=\text{CFC}_2\text{H}_5)$; 4,5,5,5-tetrafluoro-4-(trifluoromethyl)-1-pentene
 $(\text{CH}_2=\text{CHCH}_2\text{CF}(\text{CF}_3)_2)$; 1,1,1,2,5,5,5-heptafluoro-4-methyl-2-pentene
 $(\text{CF}_3\text{CF}=\text{CHCH}(\text{CF}_3)(\text{CH}_3))$; 1,1,1,3-tetrafluoro-2-(trifluoromethyl)-2-pentene
 $((\text{CF}_3)_2\text{C}=\text{CFC}_2\text{H}_5)$; 1,1,1,2,3,4,4,5,5,6,6,7,7,7-tetradecafluoro-2-heptene
25 $(\text{CF}_3\text{CF}=\text{CFCF}_2\text{CF}_2\text{C}_2\text{F}_5)$; 1,1,1,2,2,3,4,5,5,6,6,7,7,7-tetradecafluoro-3-heptene
 $(\text{CF}_3\text{CF}_2\text{CF}=\text{CFCF}_2\text{C}_2\text{F}_5)$; 1,1,1,3,4,4,5,5,6,6,7,7,7-tridecafluoro-2-heptene
 $(\text{CF}_3\text{CH}=\text{CFCF}_2\text{CF}_2\text{C}_2\text{F}_5)$; 1,1,1,2,4,4,5,5,6,6,7,7,7-tridecafluoro-2-heptene
 $(\text{CF}_3\text{CF}=\text{CHCF}_2\text{CF}_2\text{C}_2\text{F}_5)$; 1,1,1,2,2,4,5,5,6,6,7,7,7-tridecafluoro-3-heptene
 $(\text{CF}_3\text{CF}_2\text{CH}=\text{CFCF}_2\text{C}_2\text{F}_5)$; 1,1,1,2,2,3,5,5,6,6,7,7,7-tridecafluoro-3-heptene
30 $(\text{CF}_3\text{CF}_2\text{CF}=\text{CHCF}_2\text{C}_2\text{F}_5)$; $\text{CF}_2=\text{CFOCF}_2\text{CF}_3$ (PEVE) and $\text{CF}_2=\text{CFOCF}_3$ (PMVE); and at least one ionic liquid comprising pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, triazolium, phosphonium, or ammonium as the cation and an anion selected from the group consisting of $[\text{CH}_3\text{CO}_2]^-$, $[\text{HSO}_4]^-$, $[\text{CH}_3\text{OSO}_3]^-$, $[\text{C}_2\text{H}_5\text{OSO}_3]^-$, $[\text{AlCl}_4]^-$,

$[\text{CO}_3]^{2-}$, $[\text{HCO}_3]^-$, $[\text{NO}_2]^-$, $[\text{NO}_3]^-$, $[\text{SO}_4]^{2-}$, $[\text{PO}_4]^{3-}$, $[\text{HPO}_4]^{2-}$, $[\text{H}_2\text{PO}_4]^-$, $[\text{HSO}_3]^-$,
 $[\text{CuCl}_2]^-$, Cl^- , Br^- , I^- , SCN^- , $[\text{BF}_4]^-$, $[\text{PF}_6]^-$, $[\text{SbF}_6]^-$, $[\text{CF}_3\text{SO}_3]^-$, $[\text{HCF}_2\text{CF}_2\text{SO}_3]^-$,
 $[\text{CF}_3\text{HFCCF}_2\text{SO}_3]^-$, $[\text{HCClFCF}_2\text{SO}_3]^-$, $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CF}_2\text{SO}_2)_2\text{N}]^-$,
 $[(\text{CF}_3\text{SO}_2)_3\text{C}]^-$, $[\text{CF}_3\text{CO}_2]^-$, $[\text{CF}_3\text{OCFHC}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCFHC}_2\text{SO}_3]^-$,
5 $[\text{CF}_3\text{CFHOC}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{HCF}_2\text{OC}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{ICF}_2\text{OC}_2\text{CF}_2\text{SO}_3]^-$,
 $[\text{CF}_3\text{CF}_2\text{OC}_2\text{CF}_2\text{SO}_3]^-$, $[(\text{CF}_2\text{HCF}_2\text{SO}_2)_2\text{N}]^-$, and $[(\text{CF}_3\text{CFHCF}_2\text{SO}_2)_2\text{N}]^-$.

Vapor compression systems may also utilize at least one refrigerant comprising pentafluoroethane (R-125), 1,1,1,2-tetrafluoroethane (R-134a), and at
 10 least two hydrocarbons each having eight or fewer carbon atoms, wherein said at least two hydrocarbons can be C_4 to C_8 hydrocarbons, and more specifically n-butane and n-pentane; and at least one ionic liquid comprising pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, triazolium, phosphonium, or ammonium as the cation and an anion
 15 selected from the group consisting of $[\text{CH}_3\text{CO}_2]^-$, $[\text{HSO}_4]^-$, $[\text{CH}_3\text{OSO}_3]^-$, $[\text{C}_2\text{H}_5\text{OSO}_3]^-$, $[\text{AlCl}_4]^-$, $[\text{CO}_3]^{2-}$, $[\text{HCO}_3]^-$, $[\text{NO}_2]^-$, $[\text{NO}_3]^-$, $[\text{SO}_4]^{2-}$, $[\text{PO}_4]^{3-}$, $[\text{HPO}_4]^{2-}$, $[\text{H}_2\text{PO}_4]^-$, $[\text{HSO}_3]^-$, $[\text{CuCl}_2]^-$, Cl^- , Br^- , I^- , SCN^- , $[\text{BF}_4]^-$, $[\text{PF}_6]^-$, $[\text{SbF}_6]^-$, $[\text{CF}_3\text{SO}_3]^-$, $[\text{HCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{HFCCF}_2\text{SO}_3]^-$, $[\text{HCClFCF}_2\text{SO}_3]^-$, $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CF}_2\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{SO}_2)_3\text{C}]^-$, $[\text{CF}_3\text{CO}_2]^-$, $[\text{CF}_3\text{OCFHC}_2\text{SO}_3]^-$,
 20 $[\text{CF}_3\text{CF}_2\text{OCFHC}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CFHOC}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{HCF}_2\text{OC}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{ICF}_2\text{OC}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OC}_2\text{CF}_2\text{SO}_3]^-$, $[(\text{CF}_2\text{HCF}_2\text{SO}_2)_2\text{N}]^-$, and $[(\text{CF}_3\text{CFHCF}_2\text{SO}_2)_2\text{N}]^-$.

Vapor compression systems may also utilize at least one refrigerant
 25 selected from the group consisting of trifluoromethane (HFC-23), difluoromethane (HFC-32), pentafluoroethane (HFC-125), 1,1,2,2-tetrafluoroethane (HFC-134), 1,1,1,2-tetrafluoroethane (HFC-134a), 1,1,1-trifluoroethane (HFC-143a), 1,1-difluoroethane (HFC-152a), fluoroethane (HFC-161), R-404A, R-407C, HFC-410A, and CO_2 ; and at least one ionic liquid
 30 selected from the group consisting of 1-butyl-3-methylimidazolium hexafluorophosphate $[\text{bmim}][\text{PF}_6]$, 1-butyl-3-methylimidazolium tetrafluoroborate $[\text{bmim}][\text{BF}_4]$, 1,2-dimethyl-3-propylimidazolium tris(trifluoromethylsulfonyl)methide $[\text{dmpim}][\text{TMeM}]$, 1-octyl-3-methylimidazolium iodide $[\text{omim}][\text{I}]$, 1,3-dioctylimidazolium iodide $[\text{doim}][\text{I}]$, 1-

ethyl-3-methylimidazolium bis(pentafluoroethylsulfonyl)imide [emim][BEI], 1,2-dimethyl-3-propylimidazolium bis(trifluoromethylsulfonyl)imide [dmpim][BMeI], 3-methyl-1-propylpyridinium bis(trifluoromethylsulfonyl)imide [pmpy][BMeI], 1-ethyl-3-methylimidazolium hexafluorophosphate [emim][PF₆],

5 1-ethyl-3-methylimidazolium bis(trifluoroethylsulfonyl)imide [emim][BMeI], 1-butyl-3-methylpyridinium bis(trifluoromethylsulfonyl)imide [bmpy][BMeI], 1-ethyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate [emim][TFES], 1-butyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate [bmim][TFES], 1-dodecyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate [dmim][TFES],

10 1-heptyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate [hmim][TFES], 1-butyl-3-methylimidazolium acetate [bmim][Ac], 1-butyl-3-methylimidazolium 2-(1,2,2,2-tetrafluoroethoxy)-1,1,2,2-tetrafluoroethanesulfonate [bmim][FS], 1-butyl-3-methylimidazolium 1, 1, 2, 3, 3, 3-hexafluoropropanesulfonate [bmim][HFPS], 1-butyl-3-methylimidazolium methyl sulfonate [bmim][MeSO₄],

15 1-butyl-3-methylimidazolium thiocyanate [bmim][SCN], 1-butyl-3-methylimidazolium 1, 1, 2-trifluoro-2-(perfluoroethoxy)ethanesulfonate [bmim][TPES], 1-butyl-3-methylimidazolium 1, 1, 2-trifluoro-2-(trifluoromethoxy)ethanesulfonate [bmim][TTES], 1-butyl-3-methylimidazolium 1,1,2-trifluoro-2-(trifluoromethoxy)ethanesulfonate [bmim][TTES], 1-butyl-3-

20 methylimidazolium 1,1,2-trifluoro-2-(perfluoroethoxy)ethanesulfonate [bmim][TPES], 1-ethyl-3-methylimidazolium bis(pentafluoroethylsulfonyl)imide [emim][BEI], 1-butyl-3-methylimidazolium 1,1,2,3,3-hexafluoropropanesulfonate [bmim][HFPS], tetradecyl(trihexyl) phosphonium 1,1,2-trifluoro-2-(perfluoroethoxy) ethanesulfonate [6,6,6,14-P][TPES], and

25 tributyl(tetradecyl)phosphonium 1,1,2,3,3,3-hexafluoropropanesulfonate [4,4,4,14-P][HFPS].

In another embodiment, combinations of refrigerant and lubricant useful for vapor compression systems may include difluoromethane (HFC-32) and 1-

30 butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆], difluoromethane (HFC-32) and 1-butyl-3-methylimidazolium tetrafluoroborate [bmim][BF₄], pentafluoroethane (HFC-125) and 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆], 1,1,1,2-tetrafluoroethane (HFC-134a) and 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆], 1,1,1-

trifluoroethane (HFC-143a) and 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆], 1,1-difluoroethane (HFC-152a) and 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆], difluoromethane (HFC-32) and 1,2-dimethyl-3-propylimidazolium tris(trifluoromethylsulfonyl)methide

5 [dmpim][TMeM], difluoromethane (HFC-32) and 1-octyl-3-methylimidazolium iodide [omim][I], difluoromethane (HFC-32) and 1,3-dioctylimidazolium iodide [doim][I], difluoromethane (HFC-32) and 1-ethyl-3-methylimidazolium bis(pentafluoroethylsulfonyl)imide [emim][BEI], difluoromethane (HFC-32) and 1,2-dimethyl-3-propylimidazolium bis(trifluoromethylsulfonyl)imide

10 [dmpim][BMeI], difluoromethane (HFC-32) and 3-methyl-1-propylpyridinium bis(trifluoromethylsulfonyl)imide [pmpy][BMeI], trifluoromethane (HFC-23) and 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆], trifluoromethane (HFC-23) and 1-ethyl-3-methylimidazolium hexafluorophosphate [emim][PF₆], difluoromethane (HFC-32) and 1-ethyl-3-

15 methylimidazolium bis(trifluoroethylsulfonyl)imide [emim][BMeI], difluoromethane (HFC-32) and 1-butyl-3-methylpyridinium bis(trifluoromethylsulfonyl)imide [bmpy][BMeI], difluoromethane (HFC-32) and 1-ethyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate [emim][TFES], difluoromethane (HFC-32) and 1-butyl-3-methylimidazolium 1,1,2,2-

20 tetrafluoroethanesulfonate [bmim][TFES], difluoromethane (HFC-32) and 1-dodecyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate [dmim][TFES], difluoromethane (HFC-32) and 1-heptyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate [hmim][TFES], difluoromethane (HFC-32) and 1-

butyl-3-methylimidazolium acetate [bmim][Ac], (HFC-32) and 1-butyl-3-

25 methylimidazolium 2-(1,2,2,2-tetrafluoroethoxy)-1,1,2,2-tetrafluoroethanesulfonate [bmim][FS], difluoromethane (HFC-32) and 1-butyl-3-methylimidazolium 1, 1, 2, 3, 3, 3-hexafluoropropanesulfonate [bmim][HFPS], difluoromethane (HFC-32) and 1-butyl-3-methylimidazolium methyl sulfonate [bmim][MeSO₄], difluoromethane (HFC-32) and 1-butyl-3-methylimidazolium

30 thiocyanate [bmim][SCN], difluoromethane (HFC-32) and 1-butyl-3-methylimidazolium 1, 1, 2-trifluoro-2-(perfluoroethoxy)ethanesulfonate [bmim][TPES], difluoromethane (HFC-32) and 1-butyl-3-methylimidazolium 1, 1, 2-trifluoro-2-(trifluoromethoxy)ethanesulfonate [bmim][TTES], 1,1,1,2-tetrafluoroethane (HFC-134a) and 1-butyl-3-methylimidazolium 1,1,2-trifluoro-2-

(trifluoromethoxy)ethanesulfonate [bmim][TTES], 1,1,1,2-tetrafluoroethane (HFC-134a) and 1-butyl-3-methylimidazolium 1,1,2-trifluoro-2-(perfluoroethoxy)ethanesulfonate [bmim][TPES], 1,1,1,2-tetrafluoroethane (HFC-134a) and 1-ethyl-3-methylimidazolium bis(pentafluoroethylsulfonyl)imide [emim][BEI], 1,1,1,2-tetrafluoroethane (HFC-134a) and 1-butyl-3-methylimidazolium 1,1,2,3,3-hexafluoropropanesulfonate [bmim][HFPS], 1,1,1,2-tetrafluoroethane (HFC-134a) and tetradecyl(trihexyl) phosphonium 1,1,2-trifluoro-2-(perfluoroethoxy) ethanesulfonate [6,6,6,14-P][TPES], 1,1,1,2-tetrafluoroethane (HFC-134a) and tributyl(tetradecyl)phosphonium 1,1,2,3,3,3-hexafluoropropanesulfonate [4,4,4,14-P][HFPS], carbon dioxide (CO₂) and 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆], carbon dioxide (CO₂) and 1-butyl-3-methylimidazolium tetrafluoroborate [bmim][BF₄].

In alternative embodiments of this invention, a refrigerant may be any one or more of all of the members of the total group of refrigerants disclosed herein. In those embodiments, the refrigerant may also, however, be any one or more of those members of a subgroup of the total group of refrigerants disclosed herein, where the subgroup is formed by excluding any one or more other members from the total group. As a result, the refrigerant in those embodiments may not only be any one or more of the refrigerants in any subgroup of any size that may be selected from the total group of refrigerants in all the various different combinations of individual members of the total group, but the members in any subgroup may thus be used in the absence of one or more of the members of the total group that have been excluded to form the subgroup. The subgroup formed by excluding various members from the total group of refrigerants may, moreover, be an individual member of the total group such that that refrigerant is used in the absence of all other members of the total group except the selected individual member.

Correspondingly, in further alternative embodiments of this invention, an ionic liquid may be any one or more of all of the members of the total group of ionic liquids disclosed herein. In those embodiments, the liquid may also, however, be any one or more of those members of a subgroup of the total group of ionic liquids disclosed herein, where the subgroup is formed by excluding any one

or more other members from the total group. As a result, the ionic liquid in those embodiments may not only be any one or more of the ionic liquids in any subgroup of any size that may be selected from the total group of ionic liquids in all the various different combinations of individual members of the total group, 5 but the members in any subgroup may thus be used in the absence of one or more of the members of the total group that have been excluded to form the subgroup. The subgroup formed by excluding various members from the total group of ionic liquids may, moreover, be an individual member of the total group such that that ionic liquid is used in the absence of all other members of the total group except 10 the selected individual member.

As a result, in yet other alternative embodiments of this invention, pairings of one or more particular refrigerants and one or more particular ionic liquids may be formed from (i) any one or more of all of the members of the total group of 15 refrigerants disclosed herein, selected as described above as a single member or any subgroup of any size taken from the total group of refrigerants in all the various different combinations of individual members of that total group, together with (ii) any one or more of all of the members of the total group of ionic liquids disclosed herein, selected as described above as a single member or any subgroup 20 of any size taken from the total group of ionic liquids in all the various different combinations of individual members of that total group.

The following examples are presented to illustrate the advantages of the present invention and to assist one of ordinary skill in making and using the same. These examples are not intended in any way otherwise to limit the scope of the disclosure or the appended claims. The operation of this invention is illustrated in terms of the extent of solubility of various refrigerants and various ionic liquids in or with each other.

GENERAL METHODS AND MATERIALS

10

1-Butyl-3-methylimidazolium hexafluorophosphate ([bmim][PF₆], C₈H₁₅N₂F₆P, molecular weight 284 g mol⁻¹), 1-butyl-3-methylimidazolium tetrafluoroborate ([bmim][BF₄], C₈H₁₅N₂F₄B, molecular weight 226 g mol⁻¹), 1,2-dimethyl-3-propylimidazolium tris(trifluoromethylsulfonyl)methide ([dmpim][tTFMSmethide], C₁₂H₁₅N₂F₉O₆S₃, molecular weight 550 g mol⁻¹), 1,2-dimethyl-3-propylimidazolium bis(trifluoromethylsulfonyl)imide ([dmpim][bTFMSimide], C₁₀H₁₅N₃F₆O₄S₂, molecular weight 419 g mol⁻¹), 1-ethyl-3-methylimidazolium bis(pentafluoroethylsulfonyl)imide ([emim][bPFESimide], C₁₀H₁₁N₃F₁₀O₄S₂, molecular weight 491.33 g mol⁻¹), 1-propyl-3-methylpyridinium bis(trifluoromethylsulfonyl)imide ([pmpy][bTFMSimide], C₁₁H₁₄N₂F₆O₄S₂, molecular weight 416.36 g mol⁻¹), 1-ethyl-3-methylimidazolium hexafluorophosphate ([emim][PF₆], C₆H₁₁F₆N₂P, molecular weight 265.13 g mol⁻¹), 1-ethyl-3-methylimidazolium bis(trifluoromethylsulfonyl)imide ([emim][BMeI], C₈H₁₁F₆N₃O₄S₂, molecular weight 197.98 g mol⁻¹), 1-butyl-3-methylpyridinium bis(trifluoromethylsulfonyl)imide ([BMPy][bTFMSimide], C₁₂H₁₆F₆N₂O₄S₂, molecular weight 430.39 g mol⁻¹) were each obtained from Fluka Chemika (may be obtained from Sigma-Aldrich, St. Louis, Missouri) with a purity of >96 to 97% each. Trifluoromethane (HFC-23), difluoromethane (HFC-32, CH₂F₂, molecular weight 52.02 g mol⁻¹), pentafluoroethane (HFC-125, C₂HF₅, molecular weight 120.02 g mol⁻¹), 1,1,1,2-tetrafluoroethane (HFC-134a, C₂H₂F₄, molecular weight 102.03 g mol⁻¹), 1,1,1-trifluoroethane (HFC-143a, C₂H₃F₃, molecular weight 82.04 g mol⁻¹), and 1,1-difluoroethane (HFC-152a, C₂H₄F₂, molecular weight 66.05 g mol⁻¹) were obtained from DuPont Fluorochemicals (Wilmington,

Delaware), with a minimum purity of 99.99%. A molecular sieve trap was installed to remove trace amounts of water from the gases and each of the ionic liquids tested were degassed prior to making solubility measurements.

5 The following nomenclature and abbreviations are used:

- C = concentration ($\text{mol}\cdot\text{m}^{-3}$)
- C_b = buoyancy force (N)
- C_f = correction factor (kg)
- 10 C_0 = initial concentration ($\text{mol}\cdot\text{m}^{-3}$)
- C_s = saturation concentration ($\text{mol}\cdot\text{m}^{-3}$)
- $\langle C \rangle$ = space-averaged concentration ($\text{mol}\cdot\text{m}^{-3}$)
- D = diffusion constant ($\text{m}^2\cdot\text{s}^{-1}$)
- g = gravitational acceleration ($9.80665 \text{ m}\cdot\text{s}^{-2}$)
- 15 L = length (m)
- m_a = mass absorbed (kg)
- m_i = mass of i -th species on sample side of balance (kg)
- m_j = mass of j -th species on counterweight side of balance (kg)
- m_{il} = mass of ionic liquid sample (kg)
- 20 MW_i = molecular weight of i -th species ($\text{kg}\cdot\text{mol}^{-1}$)
- N = n -th number component
- P = pressure (MPa)
- P_0 = initial pressure (MPa)
- t = time (s)
- 25 T_{ci} = critical temperature of i -th species (K)
- T_i = temperature of i -th species (K)
- T_j = temperature of j -th species (K)
- T_s = temperature of sample (K)
- V_i = volume of i -th species (m^3)
- 30 V_{il} = volume of ionic liquid (m^3)

- V_m = liquid sample volume (m^3)
 $\tilde{V}_g$ = molar volume of gas ($\text{m}^3 \cdot \text{mol}^{-1}$)
 $\tilde{V}_i$ = molar volume of i -th species ($\text{m}^3 \cdot \text{mol}^{-1}$)
 $\tilde{V}_{IL}$ = molar volume of ionic liquid ($\text{m}^3 \cdot \text{mol}^{-1}$)
 5 $\tilde{V}_m$ = molar volume of mixture ($\text{m}^3 \cdot \text{mol}^{-1}$)
 $\tilde{V}_0$ = initial molar volume ($\text{m}^3 \cdot \text{mol}^{-1}$)
 $\Delta \tilde{V}$ = change in molar volume ($\text{m}^3 \cdot \text{mol}^{-1}$)
 x_i = mole fraction of i -th species
 z = depth (m)
 10 λ_n = eigenvalue (m^{-1})
 ρ_g = density of gas ($\text{kg} \cdot \text{m}^{-3}$)
 ρ_i = density of i -th component on sample side of balance ($\text{kg} \cdot \text{m}^{-3}$)
 ρ_j = density of j -th component on counter weight side of balance ($\text{kg} \cdot \text{m}^{-3}$)
 ρ_{air} = density of air ($\text{kg} \cdot \text{m}^{-3}$)
 15 ρ_s = density of sample ($\text{kg} \cdot \text{m}^{-3}$)

Units

- Pa $\equiv$ Pascal
 20 MPa $\equiv$ Mega Pascal
 mol $\equiv$ mole
 m $\equiv$ meter
 cm $\equiv$ centimeter
 K $\equiv$ Kelvin
 25 N $\equiv$ Newton
 J $\equiv$ Joule
 kJ $\equiv$ kilojoule
 kg $\equiv$ kilogram
 mg $\equiv$ milligram

μg $\equiv$ microgram

T $\equiv$ temperature

P $\equiv$ pressure

mbar $\equiv$ millibar

5 min $\equiv$ minute

$^{\circ}\text{C}$ or C $\equiv$ degrees Centigrade

$^{\circ}\text{F}$ $\equiv$ degrees Fahrenheit

sec $\equiv$ second

kW $\equiv$ kilowatt

10 kg/s $\equiv$ kilogram/second

In the following description, (A)-(D) provide syntheses for anions of ionic liquids that are useful as lubricants for the invention, and (E) -(W) provide syntheses for ionic liquids useful as lubricants for the invention.

15

Preparation of Anions Not Generally Available Commercially

(A) Synthesis of potassium 1,1,2,2-tetrafluoroethanesulfonate (TFES-K) ([HCF₂CF₂SO₃]⁻):

A 1-gallon Hastelloy® C276 reaction vessel was charged with a solution of potassium sulfite hydrate (176 g, 1.0 mol), potassium metabisulfite (610 g, 2.8 mol) and deionized water (2000 ml). The pH of this solution was 5.8. The vessel was cooled to 18 degrees C, evacuated to 0.10 MPa, and purged with nitrogen. The evacuate/purge cycle was repeated two more times. To the vessel was then added tetrafluoroethylene (TFE, 66 g), and it was heated to 100 degrees C at which time the inside pressure was 1.14 MPa. The reaction temperature was increased to 125 degrees C and kept there for 3 h. As the TFE pressure decreased due to the reaction, more TFE was added in small aliquots (20-30 g each) to maintain operating pressure roughly between 1.14 and 1.48 MPa. Once 500 g (5.0 mol) of TFE had been fed after the initial 66 g precharge, the vessel was vented and cooled to 25 degrees C. The pH of the clear light yellow reaction solution was 10-11. This solution was buffered to pH 7 through the addition of potassium metabisulfite (16 g).

The water was removed *in vacuo* on a rotary evaporator to produce a wet solid. The solid was then placed in a freeze dryer (Virtis Freezemobile 35x1; Gardiner, NY) for 72 hr to reduce the water content to approximately 1.5 wt % (1387 g crude material). The theoretical mass of total solids was 1351 g. The mass balance was very close to ideal and the isolated solid had slightly higher mass due to moisture. This added freeze drying step had the advantage of producing a free-flowing white powder whereas treatment in a vacuum oven resulted in a soapy solid cake that was very difficult to remove and had to be chipped and broken out of the flask.

The crude TFES-K can be further purified and isolated by extraction with reagent grade acetone, filtration, and drying.

^{19}F NMR (D_2O) δ , -122.0 (dt, $J_{\text{FH}} = 6$ Hz, $J_{\text{FF}} = 6$ Hz, 2F); -136.1 (dt, $J_{\text{FH}} = 53$ Hz, 2F).

^1H NMR (D_2O) δ , 6.4 (tt, $J_{\text{FH}} = 53$ Hz, $J_{\text{FH}} = 6$ Hz, 1H).

% Water by Karl-Fisher titration: 580 ppm.

Analytical calculation for $\text{C}_2\text{HO}_3\text{F}_4\text{SK}$: C, 10.9: H, 0.5: N, 0.0 Experimental results: C, 11.1: H, 0.7: N, 0.2.

Mp (DSC): 242 degrees C.

TGA (air): 10% wt. loss @ 367 degrees C, 50% wt. loss @ 375 degrees C.

TGA (N_2): 10% wt. loss @ 363 degrees C, 50% wt. loss @ 375 degrees C.

(B) Synthesis of potassium-1,1,2-trifluoro-2-(perfluoroethoxy)ethanesulfonate (TPES-K):

A 1-gallon Hastelloy® C276 reaction vessel was charged with a solution of potassium sulfite hydrate (88 g, 0.56 mol), potassium metabisulfite (340 g, 1.53 mol) and deionized water (2000 ml). The vessel was cooled to 7 degrees C, evacuated to 0.05 MPa, and purged with nitrogen. The evacuate/purge cycle was repeated two more times. To the vessel was then added perfluoro(ethylvinyl ether) (PEVE, 600 g, 2.78 mol), and it was heated to 125 degrees C at which time

the inside pressure was 2.31 MPa. The reaction temperature was maintained at 125 degrees C for 10 hr. The pressure dropped to 0.26 MPa at which point the vessel was vented and cooled to 25 degrees C. The crude reaction product was a white crystalline precipitate with a colorless aqueous layer (pH = 7) above it.

