Systems and methods for determining concentrations of components of a multiphase fluid. A first flowmeter receives a fluid flow and generates measurements of the fluid flow. After at least partially separating the fluid flow into mixtures of its components, second and third flowmeters generate measurements of the mixtures. A data processing apparatus operatively coupled to the flowmeters receives the measurements and determines concentrations of the components of the fluid flow. In an embodiment, the measurements are mass flow and density readings.

10 Claims, 4 Drawing Sheets
FIG. 1

Pressure Sensor and Transmitter

Flowmeter 1

Conductivity Sensor and Transmitter

Valve 1

Valve 2

Flowmeter 2

Isobutane

HF Mix

Flowmeter 3

Process return

Process Return Point

Sample in

Calibration Bypass 1

Calibration Bypass 2

3-way Valve

HF Mix

Isobutane

Sample Point

102

104

106

110

112

114

116

120

108

118

122

FIG. 1
Receive sample volume that includes HF mix and iC₄ through a first flowmeter

Separate HF mix from iC₄

Flow iC₄ through second flowmeter

Flow pure HF mix through conductivity sensor, third flowmeter, and pressure sensor

Compare mass flow rate through first flowmeter (M₁) with mass flow rate through second and third flowmeters (M₂ + M₃)

Increase valve opening through which iC₄ flows through second flowmeter

Lesser

Measure conductivity of pure HF mix

Determine concentration proportion of water based on conductivity

Measure mass flow and density of fluid

Determine concentration of either HF or ASO based on measured density

Determine concentration of remaining component as a difference, from 100%, of the sum of the concentration of water and either HF or ASO

Decrease valve opening through which iC₄ flows through second flowmeter

Greater

Determine concentration of iC₄ based on second and third flowmeter readings

Determine concentration of each component of the sample volume

FIG. 2A
Determine average density readings of first, second, and third flowmeters

Determine percentage drop in density at second flowmeter as a percentage of a difference between the density of HF and the average density reading of the second flowmeter to an average density reading of the first flowmeter

Determine percentage drop in density at third flowmeter as a percentage of a difference between the density of HF and the average density reading of the third flowmeter to an average density reading of the first flowmeter

Determine a K factor to convert density drop into volumetric ratio for iC₄

Determine volumetric proportion of iC₄ for the entire flow as a ratio of the percentage drop in density at the second flowmeter to the K factor

Determine volumetric flowrate of iC₄ as a product of an average volumetric flow rate through the second flowmeter and the volumetric proportion of iC₄ for the entire flow

Determine a ratio of the percentage drop in density at the third flowmeter to the K factor

Determine volumetric flowrate of iC₄ as a product of an average volumetric flow rate through the third flowmeter and the ratio of the percentage drop in density at the third flowmeter to the K factor

Determine a concentration of HF, ASO, and water in the HF mix

Determine a concentration of HF, ASO, and water in the sample volume
Set 3-way valve in a configuration in which no sample volume flows through the calibration bypass portions to the first and second flowmeters

Close the second valve and open the first valve

Compare mass flow and density readings from the first and second flowmeters, respectively

Substantially equal?

Adjust one or both flowmeters

End

FIG. 3
DETERMINING CONCENTRATIONS OF COMPONENTS OF A MIXTURE

CROSS REFERENCE TO RELATED APPLICATIONS

This application is a continuation of U.S. Ser. No. 13/302,780 filed Nov. 22, 2011, now U.S. Pat. No. 8,726,722, which is the nonprovisional of U.S. Provisional Application Ser. No. 61/418,659 filed Dec. 1, 2010 the contents of which are expressly incorporated herein by reference in their entirety including the contents and teachings of any references contained therein.


TECHNICAL FIELD

This document relates to determining concentrations of components in a multiphase flow.

BACKGROUND

Many industrial processes involve a mixture of a number of different fluids having different properties, for example, density or concentration. In some cases, knowing the concentration of the components of the mixture is useful for controlling the process. For example, the synthesis of isooctane (i.e., octane) for blending in a dynamic gasoline pool is a step in petroleum refining. One process for producing octane is HF Alkylation. In this process, Hydrogen Fluoride (HF, hydrofluoric acid) serves as a catalyst for the reaction of isobutane and C4 olefins to form octane or “alkylate.” A catalyst is a substance that increases a rate of a chemical reaction without itself becoming consumed in the process. The alkylation process turns low-molecular weight hydrocarbons, which formerly were waste, into a component of gasoline.

HF catalyst includes three components—HF (approximately 90%), water (approximately 1%), and acid-soluble organics (ASO) as the remainder. The concentration of each of the components affects the alkylation process. For example, a small concentration of water improves reaction efficiency, whereas a large concentration increases corrosiveness of the catalyst. Further, the concentration of HF can affect its role as a catalyst. For example, in acid runaway, HF’s role as a catalyst is compromised when the HF concentration falls below a threshold concentration and begins to participate as a reactant in side reactions that consume HF, but do not produce octane.

Acid runaway can be avoided by knowing HF and ASO concentrations. In particular, to avoid acid runaway, an HF concentration margin can be maintained over the critical concentration. Uncertainty around the HF and ASO concentrations can necessitate a greater margin over the critical concentration and cause a decrease in efficiency. In general, to optimize the alkylation process, the concentrations of the components of the HF catalyst are controlled. Minimizing a net consumption of HF by fine-tuning the operation of the alkylation unit can increase safety and profitability of the unit. For example, large output of octane barrels can be obtained at minimum cost by operating the unit at the lowest possible HF concentration without risking acid runaway. Grab-sampling extraction for laboratory analysis and online analysis are two methods for monitoring the concentrations of the components of HF catalyst (henceforth referred to as “HF mix”).

SUMMARY

This document describes technologies related to determining the concentrations of components in a multiphase flow. For example, this document describes technologies relating to an acid catalyst analyzer that can determine concentrations of the components of a sample mix that includes isobutane and HF mix.

Particular implementations of the subject matter described in this document can be implemented so as to realize one or more of the following advantages. Knowing HF and ASO levels can allow operation of the HF rerun in a manner that optimizes octane barrels without the risk of acid runaway, leading to improvement in profitability and safety. By calculating, rather than estimating, a concentration of iC4 in a sample volume that includes iC4 and HF mix, the concentration of each component of the sample volume (i.e., iC4, HF, ASO, and water) can