5

The ^{19}F NMR spectrum of the white solid showed pure desired product, while the spectrum of the aqueous layer showed a small but detectable amount of a fluorinated impurity. The desired isomer is less soluble in water so it precipitated in isomerically pure form.

10

The product slurry was suction filtered through a fritted glass funnel, and the wet cake was dried in a vacuum oven (60 degrees C, 0.01 MPa) for 48 hr. The product was obtained as off-white crystals (904 g, 97% yield).

15 ^{19}F NMR (D_2O) δ -86.5 (s, 3F); -89.2, -91.3 (subsplit ABq, $J_{\text{FF}} = 147$ Hz, 2F); -119.3, -121.2 (subsplit ABq, $J_{\text{FF}} = 258$ Hz, 2F); -144.3 (dm, $J_{\text{FH}} = 53$ Hz, 1F).
 ^1H NMR (D_2O) δ 6.7 (dm, $J_{\text{FH}} = 53$ Hz, 1H).

Mp (DSC) 263 degrees C.

Analytical calculation for $\text{C}_4\text{HO}_4\text{F}_8\text{SK}$: C, 14.3; H, 0.3 Experimental results: C, 14.1; H, 0.3.

20

TGA (air): 10% wt. loss @ 359 degrees C, 50% wt. loss @ 367 degrees C.

TGA (N_2): 10% wt. loss @ 362 degrees C, 50% wt. loss @ 374 degrees C.

25

(C) Synthesis of potassium-1,1,2-trifluoro-2-(trifluoromethoxy)ethanesulfonate (TTES-K)

A 1-gallon Hastelloy® C276 reaction vessel was charged with a solution of potassium sulfite hydrate (114 g, 0.72 mol), potassium metabisulfite (440 g, 1.98 mol) and deionized water (2000 ml). The pH of this solution was 5.8. The vessel was cooled to -35 degrees C, evacuated to 0.08 MPa, and purged with nitrogen. The evacuate/purge cycle was repeated two more times. To the vessel was then added perfluoro(methylvinyl ether) (PMVE, 600 g, 3.61 mol) and it was heated to 125 degrees C at which time the inside pressure was 3.29 MPa. The

35

reaction temperature was maintained at 125 degrees C for 6 hr. The pressure dropped to 0.27 MPa at which point the vessel was vented and cooled to 25 degrees C. Once cooled, a white crystalline precipitate of the desired product formed leaving a colorless clear aqueous solution above it (pH = 7).

5

The ^{19}F NMR spectrum of the white solid showed pure desired product, while the spectrum of the aqueous layer showed a small but detectable amount of a fluorinated impurity.

10

The solution was suction filtered through a fritted glass funnel for 6 hr to remove most of the water. The wet cake was then dried in a vacuum oven at 0.01 MPa and 50 degrees C for 48 h. This gave 854 g (83% yield) of a white powder. The final product was isomerically pure (by ^{19}F and ^1H NMR) since the undesired isomer remained in the water during filtration.

15

^{19}F NMR (D_2O) δ -59.9 (d, $J_{\text{FH}} = 4$ Hz, 3F); -119.6, -120.2 (subsplit ABq, $J = 260$ Hz, 2F); -144.9 (dm, $J_{\text{FH}} = 53$ Hz, 1F).

^1H NMR (D_2O) δ 6.6 (dm, $J_{\text{FH}} = 53$ Hz, 1H).

% Water by Karl-Fisher titration: 71 ppm.

20

Analytical calculation for $\text{C}_3\text{HF}_6\text{SO}_4\text{K}$: C, 12.6: H, 0.4: N, 0.0 Experimental results: C, 12.6: H, 0.0: N, 0.1.

Mp (DSC) 257 degrees C.

TGA (air): 10% wt. loss @ 343 degrees C, 50% wt. loss @ 358 degrees C.

25

TGA (N_2): 10% wt. loss @ 341 degrees C, 50% wt. loss @ 357 degrees C.

(D) Synthesis of sodium 1,1,2,3,3,3-hexafluoropropanesulfonate (HFPS-Na)

30

A 1-gallon Hastelloy® C reaction vessel was charged with a solution of anhydrous sodium sulfite (25 g, 0.20 mol), sodium bisulfite 73 g, (0.70 mol) and of deionized water (400 ml). The pH of this solution was 5.7. The vessel was cooled to 4 degrees C, evacuated to 0.08 MPa, and then charged with hexafluoropropene (HFP, 120 g, 0.8 mol, 0.43 MPa). The vessel was heated with

agitation to 120 degrees C and kept there for 3 hr. The pressure rose to a maximum of 1.83 MPa and then dropped down to 0.27 MPa within 30 minutes. At the end, the vessel was cooled and the remaining HFP was vented, and the reactor was purged with nitrogen. The final solution had a pH of 7.3.

5

The water was removed *in vacuo* on a rotary evaporator to produce a wet solid. The solid was then placed in a vacuum oven (0.02 MPa, 140 degrees C, 48 hr) to produce 219 g of white solid, which contained approximately 1 wt % water. The theoretical mass of total solids was 217 g.

10

The crude HFPS-Na can be further purified and isolated by extraction with reagent grade acetone, filtration, and drying.

^{19}F NMR (D_2O) δ -74.5 (m, 3F); -113.1, -120.4 (ABq, $J = 264$ Hz, 2F); -211.6 (dm, 1F).

15

^1H NMR (D_2O) δ 5.8 (dm, $J_{\text{FH}} = 43$ Hz, 1H).

Mp (DSC) 126 degrees C.

TGA (air): 10% wt. loss @ 326 degrees C, 50% wt. loss @ 446 degrees C.

TGA (N_2): 10% wt. loss @ 322 degrees C, 50% wt. loss @ 449 degrees C.

20

Preparation of Ionic Liquids

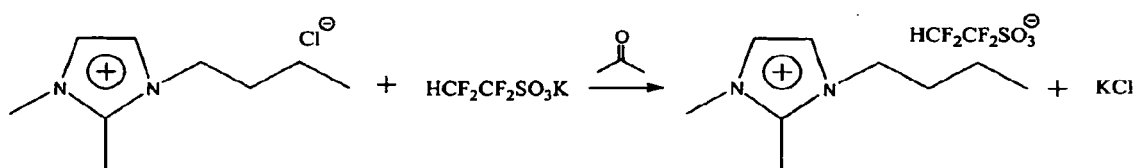
E) Synthesis of 1-butyl-2,3-dimethylimidazolium 1,1,2,2-tetrafluoroethanesulfonate

25 1-Butyl-2,3-dimethylimidazolium chloride (22.8 g, 0.121 moles) was mixed with reagent-grade acetone (250 ml) in a large round-bottomed flask and stirred vigorously. Potassium 1,1,2,2-tetrafluoroethanesulfonate (TFES-K, 26.6 g, 0.121 moles), was added to reagent grade acetone (250 ml) in a separate round-bottomed flask, and this solution was carefully added to the 1-butyl-2,3-dimethylimidazolium chloride solution. The large flask was lowered into an oil bath and heated at 60 degrees C under reflux for 10 hours. The reaction mixture was then filtered using a large frit glass funnel to remove the white KCl

30

precipitate formed, and the filtrate was placed on a rotary evaporator for 4 hours to remove the acetone.

5 The reaction scheme is shown below:



10 F) Synthesis of 1-butyl-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate

1-Butyl-3-methylimidazolium chloride (60.0 g) and high purity dry acetone (>99.5%, Aldrich, 300 ml) were combined in a 1l flask and warmed to reflux with magnetic stirring until the solid completely dissolved. At room temperature in a separate 1l flask, potassium-1,1,2,2-tetrafluoroethanesulfonate (TFES-K, 75.6 g) was dissolved in high purity dry acetone (500 ml). These two solutions were combined at room temperature and allowed to stir magnetically for 2 hr under positive nitrogen pressure. The stirring was stopped and the KCl precipitate was allowed to settle, then removed by suction filtration through a fritted glass funnel with a celite pad. The acetone was removed *in vacuo* to give a yellow oil. The oil was further purified by diluting with high purity acetone (100 ml) and stirring with decolorizing carbon (5 g). The mixture was again suction filtered and the acetone removed *in vacuo* to give a colorless oil. This was further dried at 4 Pa and 25 degrees C for 6 hr to provide 83.6 g of product.

25 ^{19}F NMR (DMSO- d_6) δ -124.7 (dt, $J = 6$ Hz, $J = 8$ Hz, 2F); -136.8 (dt, $J = 53$ Hz, 2F).

^1H NMR (DMSO- d_6) δ 0.9 (t, $J = 7.4$ Hz, 3H); 1.3 (m, 2H); 1.8 (m, 2H); 3.9 (s, 3H); 4.2 (t, $J = 7$ Hz, 2H); 6.3 (dt, $J = 53$ Hz, $J = 6$ Hz, 1H); 7.4 (s, 1H); 7.5 (s, 1H); 8.7 (s, 1H).

30 % Water by Karl-Fisher titration: 0.14 %.

Analytical calculation for $C_9H_{12}F_6N_2O_3S$: C, 37.6; H, 4.7; N, 8.8. Experimental Results: C, 37.6; H, 4.6; N, 8.7.

TGA (air): 10% wt. loss @ 380 degrees C, 50% wt. loss @ 420 degrees C.

5 TGA (N_2): 10% wt. loss @ 375 degrees C, 50% wt. loss @ 422 degrees C.

G) Synthesis of 1-ethyl-3-methylimidazolium 1,1,2,2-tetrafluoroethane sulfonate

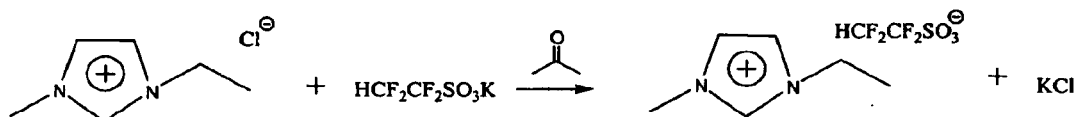
10 To a 500 ml round bottom flask was added 1-ethyl-3-methylimidazolium chloride (Emim-Cl, 98%, 61.0 g) and reagent grade acetone (500 ml). The mixture was gently warmed (50 degrees C) until almost all of the Emim-Cl dissolved. To a separate 500 ml flask was added potassium 1,1,2,2-tetrafluoroethanesulfonate (TFES-K, 90.2 g) along with reagent grade acetone
15 (350 ml). This second mixture was stirred magnetically at 24 degrees C until all of the TFES-K dissolved.

These solutions were combined in a 1l flask producing a milky white suspension. The mixture was stirred at 24 degrees C for 24 hrs. The KCl
20 precipitate was then allowed to settle leaving a clear green solution above it.

The reaction mixture was filtered once through a celite/acetone pad and again through a fritted glass funnel to remove the KCl. The acetone was removed *in vacuo* first on a rotovap and then on a high vacuum line (4 Pa, 25 degrees C)
25 for 2 hr. The product was a viscous light yellow oil (76.0 g, 64% yield).

The reaction scheme is shown below:

30



^{19}F NMR ($DMSO-d_6$) δ -124.7 (dt, $J_{FH} = 6$ Hz, $J_{FF} = 6$ Hz, 2F); -138.4 (dt, $J_{FH} = 53$ Hz, 2F).

¹H NMR (DMSO-d₆) δ 1.3 (t, *J* = 7.3 Hz, 3H); 3.7 (s, 3H); 4.0 (q, *J* = 7.3 Hz, 2H);

6.1 (tt, *J*_{FH} = 53 Hz, *J*_{FH} = 6 Hz, 1H); 7.2 (s, 1H); 7.3 (s, 1H); 8.5 (s, 1H).

% Water by Karl-Fisher titration: 0.18 %.

5 Analytical calculation for C₈H₁₂N₂O₃F₄S: C, 32.9: H, 4.1: N, 9.6 Found: C, 33.3: H, 3.7: N, 9.6.

Mp 45-46 degrees C.

TGA (air): 10% wt. loss @ 379 degrees C, 50% wt. loss @ 420 degrees C.

TGA (N₂): 10% wt. loss @ 378 degrees C, 50% wt. loss @ 418 degrees C.

10

H) Synthesis of 1-ethyl-3-methylimidazolium 1,1,2,3,3,3-hexafluoropropanesulfonate

To a 1l round bottom flask was added 1-ethyl-3-methylimidazolium
15 chloride (Emim-Cl, 98%, 50.5 g) and reagent grade acetone (400 ml). The mixture was gently warmed (50 degrees C) until almost all of the Emim-Cl dissolved. To a separate 500 ml flask was added potassium 1,1,2,3,3,3-hexafluoropropanesulfonate (HFPS-K, 92.2 g) along with reagent grade acetone (300 ml). This second mixture was stirred magnetically at room temperature until
20 all of the HFPS-K dissolved.

These solutions were combined and stirred under positive N₂ pressure at 26 degrees C for 12 hr producing a milky white suspension. The KCl precipitate was allowed to settle overnight leaving a clear yellow solution above it.

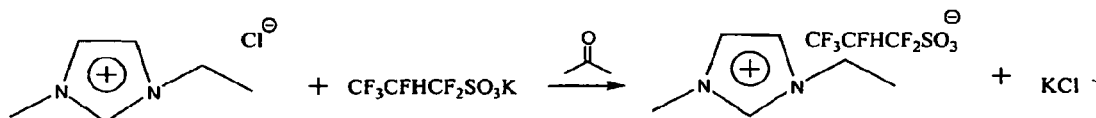
25

The reaction mixture was filtered once through a celite/acetone pad and again through a fritted glass funnel. The acetone was removed *in vacuo* first on a rotovap and then on a high vacuum line (4 Pa, 25 degrees C) for 2 hr. The product was a viscous light yellow oil (103.8 g, 89% yield).

30

35

The reaction scheme is shown below:



5 ^{19}F NMR (DMSO- d_6) δ -73.8 (s, 3F); -114.5, -121.0 (ABq, J = 258 Hz, 2F); -210.6 (m, 1F, J_{HF} = 41.5 Hz).

^1H NMR (DMSO- d_6) δ 1.4 (t, J = 7.3 Hz, 3H); 3.9 (s, 3H); 4.2 (q, J = 7.3 Hz, 2H);

5.8 (m, J_{HF} = 41.5 Hz, 1H); 7.7 (s, 1H); 7.8 (s, 1H); 9.1 (s, 1H).

10 % Water by Karl-Fisher titration: 0.12 %.

Analytical calculation for $\text{C}_9\text{H}_{12}\text{N}_2\text{O}_3\text{F}_6\text{S}$: C, 31.5; H, 3.5; N, 8.2. Experimental Results: C, 30.9; H, 3.3; N, 7.8.

TGA (air): 10% wt. loss @ 342 degrees C, 50% wt. loss @ 373 degrees C.

TGA (N_2): 10% wt. loss @ 341 degrees C, 50% wt. loss @ 374 degrees C.

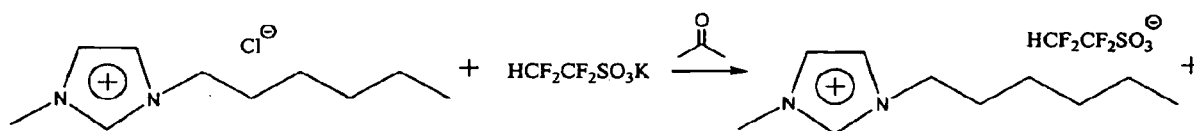
15

I) Synthesis of 1-hexyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate

1-Hexyl-3-methylimidazolium chloride (10 g, 0.0493 moles) was mixed
 20 with reagent-grade acetone (100 ml) in a large round-bottomed flask and stirred vigorously under a nitrogen blanket. Potassium 1,1,2,2-tetrafluoroethane sulfonate (TFES-K, 10 g, 0.0455 moles) was added to reagent grade acetone (100 ml) in a separate round-bottomed flask, and this solution was carefully added to the 1-hexyl-3-methylimidazolium chloride/acetone mixture. The mixture was left
 25 to stir overnight. The reaction mixture was then filtered using a large frit glass funnel to remove the white KCl precipitate formed, and the filtrate was placed on a rotary evaporator for 4 hours to remove the acetone.

30

The reaction scheme is shown below:



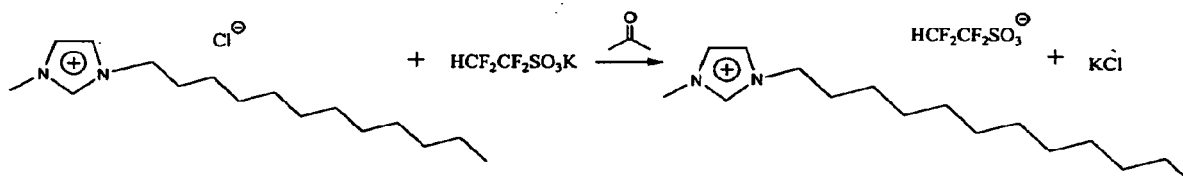
5

J) Synthesis of 1-dodecyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate

1-Dodecyl-3-methylimidazolium chloride (34.16 g, 0.119 moles) was partially dissolved in reagent-grade acetone (400 ml) in a large round-bottomed flask and stirred vigorously. Potassium 1,1,2,2-tetrafluoroethanesulfonate (TFES-K, 26.24 g, 0.119 moles) was added to reagent grade acetone (400 ml) in a separate round-bottomed flask, and this solution was carefully added to the 1-dodecyl-3-methylimidazolium chloride solution. The reaction mixture was heated at 60 degrees C under reflux for approximately 16 hours. The reaction mixture was then filtered using a large frit glass funnel to remove the white KCl precipitate formed, and the filtrate was placed on a rotary evaporator for 4 hours to remove the acetone.

The reaction scheme is shown below:

20

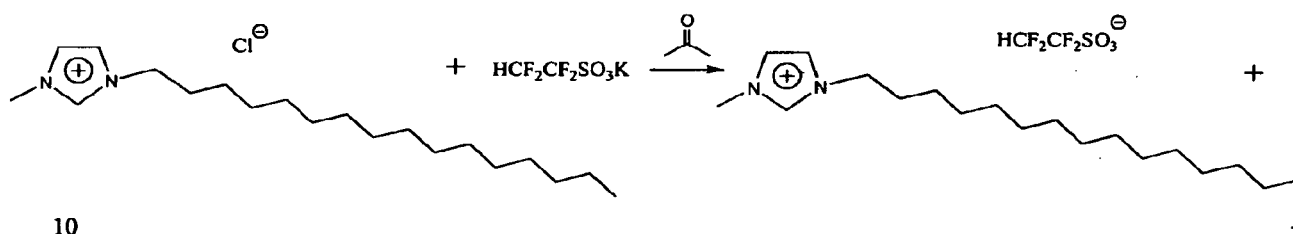


K) Synthesis of 1-hexadecyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate

1-Hexadecyl-3-methylimidazolium chloride (17.0 g, 0.0496 moles) was partially dissolved in reagent-grade acetone (100 ml) in a large round-bottomed flask and stirred vigorously. Potassium 1,1,2,2-tetrafluoroethanesulfonate (TFES-

K, 10.9 g, 0.0495 moles) was added to reagent grade acetone (100 ml) in a separate round-bottomed flask, and this solution was carefully added to the 1-hexadecyl-3-methylimidazolium chloride solution. The reaction mixture was heated at 60 degrees C under reflux for approximately 16 hours. The reaction mixture was then filtered using a large frit glass funnel to remove the white KCl precipitate formed, and the filtrate was placed on a rotary evaporator for 4 hours to remove the acetone.

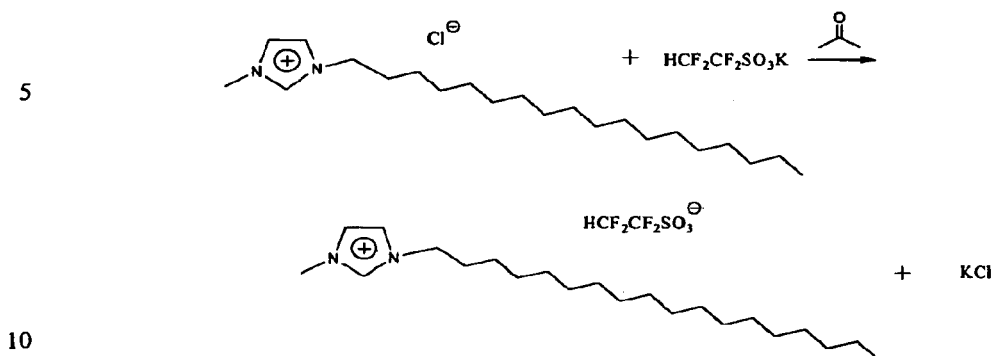
The reaction scheme is shown below:



L) Synthesis of 1-octadecyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate

1-Octadecyl-3-methylimidazolium chloride (17.0 g, 0.0458 moles) was partially dissolved in reagent-grade acetone (200 ml) in a large round-bottomed flask and stirred vigorously. Potassium 1,1,2,2-tetrafluoroethanesulfonate (TFES-K, 10.1 g, 0.0459 moles), was added to reagent grade acetone (200 ml) in a separate round-bottomed flask, and this solution was carefully added to the 1-octadecyl-3-methylimidazolium chloride solution. The reaction mixture was heated at 60 degrees C under reflux for approximately 16 hours. The reaction mixture was then filtered using a large frit glass funnel to remove the white KCl precipitate formed, and the filtrate was placed on a rotary evaporator for 4 hours to remove the acetone.

The reaction scheme is shown below:



M) Synthesis of 1-propyl-3-(1,1,2,2-TFES) imidazolium 1,1,2,2-tetrafluoroethanesulfonate

15 Imidazole (19.2 g) was added to of tetrahydrofuran (80 mls). A glass shaker tube reaction vessel was filled with the THF-containing imidazole solution. The vessel was cooled to 18°C, evacuated to 0.08 MPa, and purged with nitrogen. The evacuate/purge cycle was repeated two more times. Tetrafluoroethylene (TFE, 5 g) was then added to the vessel, and it was heated to 100 degrees C, at
 20 which time the inside pressure was about 0.72 MPa. As the TFE pressure decreased due to the reaction, more TFE was added in small aliquots (5 g each) to maintain operating pressure roughly between 0.34 MPa and 0.86 MPa. Once 40 g of TFE had been fed, the vessel was vented and cooled to 25 degrees C. The THF was then removed under vacuum and the product was vacuum distilled at 40
 25 degrees C to yield pure product as shown by ¹H and ¹⁹F NMR (yield 44 g). Iodopropane (16.99 g) was mixed with 1-(1,1,2,2-tetrafluoroethyl)imidazole (16.8 g) in dry acetonitrile (100 ml), and the mixture was refluxed for 3 days. The solvent was removed *in vacuo*, yielding a yellow waxy solid (yield 29 g). The product, 1-propyl-3-(1,1,2,2-tetrafluoroethyl)imidazolium iodide was confirmed
 30 by ¹H NMR (in CD₃CN) [0.96 (t, 3H); 1.99 (m, 2H); 4.27 (t, 2H); 6.75 (t, 1H); 7.72 (d, 2H); 9.95 (s, 1H)].

Iodide (24 g) was then added to 60 ml of dry acetone, followed by 15.4 g of potassium 1,1,2,2-tetrafluoroethanesulfonate in 75 ml of dry acetone. The

mixture was heated at 60 degrees C overnight and a dense white precipitate was formed (potassium iodide). The mixture was cooled, filtered, and the solvent from the filtrate was removed using a rotary evaporator. Some further potassium iodide was removed under filtration. The product was further purified by adding
 5 50 g of acetone, 1 g of charcoal, 1 g of celite and 1 g of silica gel. The mixture was stirred for 2 hours, filtered and the solvent removed. This yielded 15 g of a liquid, shown by NMR to be the desired product.

10 N) Synthesis of 1-butyl-3-methylimidazolium 1,1,2,3,3,3-hexafluoropropanesulfonate (Bmim-HFPS)

1-Butyl-3-methylimidazolium chloride (Bmim-Cl, 50.0 g) and high purity dry acetone (>99.5%, 500 ml) were combined in a 1l flask and warmed to reflux with magnetic stirring until the solid all dissolved. At room temperature in a
 15 separate 1l flask, potassium-1,1,2,3,3,3-hexafluoropropanesulfonate (HFPS-K) was dissolved in high purity dry acetone (550 ml). These two solutions were combined at room temperature and allowed to stir magnetically for 12 hr under positive nitrogen pressure. The stirring was stopped, and the KCl precipitate was allowed to settle. This solid was removed by suction filtration through a fritted
 20 glass funnel with a celite pad. The acetone was removed *in vacuo* to give a yellow oil. The oil was further purified by diluting with high purity acetone (100 ml) and stirring with decolorizing carbon (5 g). The mixture was suction filtered and the acetone removed *in vacuo* to give a colorless oil. This was further dried at 4 Pa and 25 degrees C for 2 hr to provide 68.6 g of product.

25

¹⁹F NMR (DMSO-d₆) δ -73.8 (s, 3F); -114.5, -121.0 (ABq, *J* = 258 Hz, 2F); -210.6 (m, *J* = 42 Hz, 1F).

¹H NMR (DMSO-d₆) δ 0.9 (t, *J* = 7.4 Hz, 3H); 1.3 (m, 2H); 1.8 (m, 2H); 3.9 (s, 3H); 4.2 (t, *J* = 7 Hz, 2H); 5.8 (dm, *J* = 42 Hz, 1H); 7.7 (s, 1H); 7.8 (s, 1H); 9.1 (s,
 30 1H).

% Water by Karl-Fisher titration: 0.12 %.

Analytical calculation for C₉H₁₂F₆N₂O₃S: C, 35.7; H, 4.4; N, 7.6. Experimental Results: C, 34.7; H, 3.8; N, 7.2.

TGA (air): 10% wt. loss @ 340 degrees C, 50% wt. loss @ 367 degrees C.

TGA (N₂): 10% wt. loss @ 335 degrees C, 50% wt. loss @ 361 degrees C.

Extractable chloride by ion chromatography: 27 ppm.

5

O) Synthesis of 1-butyl-3-methylimidazolium 1,1,2-trifluoro-2-(trifluoromethoxy)ethanesulfonate (Bmim-TTES)

1-Butyl-3-methylimidazolium chloride (Bmim-Cl, 10.0 g) and deionized water (15 ml) were combined at room temperature in a 200 ml flask. At room
10 temperature in a separate 200 ml flask, potassium 1,1,2-trifluoro-2-(trifluoromethoxy)ethanesulfonate (TTES-K, 16.4 g) was dissolved in deionized water (90 ml). These two solutions were combined at room temperature and allowed to stir magnetically for 30 min. under positive nitrogen pressure to give a biphasic mixture with the desired ionic liquid as the bottom phase. The layers
15 were separated, and the aqueous phase was extracted with 2 x 50 ml portions of methylene chloride. The combined organic layers were dried over magnesium sulfate and concentrated *in vacuo*. The colorless oil product was dried at for 4 hr at 5 Pa and 25 degrees C to afford 15.0 g of product.

20 ¹⁹F NMR (DMSO-d₆) δ -56.8 (d, *J*_{FH} = 4 Hz, 3F); -119.5, -119.9 (subsplit ABq, *J* = 260 Hz, 2F); -142.2 (dm, *J*_{FH} = 53 Hz, 1F).

¹H NMR (DMSO-d₆) δ 0.9 (t, *J* = 7.4 Hz, 3H); 1.3 (m, 2H); 1.8 (m, 2H); 3.9 (s, 3H); 4.2 (t, *J* = 7.0 Hz, 2H); 6.5 (dt, *J* = 53 Hz, *J* = 7 Hz, 1H); 7.7 (s, 1H); 7.8 (s, 1H); 9.1 (s, 1H).

25 % Water by Karl-Fisher titration: 613 ppm.

Analytical calculation for C₁₁H₁₆F₆N₂O₄S: C, 34.2: H, 4.2: N, 7.3.

Experimental Results: C, 34.0: H, 4.0: N, 7.1.

TGA (air): 10% wt. loss @ 328 degrees C, 50% wt. loss @ 354 degrees C.

TGA (N₂): 10% wt. loss @ 324 degrees C, 50% wt. loss @ 351 degrees C.

30 Extractable chloride by ion chromatography: < 2 ppm.

P) Synthesis of 1-butyl-3-methylimidazolium 1,1,2-trifluoro-2-(perfluoroethoxy)ethanesulfonate (Bmim-TPES)

1-Butyl-3-methylimidazolium chloride (Bmim-Cl, 7.8 g) and dry acetone (150 ml) were combined at room temperature in a 500 ml flask. At room temperature in a separate 200 ml flask, potassium 1,1,2-trifluoro-2-(perfluoroethoxy)ethanesulfonate (TPES-K, 15.0 g) was dissolved in dry acetone (300 ml). These two solutions were combined and allowed to stir magnetically for 12 hr under positive nitrogen pressure. The KCl precipitate was then allowed to settle leaving a colorless solution above it. The reaction mixture was filtered once through a celite/acetone pad and again through a fritted glass funnel to remove the KCl. The acetone was removed *in vacuo* first on a rotovap and then on a high vacuum line (4 Pa, 25 degrees C) for 2 hr. Residual KCl was still precipitating out of the solution, so methylene chloride (50 ml) was added to the crude product, which was then washed with deionized water (2 x 50 ml). The solution was dried over magnesium sulfate, and the solvent was removed *in vacuo* to give the product as a viscous light yellow oil (12.0 g, 62% yield).

^{19}F NMR (CD_3CN) δ -85.8 (s, 3F); -87.9, -90.1 (subsplit ABq, $J_{\text{FF}} = 147$ Hz, 2F); -120.6, -122.4 (subsplit ABq, $J_{\text{FF}} = 258$ Hz, 2F); -142.2 (dm, $J_{\text{FH}} = 53$ Hz, 1F).

^1H NMR (CD_3CN) δ 1.0 (t, $J = 7.4$ Hz, 3H); 1.4 (m, 2H); 1.8 (m, 2H); 3.9 (s, 3H); 4.2 (t, $J = 7.0$ Hz, 2H); 6.5 (dm, $J = 53$ Hz, 1H); 7.4 (s, 1H); 7.5 (s, 1H); 8.6 (s, 1H).

% Water by Karl-Fisher titration: 0.461.

Analytical calculation for $\text{C}_{12}\text{H}_{16}\text{F}_8\text{N}_2\text{O}_4\text{S}$: C, 33.0; H, 3.7. Experimental Results: C, 32.0; H, 3.6.

TGA (air): 10% wt. loss @ 334 degrees C, 50% wt. loss @ 353 degrees C.

TGA (N_2): 10% wt. loss @ 330 degrees C, 50% wt. loss @ 365 degrees C.

Q) Synthesis of tetradecyl(tri-*n*-butyl)phosphonium 1,1,2,3,3,3-hexafluoropropanesulfonate ([4.4.4.14]P-HFPS)

To a 4l round bottomed flask was added the ionic liquid tetradecyl(tri-*n*-butyl)phosphonium chloride (Cyphos® IL 167, 345 g) and deionized water (1000 ml). The mixture was magnetically stirred until it was one phase. In a separate 2l flask, potassium 1,1,2,3,3,3-hexafluoropropanesulfonate (HFPS-K, 214.2 g) was dissolved in deionized water (1100 ml). These solutions were combined and stirred under positive N₂ pressure at 26 degrees C for 1 hr producing a milky white oil. The oil slowly solidified (439 g) and was removed by suction filtration and then dissolved in chloroform (300 ml). The remaining aqueous layer (pH = 2) was extracted once with chloroform (100 ml). The chloroform layers were combined and washed with an aqueous sodium carbonate solution (50 ml) to remove any acidic impurity. They were then dried over magnesium sulfate, suction filtered, and reduced *in vacuo* first on a rotovap and then on a high vacuum line (4 Pa, 100 degrees C) for 16 hr to yield the final product as a white solid (380 g, 76% yield).

¹⁹F NMR (DMSO-d₆) δ -73.7 (s, 3F); -114.6, -120.9 (ABq, *J* = 258 Hz, 2F); -210.5 (m, *J*_{HF} = 41.5 Hz, 1F).

¹H NMR (DMSO-d₆) δ 0.8 (t, *J* = 7.0 Hz, 3H); 0.9 (t, *J* = 7.0 Hz, 9H); 1.3 (br s, 20H); 1.4 (m, 16H); 2.2 (m, 8H); 5.9 (m, *J*_{HF} = 42 Hz, 1H).

% Water by Karl-Fisher titration: 895 ppm.

Analytical calculation for C₂₉H₅₇F₆O₃PS: C, 55.2: H, 9.1: N, 0.0.

Experimental Results: C, 55.1: H, 8.8: N, 0.0.

TGA (air): 10% wt. loss @ 373 degrees C, 50% wt. loss @ 421 degrees C.

TGA (N₂): 10% wt. loss @ 383 degrees C, 50% wt. loss @ 436 degrees C.

R) Synthesis of tetradecyl(tri-*n*-hexyl)phosphonium 1,1,2-trifluoro-2-(perfluoroethoxy)ethanesulfonate ([6.6.6.14]P-TPES)

To a 500 ml round bottomed flask was added acetone (Spectroscopic grade, 50 ml) and ionic liquid tetradecyl(tri-*n*-hexyl)phosphonium chloride

(Cyphos® IL 101, 33.7 g). The mixture was magnetically stirred until it was one phase. In a separate 11 flask, potassium 1,1,2-trifluoro-2-(perfluoroethoxy)ethanesulfonate (TPES-K, 21.6 g) was dissolved in acetone (400 ml). These solutions were combined and stirred under positive N₂ pressure at 5 26 degrees C for 12 hr producing a white precipitate of KCl. The precipitate was removed by suction filtration, and the acetone was removed *in vacuo* on a rotovap to produce the crude product as a cloudy oil (48 g). Chloroform (100 ml) was added, and the solution was washed once with deionized water (50 ml). It was then dried over magnesium sulfate and reduced *in vacuo* first on a rotovap and 10 then on a high vacuum line (8 Pa, 24 degrees C) for 8 hr to yield the final product as a slightly yellow oil (28 g, 56% yield).

¹⁹F NMR (DMSO-d₆) δ -86.1 (s, 3F); -88.4, -90.3 (subsplit ABq, J_{FF} = 147 Hz, 2F); -121.4, -122.4 (subsplit ABq, J_{FF} = 258 Hz, 2F); -143.0 (dm, J_{FH} = 53 Hz, 15 1F).

¹H NMR (DMSO-d₆) δ 0.9 (m, 12H); 1.2 (m, 16H); 1.3 (m, 16H); 1.4 (m, 8H); 1.5 (m, 8H); 2.2 (m, 8H); 6.3 (dm, J_{FH} = 54 Hz, 1H).

% Water by Karl-Fisher titration: 0.11.

Analytical calculation for C₃₆H₆₉F₈O₄PS: C, 55.4: H, 8.9: N, 0.0.

20 Experimental Results: C, 55.2: H, 8.2: N, 0.1.

TGA (air): 10% wt. loss @ 311 degrees C, 50% wt. loss @ 339 degrees C.

TGA (N₂): 10% wt. loss @ 315 degrees C, 50% wt. loss @ 343 degrees C.

25 S) Synthesis of tetradecyl(tri-*n*-hexyl)phosphonium 1,1,2-trifluoro-2-(trifluoromethoxy)ethanesulfonate ([6.6.6.14]P-TTES)

To a 100 ml round bottomed flask was added acetone (Spectroscopic grade, 50 ml) and ionic liquid tetradecyl(tri-*n*-hexyl)phosphonium chloride (Cyphos® IL 101, 20.2 g). The mixture was magnetically stirred until it was one 30 phase. In a separate 100 ml flask, potassium 1,1,2-trifluoro-2-(trifluoromethoxy)ethanesulfonate (TTES-K, 11.2 g) was dissolved in acetone (100 ml). These solutions were combined and stirred under positive N₂ pressure at 26 degrees C for 12 hr producing a white precipitate of KCl.

The precipitate was removed by suction filtration, and the acetone was removed *in vacuo* on a rotovap to produce the crude product as a cloudy oil. The product was diluted with ethyl ether (100 ml) and then washed once with deionized water (50 ml), twice with an aqueous sodium carbonate solution (50 ml) to remove any acidic impurity, and twice more with deionized water (50 ml). The ether solution was then dried over magnesium sulfate and reduced *in vacuo* first on a rotovap and then on a high vacuum line (4 Pa, 24 degrees C) for 8 hr to yield the final product as an oil (19.0 g, 69% yield).

^{19}F NMR (CD_2Cl_2) δ -60.2 (d, $J_{\text{FH}} = 4$ Hz, 3F); -120.8, -125.1 (subsplit ABq, $J = 260$ Hz, 2F); -143.7 (dm, $J_{\text{FH}} = 53$ Hz, 1F).

^1H NMR (CD_2Cl_2) δ 0.9 (m, 12H); 1.2 (m, 16H); 1.3 (m, 16H); 1.4 (m, 8H); 1.5 (m, 8H); 2.2 (m, 8H); 6.3 (dm, $J_{\text{FH}} = 54$ Hz, 1H).

% Water by Karl-Fisher titration: 412 ppm.

Analytical calculation for $\text{C}_{35}\text{H}_{69}\text{F}_6\text{O}_4\text{PS}$: C, 57.5: H, 9.5: N, 0.0. Experimental results: C, 57.8: H, 9.3: N, 0.0.

TGA (air): 10% wt. loss @ 331 degrees C, 50% wt. loss @ 359 degrees C.

TGA (N_2): 10% wt. loss @ 328 degrees C, 50% wt. loss @ 360 degrees C.

T) Synthesis of 1-ethyl-3-methylimidazolium 1,1,2,2-tetrafluoro-2-(pentafluoroethoxy)sulfonate (Emim-TPENTAS)

To a 500 ml round bottomed flask was added 1-ethyl-3-methylimidazolium chloride (Emim-Cl, 98%, 18.0 g) and reagent grade acetone (150 ml). The mixture was gently warmed (50 degrees C) until all of the Emim-Cl dissolved. In a separate 500 ml flask, potassium 1,1,2,2-tetrafluoro-2-(pentafluoroethoxy)sulfonate (TPENTAS-K, 43.7 g) was dissolved in reagent grade acetone (450 ml).

These solutions were combined in a 1l flask producing a white precipitate (KCl). The mixture was stirred at 24 degrees C for 8 hr. The KCl precipitate was then allowed to settle leaving a clear yellow solution above it. The KCl was

removed by filtration through a celite/acetone pad. The acetone was removed *in vacuo* to give a yellow oil, which was then diluted with chloroform (100 ml). The chloroform was washed three times with deionized water (50 ml), dried over magnesium sulfate, filtered, and reduced *in vacuo* first on a rotovap and then on a high vacuum line (4 Pa, 25 degrees C) for 8 hr. The product was a light yellow oil (22.5 g).

^{19}F NMR (DMSO- d_6) δ -82.9 (m, 2F); -87.3 (s, 3F); -89.0 (m, 2F); -118.9 (s, 2F).

^1H NMR (DMSO- d_6) δ 1.5 (t, $J = 7.3$ Hz, 3H); 3.9 (s, 3H); 4.2 (q, $J = 7.3$ Hz, 2H); 7.7 (s, 1H); 7.8 (s, 1H); 9.1 (s, 1H).

% Water by Karl-Fisher titration: 0.17 %.

Analytical calculation for $\text{C}_{10}\text{H}_{11}\text{N}_2\text{O}_4\text{F}_9\text{S}$: C, 28.2: H, 2.6: N, 6.6

Experimental results: C, 28.1: H, 2.9: N, 6.6.

TGA (air): 10% wt. loss @ 351 degrees C, 50% wt. loss @ 401 degrees C.

TGA (N_2): 10% wt. loss @ 349 degrees C, 50% wt. loss @ 406 degrees C.

U) Synthesis of tetrabutylphosphonium 1,1,2-trifluoro-2-(perfluoroethoxy)ethanesulfonate (TBP-TPES)

To a 200 ml round bottomed flask was added deionized water (100 ml) and tetra-*n*-butylphosphonium bromide (Cytec Canada Inc., 20.2 g). The mixture was magnetically stirred until the solid all dissolved. In a separate 300 ml flask, potassium 1,1,2-trifluoro-2-(perfluoroethoxy)ethanesulfonate (TPES-K, 20.0 g) was dissolved in deionized water (400 ml) heated to 70 degrees C. These solutions were combined and stirred under positive N_2 pressure at 26 degrees C for 2 hr producing a lower oily layer. The product oil layer was separated and diluted with chloroform (30 ml), then washed once with an aqueous sodium carbonate solution (4 ml) to remove any acidic impurity, and three times with deionized water (20 ml). It was then dried over magnesium sulfate and reduced *in vacuo* first on a rotovap and then on a high vacuum line (8 Pa, 24 degrees C) for 2 hr to yield the final product as a colorless oil (28.1 g, 85% yield).

^{19}F NMR (CD_2Cl_2) δ -86.4 (s, 3F); -89.0, -90.8 (subsplit ABq, $J_{\text{FF}} = 147$ Hz, 2F); -119.2, -125.8 (subsplit ABq, $J_{\text{FF}} = 254$ Hz, 2F); -141.7 (dm, $J_{\text{FH}} = 53$ Hz, 1F).

^1H NMR (CD_2Cl_2) δ 1.0 (t, $J = 7.3$ Hz, 12H); 1.5 (m, 16H); 2.2 (m, 8H); 6.3 (dm, $J_{\text{FH}} = 54$ Hz, 1H).

% Water by Karl-Fisher titration: 0.29.

Analytical calculation for $\text{C}_{20}\text{H}_{37}\text{F}_8\text{O}_4\text{PS}$: C, 43.2; H, 6.7; N, 0.0.

Experimental results: C, 42.0; H, 6.9; N, 0.1.

10 Extractable bromide by ion chromatography: 21 ppm.

(V) Preparation of 1,3-dioctylimidazolium iodide [doim][I]

15 1,3-Dioctylimidazolium iodide [ooim][I] was prepared as described by L. Xu, *et al.*, Journal of Organometallic Chemistry, 2000, 598, 409-416:

Imidazole (2.72 g; 0.04 mmol) and octyl bromide (3.1 g; 0.016 mmol) were dissolved in 55 ml of ethyl acetate. The mixture was refluxed under a nitrogen blanket. Initially, the solution was clear and colorless, however upon refluxing approximately 1 hour the mixture became cloudy with a tannish color. The mixture was allowed to reflux overnight. The mixture was then cooled to room temperature (RT) upon which a white precipitate formed. The mixture was extracted with water (2x: 30ml). After drying the solvent with magnesium sulfate, the solvent was removed using a vacuum, yielding a tannish oil.

To the oily residue was added 60 ml of toluene followed by 1-iodooctane (4.8 g; 0.02). The mixture was refluxed overnight under a nitrogen blanket, resulting in a dark yellow mixture. The yellow oil was collected via a separation funnel rinsed with toluene (2x: 20 ml) and dried under vacuum.

(W) Preparation of 1-methyl-3-octylimidazolium iodide [omim][I]

1-Methyl-3-octylimidazolium iodide [omim][I] was prepared as described by L. Xu, *et al.* (Journal of Organometallic Chemistry, 2000, 598, 409-416):

5 1-Methylimidazole (1.65 g; 0.02 mmol) and 1-iodooctane (5.31 g; 0.022 mmol) were dissolved in 30 ml of toluene. The reaction was refluxed, whereupon the mixture immediately became yellow in color and cloudy. The mixture was refluxed overnight, during which a yellowish oily precipitate formed. The yellowish oil was collected and dried under vacuum.

10 Gravimetric Microbalance

The gas solubility and diffusivity measurements were made using a gravimetric microbalance (Hiden Isochema Ltd, IGA 003, Warrington, UK). The IGA design integrates precise computer-control and measurement of weight
15 change, pressure and temperature to enable fully automatic and reproducible determination of gas adsorption-desorption isotherms and isobars. The microbalance consists of an electrobalance with sample and counterweight components inside a stainless steel pressure-vessel as shown in Figure 18 and described in Example 38, Table 31. The balance has a weigh range of 0-100 mg
20 with a resolution of 0.1 μ g. An enhanced pressure stainless steel (SS316LN) reactor capable of operation to 20.0 bar and 100 °C was installed. Approximately 60 mg of ionic liquid sample was added to the sample container and the reactor was sealed. The sample was dried and degassed by first pulling a course vacuum on the sample with a diaphragm pump (Pfeiffer, model MVP055-3, Asslar,
25 Germany) and then fully evacuating the reactor to 10^{-8} bar with a turbopump (Pfeiffer, model TSH-071). While under deep vacuum, the sample was heated to 75 °C for 10 hr with an external water jacket connected to a remote-controlled constant-temperature bath (Huber Ministat, model cc-S3, Offenburg, Germany). A 30 percent ethylene glycol and 70 percent water mixture by volume was used as
30 the recirculating fluid with a temperature range of 5 to 90 °C. The sample mass slowly decreased as residual water and gases were removed. Once the mass had stabilized for at least 60 min, the sample dry mass was recorded. The percent weight loss for the various ionic liquids tested was in the range of 1 to 3%.

The IGA003 can operate in both dynamic and static mode. Dynamic mode operation provides a continuous flow of gas (max. 500 cm³ min⁻¹) past the sample and the exhaust valve controls the set-point pressure. Static mode operation introduces gas into the top of the balance away from the sample and both the admittance and exhaust valves control the set-point pressure. All absorption measurements were performed in static mode. The sample temperature was measured with a type K thermocouple with an accuracy of $\pm 0.1^{\circ}\text{C}$. The thermocouple was located inside the reactor next to the sample container. The water jacket maintained the set-point temperature automatically to within a typical regulation accuracy of $\pm 0.1^{\circ}\text{C}$. Four isotherms (at 10, 25, 50, and 75 $^{\circ}\text{C}$) were measured beginning with 10 $^{\circ}\text{C}$. Once the desired temperature was achieved and stable, the admittance and exhaust valves automatically opened and closed to adjust the pressure to the first set-point. Pressures from 10^{-9} to 10^{-1} bar were measured using a capacitance manometer (Pfeiffer, model PKR251), and pressures from 10^{-1} to 20.0 bar were measured using a piezo-resistive strain gauge (Druck, model PDCR4010, New Fairfield, CT). Regulation maintained the reactor pressure set-point to within ± 4 to 8 mbar. The pressure ramp rate was set at 200 mbar min⁻¹ and the temperature ramp rate was set at 1 $^{\circ}\text{C}$ min⁻¹. The upper pressure limit of the stainless steel reactor was 20.0 bar, and several isobars up to 10 bar (i.e., 0.1, 0.5, 1, 4, 7, 10 bar) were measured. To ensure sufficient time for gas-liquid equilibrium, the ionic liquid samples were maintained at set-point for a minimum of 3 hr with a maximum time-out of 8 hr.

The IGA method exploits the relaxation behavior following pressure and temperature changes to simultaneously evaluate the time-dependent absorption and asymptotic uptake. The real-time processor was used to determine the end-point for each isotherm. The percent relaxation used as an end point for the real-time analysis was 99 percent. The minimum weight change for real-time analysis was set at 1 μg , the acceptable average deviation of the model from the acquired data was set at 7 μg , and the target interval for weight acquisition was set at a typical value of 1 μg . The temperature variation during an isotherm was maintained less than 0.1 $^{\circ}\text{C}$ min⁻¹.

Safety features of the IGA003 included a pressure relief valve and over-temperature control for the reactor. The factory-installed relief valve was replaced with a DuPont guideline relief valve (Circle-Seal, set-point pressure 24.5 bar; DuPont, Wilmington, Delaware). To further protect the microbalance system
5 from over-pressure, additional relief valves were installed on the custom gas manifold and on each gas cylinder; these relief valves were set to open if the pressure exceeded 25 bar. The reactor over-temperature interlock controller that comes standard on the IGA003 was set to turn off the water bath if the temperature exceeded 100 °C. Due to the fact that some of the gases tested were
10 flammable (i.e. HFC-32, HFC-143a, and HFC-152a), the IGA003 was mounted inside a custom stainless steel cabinet purged with nitrogen that would minimize the possibility of a flame.

Thermogravimetric measurements were corrected for a number of
15 gravitational balance forces introduced at high pressure as described by Pinkerton, E. P., *et al* (High-pressure gravimetric measurement of hydrogen capacity in vapor-grown carbon nanofibers and related materials. Proceedings of the 11th Canadian Hydrogen Conference, Victoria, BC (2001) pages 633-642). These included:

- 20 (1) Changes in the buoyant forces due to changes in pressure and temperature.
- (2) Aerodynamic drag forces created by the flow of gases.
- (3) Changes in the balance sensitivity due to changes in temperature and pressure.
- (4) Volumetric changes in the samples due to expansivity.

25

The gravitational balance forces previously described are often of the same order of magnitude (0.1 to 5 mg) as the overall weight change in the sample and can lead to inaccurate results if not accounted for precisely. Distinguishing mass changes with an accuracy of 0.01 wt.% on small and sometimes limited sample
30 quantities requires knowledge of the sample weight to within about 5 to 10 µg.

The buoyancy correction follows from Archimedes' principal: there is an upward force exerted on an object equivalent to the mass of fluid displaced. The

upward force (C_b) due to buoyancy is calculated using eq 1 where the mass of the gas displaced is equivalent to the volume of the submersed object (V_i) times the density (ρ_g) of the gas at a given (T, P) and the gravitational acceleration (g). If the volume of the object remains constant, V_i can be calculated by knowing the mass (m_i) and density (ρ_i) of the object.

$$C_b = \text{Buoyancy} = gV_i\rho_g(T, P) = g\frac{m_i}{\rho_i}\rho_g(T, P) \quad (1)$$

Instead of using the gas densities provided in the Hiden Isochema IGA software, the gas density for each gas was calculated using a computer program (REFPROP v.7) developed by the National Institute of Standards and Technology (NIST) (Lemmon EW, *et al.* [NIST reference fluid thermodynamic and transport properties – REFPROP, version 7.0 user's guide, U.S. Department of Commerce, Technology Administration, National Institute of Standards and Technology, Standard Reference Data Program, Gaithersburg, Maryland, 2002]).

The buoyancy correction using the IGA003 system involves many additional objects for weighing the sample. Table 31 provides a list of each critical component along with the objects weight, material, density, and temperature. The component arrangement in Figure 18 leads to a mass balance as shown by eq 2. This expression accounts for the summation of all components as well as the contribution of the absorbed gas mass (m_a) and a correction factor (C_f) which accounts for the balance sensitivity to T, P . The density of air (ρ_{air}) at ambient temperature and pressure was subtracted from ρ_i and ρ_j because the components were initially weighed in air.

$$\begin{aligned} & \sum_{i=1} m_i - \sum_{j=1} m_j - \sum_{i=1} \frac{m_i}{\rho_i} \rho_g(T_i, P) + \sum_{j=1} \frac{m_j}{\rho_j} \rho_g(T_j, P) + m_{IL} + m_a \\ & - \frac{m_{IL}}{\rho_s(T_s)} \rho_g(T_s, P) - \frac{m_a}{\rho_a(T_s)} \rho_g(T_s, P) - C_f(T_s, P) = \text{reading} \end{aligned} \quad (2)$$

The largest contributions in eq 2 are typically those of the sample container, sample, and counter weight; the other referenced objects in Table 31 contribute less because of their larger densities (denominators in eq 2). Physical densities of ionic liquids were measured using a Micromeritics Accupyc 1330 helium

pycnometer with an accuracy of $\pm 0.001 \text{ g cm}^{-3}$ (Micromeritics Instrument Corp., Norcross, GA). Initially, the volume (V_{IL}) of each sample was calculated from its pycnometric density (ρ_s) and dry mass sample weight (ρ_s), but volumetric expansion ($\Delta\tilde{V}/\tilde{V}_0$) due to the gas absorption was later taken into account as
 5 described below to more accurately determine the buoyancy effect.

The system was operated in static mode that essentially eliminates any aerodynamic drag forces due to flowing gases. Electrobalances are sensitive to temperature and pressure fluctuations on the beam arm and internal electronics.
 10 To minimize this effect, the balance electronics are heated externally with a band heater to a temperature of $45 \pm 0.1 \text{ }^\circ\text{C}$. In addition, the component temperatures provided in Table 31 are measured for the sample (T_s) and all others are estimated. Therefore, a correction factor (C_f) was determined as a function of T , P by measuring the buoyancy effect without a sample and calculating a least-squares fit to tare the balance. The correction factor was on the order of 0.1 to 0.3
 15 mg and increased as expected with decreasing temperature and increasing pressure.

Initially the ionic liquid sample volume was considered to be constant and
 20 the mole fraction solubility calculated without taking into account buoyancy effects due to sample expansivity. In order to make a proper buoyancy correction due to the liquid volume change, a simple mole fraction average for the molar volume, $\tilde{V}_m$, was used.

$$\tilde{V}_m(T, P) = \tilde{V}_{IL}(1 - x) + \tilde{V}_g x, \quad (3)$$
 25

where $\tilde{V}_i = MW_i/\rho_i$ and x represents the molar fraction of gas in the solution.

$$V_m(T, P) = \tilde{V}_m(T, P) \left[\left(\frac{m_{IL}}{MW_{IL}} \right) + \left(\frac{m_g}{MW_g} \right) \right] \quad (4)$$

$$\frac{m_s}{\rho_s(T_s)} \rho_g(T_s, P) + \frac{m_a}{\rho_a(T_s)} \rho_g(T_s, P) = V_m(T, P) \rho_g(T, P) \quad (5)$$

As a first approximation, eqs 3 and 4 were used to estimate the change in the liquid sample volume, V_m , at the measured T , P conditions. Eq 5 can be substituted into eq 2 to account for the buoyancy change with respect to sample expansivity.

5

Besides the equilibrium solubility, time-dependent absorption data were also gathered using the Hiden gravimetric microbalance for each T , P set-point. In order to understand the time-dependent behavior of gas dissolving in liquid, we applied a mathematical model based on a simplified mass diffusion process.

10 Imagine a flat-bottom sample container filled with ionic liquid at a certain liquid level height (L). The height is determined by knowing the cylindrical geometry of the sample container, dry sample weight after evacuation and heating, and the ionic liquid density at the proper temperature. After evacuation, the gas is introduced into the Pyrex[®] sample container with a constant pressure at a given

15 temperature. A small amount of gas will start dissolving into the ionic liquid, and after a sufficient time it will reach a thermodynamic equilibrium, that is the solubility limit of the gas in the ionic liquid at the given T and P . This transient behavior with time is modeled as described by Shiflett, M. B. and Yokozeki, A. (*Ind. Eng. Chem. Research*, 2005, 44, 4453-4464) and Yokozeki, A. (*Int. J. Refrigeration*, 2002, 22, 695-704).

20

Processes of gas dissolving in liquid may be highly complex phenomena because of a possible evolution of heat of mixing, the subsequent liquid convection due to the local temperature difference, as well as the free convection

25 due to the density difference, and the possible change in thermophysical properties of the liquid. The following assumptions were made for the dissolving gas (Shiflett, M. B., and Yokozeki, A. (*supra*); and Yokozeki, A. (Time-dependent behavior of gas absorption in lubricant oil [*Int. J. Refrigeration* (2002), 22, 695-704])):

- 30 (1) Gas dissolves through a one-dimensional (vertical) diffusion process, in which there is no convective flow in the liquid.
- (2) A thin boundary layer between the gas and liquid phases exists, where the thermodynamic equilibrium is instantly established with the saturation

concentration (C_s), and where the concentration is constant all the time at a given temperature and pressure.

(3) Temperature and pressure are kept constant.

(4) The gas-dissolved liquid is a highly dilute solution, and so the relevant thermophysical properties of the solution do not change.

The process may then be described by one-dimensional mass diffusion due to the local concentration difference. The governing differential equations are:

$$\frac{\partial C}{\partial t} = D \frac{\partial^2 C}{\partial z^2} \quad (6)$$

$$\text{Initial Condition: } C = C_0 \text{ when } t = 0 \text{ and } 0 < z < L \quad (7)$$

$$\text{Boundary Conditions: } C = C_s \text{ when } t > 0 \text{ and } z = 0 \quad (8)$$

$$\frac{\partial C}{\partial z} = 0 \text{ at } z = L \quad (9)$$

where C is the concentration of a dissolving substance in ionic liquid as a function of time, t and vertical location, z , where L is the depth of ionic liquid in the container, and $z = 0$ corresponds to the vapor-liquid boundary. C_0 is an initial homogenous concentration of the dissolving gas, and is zero (initially) or a small finite amount at $t > 0$. D is the diffusion coefficient that is assumed to be constant.

Eq 6 can be solved analytically for the initial and boundary conditions eqs 7 - 9 by a standard method such as separation variables or Laplace transform and yields:

$$C = C_s \left[1 - 2 \left(1 - \frac{C_0}{C_s} \right) \sum_{n=0}^{\infty} \frac{\exp(-\lambda_n^2 D t) \sin \lambda_n z}{L \lambda_n} \right], \quad (10)$$

$$\text{where } \lambda_n = \left(n + \frac{1}{2} \right) \frac{\pi}{L}.$$

An experimentally observed quantity at a specified time is the total concentration (or mass) of dissolved gas in ionic liquid, and not the concentration profile in z . This space-averaged concentration at a given time, $\langle C \rangle$, can be calculated from eq 11.

$$\langle C \rangle = \int_0^L C dz / L \quad (11)$$

$$\langle C \rangle = C_s \left[1 - 2 \left(1 - \frac{C_0}{C_s} \right) \sum_{n=0}^{\infty} \frac{\exp(-\lambda_n^2 D t)}{L^2 \lambda_n^2} \right] \quad (12)$$

Although eq 12 contains an infinite summation, only the first few terms, except for initial small time periods, are sufficient in practical applications. In this work, the summation was terminated after ten terms when the numerical contribution to the summation in $\langle C \rangle$ became less than 10^{-12} . By analyzing experimental data with this equation, we obtained the saturation concentration (C_s) and diffusion constant (D) at given T and P , when C_0 was known.

10

Properties of lubricants and lubricant/refrigerant mixtures

For vapor compression refrigeration and air-conditioning applications, the viscosity, density, and molecular weight of the lubricating oil are important properties. The viscosity of the oil and the solubility of the refrigerant in the oil are the two key factors in good oil return to the compressor. The viscosity of the oil cannot be so high that the oil becomes viscous at lower temperatures or so low that it does not lubricate the compressor properly at high temperatures. A viscosity of about 150 Saybolt universal seconds (SUS) or about 30–35 centipoise (cP) at 37.8°C (100°F) is generally used for low and medium temperature refrigeration applications. A viscosity of about 300 SUS or about 50–60 cP at 37.8°C (100°F) is generally used for higher temperature air-conditioning applications. The division, however, is not exact and both viscosities can be used for low temperature applications. Even higher viscosity oils such as 500 SUS are used in mobile air-conditioning applications. Examples of mineral oils that were commonly used with chlorofluorocarbon (CFC) refrigerants are Suniso 3GS, 4GS, and 5GS (Sun Oil Company) with viscosities of 150, 300, and 500 SUS, respectively.

The densities of the three Suniso oils are about 0.91 to 0.92 g cm⁻³ at 21.1°C (70°F). The molecular weight can vary from about 300 to 330 g mol⁻¹. The properties of these products can vary since they are natural or mineral-based

oils (MO) derived from underground petroleum sources which are a complex mixture of chemical compounds; the composition may also vary with drilling location and time (Downing, R.C. Fluorocarbon Refrigerants Handbook, Prentice-Hall, 1988).

5 Many ionic liquids have the proper viscosity, density, and molecular weight range to be used as lubricants as shown in Example 1, Table 1. For vapor compression heating or cooling systems the lubricant must be soluble in the refrigerant such that the lubricant that escapes the compressor and becomes
10 entrained with the refrigerant is returned to the compressor. In addition, the mixture properties of the refrigerant and lubricant must adequately perform as a lubricant in the compressor.

The viscosity of oil with dissolved refrigerant is the key factor in oil return
15 since the oil is moved by the force of the flowing refrigerant and thin oil moves better than thick or viscous oil. Temperature is also an important factor, but ultimately the mixture viscosity properties of the oil and refrigerant dominate the ability of the oil to return to the compressor.

20 Several measurements of the viscosity of oil-refrigerant solutions have been reported (Downing, R.C. Fluorocarbon Refrigerants Handbook, Prentice-Hall, 1988). In order to have the proper lubrication in the compressor, the mixture viscosity should be similar to the mixture viscosity of the traditional refrigerant-oil working fluid. For illustration, the viscosity of the mixtures of the refrigerant
25 chlorodifluoromethane (HCFC-22) with several SUNISO lubricants were compared to that of refrigerant-ionic liquid working fluids useful for the invention. (see Example 2, Table 2).

Example 1

Viscosity, density, and molecular weight comparison of ionic liquids with SUNISO 3GS, 4GS, and 5GS mineral oils are shown in Table 1 (Downing, R. C., *supra*).

5

Table 1.

Ionic Liquid	Viscosity at 40°C (cP or mPa.s)	Density at 21°C (g cm ⁻³)	Molecular Weight (g mol ⁻¹)
SUNISO 3GS	27	0.91	300-330
SUNISO 4GS	51	0.92	300-330
SUNISO 5GS	87	0.92	300-330
[bmim][PF ₆]	147	1.37	284.2
[bmim][BF ₄]	59	1.21	226.0
[dmpim][TMeM]	239	1.60	551.5
[emim][BEI]	44	1.59	491.3
[emim][BMeI]	20	1.52	391.3
[dmpim][BMeI]	-	-	419.4
[pmpy][BMeI]	31	1.45	416.4
[bmpy][BMeI]	-	1.42	430.4
[emim][TFES]	-	1.50*	292.3
[bmim][TFES]	-	1.32*	320.3
[dmim][TFES]	-	1.14*	432.5
[hmim][TFES]	-	1.27*	362.4
[bmim][Ac]	-	1.05**	198.3
[bmim][MeSO ₄]	-	1.21**	250.3
[bmim][SCN]	-	1.07**	197.3
[bmim][FS]	-	1.45	436.3
[bmim][HFPS]	108	1.41	370.3
[bmim][TPES]	119	1.43	436.3
[bmim][TTES]	93	1.40	386.3
[6,6,6,14-P][TPES]	191	1.07	780.0
[4,4,4,14-P][HFPS]	289	1.07	629.4

*T = 28.3°C, **T = 25-26°C

EXAMPLE 2

The concentration of the refrigerant in the ionic liquid (or oil) and viscosity of the refrigerant and ionic liquid (or oil) mixtures were calculated and compared with chlorodifluoromethane (HCFC-22) and the three SUNISO lubricants (3GS, 4GS, and 5GS) (see Downing, R. C., (*supra*)). The temperature in all cases was 40°C and the lubrication effectiveness was evaluated as excellent, good, average, or poor based on how close the viscosity of the mixture compared with HCFC-22 and the SUNISO lubricants as shown in Table 2. Five mixtures were rated as "excellent": HFC-134a and [bmim][PF₆], HFC-152a and [bmim][PF₆], HFC-134a and [bmim][TPES], HFC-134a and [bmim][TTES], and HFC-134a and [4,4,4,14-P][HFPS]. Three refrigerant-ionic liquid mixtures were evaluated with a "good" lubrication performance: HFC-143a and [bmim][PF₆], HFC-134a and [bmim][HFPS], and HFC-134a and [6,6,6,14-P][TPES]. In addition, four refrigerant-ionic liquid mixtures were evaluated to have "average" lubrication performance: HFC-134a and [emim][BEI], HFC-32 and [bmim][BF₄], HFC-125 and [bmim][PF₆], and HFC-32 and [bmim]PF₆. Finally, several mixtures were found to have "poor" lubrication performance because the viscosity of the mixture was too low compared with HCFC-22 and the SUNISO lubricants. Those mixtures included HFC-32 with [dmpim][TMeM], [emim][BEI], [pmpy][BMeI], [emim][BMeI], [bmim][HFPS], [bmim][TPES], and [bmim][TTES]. Although HFC-32 has a high solubility in the ionic liquid which reduces the viscosity of the mixture, it also most not decrease the viscosity so much that the mixture viscosity is too low for providing adequate lubrication in the compressor. Therefore, ionic liquids with higher molecular weights may work better with the HFC-32 refrigerant for lubricating the compressor.

Table 2.

Refrigerant	Lubricant	Refrigerant in Lubricant	Mixture Viscosity	Lubrication Effectiveness
		(mass %)	(cp)	
HCFC-22	SUNISO 3GS	10	10	Excellent
HCFC-22	SUNISO 4GS	10	16	Excellent
HCFC-22	SUNISO 5GS	10	23	Excellent
HFC-32	[bmim][PF ₆]	20	4	Average
HFC-125	[bmim][PF ₆]	10	48	Average
HFC-134a	[bmim][PF ₆]	15	26	Excellent
HFC-143a	[bmim][PF ₆]	10	31	Good
HFC-152a	[bmim][PF ₆]	15	13	Excellent
HFC-32	[bmim][BF ₄]	20	4	Average
HFC-32	[dmpim][TMeM]	20	1	Poor
HFC-32	[emim][BEI]	20	1	Poor
HFC-32	[pmpy][BMeI]	20	1	Poor
HFC-32	[emim][BMeI]	20	1	Poor
HFC-32	[bmim][HFPS]	20	2	Poor
HFC-32	[bmim][TPES]	20	2	Poor
HFC-32	[bmim][TTES]	20	2	Poor
HFC-134a	[emim][BEI]	15	6	Average
HFC-134a	[bmim][HFPS]	15	15	Good
HFC-134a	[bmim][TPES]	15	12	Excellent
HFC-134a	[bmim][TTES]	15	13	Excellent
HFC-134a	[6,6,6,14-P][TPES]	15	6	Good
HFC-134a	[4,4,4,14-P][HFPS]	15	10	Excellent

Examples 3-35 provide solubility and diffusivity results for several hydrofluorocarbon refrigerants. These data are used for solubility with lubricant in Examples 1 and 2.

Examples 3 – 7 and Figures 3 – 7 show solubility and diffusivity results for several hydrofluorocarbons (HFC-32, HFC-125, HFC-134a, HFC-143a, and HFC-152a) in one ionic liquid, [bmim][PF₆], at 10, 25, 50, and 75 °C. Compositions were prepared that consisted of HFC-32 and [bmim][PF₆] from about 0.3 to about 81.2 mole percent of HFC-32 over a temperature range from 10 to 75 °C at a pressure from about 0.1 to 10 bar. Compositions were prepared that consisted of HFC-125 and [bmim][PF₆] from about 0.1 to about 65.1 mole percent of HFC-125 over a temperature range from 10 to 75 °C at a pressure from about 0.1 to 10 bar. Compositions were prepared that consisted of HFC-134a and [bmim][PF₆] from about 0.1 to about 72.1 mole percent of HFC-134a over a temperature range from 10 to 75 °C at a pressure from about 0.1 to 3.5 bar. Compositions were prepared that consisted of HFC-143a and [bmim][PF₆] from about 0.1 to about 26.5 mole percent of HFC-143a over a temperature range from 10 to 75 °C at a pressure from about 0.1 to 7.5 bar. Compositions were prepared that consisted of HFC-152a and [bmim][PF₆] from about 0.5 to about 79.7 mole percent of HFC-152a over a temperature range from 10 to 75 °C at a pressure from about 0.1 to 4.5 bar.

Examples 8 – 14 and 17 – 29 and Figures 8, 11 - 16 show solubility and diffusivity results for HFC-32 in several additional ionic liquids. Examples 15 and 16 and Figures 9 and 10 show solubility and diffusivity results for HFC-23 in the ionic liquids [bmim][PF₆] and [emim][PF₆].

Examples 30-35 show solubility and diffusivity results for HFC-134a in several ionic liquids.

Compositions were prepared that consisted of HFC-32 and [bmim][BF₄] from about 0.1 to about 76.5 mole percent of HFC-32 over a temperature range from 10 to 75 °C at a pressure from about 0.1 to 10 bar. Compositions were prepared that consisted of HFC-32 and [dmpim][TMeM] from about 0.9 to about 66 mole percent of HFC-32 at a temperature of 25 °C and a pressure from about 0.1 to 10 bar. Compositions were prepared that consisted of HFC-32 and

- [omim][I] from about 0.4 to about 41.6 mole percent of HFC-32 at a temperature of 25 °C and a pressure from about 0.1 to 10 bar. Compositions were prepared that consisted of HFC-32 and [doim][I] from about 0.7 to about 46.8 mole percent of HFC-32 at a temperature of 25 °C and a pressure from about 0.1 to 10 bar.
- 5 Compositions were prepared that consisted of HFC-32 and [emim][BEI] from about 1.0 to about 66.6 mole percent of HFC-32 at a temperature of 25 °C and a pressure from about 0.1 to 10 bar. Compositions were prepared that consisted of HFC-32 and [dmpim][TMeM] from about 0.8 to about 64.5 mole percent of HFC-32 at a temperature of 25 °C and a pressure from about 0.1 to 10 bar.
- 10 Compositions were prepared that consisted of HFC-32 and [pmpy][BMeI] from about 1.0 to about 63.9 mole percent of HFC-32 at a temperature of 25 °C and a pressure from about 0.1 to 10 bar. Compositions were prepared that consisted of HFC-32 and [emim][BMeI] from about 0.1 to about 78.5 mole percent of HFC-32 over a temperature range from 10 to 75 °C at a pressure from about 0.1 to 10 bar.
- 15 Compositions were prepared that consisted of HFC-32 and [bmpy][BMeI] from about 1.0 to about 64.8 mole percent of HFC-32 at a temperature of 25 °C and a pressure from about 0.1 to 10 bar. Compositions were prepared that consisted of HFC-32 and [emim][TFES] from about 1.0 to about 47.1 mole percent of HFC-32 at a temperature of 25 °C and a pressure from about 0.1 to 10 bar. Compositions
- 20 were prepared that consisted of HFC-32 and [bmim][TFES] from about 1.0 to about 55.0 mole percent of HFC-32 at a temperature of 25 °C and a pressure from about 0.1 to 10 bar. Compositions were prepared that consisted of HFC-32 and [odmim][TFES] from about 1.0 to about 56.2 mole percent of HFC-32 at a temperature of 25 °C and a pressure from about 0.1 to 10 bar. Compositions were
- 25 prepared that consisted of HFC-32 and [hmim][TFES] from about 1.0 to about 58.6 mole percent of HFC-32 at a temperature of 25 °C and a pressure from about 0.1 to 10 bar. Compositions were prepared that consisted of HFC-23 and [bmim][PF₆] from about 0.1 to about 52.8 mole percent of HFC-23 over a temperature range from 10 to 75 °C at a pressure from about 0.1 to 20 bar.
- 30 Compositions were prepared that consisted of HFC-23 and [emim][PF₆] from about 0.1 to about 15.1 mole percent of HFC-23 over a temperature range from 60 to 75 °C at a pressure from about 0.1 to 20 bar.

Figure 17 shows measured isothermal solubility data (in mole fraction) at 10 °C of the systems HFC-32, HFC-152a, HFC-134a, HFC-125, and HFC-143a + [bmim][PF₆] in terms of absolute pressure divided by the gas saturation pressure (P_0) at 10 °C shown by ratio (P/P_0). The saturation pressures for HFC-32, HFC-125, HFC-134a, HFC-143a, and HFC-152a at 10 °C are $P_0 = 11.069$ bar, $P_0 = 3.7277$ bar, $P_0 = 4.1461$ bar, $P_0 = 9.0875$, and $P_0 = 8.3628$ bar, respectively. Negative deviations from Raoult's law (i.e. curvature below the dashed line) indicate strong interaction between the refrigerant and the ionic liquid, which indicates high solubility. In particular HFC-32 has negative deviation from Raoult's law as shown in Figure 17. Compositions comprise HFC-32 and [bmim][PF₆] from about 0.1 to 63 mole percent of HFC-32 at 10 °C and P/P_0 from about 0.1 to about 0.63. Strong positive deviations from Raoult's law (i.e. curvature above the dashed line) are more typical and indicate refrigerant and ionic liquids are less soluble and eventually may form a liquid-liquid phase separation. Compositions comprise HFC-152a and [bmim][PF₆] from about 0.1 to about 80 mole percent of HFC-152a at 10 °C and P/P_0 from 0.1 to about 0.86. Compositions comprise HFC-134a and [bmim][PF₆] from about 0.1 to about 72 mole percent of HFC-134a at 10 °C and P/P_0 from about 0.1 to about 0.84. Compositions comprise HFC-125 and [bmim][PF₆] from about 0.1 mole to about 65 mole percent of HFC-125 at 10 °C and P/P_0 from about 0.1 to about 0.88. Compositions comprise HFC-143a and [bmim][PF₆] from about 0.1 to about 25 mole percent at 10 °C and P/P_0 from about 0.1 to about 0.90.

Example 36 provides details about the components of the microbalance.

Examples 37-38 show solubility and diffusivity results for CO₂ in two different ionic liquids.

EXAMPLE 3

Solubility of difluoromethane (HFC-32) in 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆]

A solubility and diffusivity study was made at temperatures of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were

calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility ($X_{calc.}$) are also provided.

Tables 4a, 4b, 4c and 4d provide data for C_o , C_s , D , $X_{calc.}$ and $X_{meas.}$ at
5 temperatures of 10, 25, 50 and 75 °C, respectively.

Table 4a

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m ² /sec)	$X_{calc.}$ (mol. fraction)	$X_{meas.}$ (mol. fraction)
10.0	0.0979	0.52	0.54	1.54E-09	0.029	0.026
10.0	0.9957	0.82	2.53	1.94E-11	0.124	0.106
10.0	2.4967	3.32	7.56	1.71E-11	0.309	0.270
10.0	3.9964	8.18	12.38	3.65E-11	0.436	0.426
10.0	5.4975	14.44	18.71	6.34E-11	0.557	0.555
10.0	6.9965	22.12	27.85	7.42E-11	0.678	0.676
10.0	8.4954	--	--	--	--	0.812

10 **Table 4b**

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m ² /sec)	$X_{calc.}$ (mol. fraction)	$X_{meas.}$ (mol. fraction)
24.9	0.0965	0.16	0.21	1.84E-10	0.012	0.018
25.0	0.9952	0.49	1.69	2.45E-11	0.086	0.076
25.0	2.4965	2.22	4.53	2.44E-11	0.206	0.189
25.0	3.9979	5.05	7.37	3.51E-11	0.303	0.295
24.9	5.4969	8.23	10.47	5.41E-11	0.390	0.387
24.9	6.9950	11.82	14.09	6.75E-11	0.473	0.471
25.0	8.5012	15.75	18.26	8.33E-11	0.550	0.548
24.9	9.9994	20.38	23.31	8.84E-11	0.624	0.622

Table 4c

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
49.6	0.0992	0.00	0.12	4.76E-11	0.007	0.006
49.9	0.9954	0.33	0.92	5.28E-11	0.048	0.047
49.9	2.4963	1.43	2.31	5.29E-11	0.115	0.113
49.9	3.9949	2.84	3.72	5.98E-11	0.174	0.173
49.9	5.4966	4.41	5.22	5.99E-11	0.231	0.229
49.9	6.9965	5.81	6.72	7.69E-11	0.282	0.282
50.0	8.4959	7.37	8.32	8.54E-11	0.331	0.331
50.0	9.9959	9.78	10.05	4.04E-11	0.379	0.377

5

Table 4d

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
75.0	0.0988	0.00	0.06	7.12E-11	0.003	0.003
75.0	0.9968	0.30	0.56	8.19E-11	0.030	0.029
75.0	2.4950	0.96	1.38	8.14E-11	0.071	0.069
75.0	3.9944	1.74	2.19	9.82E-11	0.109	0.108
74.9	5.4983	2.60	3.03	9.70E-11	0.146	0.145
74.9	6.9966	3.42	3.89	9.58E-11	0.181	0.180
75.0	8.4958	4.28	4.77	9.56E-11	0.215	0.212
75.0	9.9989	5.12	5.62	1.18E-10	0.245	0.244

EXAMPLE 4**Solubility of pentafluoroethane (HFC-125) in 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆]**

5 A solubility and diffusivity study was made at temperatures of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility

10 ($X_{\text{calc.}}$) are also provided.

Tables 5a, 5b, 5c and 5d provide data for C_0 , C_s , D , $X_{\text{calc.}}$, and $X_{\text{meas.}}$ at temperatures of 10, 25, 50 and 75 °C, respectively.

Table 5a

15

T (°C)	P (bar)	C_0 (mass %)	C_s (mass%)	D (m ² /sec)	$X_{\text{calc.}}$ (mol. fraction)	$X_{\text{meas.}}$ (mol. fraction)
9.9	0.0992	0.0	0.12	2.52E-12	0.003	0.013
10.0	0.9964	0.73	1.50	1.83E-11	0.035	0.034
10.1	1.9959	1.72	3.96	6.36E-12	0.089	0.074
10.0	2.9960	3.55	6.25	9.31E-12	0.136	0.125
10.1	3.9964	6.03	8.88	1.56E-11	0.187	0.182
9.9	4.9965	9.10	12.52	2.44E-11	0.253	0.250
10.0	5.9965	13.18	17.56	4.05E-11	0.335	0.336
9.9	6.9962	19.19	26.04	6.12E-11	0.455	0.454
10.0	7.9979	--	--	--	--	0.651

20

Table 5b

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0977	0.0	0.09	3.29E-12	0.002	0.003
25.0	0.9963	0.23	0.09	1.81E-11	0.002	0.023
25.0	1.9982	1.05	2.12	1.50E-11	0.049	0.050
24.9	2.9949	2.13	3.11	2.15E-11	0.071	0.079
25.0	3.9982	3.50	4.71	2.03E-11	0.105	0.109
25.0	4.9947	4.84	6.18	2.39E-11	0.135	0.140
25.0	5.9951	6.38	7.91	2.65E-11	0.169	0.176
25.0	7.9955	8.96	12.10	4.81E-11	0.246	0.254
24.9	9.9977	14.20	18.16	7.82E-11	0.344	0.352

Table 5c

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
49.9	0.1003	0.0	0.02	1.96E-10	0.000	0.000
49.9	0.9963	0.18	0.55	4.29E-11	0.013	0.013
49.9	1.9983	0.73	1.17	4.59E-11	0.027	0.027
50.0	2.9996	1.34	1.78	5.19E-11	0.041	0.041
49.9	3.9969	1.96	2.44	4.75E-11	0.056	0.056
50.0	4.9993	2.60	3.10	5.38E-11	0.070	0.070
49.9	5.9961	3.29	3.80	5.14E-11	0.086	0.085
49.9	7.9970	4.38	5.25	5.55E-11	0.116	0.116
49.9	9.9958	5.85	6.82	5.87E-11	0.148	0.148

5

10

Table 5d

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
75.0	0.1021	0.0	0.03	6.85E-10	0.001	0.001
74.9	0.9965	0.07	0.28	7.49E-11	0.007	0.007
75.0	1.9961	0.36	0.60	9.46E-11	0.014	0.016
75.1	2.9967	0.70	0.93	7.04E-11	0.022	0.025
75.0	3.9971	1.04	1.27	7.96E-11	0.030	0.033
75.0	4.9983	1.36	1.61	9.86E-11	0.037	0.042
75.0	5.9980	1.75	1.97	7.12E-11	0.045	0.052
75.1	7.9997	2.26	2.65	1.14E-10	0.061	0.068
75.0	9.9959	3.00	3.33	8.89E-11	0.075	0.085

5

EXAMPLE 5**Solubility of 1,1,1,2-tetrafluoroethane (HFC-134a) in 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆]**

- 10 A solubility and diffusivity study was made at temperatures of 10, 25, 50, and 75 °C over a pressure range from 0 to 3.5 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility
- 15 (X_{calc.}) are also provided.

Tables 6a, 6b, 6c and 6d provide data for C_o , C_s , D , X_{calc} , and X_{meas} at temperatures of 10, 25, 50 and 75 °C, respectively.

Table 6a

5

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
9.8	0.0999	0.0	0.23	4.21E-12	0.006	0.003
10.0	0.4981	0.35	2.20	6.46E-12	0.059	0.050
9.9	0.9986	2.25	5.73	5.78E-12	0.145	0.126
9.9	1.4981	5.40	9.15	1.01E-11	0.219	0.212
9.9	2.0024	9.50	13.64	1.48E-11	0.306	0.303
9.9	2.4907	14.39	19.36	2.67E-11	0.401	0.402
9.9	2.9974	20.96	27.51	5.33E-11	0.514	0.516
9.9	3.4900	--	--	--	--	0.721

Table 6b

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.1002	0.17	0.29	4.36E-12	0.008	0.011
24.9	0.4981	0.57	1.52	1.89E-11	0.041	0.042
25.0	0.9972	1.82	3.26	1.71E-11	0.086	0.085
25.0	1.4987	3.60	5.09	2.00E-11	0.130	0.130
25.0	1.9930	5.43	7.09	2.27E-11	0.175	0.175
24.9	2.4996	7.53	9.31	2.59E-11	0.222	0.222
25.0	2.9952	9.78	11.82	2.82E-11	0.272	0.273
24.9	3.5000	12.51	14.62	3.99E-11	0.323	0.323

Table 6c

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
49.9	0.0992	0.07	0.13	2.44E-11	0.004	0.004
50.0	0.4984	0.25	0.75	4.39E-11	0.021	0.021
49.9	0.9971	1.00	1.57	3.94E-11	0.043	0.043
49.9	1.4989	1.79	2.42	4.48E-11	0.064	0.065
50.0	1.9895	2.65	3.28	4.38E-11	0.086	0.086
50.0	2.4900	3.75	4.23	2.33E-11	0.110	0.108
50.0	2.9897	4.43	5.10	4.90E-11	0.130	0.130
50.0	3.4933	5.39	6.06	5.00E-11	0.152	0.152

5 Table 6d

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
75.0	0.0970	0.00	0.03	6.45E-11	0.001	0.001
74.9	0.4984	0.09	0.32	7.49E-11	0.009	0.009
74.9	0.9934	0.51	0.79	7.93E-11	0.022	0.022
74.9	1.5010	0.98	1.27	7.78E-11	0.035	0.035
75.0	1.9983	1.44	1.73	8.37E-11	0.047	0.046
75.0	2.5014	1.89	2.21	8.37E-11	0.059	0.059
75.0	3.0022	2.39	2.71	8.26E-11	0.072	0.072
75.0	3.4897	2.95	3.21	5.53E-11	0.085	0.084

EXAMPLE 6**Solubility of 1,1,1-trifluoroethane (HFC-143a) in 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆]**

5 A solubility and diffusivity study was made at temperatures of 10, 25, 50, and 75 °C over a pressure range from 0 to 7.5 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility

10 ($X_{\text{calc.}}$) are also provided.

Tables 7a, 7b, 7c and 7d provide data for C_o , C_s , D , $X_{\text{calc.}}$, and $X_{\text{meas.}}$ at temperatures of 10, 25, 50 and 75 °C, respectively.

Table 7a

15

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
11.7	0.0956	0.03	0.10	8.10E-12	0.003	0.003
12.0	0.9970	0.22	0.92	8.51E-12	0.031	0.029
11.9	1.9830	0.99	1.93	8.11E-12	0.064	0.060
12.0	2.9740	1.95	2.39	3.21E-12	0.078	0.093
12.3	3.9808	3.06	4.03	1.04E-11	0.127	0.124
12.0	4.9975	4.16	5.23	1.10E-11	0.161	0.156
12.0	5.9821	5.30	6.42	1.44E-11	0.192	0.188
12.2	6.9975	6.54	7.63	1.94E-11	0.223	0.219
12.2	7.4832	7.80	8.31	2.03E-11	0.239	0.235

Table 7b

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0951	0.00	0.01	1.53E-11	0.001	0.004
24.9	0.9970	0.24	0.69	2.05E-11	0.023	0.023
24.9	2.0054	0.84	1.33	2.56E-11	0.045	0.042
24.9	2.9895	1.40	2.10	1.83E-11	0.069	0.068
24.9	4.0147	2.26	2.89	1.77E-11	0.093	0.090
24.9	4.9886	2.95	3.60	2.24E-11	0.114	0.112
24.9	5.9855	3.71	4.33	2.73E-11	0.136	0.134
24.9	7.0019	4.47	5.12	2.83E-11	0.157	0.155
24.9	7.5011	5.14	5.53	3.61E-11	0.169	0.165

Table 7c

5

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
49.9	0.1050	0.00	0.03	1.51E-10	0.000	0.001
49.9	1.0023	0.16	0.40	4.47E-11	0.014	0.013
50.1	2.0045	0.61	0.84	3.41E-11	0.028	0.027
50.0	3.0002	1.03	1.26	2.90E-11	0.042	0.040
50.0	4.0021	1.39	1.65	5.08E-11	0.055	0.054
50.0	5.0046	1.81	2.08	4.10E-11	0.069	0.067
50.0	6.0039	2.29	2.50	3.75E-11	0.082	0.079
50.0	7.0029	2.63	2.90	5.57E-11	0.094	0.092
50.0	10.0030	3.56	4.16	5.51E-11	0.131	0.127

Table 7d

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
75.0	0.0995	0.00	0.01	3.86E-12	0.000	0.001
74.9	1.0005	0.18	0.26	7.38E-11	0.009	0.009
74.8	1.9960	0.38	0.54	1.04E-10	0.018	0.018
74.9	3.0001	0.67	0.81	1.07E-10	0.028	0.027
74.9	4.0015	0.91	1.08	1.32E-10	0.037	0.036
74.9	5.0027	1.18	1.36	1.20E-10	0.045	0.044
75.0	5.9979	1.44	1.63	1.40E-10	0.054	0.053
75.0	7.0026	1.92	1.94	3.79E-09	0.064	0.061
74.9	10.0035	2.65	2.76	1.90E-09	0.089	0.083

EXAMPLE 7

5 **Solubility of 1,1-difluoroethane (HFC-152a) in 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆]**

A solubility and diffusivity study was made at temperatures of 10, 25, 50, and 75 °C over a pressure range from 0 to 4.5 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were
 10 calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided.

Tables 8a, 8b, 8c and 8d provide data for C_o, C_s, D, X_{calc.}, and X_{meas.} at
 15 temperatures of 10, 25, 50 and 75 °C, respectively.

Table 8a

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
10.0	0.0973	0.10	0.73	2.13E-12	0.031	0.021
10.0	0.4994	1.23	2.90	1.14E-11	0.114	0.103
10.0	0.9933	3.58	6.11	1.56E-11	0.219	0.210
10.0	1.4985	6.91	9.60	3.09E-11	0.314	0.301
9.9	2.0011	10.40	14.00	3.60E-11	0.412	0.407
9.9	2.4952	15.52	20.42	6.44E-11	0.525	0.521
9.9	3.1963	--	--	--	--	0.797

5 Table 8b

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.1002	0.16	0.66	2.00E-11	0.028	0.030
25.0	0.5006	1.02	1.92	2.01E-11	0.078	0.077
24.9	0.9982	2.34	3.55	2.64E-11	0.137	0.136
25.0	1.4924	4.20	5.35	2.89E-11	0.196	0.194
25.0	2.4969	6.74	9.52	4.96E-11	0.312	0.311
25.0	3.4818	11.59	15.05	7.73E-11	0.433	0.432
25.0	4.5051	18.83	23.81	1.04E-10	0.573	0.574

Table 8c

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
50.1	0.9921	0.03	0.15	5.73E-11	0.007	0.007
50.0	1.0017	0.88	1.46	5.52E-11	0.060	0.060
50.0	1.5020	1.63	2.22	5.94E-11	0.089	0.089
50.0	2.4969	2.72	3.81	6.43E-11	0.145	0.145
50.0	4.5051	6.31	7.33	7.88E-11	0.254	0.254

Table 8d

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
74.9	0.1032	0.04	0.11	1.38E-10	0.005	0.005
74.9	0.5019	0.19	0.42	1.25E-10	0.018	0.018
74.9	1.0023	0.57	0.84	1.21E-10	0.035	0.035
74.9	1.5014	0.99	1.27	1.25E-10	0.052	0.052
75.0	2.4964	1.63	2.12	1.42E-10	0.085	0.085
75.0	3.4970	2.57	2.98	1.48E-10	0.117	0.117
74.8	4.5003	3.51	3.89	1.21E-10	0.148	0.149

5

EXAMPLE 8**Solubility of difluoromethane (HFC-32) in 1-butyl-3-methylimidazolium tetrafluoroborate [bmim][BF₄]**

10 A solubility and diffusivity study was made at temperatures of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided.

15

Tables 9a, 9b, 9c and 9d provide data for C_o, C_s, D, X_{calc.}, and X_{meas.} at temperatures of 10, 25, 50 and 75 °C, respectively.

Table 9a

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
9.9	0.1002	8.35	9.20	1.76E-11	0.008	0.009
9.9	0.9985	10.08	13.74	1.72E-11	0.100	0.108
10.0	2.4995	15.10	18.94	3.29E-11	0.239	0.254
10.0	3.9954	21.28	25.08	4.53E-11	0.376	0.396
9.8	5.4992	28.16	33.17	8.48E-11	0.499	0.519
9.9	6.9988	37.79	46.86	1.08E-10	0.625	0.636
9.9	8.4966	52.61	52.61	1.01E-10	0.766	0.765

Table 9b

5

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0969	0.01	0.15	3.37E-11	0.007	0.006
25.0	0.9968	0.59	1.81	3.36E-11	0.074	0.070
25.0	2.4955	2.75	4.79	3.70E-11	0.180	0.174
25.0	3.9989	5.87	7.95	4.62E-11	0.273	0.270
25.0	5.4977	9.23	11.36	5.98E-11	0.358	0.356
25.0	6.9955	12.90	15.12	7.44E-11	0.436	0.434
25.0	8.4945	17.08	19.33	9.10E-11	0.510	0.510
25.0	9.9985	21.83	24.46	9.94E-11	0.585	0.583

Table 9c

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
50.0	0.0977	0.01	0.07	8.71E-11	0.003	0.003
49.9	0.9961	0.37	0.95	7.56E-11	0.040	0.039
50.0	2.4967	1.67	2.47	7.40E-11	0.099	0.099
50.0	3.9964	3.16	4.01	8.23E-11	0.154	0.153
49.9	5.4956	4.75	5.59	8.95E-11	0.205	0.204
49.9	6.9953	6.38	7.22	9.88E-11	0.253	0.253
49.8	8.4986	8.05	8.91	1.06E-10	0.298	0.298
50.0	9.9963	9.75	10.64	1.11E-10	0.341	0.341

Table 9d

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
75.0	0.0971	0.0	0.03	1.26E-10	0.001	0.001
74.9	0.9956	0.26	0.54	1.28E-10	0.023	0.023
74.9	2.4948	1.03	1.40	1.25E-10	0.058	0.058
75.0	3.9950	1.92	2.27	1.22E-10	0.092	0.091
74.9	5.4951	2.75	3.14	1.45E-10	0.124	0.123
75.0	6.9955	3.64	4.03	1.59E-10	0.154	0.154
74.9	8.4964	4.54	4.94	1.42E-10	0.184	0.183
74.9	9.9994	5.44	5.82	1.89E-10	0.212	0.212

EXAMPLE 9

5 Solubility of difluoromethane (HFC-32) in 1,2-dimethyl-3-propylimidazolium tris(trifluoromethylsulfonyl)methide [dmpim][TMeM]

A solubility and diffusivity study was made at temperatures of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided.

Tables 10a, 10b, 10c and 10d provide data for C_o, C_s, D, X_{calc.}, and X_{meas.} at temperatures of 10, 25, 50 and 75 °C, respectively.

Table 10a

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
10.0	0.1010	0.03	0.11	1.71E-11	0.012	0.012
10.0	0.9964	0.43	1.44	1.39E-11	0.134	0.136
10.0	2.4970	2.39	4.13	2.52E-11	0.313	0.311
10.0	3.9969	5.57	7.39	5.04E-11	0.458	0.457
10.0	5.4947	9.70	11.67	8.93E-11	0.583	0.583
10.0	6.9966	15.43	17.70	1.37E-10	0.695	0.696
10.0	8.4959	24.33	28.09	1.56E-10	0.805	0.802

5 Table 10b

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
24.9	0.0998	0.01	0.09	2.71E-11	0.010	0.010
24.9	0.9997	0.42	1.01	2.52E-11	0.098	0.096
24.9	2.4956	--	--	--	--	0.225
24.9	3.9958	3.61	4.55	5.46E-11	0.336	0.335
24.9	5.4927	5.76	6.69	7.98E-11	0.432	0.431
24.9	6.9955	8.15	9.13	1.10E-10	0.516	0.515
24.9	8.4948	11.02	12.07	1.34E-10	0.593	0.593
24.9	10.0000	14.52	15.59	1.83E-10	0.662	0.662

Table 10c

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
50.0	0.0991	0.21	0.04	6.45E-11	0.004	0.004
50.0	0.9995	0.29	0.57	6.75E-11	0.058	0.057
50.0	2.4945	1.11	1.52	7.87E-11	0.141	0.141
50.0	3.9947	2.10	2.50	9.56E-11	0.213	0.213
50.0	5.4954	3.15	3.51	1.15E-10	0.278	0.278
50.0	6.9968	4.24	4.59	1.33E-10	0.338	0.338
50.0	8.4944	5.37	5.73	1.51E-10	0.392	0.392
50.0	9.9952	6.61	6.96	1.68E-10	0.442	0.442

5 Table 10d

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
75.0	0.0940	0.0	0.0	5.75E-11	0.000	0.000
74.9	1.0018	0.06	0.31	6.06E-11	0.032	0.031
75.0	2.5040	0.71	0.89	1.23E-10	0.087	0.087
74.9	3.9958	1.32	1.49	1.26E-10	0.138	0.138
74.9	5.4938	1.92	2.09	1.59E-10	0.184	0.184
74.9	7.0051	2.58	2.72	1.35E-10	0.229	0.229
74.9	8.4954	3.24	3.37	1.19E-10	0.270	0.268
74.9	10.0046	3.89	4.05	2.10E-10	0.309	0.308

EXAMPLE 10**Solubility of difluoromethane (HFC-32) in 1-octyl-3-methylimidazolium iodide [omim][I]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided.

Table 11 provides data for C_o , C_s , D, $X_{\text{calc.}}$, and $X_{\text{meas.}}$ at a temperature of 25 °C.

Table 11

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	$X_{\text{calc.}}$ (mol. fraction)	$X_{\text{meas.}}$ (mol. fraction)
25.0	0.1007	0.01	0.06	1.75E-11	0.004	0.004
25.2	1.0021	0.23	0.80	1.77E-11	0.048	0.048
25.0	2.4971	1.20	2.13	1.86E-11	0.119	0.118
25.0	3.9999	2.58	3.55	2.27E-11	0.186	0.185
25.0	5.5008	4.07	5.04	3.13E-11	0.247	0.246
25.0	6.9964	5.64	6.64	3.81E-11	0.306	0.306
25.0	8.5027	7.52	8.33	2.86E-11	0.360	0.362
25.0	10.0022	9.27	10.35	6.37E-11	0.417	0.416

EXAMPLE 11**Solubility of difluoromethane (HFC-32) in 1,3-dioctylimidazolium iodide [doim][I]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided.

Table 12 provides data for C_o , C_s , D , X_{calc} , and X_{meas} at a temperature of 25 °C.

Table 12

5

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc} (mol.fraction)	X_{meas} (mol. fraction)
25.0	0.1002	0.03	0.11	1.78E-11	0.009	0.007
25.0	1.0010	0.29	0.87	2.11E-11	0.066	0.064
25.0	2.5003	1.29	2.17	2.35E-11	0.152	0.150
25.0	4.0024	2.62	3.51	2.91E-11	0.227	0.225
25.0	5.5024	4.03	4.93	3.54E-11	0.295	0.293
25.0	7.0010	5.51	6.43	4.25E-11	0.357	0.355
24.9	8.4988	7.12	8.07	5.00E-11	0.415	0.413
25.0	10.0024	8.83	9.85	5.77E-11	0.469	0.468

EXAMPLE 12

Solubility of difluoromethane (HFC-32) in 1-ethyl-3-methylimidazolium

10 bis(pentafluoroethylsulfonyl)imide [emim][BEI]

A solubility and diffusivity study was made at a temperature of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial
15 concentration (C_o), final saturation concentration (C_s), and calculated solubility (X_{calc}) are also provided.

Tables 13a, 13b, 13c and 13d provide data for C_o , C_s , D , X_{calc} , and X_{meas} at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

Table 13a

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
10.0	0.101	0.06	0.15	3.79E-11	0.014	0.014
10.0	1.000	1.06	1.78	4.78E-11	0.146	0.144
10.0	2.495	3.58	4.83	7.37E-11	0.324	0.323
10.0	3.995	7.14	8.52	1.17E-10	0.468	0.467
10.0	5.496	11.75	13.23	1.51E-10	0.590	0.590
10.0	6.994	17.76	19.75	1.72E-10	0.699	0.699
10.0	8.505	26.95	30.37	1.67E-10	0.805	0.799

Table 13b

5

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.096	0.03	0.11	7.5E-11	0.010	0.010
25.0	0.997	0.71	1.22	7.9E-11	0.104	0.104
25.0	2.496	2.49	3.19	1.1E-10	0.237	0.237
25.0	3.996	4.61	5.33	1.3E-10	0.347	0.347
25.0	5.493	7.03	7.75	1.6E-10	0.443	0.442
25.0	6.993	9.70	10.49	1.8E-10	0.525	0.525
25.0	8.503	12.87	13.71	2.1E-10	0.600	0.598
25.0	10.005	16.49	17.56	1.7E-10	0.668	0.666

Table 13c

10

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
50.0	0.100	0.00	0.04	1.66E-10	0.004	0.004
50.0	0.997	0.49	0.65	1.34E-10	0.058	0.059
50.0	2.497	1.46	1.73	1.79E-10	0.142	0.145
50.0	3.996	2.61	2.83	1.92E-10	0.216	0.219
50.0	5.495	3.82	3.98	2.19E-10	0.281	0.285
50.0	6.995	4.92	5.19	2.28E-10	0.341	0.345
50.0	8.504	6.20	6.46	2.73E-10	0.395	0.399
50.0	9.993	7.54	7.81	1.62E-10	0.444	0.449

Table 13d

T (°C)	P (bar)	C _o (mass%)	C _s (mass%)	D (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
74.9	0.101	0.00	0.01	3.92E-10	0.001	0.001
74.9	1.000	0.32	0.41	2.60E-10	0.038	0.038
74.9	2.501	0.99	1.10	3.32E-10	0.095	0.095
74.9	3.992	1.72	1.79	3.96E-10	0.147	0.146
74.9	5.496	2.39	2.49	3.53E-10	0.194	0.194
74.9	6.996	3.08	3.22	3.41E-10	0.239	0.239
74.9	8.504	3.87	3.96	3.48E-10	0.280	0.280
74.9	9.994	4.55	4.70	1.92E-10	0.318	0.317

5 EXAMPLE 13

Solubility of difluoromethane (HFC-32) in 1,2-dimethyl-3-propylimidazolium bis(trifluoromethylsulfonyl)imide [dmpim][BMeI]

A solubility and diffusivity study was made at a temperature of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided.

Table 14 provides data for C_o, C_s, D, X_{calc.}, and X_{meas.} at a temperature of 25 °C.

Table 14

T (°C)	P (bar)	C _o (mass%)	C _s (mass%)	D (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
24.9	0.0989	0.02	0.11	6.31E-11	0.008	0.008
25.0	0.9951	0.65	1.22	6.60E-11	0.091	0.090
25.0	2.4949	2.44	3.25	8.94E-11	0.213	0.212
25.0	3.9762	4.62	5.46	1.21E-10	0.317	0.317
25.0	5.5013	7.08	8.00	1.46E-10	0.412	0.412
25.0	7.0174	10.02	10.92	1.75E-10	0.497	0.496
25.0	8.5131	13.56	14.29	2.23E-10	0.573	0.573
25.0	10.0108	17.55	18.41	2.33E-10	0.645	0.645

EXAMPLE 14**Solubility of difluoromethane (HFC-32) in 3-methyl-1-propylpyridinium bis(trifluoromethylsulfonyl)imide [pmpy][BMeI]**

A solubility and diffusivity study was made at a temperature of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided.

Tables 15a, 15b, 15c, and 15d provide data for C_0 , C_s , D , $X_{\text{calc.}}$, and $X_{\text{meas.}}$ at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

Table 15a

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
10.0	0.1021	0.08	0.02	5.76E-11	0.002	0.015
10.0	1.0001	1.03	2.01	5.72E-11	0.141	0.140
10.0	2.4942	3.95	5.31	1.05E-10	0.310	0.311
10.0	3.9963	7.78	9.35	1.28E-10	0.452	0.452
10.0	5.4935	12.68	14.05	2.89E-10	0.567	0.570
10.0	6.9960	18.73	20.79	2.01E-10	0.678	0.679
10.0	8.4951	27.80	30.88	2.71E-10	0.781	0.778

15

Table 15b

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
24.9	0.0951	0.02	0.12	9.96E-11	0.010	0.010
24.9	1.0020	0.74	1.32	1.00E-10	0.097	0.096
24.9	2.5034	2.67	3.44	1.20E-10	0.222	0.221
24.9	3.9959	4.93	5.73	1.52E-10	0.327	0.328
24.9	5.4973	7.52	8.30	1.92E-10	0.420	0.419
24.9	6.9923	10.35	11.16	2.20E-10	0.501	0.502
24.9	8.4965	13.61	14.48	2.41E-10	0.575	0.575
24.9	10.0044	17.35	18.06	6.21E-10	0.638	0.639

Table 15c

T (°C)	P (bar)	C ₀ (mass%)	C _s (mass%)	D (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
50.0	0.1025	0.04	0.08	2.10E-10	0.007	0.007
50.0	1.0031	0.59	0.76	1.86E-10	0.058	0.058
50.0	2.4979	1.64	1.93	2.01E-10	0.136	0.137
50.0	4.0004	2.82	3.11	2.80E-10	0.205	0.206
50.0	5.4945	4.05	4.36	2.37E-10	0.268	0.270
50.0	6.9935	5.39	5.64	3.50E-10	0.323	0.326
50.0	8.5031	6.71	6.97	3.95E-10	0.375	0.378
50.0	9.9939	8.06	8.44	2.30E-10	0.425	0.427

5 Table 15d

T (°C)	P (bar)	C ₀ (mass%)	C _s (mass%)	D (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
74.9	0.1026	0.03	0.04	3.94E-10	0.003	0.003
74.9	1.0023	0.04	0.46	3.89E-10	0.036	0.037
74.9	2.5020	1.06	1.19	3.96E-10	0.088	0.089
74.9	4.0021	1.77	1.91	4.00E-10	0.135	0.138
74.9	5.4931	2.53	2.65	3.62E-10	0.179	0.183
74.9	7.0026	3.27	3.39	4.62E-10	0.219	0.223
74.9	8.4935	4.04	4.15	4.76E-10	0.257	0.262
74.9	10.0019	4.76	4.91	5.48E-10	0.293	0.300

EXAMPLE 15

10 **Solubility of trifluoromethane (HFC-23) in 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆]**

A solubility and diffusivity study was made at a temperature of 10, 25, 50, and 75 °C over a pressure range from 0 to 20 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were
 15 calculated using a one-dimensional diffusion model analysis. The initial concentration (C₀), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided.

Tables 16a, 16b, 16c, and 16d provide data for C₀, C_s, D, X_{calc.}, and X_{meas.} at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

Table 16a

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
9.4	0.0962	--	--	--	--	0.000
9.4	0.5000	0.00	0.25	1.54E-11	0.010	0.010
9.6	1.0979	--	--	--	--	0.028
9.5	4.0003	1.56	3.05	1.54E-11	0.113	0.113
9.4	7.0000	4.14	5.76	2.17E-11	0.199	0.198
9.5	9.9934	7.15	8.81	2.89E-11	0.282	0.281
9.5	12.9972	10.59	12.22	4.26E-11	0.361	0.361
9.5	14.9964	13.48	14.81	5.68E-11	0.414	0.414
10.0	20.0017	--	--	--	--	0.528

Table 16b

5

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
24.9	0.0991	--	--	--	--	0.000
24.9	0.4972	0.03	0.19	2.56E-11	0.008	0.008
24.9	0.9994	0.24	0.44	3.22E-11	0.018	0.018
24.9	3.9934	1.17	2.08	2.37E-11	0.080	0.079
24.9	6.9953	2.86	3.79	3.01E-11	0.138	0.137
24.9	10.0041	4.68	5.59	3.95E-11	0.194	0.193
24.9	13.0056	6.66	7.52	3.89E-11	0.248	0.247
25.0	15.0000	8.09	8.80	5.73E-11	0.281	0.281
24.9	19.9990	11.36	12.49	7.12E-11	0.367	0.367

Table 16c

10

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
50.0	0.0981	0.00	0.01	6.34E-11	0.000	0.000
50.0	0.4984	0.03	0.11	6.26E-11	0.005	0.005
50.0	0.9961	0.17	0.27	7.35E-11	0.011	0.011
50.0	3.9965	0.89	1.27	5.88E-11	0.049	0.049
50.0	7.0036	1.90	2.25	6.74E-11	0.085	0.085
50.0	10.0041	2.92	3.27	8.02E-11	0.121	0.120
50.0	12.9931	3.95	4.29	7.47E-11	0.154	0.154
50.0	15.0015	4.69	5.01	1.16E-10	0.176	0.176
50.0	19.9932	6.41	6.78	1.08E-10	0.228	0.227

Table 16d

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
75.0	0.0965	--	--	--	--	0.001
74.9	0.4973	0.03	0.08	8.13E-11	0.003	0.003
74.9	0.9975	0.12	0.21	1.22E-10	0.008	0.008
74.9	3.9971	0.63	0.84	1.04E-10	0.033	0.033
74.9	7.0016	1.45	1.42	2.86E-12	0.055	0.057
75.0	9.9934	1.92	2.08	1.08E-10	0.079	0.080
74.9	13.0031	2.55	2.72	2.23E-10	0.102	0.103
74.9	14.9943	2.98	3.17	1.09E-10	0.117	0.118
74.9	19.9998	4.00	4.22	2.31E-10	0.152	0.146

EXAMPLE 16

5

Solubility of trifluoromethane (HFC-23) in 1-ethyl-3-methylimidazolium hexafluorophosphate [emim][PF₆]

A solubility and diffusivity study was made at a temperature of 60, and 75 °C over a pressure range from 0 to 20 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were
10 calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided.

Tables 17a, and 17b provide data for C_o, C_s, D, X_{calc.}, and X_{meas.} at a
15 temperature of 60°C, and 75°C, respectively.

Table 17a

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
59.9	0.0992	--	--	--	--	0.000
59.9	0.4997	0.03	0.09	1.23E-10	0.003	0.003
59.9	0.9973	0.13	0.20	1.28E-10	0.007	0.007
59.9	4.0026	0.76	0.86	1.21E-10	0.031	0.030
59.9	6.9974	1.30	1.50	1.58E-10	0.053	0.053
59.9	10.0001	2.02	2.18	1.12E-10	0.075	0.076
60.0	12.9920	2.71	2.86	2.55E-10	0.097	0.098
59.9	15.0002	3.20	3.35	1.68E-10	0.113	0.113
59.9	19.9990	4.39	4.54	3.12E-10	0.148	0.151

20

Table 17b

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
75.0	0.0965	0.02	0.02	1.12E-10	0.001	0.001
74.9	0.4982	--	--	--	--	0.002
74.9	0.9998	0.12	0.16	1.94E-10	0.006	0.006
74.9	4.0035	0.56	0.65	2.18E-10	0.023	0.024
74.9	6.9933	1.06	1.14	1.17E-10	0.040	0.040
74.9	10.0041	1.56	1.65	2.73E-10	0.058	0.057
75.0	12.9969	2.00	2.16	1.02E-10	0.075	0.074
74.9	15.0041	2.47	2.49	7.22E-10	0.085	0.083
75.0	19.9939	--	--	--	--	0.116

5

EXAMPLE 17**Solubility of difluoromethane (HFC-32) in 1-ethyl-3-methylimidazolium bis(trifluoroethylsulfonyl)imide [emim][BMeI]**

5 A solubility and diffusivity study was made at a temperature of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility

10 ($X_{\text{calc.}}$) are also provided.

Tables 18a, 18b, 18c, and 18d provide data for C_0 , C_s , D , $X_{\text{calc.}}$, and $X_{\text{meas.}}$ at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

15 **Table 18a**

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
10.0	0.1015	0.11	0.19	6.94E-11	0.014	0.014
10.0	1.0012	1.12	2.06	8.72E-11	0.137	0.136
10.0	2.5030	4.25	5.55	1.18E-10	0.306	0.305
10.0	3.9929	8.20	9.58	1.50E-10	0.444	0.446
10.0	5.4925	13.38	14.83	1.78E-10	0.567	0.567
10.0	7.0043	19.75	21.63	2.36E-10	0.675	0.668
10.0	8.4935	27.92	31.92	1.24E-10	0.779	0.785

Table 18b

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0959	0.09	0.13	8.36E-11	0.010	0.010
25.0	0.9981	0.86	1.38	1.22E-10	0.095	0.095
25.0	2.5024	2.88	3.56	1.61E-10	0.217	0.217
25.0	3.9937	5.27	5.97	1.56E-10	0.323	0.323
25.0	5.4940	7.90	8.60	2.00E-10	0.414	0.414
25.0	6.9946	10.77	11.53	2.33E-10	0.495	0.495
25.0	8.4952	14.06	14.80	3.24E-10	0.566	0.565
25.0	9.9967	17.74	18.58	3.20E-10	0.632	0.637

20

Table 18c

T (°C)	P (bar)	C _o (mass%)	C _s (mass%)	D (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
50.0	0.1022	0.04	0.07	3.03E-10	0.005	0.005
50.0	1.0029	0.55	0.77	2.18E-10	0.055	0.055
50.0	2.4972	1.71	1.98	2.19E-10	0.132	0.132
50.0	4.0011	2.95	3.21	2.86E-10	0.199	0.199
50.0	5.4949	4.22	4.50	2.47E-10	0.261	0.262
50.0	7.0033	5.52	5.80	3.97E-10	0.316	0.316
50.0	8.5044	6.93	7.20	2.90E-10	0.368	0.364
50.0	10.0038	8.22	8.51	3.43E-10	0.411	0.412

Table 18d

T (°C)	P (bar)	C _o (mass%)	C _s (mass%)	D (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
74.9	0.1028	0.01	0.03	6.36E-10	0.002	0.002
74.9	0.9981	0.36	0.46	3.41E-10	0.034	0.034
74.9	2.4971	1.09	1.21	4.21E-10	0.084	0.084
74.9	3.9948	1.82	1.96	5.11E-10	0.130	0.130
74.9	5.5026	2.60	2.71	5.24E-10	0.173	0.173
74.9	6.9919	3.37	3.49	3.22E-10	0.213	0.213
74.9	8.5039	4.16	4.28	4.63E-10	0.252	0.251
74.9	10.0045	5.10	5.10	4.75E-09	0.288	0.284

EXAMPLE 18**Solubility of difluoromethane (HFC-32) in 1-butyl-3-methylpyridinium****10 bis(trifluoromethylsulfonyl)imide [bmpy][BMeI]**

A solubility and diffusivity study was made at a temperature of 25°C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided in Table 19.

Table 19.

T (°C)	P (bar)	C ₀ (mass%)	C _s (mass%)	D (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
25.0	0.0961	0.04	0.12	6.81E-11	0.010	0.010
25.0	0.9950	0.66	1.32	7.82E-11	0.097	0.100
25.0	2.4949	2.58	3.38	1.21E-10	0.219	0.223
25.0	3.9948	4.76	5.59	1.49E-10	0.321	0.329
25.0	5.4962	7.25	8.10	1.53E-10	0.414	0.424
25.0	7.0055	--	--	--	--	0.505
25.0	8.5057	13.03	14.47	1.15E-11	0.575	0.580
25.0	10.0002	17.06	18.28	2.31E-10	0.642	0.648

EXAMPLE 19

5

**Solubility of difluoromethane (HFC-32) in 1-ethyl-3-methylimidazolium
1,1,2,2-tetrafluoroethanesulfonate [emim][TFES]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C₀), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided in Table 20.

15 Table 20

T (°C)	P (bar)	C ₀ (mass%)	C _s (mass%)	D (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
25.0	0.0987	0.01	0.10	4.12E-11	0.006	0.006
24.9	0.9910	0.40	1.03	3.25E-11	0.055	0.054
24.9	2.4841	2.48	2.65	2.94E-11	0.133	0.132
24.9	3.9945	3.66	4.45	4.93E-11	0.207	0.207
24.9	5.4957	5.78	6.37	5.92E-11	0.276	0.277
24.9	7.0221	--	--	--	--	0.344
24.9	8.4832	9.79	10.90	1.04E-10	0.407	0.407
24.9	10.0160	12.55	13.66	1.21E-10	0.470	0.471

EXAMPLE 20**Solubility of difluoromethane (HFC-32) in 1-butyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate [bmim][TFES]**

5 A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided in

10 Table 21.

Table 21

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0967	0.02	0.12	2.37E-11	0.007	0.007
25.0	0.9986	0.99	1.29	1.47E-11	0.075	0.072
25.0	2.4997	2.19	3.31	2.67E-11	0.174	0.171
25.0	3.9716	4.33	5.40	3.95E-11	0.260	0.261
25.0	5.4838	6.84	7.78	4.76E-11	0.342	0.342
25.0	6.9946	8.98	10.39	7.75E-11	0.416	0.416
25.0	8.4811	11.98	13.27	8.73E-11	0.485	0.485
25.0	9.9886	15.07	16.62	1.35E-10	0.551	0.550

EXAMPLE 21

15 **Solubility of difluoromethane (HFC-32) in 1-dodecyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate [dmim][TFES]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a

20 one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided in Table 22.

Table 22

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0963	0.00	0.06	5.01E-11	0.005	0.006
25.0	0.9950	0.35	0.95	4.72E-11	0.072	0.074
25.0	2.5100	1.63	2.56	5.06E-11	0.175	0.178
25.0	3.9971	4.15	4.30	3.01E-11	0.266	0.271
25.0	5.4807	6.06	6.16	4.74E-11	0.346	0.353
25.0	7.0007	7.98	8.29	6.81E-11	0.421	0.429
25.0	8.5003	10.50	10.66	8.17E-11	0.490	0.497
25.0	10.0101	12.09	13.39	1.25E-10	0.555	0.562

EXAMPLE 22

5 **Solubility of difluoromethane (HFC-32) in 1-heptyl-3-methylimidazolium 1,1,2,2-tetrafluoroethanesulfonate [hmim][TFES]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C₀), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided in Table 23.

Table 23

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0988	0.01	0.11	3.86E-11	0.008	0.008
25.0	1.0023	0.47	1.25	3.87E-11	0.081	0.081
25.0	2.5100	2.18	3.30	4.35E-11	0.192	0.190
25.0	3.9884	4.39	5.44	5.84E-11	0.286	0.286
25.0	5.4973	7.25	7.82	6.41E-11	0.371	0.371
25.0	6.9871	9.99	10.43	9.01E-11	0.448	0.448
25.0	8.4785	12.28	13.40	1.30E-10	0.518	0.518
25.0	9.9795	15.45	16.83	1.56E-10	0.585	0.586

EXAMPLE 23**Solubility of difluoromethane (HFC-32) in 1-butyl-3-methylimidazolium acetate [bmim][Ac]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided in Table 24.

Table 24

T (°C)	P (bar)	C_0 (mass%)	C_s (mass%)	D (m²/sec)	$X_{\text{calc.}}$ (mol. fraction)	$X_{\text{meas.}}$ (mol. fraction)
25.1	0.0985	0.09	0.25	2.19E-11	0.010	0.010
25.0	0.9968	0.72	2.17	2.64E-11	0.078	0.077
25.0	2.4979	3.25	5.30	4.05E-11	0.176	0.174
25.0	4.0040	6.59	8.59	5.64E-11	0.264	0.258
25.0	5.4984	9.83	11.70	1.02E-10	0.335	0.333
25.0	6.9974	13.24	15.00	1.46E-10	0.402	0.397
24.9	8.5016	16.74	18.36	1.83E-10	0.462	0.456
25.0	10.0044	20.30	21.89	2.10E-10	0.516	0.511

EXAMPLE 24**Solubility of difluoromethane (HFC-32) in 1-butyl-3-methylimidazolium 2-(1,2,2,2-tetrafluoroethoxy)-1,1,2,2-tetrafluoroethanesulfonate [bmim][FS]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided in Table 25.

Table 25

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0999	0.02	0.11	4.30E-11	0.009	0.009
25.0	0.9966	0.82	1.20	4.29E-11	0.092	0.092
25.0	2.5009	2.29	3.17	5.44E-11	0.215	0.213
25.0	4.0040	4.16	5.26	9.11E-11	0.318	0.317
25.0	5.4999	6.53	7.68	1.04E-10	0.411	0.411
25.0	6.9963	9.19	10.36	1.49E-10	0.492	0.493
25.0	8.4944	12.24	13.24	1.26E-09	0.561	0.565
25.0	10.0048	15.74	17.00	2.78E-10	0.632	0.632

EXAMPLE 25

5 **Solubility of difluoromethane (HFC-32) in 1-butyl-3-methylimidazolium 1, 1, 2, 3, 3, 3-hexafluoropropanesulfonate [bmim][HFPS]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided in Table 26.

Table 26

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0945	0.02	0.11	3.33E-11	0.010	0.010
25.0	0.9999	0.56	1.25	3.17E-11	0.106	0.104
25.0	2.4976	2.29	3.29	3.90E-11	0.242	0.241
25.0	3.9945	4.34	5.40	6.98E-11	0.349	0.347
25.0	5.4949	6.56	7.79	6.98E-11	0.443	0.443
25.0	6.9975	9.29	10.45	1.11E-10	0.523	0.523
25.0	8.4943	12.16	13.60	1.04E-10	0.597	0.599
25.0	10.0042	15.98	17.43	1.67E-10	0.665	0.664

EXAMPLE 26**Solubility of difluoromethane (HFC-32) in 1-butyl-3-methylimidazolium methyl sulfonate [bmim][MeSO₄]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided in Table 27.

Table 27

T (°C)	P (bar)	C_0 (mass%)	C_s (mass%)	D (m ² /sec)	$X_{\text{calc.}}$ (mol. fraction)	$X_{\text{meas.}}$ (mol. fraction)
25.0	0.0993	0.12	0.24	2.08E-11	0.012	0.012
25.0	1.0010	0.53	1.48	2.67E-11	0.068	0.068
25.0	2.4982	2.15	3.65	3.04E-11	0.154	0.155
25.0	3.9954	4.41	5.87	4.15E-11	0.231	0.232
25.1	5.5009	6.87	8.16	5.23E-11	0.299	0.302
25.0	6.9953	9.24	10.77	6.24E-11	0.367	0.369
25.0	8.5005	11.97	13.33	9.89E-11	0.425	0.427
25.0	10.0059	14.75	16.32	1.20E-10	0.484	0.482

EXAMPLE 27**Solubility of difluoromethane (HFC-32) in 1-butyl-3-methylimidazolium thiocyanate [bmim][SCN]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided in Table 28.

Table 28

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0947	0.02	0.10	8.08E-11	0.004	0.004
25.0	1.0031	0.45	1.11	8.57E-11	0.041	0.041
25.0	2.5033	1.90	2.84	1.03E-10	0.100	0.099
25.0	3.9958	3.66	4.68	1.11E-10	0.157	0.156
25.0	5.4999	--	--	--	--	0.212
25.0	6.9966	7.62	8.73	1.42E-10	0.266	0.267
25.0	8.4947	9.93	11.01	1.83E-10	0.319	0.320
25.0	9.9919	12.30	13.55	2.05E-10	0.373	0.373

EXAMPLE 28

5 **Solubility of difluoromethane (HFC-32) in 1-butyl-3-methylimidazolium 1, 1, 2-trifluoro-2-(perfluoroethoxy)ethanesulfonate [bmim][TPES]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided in Table 29.

Table 29

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0951	0.02	0.12	4.46E-11	0.010	0.010
25.0	1.0007	0.58	1.35	5.27E-11	0.103	0.102
25.0	2.4964	2.43	3.56	6.70E-11	0.236	0.236
25.0	3.9947	4.81	5.94	9.64E-11	0.346	0.346
25.0	5.4938	7.52	8.62	1.20E-10	0.442	0.442
25.0	6.9941	10.49	11.65	1.49E-10	0.525	0.525
25.0	8.4946	13.93	15.15	1.78E-10	0.600	0.599
25.0	9.9937	18.00	19.36	2.06E-10	0.668	0.668

EXAMPLE 29**Solubility of difluoromethane (HFC-32) in 1-butyl-3-methylimidazolium 1, 1, 2-trifluoro-2-(trifluoromethoxy)ethanesulfonate [bmim][TTES]**

A solubility and diffusivity study was made at a temperature of 25 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided in Table 30.

10

Table 30

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
25.0	0.0947	0.02	0.13	4.26E-11	0.010	0.010
25.0	1.0031	0.57	1.42	4.51E-11	0.097	0.096
25.0	2.5033	2.40	3.71	5.83E-11	0.222	0.222
25.0	3.9958	4.92	6.28	7.11E-11	0.332	0.332
25.0	5.4999	7.79	9.04	9.96E-11	0.425	0.424
25.0	6.9966	10.71	12.12	1.23E-10	0.506	0.506
25.0	8.4947	14.21	15.63	1.59E-10	0.579	0.578
25.0	9.9919	18.20	19.62	2.51E-10	0.644	0.644

EXAMPLE 30

15

Solubility of 1,1,1,2-tetrafluoroethane (HFC-134a) in 1-butyl-3-methylimidazolium 1,1,2-trifluoro-2-(trifluoromethoxy)ethanesulfonate [bmim][TTES]

A solubility and diffusivity study was made at a temperature of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided.

20

Tables 18a, 18b, 18c, and 18d provide data for C_o , C_s , D , X_{calc} , and X_{meas} at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

Table 31a

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	$D_{eff.}$ (m ² /sec)	$X_{calc.}$ (mol. fraction)	$X_{meas.}$ (mol. fraction)
10.0	0.1025	0.08	0.66	1.04E-11	0.025	0.026
10.0	0.5002	0.97	3.29	1.25E-11	0.114	0.117
10.0	1.0027	4.03	7.05	1.62E-11	0.223	0.225
10.0	1.5018	7.93	11.31	2.16E-11	0.326	0.326
9.9	2.0022	12.23	16.25	3.26E-11	0.424	0.424
10.0	2.5048	17.58	22.11	5.31E-11	0.518	0.514
10.0	2.9946	23.87	30.15	5.28E-11	0.620	0.628
10.0	3.5047	36.32	44.43	7.71E-11	0.752	0.745

5

Table 31b

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	$D_{eff.}$ (m ² /sec)	$X_{calc.}$ (mol. fraction)	$X_{meas.}$ (mol. fraction)
24.9	0.1018	1.51	0.35	1.19E-11	0.013	0.017
24.9	0.5032	0.77	2.07	2.17E-11	0.074	0.075
25.1	1.0024	2.52	4.22	2.60E-11	0.143	0.143
24.8	1.5015	4.77	6.52	3.00E-11	0.209	0.208
25.0	2.0032	7.17	9.00	3.27E-11	0.272	0.271
25.0	2.5035	9.59	11.56	4.43E-11	0.331	0.331
24.9	3.0013	12.31	14.44	5.05E-11	0.390	0.389
24.8	3.5010	15.87	17.69	4.50E-11	0.449	0.450

Table 31c

10

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	$D_{eff.}$ (m ² /sec)	$X_{calc.}$ (mol. fraction)	$X_{meas.}$ (mol. fraction)
50.0	0.1048	0.17	0.25	5.76E-11	0.009	0.009
50.0	0.5031	0.47	1.06	5.35E-11	0.039	0.039
50.0	1.0023	1.37	2.11	5.79E-11	0.076	0.076
50.0	1.5021	2.43	3.19	6.35E-11	0.111	0.111
50.0	2.0026	3.50	4.28	6.64E-11	0.145	0.145
50.0	2.5033	4.67	5.41	6.97E-11	0.178	0.179
50.0	3.0035	5.81	6.58	7.24E-11	0.211	0.211
50.0	3.5016	7.22	7.78	6.89E-11	0.242	0.243

Table 31d

T (°C)	P (bar)	C ₀ (mass%)	C _s (mass%)	D _{eff.} (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
75.0	0.1031	0.06	0.13	1.04E-10	0.005	0.005
74.9	0.5054	0.31	0.62	1.18E-10	0.023	0.023
74.9	1.0049	0.85	1.23	1.22E-10	0.045	0.045
74.9	1.5029	1.49	1.85	1.21E-10	0.067	0.067
74.9	2.0041	2.10	2.46	1.25E-10	0.087	0.087
74.9	2.5042	2.71	3.08	1.26E-10	0.107	0.108
74.9	3.0024	3.33	3.72	1.38E-10	0.128	0.128
74.9	3.5039	4.19	4.36	1.09E-10	0.147	0.147

EXAMPLE 31

5

Solubility of 1,1,1,2-tetrafluoroethane (HFC-134a) in 1-butyl-3-methylimidazolium 1,1,2-trifluoro-2-(perfluoroethoxy)ethanesulfonate [bmim][TPES]

A solubility and diffusivity study was made at a temperature of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C₀), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided.

Tables 32a, 32b, 32c, and 32d provide data for C₀, C_s, D, X_{calc.}, and X_{meas.} at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

Table 32a

T (°C)	P (bar)	C ₀ (mass%)	C _s (mass%)	D _{eff.} (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
10.0	0.1024	0.06	0.66	9.33E-12	0.028	0.028
10.0	0.5038	1.01	3.39	1.15E-11	0.131	0.132
10.0	1.0043	4.05	7.26	1.71E-11	0.251	0.253
9.9	1.5033	8.17	11.65	2.53E-11	0.361	0.362
10.0	2.0022	12.78	16.90	3.67E-11	0.465	0.464
10.0	2.5024	18.33	23.30	5.37E-11	0.565	0.566
10.0	3.0041	25.90	32.36	7.06E-11	0.672	0.670
9.9	3.5039	38.42	47.48	6.49E-11	0.794	0.796

20

Table 32b

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
24.9	0.1026	0.11	0.45	1.80E-11	0.019	0.018
24.9	0.5031	0.72	2.09	2.32E-11	0.084	0.084
24.9	1.0018	2.62	4.33	2.59E-11	0.162	0.162
24.9	1.5015	4.92	6.70	3.23E-11	0.235	0.235
24.9	2.0029	7.33	9.23	4.14E-11	0.303	0.303
24.9	2.5038	9.92	11.93	4.99E-11	0.367	0.366
24.9	3.0034	12.73	14.93	5.74E-11	0.429	0.428
24.9	3.5012	16.44	18.40	4.94E-11	0.491	0.490

5 Table 32c

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
50.0	0.1036	0.20	0.26	8.37E-11	0.011	0.011
50.0	0.5032	0.47	1.10	5.99E-11	0.045	0.045
50.0	1.0023	1.52	2.20	5.66E-11	0.088	0.087
50.0	1.5021	2.55	3.32	6.59E-11	0.128	0.128
50.0	2.0025	3.69	4.47	6.87E-11	0.167	0.167
50.0	2.5035	4.90	5.66	7.37E-11	0.204	0.204
50.0	3.0042	6.08	6.87	8.56E-11	0.240	0.240
50.0	3.5035	7.49	8.10	8.02E-11	0.274	0.274

Table 32d

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
74.9	0.1051	0.11	0.15	1.09E-10	0.006	0.006
74.9	0.5052	0.34	0.65	1.19E-10	0.027	0.027
74.9	1.0054	0.92	1.29	1.22E-10	0.053	0.053
75.0	1.5046	1.90	1.93	1.93E-09	0.078	0.078
74.7	2.0056	2.25	2.59	1.05E-10	0.102	0.102
74.9	2.5053	2.88	3.22	1.50E-10	0.124	0.125
74.9	3.0041	3.56	3.90	1.30E-10	0.148	0.148
74.9	3.5051	4.34	4.56	1.42E-10	0.170	0.170

EXAMPLE 32**Solubility of 1,1,1,2-tetrafluoroethane (HFC-134a) in 1-ethyl-3-methylimidazolium bis(pentafluoroethylsulfonyl)imide [emim][BEI]**

5 A solubility and diffusivity study was made at a temperature of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_0), final saturation concentration (C_s), and calculated solubility
 10 ($X_{\text{calc.}}$) are also provided.

Tables 33a, 33b, 33c, and 33d provide data for C_0 , C_s , D , $X_{\text{calc.}}$, and $X_{\text{meas.}}$ at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

Table 33a

15

T (°C)	P (bar)	C_0 (mass%)	C_s (mass%)	$D_{\text{eff.}}$ (m ² /sec)	$X_{\text{calc.}}$ (mol. fraction)	$X_{\text{meas.}}$ (mol. fraction)
10.0	0.1031	0.09	0.61	1.92E-11	0.029	0.024
10.0	0.5039	1.21	2.51	4.25E-07	0.110	0.120
10.0	1.0027	4.05	6.65	2.95E-11	0.255	0.239
10.0	1.5024	7.74	10.72	3.68E-11	0.366	0.354
10.0	2.0011	12.01	15.61	4.88E-11	0.471	0.464
10.0	2.5009	17.79	21.74	6.58E-11	0.572	0.569
10.0	3.0043	24.67	30.25	8.67E-11	0.676	0.668
10.0	3.5049	37.47	44.30	6.14E-11	0.793	0.793

Table 33b

T (°C)	P (bar)	C_0 (mass%)	C_s (mass%)	$D_{\text{eff.}}$ (m ² /sec)	$X_{\text{calc.}}$ (mol. fraction)	$X_{\text{meas.}}$ (mol. fraction)
24.9	0.1054	0.21	0.42	2.60E-11	0.020	0.019
24.9	0.5052	0.82	1.92	3.76E-11	0.086	0.086
24.9	1.0046	2.55	3.90	4.22E-11	0.163	0.163
24.9	1.5040	4.69	6.02	4.77E-11	0.236	0.235
24.9	2.0037	6.73	8.29	5.70E-11	0.303	0.304
24.9	2.5031	9.15	10.79	6.65E-11	0.368	0.368
24.9	3.0043	11.73	13.53	7.90E-11	0.430	0.429
24.9	3.5054	15.15	16.56	7.29E-11	0.489	0.488

Table 33c

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
50.0	0.1046	0.14	0.23	5.84E-11	0.011	0.011
50.0	0.5050	0.58	1.00	6.72E-11	0.046	0.046
50.0	1.0043	1.42	1.99	8.15E-11	0.089	0.089
50.0	1.5046	2.48	3.00	7.67E-11	0.130	0.130
50.0	2.0037	3.46	4.04	8.44E-11	0.168	0.168
50.0	2.5033	4.51	5.10	8.82E-11	0.205	0.205
50.0	3.0034	5.57	6.19	9.36E-11	0.241	0.241
50.0	3.5040	6.98	7.32	8.24E-11	0.275	0.276

5 Table 33d

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
74.9	0.1044	0.10	0.13	1.30E-10	0.006	0.006
74.9	0.5057	0.37	0.58	1.36E-10	0.027	0.027
74.9	1.0042	0.87	1.16	1.35E-10	0.053	0.053
74.9	1.5043	1.48	1.73	1.32E-10	0.078	0.078
74.9	2.0041	2.01	2.30	1.49E-10	0.102	0.102
74.9	2.4957	2.60	2.88	1.42E-10	0.125	0.125
74.9	3.0049	3.22	3.47	1.69E-10	0.148	0.147
74.9	3.5027	3.89	4.06	1.17E-10	0.169	0.169

EXAMPLE 33**10 Solubility of 1,1,1,2-tetrafluoroethane (HFC-134a) in 1-butyl-3-methylimidazolium 1,1,2,3,3-hexafluoropropanesulfonate [bmim][HFPS]**

A solubility and diffusivity study was made at a temperature of 10, 25, 50, and 75 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C₀), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided.

Tables 34a, 34b, 34c, and 34d provide data for C_o , C_s , D , X_{calc} , and X_{meas} at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

5 **Table 34a**

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
10.0	0.0993	0.00	0.41	1.09E-11	0.015	0.015
9.9	0.5012	0.62	2.43	8.91E-12	0.083	0.082
10.0	1.0001	2.78	5.36	1.13E-11	0.170	0.172
10.0	1.4989	5.94	8.89	1.38E-11	0.261	0.264
9.9	1.9997	9.63	12.82	2.42E-11	0.348	0.350
10.0	2.4950	13.70	18.23	2.42E-11	0.447	0.447
10.0	3.0010	19.60	24.78	4.81E-11	0.545	0.550
10.1	3.4937	27.72	36.37	7.13E-11	0.675	0.677

Table 34b

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
24.9	0.1007	-0.02	0.26	1.61E-11	0.009	0.011
24.9	0.5000	0.50	1.75	2.46E-11	0.061	0.055
24.9	1.0002	1.80	3.22	1.51E-10	0.108	0.109
24.9	1.4995	3.60	5.07	1.50E-11	0.162	0.163
24.9	1.9931	5.36	7.12	1.78E-11	0.218	0.220
25.0	2.5041	7.52	9.10	2.66E-11	0.267	0.269
24.9	3.0042	9.65	11.44	2.46E-11	0.319	0.322
24.9	3.5020	12.23	13.92	3.10E-11	0.370	0.374

Table 34c

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
50.0	0.1007	0.01	0.16	3.94E-11	0.006	0.006
50.0	0.5006	0.28	0.81	3.51E-11	0.029	0.029
50.0	0.9997	1.11	1.69	2.84E-11	0.059	0.059
50.0	1.4987	1.93	2.58	3.30E-11	0.088	0.088
50.0	1.9941	2.87	3.53	2.73E-11	0.117	0.118
50.0	2.5040	3.73	4.42	4.20E-11	0.144	0.145
50.0	2.9997	4.65	5.37	4.79E-11	0.171	0.172
50.0	3.5040	5.64	6.32	4.79E-11	0.197	0.198

5 Table 34d

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
74.9	0.0989	0.04	0.10	5.08E-11	0.003	0.004
74.9	0.5015	0.21	0.46	2.62E-10	0.016	0.018
74.9	1.0009	0.69	1.01	6.65E-11	0.036	0.036
74.9	1.5002	1.17	1.51	7.55E-11	0.053	0.053
74.9	2.0006	1.67	2.03	6.73E-11	0.070	0.070
74.9	2.4996	2.18	2.53	8.11E-11	0.086	0.087
74.9	3.0020	2.70	3.06	8.14E-11	0.103	0.104

EXAMPLE 34

- 10 **Solubility of 1,1,1,2-tetrafluoroethane (HFC-134a) in tetradecyl(trihexyl) phosphonium 1,1,2-trifluoro-2-(perfluoroethoxy) ethanesulfonate [6,6,6,14-P][TPES]**

A solubility and diffusivity study was made at a temperature of 10, 25, 50,
 15 and 75 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.}) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility (X_{calc.}) are also provided.

Tables 35a, 35b, 35c, and 35d provide data for C_o , C_s , D , X_{calc} , and X_{meas} at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

Table 35a

5

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	$D_{eff.}$ (m ² /sec)	$X_{calc.}$ (mol. fraction)	$X_{meas.}$ (mol. Fraction)
10.0	0.0993	0.10	0.52	1.65E-11	0.038	0.038
9.7	0.5001	0.87	2.99	2.04E-11	0.190	0.190
9.9	1.0005	3.55	6.26	2.72E-11	0.338	0.338
9.8	1.4988	7.01	9.95	3.28E-11	0.458	0.452
10.1	1.9940	10.46	13.72	5.63E-11	0.549	0.551
9.8	2.4956	14.69	18.30	1.01E-10	0.631	0.634
9.7	2.9998	19.78	24.52	1.23E-10	0.713	0.718
9.6	3.4947	26.93	34.29	2.24E-10	0.800	0.799

Table 35b

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	$D_{eff.}$ (m ² /sec)	$X_{calc.}$ (mol. fraction)	$X_{meas.}$ (mol. Fraction)
24.9	0.1000	-0.01	0.26	2.82E-11	0.019	0.018
24.9	0.5002	0.50	1.75	4.18E-11	0.120	0.121
25.0	0.9998	2.14	3.73	4.58E-11	0.229	0.228
24.9	1.4991	4.13	5.79	5.46E-11	0.320	0.320
24.9	2.0001	6.22	7.90	6.55E-11	0.396	0.397
24.9	2.5034	8.35	10.05	8.92E-11	0.461	0.462
24.9	3.0041	10.54	12.31	9.57E-11	0.518	0.520
24.9	3.5040	12.92	14.84	1.11E-10	0.571	0.574

10 **Table 35c**

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	$D_{eff.}$ (m ² /sec)	$X_{calc.}$ (mol. fraction)	$X_{meas.}$ (mol. fraction)
50.0	0.1013	0.21	0.09	1.08E-11	0.007	0.011
50.0	0.5011	0.34	0.94	9.52E-11	0.068	0.068
50.0	1.0012	1.24	1.97	9.91E-11	0.133	0.134
50.0	1.4996	2.29	3.01	1.07E-10	0.192	0.193
50.0	2.0006	3.37	4.07	9.79E-11	0.245	0.246
50.0	2.5005	4.37	5.10	1.22E-10	0.291	0.294
50.0	2.9997	5.44	6.19	1.19E-10	0.335	0.339
50.1	3.4970	6.68	7.33	1.14E-10	0.377	0.381

Table 35d

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
74.9	0.1011	0.00	0.03	1.84E-10	0.002	0.003
74.9	0.5019	0.22	0.52	1.81E-10	0.039	0.039
74.9	1.0009	0.77	1.16	1.97E-10	0.082	0.083
74.9	1.4959	1.41	1.77	2.08E-10	0.121	0.122
74.9	2.0012	2.03	2.40	2.27E-10	0.158	0.160
74.9	2.5033	2.65	3.03	2.28E-10	0.193	0.194
74.9	3.0034	3.30	3.65	2.05E-10	0.225	0.227
74.9	3.5051	3.96	4.27	2.13E-10	0.254	0.256

5 EXAMPLE 35

**Solubility of 1,1,1,2-tetrafluoroethane (HFC-134a) in
tributyl(tetradecyl)phosphonium 1,1,2,3,3,3-hexafluoropropanesulfonate
[4,4,4,14-P][HFPS]**

10

A solubility and diffusivity study was made at a temperature of 10, 25, 50,
and 75 °C over a pressure range from 0 to 10 bar where the solubilities (X_{meas.})
were measured using a gravimetric microbalance and the diffusivities (D) were
calculated using a one-dimensional diffusion model analysis. The initial
concentration (C₀), final saturation concentration (C_s), and calculated solubility
(X_{calc.}) are also provided.

15

Tables 36a, 36b, 36c, and 36d provide data for C₀, C_s, D, X_{calc.}, and X_{meas.}
at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

Table 36a

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. Fraction)
10.2	0.0991	0.08	0.49	2.23E-11	0.029	0.032
9.9	0.5001	0.72	2.95	1.30E-11	0.158	0.152
10.2	0.9998	3.17	6.30	1.74E-11	0.293	0.289
10.0	1.4999	6.59	9.78	2.67E-11	0.401	0.403
10.0	1.9996	10.48	13.80	4.77E-11	0.497	0.494
10.0	2.5034	14.41	18.75	5.41E-11	0.587	0.587
10.0	3.0020	19.66	24.79	1.49E-10	0.670	0.672
10.1	3.4928	27.70	34.01	2.02E-10	0.761	0.763

5 Table 36b

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. Fraction)
25.0	0.0998	0.05	0.34	1.70E-11	0.021	0.019
24.9	0.5001	0.50	1.83	2.56E-11	0.103	0.104
24.9	0.9994	2.11	3.76	3.19E-11	0.194	0.194
25.0	1.4988	4.06	5.79	3.71E-11	0.275	0.273
24.9	2.0017	6.03	8.06	3.60E-11	0.351	0.350
25.0	2.5003	8.43	10.48	4.88E-11	0.419	0.418
25.0	2.9990	10.82	12.84	7.38E-11	0.476	0.478
25.0	3.5021	13.55	15.47	1.01E-10	0.530	0.530

Table 36c

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m ² /sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. fraction)
50.0	0.1009	0.00	0.17	6.85E-11	0.010	0.010
50.0	0.5001	0.32	0.96	6.65E-11	0.056	0.056
50.0	0.9994	1.20	1.99	6.73E-11	0.111	0.110
50.0	1.4992	2.24	3.04	6.51E-11	0.162	0.161
50.0	2.0003	3.31	4.09	7.46E-11	0.208	0.209
50.0	2.4945	4.29	5.16	8.18E-11	0.251	0.254
50.0	2.9994	5.46	6.22	1.11E-10	0.290	0.293
50.0	3.4964	7.54	8.32	7.36E-11	0.359	0.333

10

Table 36d

T (°C)	P (bar)	C _o (mass%)	C _s (mass%)	D _{eff.} (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
75.0	0.1006	0.08	0.14	1.36E-10	0.009	0.009
74.9	0.5041	0.30	0.63	1.39E-10	0.037	0.037
74.9	1.0014	0.83	1.25	1.37E-10	0.072	0.072
74.9	1.5002	1.47	1.87	1.43E-10	0.105	0.105
74.9	2.0014	2.07	2.47	1.63E-10	0.135	0.136
74.9	2.5044	2.66	3.08	1.70E-10	0.164	0.165
74.9	3.0037	2.75	3.15	1.51E-10	0.167	0.194
74.9	3.5039	3.44	3.79	1.70E-10	0.196	0.221

5

EXAMPLE 36

The description of the microbalance components, shown in Figure 18, are provided below.

10

Table 37. Microbalance Components Contributing to Buoyancy Calculation

Subscript	Item	Weight (g)	Material	Density (g•cm ⁻³)	Temperature (°C)
s	Dry sample	m _s	[bmim][PF ₆] [bmim][BF ₄]	ρ _s	Sample Temp. (T _s)
a	Interacted gas	m _a	CO ₂	ρ _a	(T _s)
i ₁	Sample container	0.5986	Pyrex	2.23	(T _s)
i ₂	Wire	0.051	Tungsten	21.0	(T _s)
i ₃	Chain	0.3205	Gold	19.3	30
j ₁	Counter-weight	0.8054	Stainless Steel	7.9	25
j ₂	Hook	0.00582	Tungsten	21.0	25
j ₃	Chain	0.2407	Gold	19.3	30

15

EXAMPLE 37**Solubility of carbon dioxide (CO₂) in 1-butyl-3-methylimidazolium hexafluorophosphate [bmim][PF₆]**

5

A solubility and diffusivity study was made at a temperature of 10, 25, 50, and 75 °C over a pressure range from 0 to 20 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial concentration (C_o), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided.

Tables 38a, 38b, 38c, and 38d provide data for C_o , C_s , D , $X_{\text{calc.}}$, and $X_{\text{meas.}}$ at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

15 **Table 38a**

T (°C)	P (bar)	C_o (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. Fraction)
9.9	0.097	0.00	0.07	1.5E-11	0.006	0.004
9.9	0.501	0.10	0.24	2.0E-11	0.017	0.016
9.9	1.002	0.29	0.46	2.2E-11	0.030	0.029
10.4	3.996	0.85	1.72	2.6E-11	0.101	0.102
10.6	6.996	2.15	2.97	3.0E-11	0.165	0.165
10.5	10.000	3.43	4.22	3.6E-11	0.221	0.220
8.9	13.003	4.95	5.69	-	0.280	0.278
9.9	14.998	5.84	6.35	7.4E-11	0.306	0.302
9.9	19.998	7.51	8.35	5.2E-11	0.375	0.367

Table 38b

T (°C)	P (bar)	C _o (mass%)	C _s (mass%)	D _{eff.} (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. Fraction)
24.9	0.102	0.00	0.01	-	0.006	0.000
24.9	0.502	0.03	0.14	5.1E-11	0.011	0.009
24.9	1.002	0.16	0.29	4.8E-11	0.020	0.018
25.0	3.996	0.65	1.19	4.5E-11	0.072	0.071
25.0	7.000	1.55	2.07	5.4E-11	0.122	0.120
24.9	9.994	2.45	2.95	5.7E-11	0.167	0.164
24.9	12.999	3.31	3.80	6.8E-11	0.208	0.203
24.9	14.994	4.05	4.41	7.7E-11	0.238	0.227
24.9	19.992	5.19	5.80	8.1E-11	0.285	0.282

Table 38c

5

T (°C)	P (bar)	C _o (mass%)	C _s (mass%)	D _{eff.} (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
50.1	0.102	0.01	0.03	11.3E-11	0.002	0.002
50.0	0.503	0.04	0.09	10.8E-11	0.006	0.006
50.0	1.002	0.12	0.19	9.7E-11	0.012	0.012
50.1	3.996	0.50	0.76	10.1E-11	0.047	0.047
50.0	7.000	1.07	1.29	10.1E-11	0.078	0.078
50.0	9.998	1.59	1.81	12.6E-11	0.107	0.107
50.0	13.002	2.13	2.34	12.7E-11	0.134	0.133
50.1	15.003	2.53	2.70	13.5E-11	0.152	0.151
50.0	19.998	3.27	3.56	14.8E-11	0.192	0.191

Table 38d

T (°C)	P (bar)	C _o (mass%)	C _s (mass%)	D _{eff.} (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
75.0	0.102	0.01	0.02	9.5E-11	0.001	0.001
74.9	0.501	0.02	0.04	23.5E-11	0.002	0.002
74.9	1.000	0.07	0.11	21.3E-11	0.007	0.007
74.9	3.997	0.40	0.51	27.5E-11	0.032	0.032
74.9	7.000	0.78	0.89	23.7E-11	0.055	0.055
74.8	10.002	1.18	1.28	13.2E-11	0.077	0.077
75.0	13.003	1.50	1.64	69.5E-11	0.097	0.096
74.9	14.999	1.75	1.88	58.5E-11	0.110	0.110
75.1	19.995	2.36	2.45	28.4E-11	0.140	0.144

10

EXAMPLE 38**Solubility of carbon dioxide (CO₂) in 1-butyl-3-methylimidazolium tetrafluoroborate [bmim][BF₄]**

5

A solubility and diffusivity study was made at a temperature of 10, 25, 50, and 75 °C over a pressure range from 0 to 20 bar where the solubilities ($X_{\text{meas.}}$) were measured using a gravimetric microbalance and the diffusivities (D) were calculated using a one-dimensional diffusion model analysis. The initial
 10 concentration (C_0), final saturation concentration (C_s), and calculated solubility ($X_{\text{calc.}}$) are also provided.

Tables 39a, 39b, 39c, and 39d provide data for C_0 , C_s , D , $X_{\text{calc.}}$, and X_{meas} at a temperature of 10°C, 25 °C, 50°C, and 75°C, respectively.

15 **Table 39a**

T (°C)	P (bar)	C₀ (mass%)	C_s (mass%)	D_{eff.} (m²/sec)	X_{calc.} (mol. fraction)	X_{meas.} (mol. Fraction)
9.6	0.102	0.00	0.04	4.0E-11	0.002	0.002
9.8	0.502	0.07	0.24	3.4E-11	0.013	0.013
9.9	1.001	0.26	0.46	4.8E-11	0.023	0.023
9.9	4.001	0.97	1.88	3.8E-11	0.090	0.090
10.1	6.996	2.40	3.25	4.5E-11	0.147	0.147
9.9	9.997	3.69	4.64	5.5E-11	0.202	0.202
9.9	13.002	5.12	5.93	6.9E-11	0.249	0.259
10.0	15.001	6.28	6.79	6.7E-11	0.278	0.288
9.9	20.002	7.89	8.97	8.0E-11	0.346	0.353

Table 39b

T (°C)	P (bar)	C ₀ (mass%)	C _s (mass%)	D _{eff.} (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. Fraction)
24.8	0.097	-	-	-	-	0.002
25.0	0.500	0.08	0.19	6.6E-11	0.011	0.010
24.9	1.001	0.23	0.36	7.3E-11	0.020	0.018
25.0	3.996	0.81	1.42	6.4E-11	0.069	0.068
25.0	7.002	1.87	2.45	7.1E-11	0.116	0.114
24.8	9.997	2.88	3.46	7.7E-11	0.158	0.156
25.0	13.002	3.88	4.44	9.3E-11	0.197	0.192
24.8	15.001	4.71	5.09	8.2E-11	0.222	0.216
24.9	20.002	6.03	6.66	10.0E-11	0.277	0.268

Table 39c

5

T (°C)	P (bar)	C ₀ (mass%)	C _s (mass%)	D _{eff.} (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
50.1	0.102	0.02	0.06	20.7E-11	0.003	0.003
50.0	0.501	0.08	0.13	24.8E-11	0.006	0.006
50.0	1.001	0.18	0.23	12.2E-11	0.012	0.012
50.0	3.997	0.62	0.88	15.5E-11	0.043	0.043
50.0	6.996	1.23	1.49	17.4E-11	0.072	0.072
50.0	10.002	1.88	2.09	14.2E-11	0.099	0.099
50.0	12.997	2.48	2.69	12.9E-11	0.124	0.124
50.0	15.001	2.94	3.07	17.4E-11	0.140	0.140
50.0	20.000	3.82	4.02	14.5E-11	0.177	0.176

Table 39d

T (°C)	P (bar)	C ₀ (mass%)	C _s (mass%)	D _{eff.} (m ² /sec)	X _{calc.} (mol. fraction)	X _{meas.} (mol. fraction)
75.0	0.102	0.02	0.04	32.8E-11	0.002	0.002
75.0	0.501	0.04	0.07	34.2E-11	0.003	0.003
74.9	1.002	0.10	0.15	32.9E-11	0.008	0.007
74.9	4.002	0.48	0.60	29.7E-11	0.030	0.030
75.0	6.996	0.91	1.01	24.0E-11	0.050	0.050
74.8	10.003	1.30	1.43	32.5E-11	0.069	0.070
75.0	13.000	1.72	1.83	23.6E-11	0.087	0.087
75.0	15.002	2.01	2.11	29.2E-11	0.100	0.101
74.9	19.999	2.68	2.77	46.2E-11	0.127	0.129

10

Where the indefinite article “a” or “an” is used with respect to a statement or description of the presence of a feature, component or step in a composition, apparatus or process of this invention, it is to be understood, unless the statement or description explicitly provides to the contrary, that the use of such indefinite article does not limit the presence of the feature, component or step in the composition, apparatus or process to one in number.

Where a apparatus or process of this invention is stated or described as comprising, including, containing, having, being composed of or being constituted by certain features, components or steps, it is to be understood, unless the statement or description explicitly provides to the contrary, that one or more features, components or steps in addition to those explicitly stated or described may be present in the apparatus or process. In an alternative embodiment, however, the apparatus or process of this invention may be stated or described as consisting essentially of certain features, components or steps, in which embodiment features, components or steps that would materially alter the principle of operation or the distinguishing characteristics of the composition, apparatus or process are not present therein. In a further alternative embodiment, the apparatus or process of this invention may be stated or described as consisting of certain features, components or steps, in which embodiment features, components or steps other than as named are not present therein.

25

CLAIMS

What is claimed is:

- 5 1. An apparatus for temperature adjustment comprising
 - (a) a compressor that increases the pressure of the vapor of at least one refrigerant, wherein the compressor comprises moving parts that are lubricated by at least one ionic liquid;
 - (b) a condenser that receives refrigerant vapor that is passed out of the compressor, and condenses the vapor under pressure to a liquid;
 - 10 (c) a pressure reduction device that receives liquid refrigerant that is passed out of the condenser, and reduces the pressure of the liquid to form a mixture of refrigerant in liquid and vapor form;
 - (d) an evaporator that receives the mixture of liquid and vapor refrigerant
 - 15 that is passed out of the pressure reduction device, and evaporates the remaining liquid in the mixture to form refrigerant vapor; and
 - (e) a conduit that returns to the compressor refrigerant vapor that is passed out of the evaporator.
- 20 2. An apparatus according to Claim 1 wherein the condenser is located in proximity to an object, medium or space to be heated.
3. An apparatus according to Claim 1 wherein the evaporator is located in proximity to an object, medium or space to be cooled.
- 25 4. An apparatus according to Claim 1 wherein a refrigerant is selected from the group consisting of CHClF_2 (R-22); CHF_3 (R-23); CH_2F_2 (R-32); CH_3F (R-41); CHF_2CF_3 (R-125); CH_2FCF_3 (R-134a); $\text{CHF}_2\text{OCHF}_2$ (E-134); CH_3CClF_2 (R-142b); CH_3CF_3 (R-143a); CH_3CHF_2 (R-152a); $\text{CH}_3\text{CH}_2\text{F}$ (R-161); CH_3OCH_3 (E170); $\text{CF}_3\text{CF}_2\text{CF}_3$ (R-218); $\text{CF}_3\text{CHFCF}_3$ (R-227ea); $\text{CF}_3\text{CH}_2\text{CF}_3$ (R-236fa); $\text{CH}_2\text{FCF}_2\text{CHF}_2$ (R-245ca); $\text{CHF}_2\text{CH}_2\text{CF}_3$ (R-245fa); $\text{CH}_3\text{CH}_2\text{CH}_3$ (R-290); $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_3$ (R-600); $\text{CH}(\text{CH}_3)_2\text{CH}_3$ (R-600a); $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3$ (R-601); $(\text{CH}_3)_2\text{CHCH}_2\text{CH}_3$ (R-601a); $\text{CH}_3\text{CH}_2\text{OCH}_2\text{CH}_3$ (R-610); NH_3 ; CO_2 ; and
- 30

CH₃CH=CH₂; and combinations thereof; or a refrigerant blend selected from the group consisting R-404A; R-407A; R-407B; R-407C; R-407D; R-407E; R-410A; R-410B; R-413A; 417A; R-419A; R-420A; 80.6 % R-134a and 19.4% R-142b (by weight); R-421A; R-421B; R-422A; R-422B; R-422C; R-422D; R-423A; R-424A; R-425A; R-426A; R-427A; 2.0% R-32, 41.0% R-125, 50.0% R-143a and 7.0% R-134a (by weight); 10.0% R-32, 33.0% R-125, 36.0% R-143a and 21.0% R-134a (by weight); R-428A; and R-507A.

5. An apparatus according to Claim 2, wherein a refrigerant is selected from the group consisting of R-22, R-32, R-125, R-134a, R-404A, R-410A, R-413A, R-422A, R-422D, R-423A, R-426A, R-427A, R-507A, and combinations thereof.

6. An apparatus according to Claim 1 wherein a refrigerant comprises at least one hydrofluorocarbon selected from the group consisting of trifluoromethane (HFC-23), difluoromethane (HFC-32), pentafluoroethane (HFC-125), 1,1,2,2-tetrafluoroethane (HFC-134), 1,1,1,2-tetrafluoroethane (HFC-134a), 1,1,1-trifluoroethane (HFC-143a), 1,1-difluoroethane (HFC-152a), fluoroethane (HFC-161), R-404A, R-407C, R-410A, and combinations thereof.

7. An apparatus according to Claim 4 wherein a refrigerant comprises at least one hydrofluorocarbon selected from the group consisting of pentafluoroethane (HFC-125), 1,1,2,2-tetrafluoroethane (HFC-134), 1,1,1,2-tetrafluoroethane (HFC-134a), R-404aA, R-407C, R-410A, and combinations thereof.

8. An apparatus according to Claim 1 wherein a refrigerant comprises at least one fluoroolefin selected from the group consisting of:

- (i) fluoroolefins of the formula *E*- or *Z*-R¹CH=CHR², wherein R¹ and R² are, independently, C₁ to C₆ perfluoroalkyl groups, and wherein the total number of carbons in the compound is at least 5;

(ii) cyclic fluoroolefins of the formula cyclo-[CX=CY(CZW)_n], wherein X, Y, Z, and W, independently, are H or F, and n is an integer from 2 to 5; and

(iii) fluoroolefins selected from the group consisting of:

- 5 2,3,3-trifluoro-1-propene (CHF₂CF=CH₂); 1,1,2-trifluoro-1-propene (CH₃CF=CF₂); 1,2,3-trifluoro-1-propene (CH₂FCF=CF₂); 1,1,3-trifluoro-1-propene (CH₂FCH=CF₂); 1,3,3-trifluoro-1-propene (CHF₂CH=CHF); 1,1,1,2,3,4,4,4-octafluoro-2-butene (CF₃CF=CFCF₃); 1,1,2,3,3,4,4,4-octafluoro-1-butene (CF₃CF₂CF=CF₂); 1,1,1,2,4,4,4-heptafluoro-2-butene (CF₃CF=CHCF₃);
- 10 1,2,3,3,4,4,4-heptafluoro-1-butene (CHF=CFCF₂CF₃); 1,1,1,2,3,4,4-heptafluoro-2-butene (CHF₂CF=CFCF₃); 1,3,3,3-tetrafluoro-2-(trifluoromethyl)-1-propene ((CF₃)₂C=CHF); 1,1,3,3,4,4,4-heptafluoro-1-butene (CF₂=CHCF₂CF₃); 1,1,2,3,4,4,4-heptafluoro-1-butene (CF₂=CFCHF₂CF₃); 1,1,2,3,3,4,4-heptafluoro-1-butene (CF₂=CFCF₂CHF₂); 2,3,3,4,4,4-hexafluoro-1-butene (CF₃CF₂CF=CH₂);
- 15 1,3,3,4,4,4-hexafluoro-1-butene (CHF=CHCF₂CF₃); 1,2,3,4,4,4-hexafluoro-1-butene (CHF=CFCHF₂CF₃); 1,2,3,3,4,4-hexafluoro-1-butene (CHF=CFCF₂CHF₂); 1,1,2,3,4,4-hexafluoro-2-butene (CHF₂CF=CFCHF₂); 1,1,1,2,3,4-hexafluoro-2-butene (CH₂FCF=CFCF₃); 1,1,1,2,4,4-hexafluoro-2-butene (CHF₂CH=CFCF₃); 1,1,1,3,4,4-hexafluoro-2-butene (CF₃CH=CFCHF₂); 1,1,2,3,3,4-hexafluoro-1-butene (CF₂=CFCF₂CH₂F); 1,1,2,3,4,4-hexafluoro-1-butene (CF₂=CFCHF₂CHF₂);
- 20 3,3,3-trifluoro-2-(trifluoromethyl)-1-propene (CH₂=C(CF₃)₂); 1,1,1,2,4-pentafluoro-2-butene (CH₂FCH=CFCF₃); 1,1,1,3,4-pentafluoro-2-butene (CF₃CH=CFCH₂F); 3,3,4,4,4-pentafluoro-1-butene (CF₃CF₂CH=CH₂); 1,1,1,4,4-pentafluoro-2-butene (CHF₂CH=CHCF₃); 1,1,1,2,3-pentafluoro-2-butene (CH₃CF=CFCF₃); 2,3,3,4,4-pentafluoro-1-butene (CH₂=CFCF₂CHF₂); 1,1,2,4,4-pentafluoro-2-butene (CHF₂CF=CHCHF₂); 1,1,2,3,3-pentafluoro-1-butene (CH₃CF₂CF=CF₂); 1,1,2,3,4-pentafluoro-2-butene (CH₂FCF=CFCHF₂); 1,1,3,3,3-pentafluoro-2-methyl-1-propene (CF₂=C(CF₃)(CH₃)); 2-(difluoromethyl)-3,3,3-trifluoro-1-propene (CH₂=C(CHF₂)(CF₃)); 2,3,4,4,4-pentafluoro-1-butene (CH₂=CFCHF₂CF₃); 1,2,4,4,4-pentafluoro-1-butene (CHF=CFCH₂CF₃); 1,3,4,4,4-pentafluoro-1-butene (CHF=CHCHF₂CF₃); 1,3,3,4,4-pentafluoro-1-butene (CHF=CHCF₂CHF₂); 1,2,3,4,4-pentafluoro-1-butene (CHF=CFCHF₂CHF₂); 3,3,4,4-tetrafluoro-1-butene (CH₂=CHCF₂CHF₂); 1,1-difluoro-2-(difluoromethyl)-1-propene (CF₂=C(CHF₂)(CH₃)); 1,3,3,3-tetrafluoro-2-methyl-1-propene

- (CHF=C(CF₃)(CH₃)); 3,3-difluoro-2-(difluoromethyl)-1-propene
 (CH₂=C(CHF₂)₂); 1,1,1,2-tetrafluoro-2-butene (CF₃CF=CHCH₃); 1,1,1,3-
 tetrafluoro-2-butene (CH₃CF=CHCF₃); 1,1,1,2,3,4,4,5,5,5-decafluoro-2-pentene
 (CF₃CF=CFCF₂CF₃); 1,1,2,3,3,4,4,5,5,5-decafluoro-1-pentene
 5 (CF₂=CFCF₂CF₂CF₃); 1,1,1,4,4,4-hexafluoro-2-(trifluoromethyl)-2-butene
 ((CF₃)₂C=CHCF₃); 1,1,1,2,4,4,5,5,5-nonafluoro-2-pentene (CF₃CF=CHCF₂CF₃);
 1,1,1,3,4,4,5,5,5-nonafluoro-2-pentene (CF₃CH=CFCF₂CF₃); 1,2,3,3,4,4,5,5,5-
 nonafluoro-1-pentene (CHF=CFCF₂CF₂CF₃); 1,1,3,3,4,4,5,5,5-nonafluoro-1-
 pentene (CF₂=CHCF₂CF₂CF₃); 1,1,2,3,3,4,4,5,5-nonafluoro-1-pentene
 10 (CF₂=CFCF₂CF₂CHF₂); 1,1,2,3,4,4,5,5,5-nonafluoro-2-pentene
 (CHF₂CF=CFCF₂CF₃); 1,1,1,2,3,4,4,5,5-nonafluoro-2-pentene
 (CF₃CF=CFCF₂CHF₂); 1,1,1,2,3,4,5,5,5-nonafluoro-2-pentene
 (CF₃CF=CFCHFCF₃); 1,2,3,4,4,4-hexafluoro-3-(trifluoromethyl)-1-butene
 (CHF=CFCF(CF₃)₂); 1,1,2,4,4,4-hexafluoro-3-(trifluoromethyl)-1-butene
 15 (CF₂=CFCH(CF₃)₂); 1,1,1,4,4,4-hexafluoro-2-(trifluoromethyl)-2-butene
 (CF₃CH=C(CF₃)₂); 1,1,3,4,4,4-hexafluoro-3-(trifluoromethyl)-1-butene
 (CF₂=CHCF(CF₃)₂); 2,3,3,4,4,5,5,5-octafluoro-1-pentene (CH₂=CFCF₂CF₂CF₃);
 1,2,3,3,4,4,5,5-octafluoro-1-pentene (CHF=CFCF₂CF₂CHF₂); 3,3,4,4,4-
 pentafluoro-2-(trifluoromethyl)-1-butene (CH₂=C(CF₃)CF₂CF₃); 1,1,4,4,4-
 20 pentafluoro-3-(trifluoromethyl)-1-butene (CF₂=CHCH(CF₃)₂); 1,3,4,4,4-
 pentafluoro-3-(trifluoromethyl)-1-butene (CHF=CHCF(CF₃)₂); 1,1,4,4,4-
 pentafluoro-2-(trifluoromethyl)-1-butene (CF₂=C(CF₃)CH₂CF₃); 3,4,4,4-
 tetrafluoro-3-(trifluoromethyl)-1-butene ((CF₃)₂CFCH=CH₂); 3,3,4,4,5,5,5-
 heptafluoro-1-pentene (CF₃CF₂CF₂CH=CH₂); 2,3,3,4,4,5,5-heptafluoro-1-pentene
 25 (CH₂=CFCF₂CF₂CHF₂); 1,1,3,3,5,5,5-heptafluoro-1-butene
 (CF₂=CHCF₂CH₂CF₃); 1,1,1,2,4,4,4-heptafluoro-3-methyl-2-butene
 (CF₃CF=C(CF₃)(CH₃)); 2,4,4,4-tetrafluoro-3-(trifluoromethyl)-1-butene
 (CH₂=CFCH(CF₃)₂); 1,4,4,4-tetrafluoro-3-(trifluoromethyl)-1-butene
 (CHF=CHCH(CF₃)₂); 1,1,1,4-tetrafluoro-2-(trifluoromethyl)-2-butene
 30 (CH₂FCH=C(CF₃)₂); 1,1,1,3-tetrafluoro-2-(trifluoromethyl)-2-butene
 (CH₃CF=C(CF₃)₂); 1,1,1-trifluoro-2-(trifluoromethyl)-2-butene
 ((CF₃)₂C=CHCH₃); 3,4,4,5,5,5-hexafluoro-2-pentene (CF₃CF₂CF=CHCH₃);
 1,1,1,4,4,4-hexafluoro-2-methyl-2-butene (CF₃C(CH₃)=CHCF₃); 3,3,4,5,5,5-
 hexafluoro-1-pentene (CH₂=CHCF₂CHFCF₃); 4,4,4-trifluoro-3-(trifluoromethyl)-

- 1-butene ($\text{CH}_2=\text{C}(\text{CF}_3)\text{CH}_2\text{CF}_3$); 1,1,2,3,3,4,4,5,5,6,6,6-dodecafluoro-1-hexene
 $(\text{CF}_3(\text{CF}_2)_3\text{CF}=\text{CF}_2)$; 1,1,1,2,2,3,4,5,5,6,6,6-dodecafluoro-3-hexene
 $(\text{CF}_3\text{CF}_2\text{CF}=\text{CFCF}_2\text{CF}_3)$; 1,1,1,4,4,4-hexafluoro-2,3-bis(trifluoromethyl)-2-butene
 $((\text{CF}_3)_2\text{C}=\text{C}(\text{CF}_3)_2)$; 1,1,1,2,3,4,5,5,5-nonafluoro-4-(trifluoromethyl)-2-pentene
5 $((\text{CF}_3)_2\text{CFCF}=\text{CFCF}_3)$; 1,1,1,4,4,5,5,5-octafluoro-2-(trifluoromethyl)-2-pentene
 $((\text{CF}_3)_2\text{C}=\text{CHC}_2\text{F}_5)$; 1,1,1,3,4,5,5,5-octafluoro-4-(trifluoromethyl)-2-pentene
 $((\text{CF}_3)_2\text{CFCF}=\text{CHCF}_3)$; 3,3,4,4,5,5,6,6,6-nonafluoro-1-hexene
 $(\text{CF}_3\text{CF}_2\text{CF}_2\text{CF}_2\text{CH}=\text{CH}_2)$; 4,4,4-trifluoro-3,3-bis(trifluoromethyl)-1-butene
 $(\text{CH}_2=\text{CHC}(\text{CF}_3)_3)$; 1,1,1,4,4,4-hexafluoro-2-(trifluoromethyl)-3-methyl-2-butene
10 $((\text{CF}_3)_2\text{C}=\text{C}(\text{CH}_3)(\text{CF}_3))$; 2,3,3,5,5,5-hexafluoro-4-(trifluoromethyl)-1-pentene
 $(\text{CH}_2=\text{CFCF}_2\text{CH}(\text{CF}_3)_2)$; 1,1,1,2,4,4,5,5,5-nonafluoro-3-methyl-2-pentene
 $(\text{CF}_3\text{CF}=\text{C}(\text{CH}_3)\text{CF}_2\text{CF}_3)$; 1,1,1,5,5,5-hexafluoro-4-(trifluoromethyl)-2-pentene
 $(\text{CF}_3\text{CH}=\text{CHCH}(\text{CF}_3)_2)$; 3,4,4,5,5,6,6,6-octafluoro-2-hexene
 $(\text{CF}_3\text{CF}_2\text{CF}_2\text{CF}=\text{CHCH}_3)$; 3,3,4,4,5,5,6,6-octafluoro-1-hexene
15 $(\text{CH}_2=\text{CHCF}_2\text{CF}_2\text{CF}_2\text{CHF}_2)$; 1,1,1,4,4-pentafluoro-2-(trifluoromethyl)-2-pentene
 $((\text{CF}_3)_2\text{C}=\text{CHCF}_2\text{CH}_3)$; 4,4,5,5,5-pentafluoro-2-(trifluoromethyl)-1-pentene
 $(\text{CH}_2=\text{C}(\text{CF}_3)\text{CH}_2\text{C}_2\text{F}_5)$; 3,3,4,4,5,5,5-heptafluoro-2-methyl-1-pentene
 $(\text{CF}_3\text{CF}_2\text{CF}_2\text{C}(\text{CH}_3)=\text{CH}_2)$; 4,4,5,5,6,6,6-heptafluoro-2-hexene
 $(\text{CF}_3\text{CF}_2\text{CF}_2\text{CH}=\text{CHCH}_3)$; 4,4,5,5,6,6,6-heptafluoro-1-hexene
20 $(\text{CH}_2=\text{CHCH}_2\text{CF}_2\text{C}_2\text{F}_5)$; 1,1,1,2,3,4-heptafluoro-3-hexene
 $(\text{CF}_3\text{CF}_2\text{CF}=\text{CFC}_2\text{H}_5)$; 4,5,5,5-tetrafluoro-4-(trifluoromethyl)-1-pentene
 $(\text{CH}_2=\text{CHCH}_2\text{CF}(\text{CF}_3)_2)$; 1,1,1,2,5,5,5-heptafluoro-4-methyl-2-pentene
 $(\text{CF}_3\text{CF}=\text{CHCH}(\text{CF}_3)(\text{CH}_3))$; 1,1,1,3-tetrafluoro-2-(trifluoromethyl)-2-pentene
 $((\text{CF}_3)_2\text{C}=\text{CFC}_2\text{H}_5)$; 1,1,1,2,3,4,4,5,5,6,6,7,7,7-tetradecafluoro-2-heptene
25 $(\text{CF}_3\text{CF}=\text{CFCF}_2\text{CF}_2\text{C}_2\text{F}_5)$; 1,1,1,2,2,3,4,5,5,6,6,7,7,7-tetradecafluoro-3-heptene
 $(\text{CF}_3\text{CF}_2\text{CF}=\text{CFCF}_2\text{C}_2\text{F}_5)$; 1,1,1,3,4,4,5,5,6,6,7,7,7-tridecafluoro-2-heptene
 $(\text{CF}_3\text{CH}=\text{CFCF}_2\text{CF}_2\text{C}_2\text{F}_5)$; 1,1,1,2,4,4,5,5,6,6,7,7,7-tridecafluoro-2-heptene
 $(\text{CF}_3\text{CF}=\text{CHCF}_2\text{CF}_2\text{C}_2\text{F}_5)$; 1,1,1,2,2,4,5,5,6,6,7,7,7-tridecafluoro-3-heptene
 $(\text{CF}_3\text{CF}_2\text{CH}=\text{CFCF}_2\text{C}_2\text{F}_5)$; 1,1,1,2,2,3,5,5,6,6,7,7,7-tridecafluoro-3-heptene
30 $(\text{CF}_3\text{CF}_2\text{CF}=\text{CHCF}_2\text{C}_2\text{F}_5)$; $\text{CF}_2=\text{CFOCF}_2\text{CF}_3$ (PEVE) and $\text{CF}_2=\text{CFOCF}_3$ (PMVE).

9. An apparatus according to Claim 1 wherein a refrigerant comprises

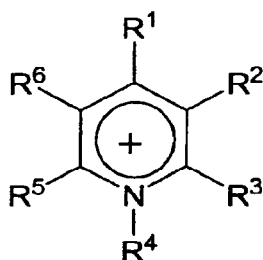
(a) pentafluoroethane (R-125), 1,1,1,2-tetrafluoroethane (R-134a), and at least two hydrocarbons each having eight or fewer carbon atoms, or

(b) pentafluoroethane (R-125), 1,1,1,2-tetrafluoroethane (R-134a), n-butane (R-600) and n-pentane (R-601).

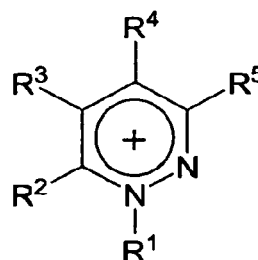
5

10. An apparatus according to Claim 1 wherein an ionic liquid comprises a cation selected from the group consisting of the following eleven cations:

10

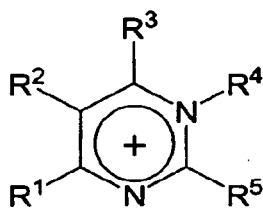


Pyridinium

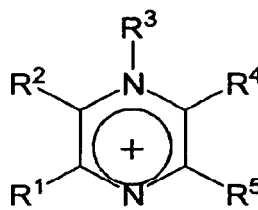


Pyridazinium

15

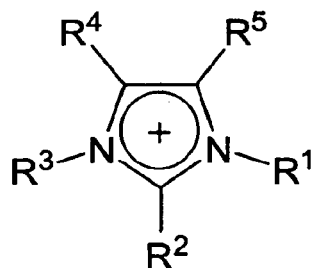


Pyrimidinium

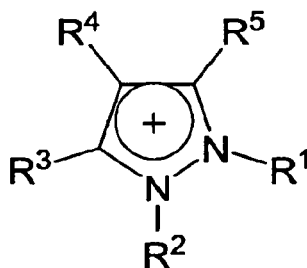


Pyrazinium

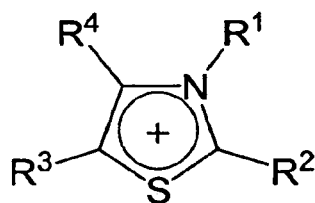
20



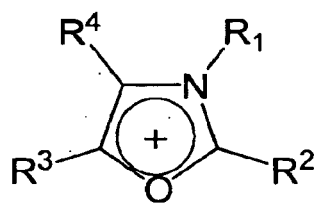
Imidazolium



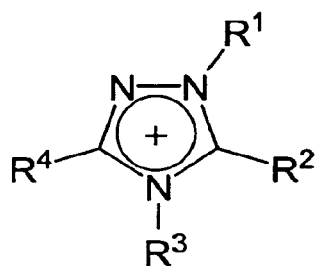
Pyrazolium



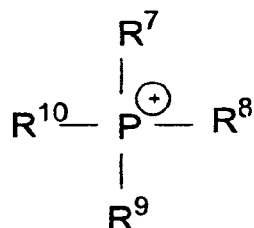
Thiazolium



Oxazolium

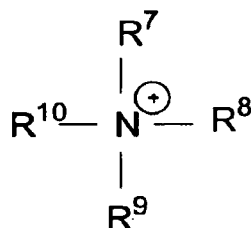


Triazolium



Phosphonium

and



Ammonium

wherein R^1 , R^2 , R^3 , R^4 , R^5 and R^6 are independently selected from the group consisting of:

- (i) H
- (ii) halogen
- (iii) $-CH_3$, $-C_2H_5$, or C_3 to C_{25} straight-chain, branched or cyclic alkane or alkene, optionally substituted with at least one member selected from the group consisting of Cl, Br, F, I, OH, NH_2 and SH;
- (iv) $-CH_3$, $-C_2H_5$, or C_3 to C_{25} straight-chain, branched or cyclic alkane or alkene comprising one to three heteroatoms selected from the group consisting of O, N, Si and S, and optionally substituted with at least one member selected from the group consisting of Cl, Br, F, I, OH, NH_2 and SH;

(v) C₆ to C₂₀ unsubstituted aryl, or C₃ to C₂₅ unsubstituted heteroaryl having one to three heteroatoms independently selected from the group consisting of O, N, Si and S; and

(vi) C₆ to C₂₅ substituted aryl, or C₃ to C₂₅ substituted heteroaryl having one to three heteroatoms independently selected from the group consisting of O, N, Si and S; and wherein said substituted aryl or substituted heteroaryl has one to three substituents independently selected from the group consisting of:

(1) -CH₃, -C₂H₅, or C₃ to C₂₅ straight-chain, branched or cyclic alkane or alkene, optionally substituted with at least one member selected from the group consisting of Cl, Br, F, I, OH, NH₂ and SH,

(2) OH,

(3) NH₂, and

(4) SH;

R⁷, R⁸, R⁹, and R¹⁰ are independently selected from the group consisting of:

(vii) -CH₃, -C₂H₅, or C₃ to C₂₅ straight-chain, branched or cyclic alkane or alkene, optionally substituted with at least one member selected from the group consisting of Cl, Br, F, I, OH, NH₂ and SH;

(viii) -CH₃, -C₂H₅, or C₃ to C₂₅ straight-chain, branched or cyclic alkane or alkene comprising one to three heteroatoms selected from the group consisting of O, N, Si and S, and optionally substituted with at least one member selected from the group consisting of Cl, Br, F, I, OH, NH₂ and SH;

(ix) C₆ to C₂₅ unsubstituted aryl, or C₃ to C₂₅ unsubstituted heteroaryl having one to three heteroatoms independently selected from the group consisting of O, N, Si and S; and

(x) C₆ to C₂₅ substituted aryl, or C₃ to C₂₅ substituted heteroaryl having one to three heteroatoms independently selected from the group consisting of O, N, Si and S; and wherein said substituted aryl or substituted heteroaryl has one to three substituents independently selected from the group consisting of:

(1) $-\text{CH}_3$, $-\text{C}_2\text{H}_5$, or C_3 to C_{25} straight-chain, branched or cyclic alkane or alkene, optionally substituted with at least one member selected from the group consisting of Cl, Br, F, I, OH, NH_2 and SH,

5

(2) OH,

(3) NH_2 , and

(4) SH; and

wherein optionally at least two of R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , and R^{10} can together form a cyclic or bicyclic alkanyl or alkenyl group.

10

11. An apparatus according to Claim 10 wherein at least one of R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , and R^{10} comprises F^- .

12. An apparatus according to Claim 1 wherein an ionic liquid comprises an anion selected from the group consisting of $[\text{CH}_3\text{CO}_2]^-$, $[\text{HSO}_4]^-$, $[\text{CH}_3\text{OSO}_3]^-$, $[\text{C}_2\text{H}_5\text{OSO}_3]^-$, $[\text{AlCl}_4]^-$, $[\text{CO}_3]^{2-}$, $[\text{HCO}_3]^-$, $[\text{NO}_2]^-$, $[\text{NO}_3]^-$, $[\text{SO}_4]^{2-}$, $[\text{PO}_4]^{3-}$, $[\text{HPO}_4]^{2-}$, $[\text{H}_2\text{PO}_4]^-$, $[\text{HSO}_3]^-$, $[\text{CuCl}_2]^-$, Cl^- , Br^- , I^- , SCN^- , and any fluorinated anion.

20

13. An apparatus according to Claim 1 wherein an ionic liquid comprises an anion selected from the group consisting of $[\text{BF}_4]^-$, $[\text{PF}_6]^-$, $[\text{SbF}_6]^-$, $[\text{CF}_3\text{SO}_3]^-$, $[\text{HCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{HFCCF}_2\text{SO}_3]^-$, $[\text{HCClFCF}_2\text{SO}_3]^-$, $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CF}_2\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{SO}_2)_3\text{C}]^-$, $[\text{CF}_3\text{CO}_2]^-$, $[\text{CF}_3\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CFHOCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{HCF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{ICF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[(\text{CF}_2\text{HCF}_2\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CFHCF}_2\text{SO}_2)_2\text{N}]^-$, and F^- .

14. An apparatus according to Claim 1 wherein an ionic liquid comprises a cation selected from the group consisting of pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, triazolium, phosphonium, or ammonium ions; and an anion selected from the group consisting of $[\text{CH}_3\text{CO}_2]^-$, $[\text{HSO}_4]^-$, $[\text{CH}_3\text{OSO}_3]^-$, $[\text{C}_2\text{H}_5\text{OSO}_3]^-$, $[\text{AlCl}_4]^-$, $[\text{CO}_3]^{2-}$, $[\text{HCO}_3]^-$, $[\text{NO}_2]^-$, $[\text{NO}_3]^-$, $[\text{SO}_4]^{2-}$, $[\text{PO}_4]^{3-}$, $[\text{HPO}_4]^{2-}$, $[\text{H}_2\text{PO}_4]^-$, $[\text{HSO}_3]^-$, $[\text{CuCl}_2]^-$, Cl^- , Br^- , I^- , SCN^- , $[\text{BF}_4]^-$, $[\text{PF}_6]^-$, $[\text{SbF}_6]^-$, $[\text{CF}_3\text{SO}_3]^-$,

30

$[\text{HCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{HFCCF}_2\text{SO}_3]^-$, $[\text{HCClFCF}_2\text{SO}_3]^-$, $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$,
 $[(\text{CF}_3\text{CF}_2\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{SO}_2)_3\text{C}]^-$, $[\text{CF}_3\text{CO}_2]^-$, $[\text{CF}_3\text{OCFHCF}_2\text{SO}_3]^-$,
 $[\text{CF}_3\text{CF}_2\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CFHOCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{HCF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$,
 $[\text{CF}_2\text{ICF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[(\text{CF}_2\text{HCF}_2\text{SO}_2)_2\text{N}]^-$, and
 5 $[(\text{CF}_3\text{CFHCF}_2\text{SO}_2)_2\text{N}]^-$.

15. An apparatus according to Claim 18 wherein an ionic liquid comprises a cation selected from the group consisting of 1-butyl-3-methylimidazolium, 1,2-dimethyl-3-propylimidazolium, 1-octyl-3-methylimidazolium, 1,3-dioctylimidazolium, 1-ethyl-3-methylimidazolium, 1-dodecyl-3-methylimidazolium, 1-heptyl-3-methylimidazolium, 3-methyl-1-propylpyridinium, 1-butyl-3-methylpyridinium, tetradecyl(trihexyl)phosphonium, or tributyl(tetradecyl)phosphonium ions; and an anion selected from the group consisting of $[\text{CH}_3\text{CO}_2]^-$, $[\text{HSO}_4]^-$, $[\text{CH}_3\text{OSO}_3]^-$, $[\text{C}_2\text{H}_5\text{OSO}_3]^-$, $[\text{AlCl}_4]^-$, $[\text{CO}_3]^{2-}$,
 15 $[\text{HCO}_3]^-$, $[\text{NO}_2]^-$, $[\text{NO}_3]^-$, $[\text{SO}_4]^{2-}$, $[\text{PO}_4]^{3-}$, $[\text{HPO}_4]^{2-}$, $[\text{H}_2\text{PO}_4]^-$, $[\text{HSO}_3]^-$, $[\text{CuCl}_2]^-$,
 Cl^- , Br^- , I^- , SCN^- , $[\text{BF}_4]^-$, $[\text{PF}_6]^-$, $[\text{SbF}_6]^-$, $[\text{CF}_3\text{SO}_3]^-$, $[\text{HCF}_2\text{CF}_2\text{SO}_3]^-$,
 $[\text{CF}_3\text{HFCCF}_2\text{SO}_3]^-$, $[\text{HCClFCF}_2\text{SO}_3]^-$, $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CF}_2\text{SO}_2)_2\text{N}]^-$,
 $[(\text{CF}_3\text{SO}_2)_3\text{C}]^-$, $[\text{CF}_3\text{CO}_2]^-$, $[\text{CF}_3\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCFHCF}_2\text{SO}_3]^-$,
 $[\text{CF}_3\text{CFHOCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{HCF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{ICF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$,
 20 $[\text{CF}_3\text{CF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[(\text{CF}_2\text{HCF}_2\text{SO}_2)_2\text{N}]^-$, and $[(\text{CF}_3\text{CFHCF}_2\text{SO}_2)_2\text{N}]^-$.

16. An apparatus according to Claim 1 wherein a refrigerant is selected from the group consisting of CHClF_2 (R-22), CHF_3 (R-23), CH_2F_2 (R-32), CH_3F (R-41), CHF_2CF_3 (R-125), CH_2FCF_3 (R-134a), $\text{CHF}_2\text{OCHF}_2$ (E-134), CH_3CClF_2 (R-142b), CH_3CF_3 (R-143a), CH_3CHF_2 (R-152a), $\text{CH}_3\text{CH}_2\text{F}$ (R-161), CH_3OCH_3 (E170), $\text{CF}_3\text{CF}_2\text{CF}_3$ (R-218), $\text{CF}_3\text{CHFCF}_3$ (R-227ea), $\text{CF}_3\text{CH}_2\text{CF}_3$ (R-236fa), $\text{CH}_2\text{FCF}_2\text{CHF}_2$ (R-245ca), $\text{CHF}_2\text{CH}_2\text{CF}_3$ (R-245fa), $\text{CH}_3\text{CH}_2\text{CH}_3$ (R-290), $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_3$ (R-600), $\text{CH}(\text{CH}_3)_2\text{CH}_3$ (R-600a), $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3$ (R-601), $(\text{CH}_3)_2\text{CHCH}_2\text{CH}_3$ (R-601a), $\text{CH}_3\text{CH}_2\text{OCH}_2\text{CH}_3$ (R-610), NH_3 , CO_2 ,
 30 $\text{CH}_3\text{CH}=\text{CH}_2$, and combinations thereof; and an ionic liquid comprises a cation selected from the group consisting of pyridinium, pyridazinium, pyrimidinium, pyrazinium, imidazolium, pyrazolium, thiazolium, oxazolium, triazolium, phosphonium, or ammonium ions; and an anion selected from the group consisting of $[\text{CH}_3\text{CO}_2]^-$, $[\text{HSO}_4]^-$, $[\text{CH}_3\text{OSO}_3]^-$, $[\text{C}_2\text{H}_5\text{OSO}_3]^-$, $[\text{AlCl}_4]^-$, $[\text{CO}_3]^{2-}$,

$[\text{HCO}_3]^-$, $[\text{NO}_2]^-$, $[\text{NO}_3]^-$, $[\text{SO}_4]^{2-}$, $[\text{PO}_4]^{3-}$, $[\text{HPO}_4]^{2-}$, $[\text{H}_2\text{PO}_4]^-$, $[\text{HSO}_3]^-$, $[\text{CuCl}_2]^-$,
 Cl^- , Br^- , I^- , SCN^- , $[\text{BF}_4]^-$, $[\text{PF}_6]^-$, $[\text{SbF}_6]^-$, $[\text{CF}_3\text{SO}_3]^-$, $[\text{HCF}_2\text{CF}_2\text{SO}_3]^-$,
 $[\text{CF}_3\text{HFCCF}_2\text{SO}_3]^-$, $[\text{HCClFCF}_2\text{SO}_3]^-$, $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$, $[(\text{CF}_3\text{CF}_2\text{SO}_2)_2\text{N}]^-$,
 $[(\text{CF}_3\text{SO}_2)_3\text{C}]^-$, $[\text{CF}_3\text{CO}_2]^-$, $[\text{CF}_3\text{OCFHCF}_2\text{SO}_3]^-$, $[\text{CF}_3\text{CF}_2\text{OCFHCF}_2\text{SO}_3]^-$,
 5 $[\text{CF}_3\text{CFHOCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{HCF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[\text{CF}_2\text{ICF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$,
 $[\text{CF}_3\text{CF}_2\text{OCF}_2\text{CF}_2\text{SO}_3]^-$, $[(\text{CF}_2\text{HCF}_2\text{SO}_2)_2\text{N}]^-$, and $[(\text{CF}_3\text{CFHCF}_2\text{SO}_2)_2\text{N}]^-$.

17. An apparatus according to Claim 1 that is fabricated as a refrigerator,
 a freezer, an ice machine, an air conditioner, an industrial cooling system, heater
 10 or heat pump.

18. An apparatus for temperature adjustment comprising
 (a) a compressor that increases the pressure of the vapor of at least one
 refrigerant;
 15 (b) a condenser that receives refrigerant vapor that is passed out of the
 compressor, and condenses the vapor under pressure to a liquid;
 (c) a pressure reduction device that receives liquid refrigerant that is
 passed out of the condenser, and reduces the pressure of the liquid to form a
 mixture of refrigerant in liquid and vapor form;
 20 (d) an evaporator that receives the mixture of liquid and vapor refrigerant
 that is passed out of the pressure reduction device, and evaporates the remaining
 liquid in the mixture to form refrigerant vapor; and
 (e) a conduit that returns to the compressor refrigerant vapor that is passed
 out of the evaporator;
 25 wherein a refrigerant is admixed with at least one ionic liquid.

19. A process for adjusting the temperature of an object, medium or a
 space comprising
 (a) providing a mechanical device having moving parts to increase the
 30 pressure of the vapor of at least one refrigerant, and providing at least one ionic
 liquid to lubricate the moving parts of the device;
 (b) condensing the refrigerant vapor under pressure to a liquid;
 (c) reducing the pressure of the liquid refrigerant to form a mixture of
 refrigerant in liquid and vapor form;

(d) evaporating the liquid refrigerant to form refrigerant vapor; and
(e) repeating step (a) to increase the pressure of the refrigerant vapor
formed in steps (c) and (d).

5 20. A process for adjusting the temperature of an object, medium or a
space comprising

(a) increasing the pressure of the vapor of at least one refrigerant;
(b) condensing the refrigerant vapor under pressure to a liquid;
(c) reducing the pressure of the liquid refrigerant to form a mixture of
10 refrigerant in liquid and vapor form;
(d) evaporating the liquid refrigerant to form refrigerant vapor;
(e) separating from the refrigerant vapor any ionic liquid present therein;
and
(f) repeating step (a) to increase the pressure of the refrigerant vapor
15 formed in steps (c) and (d).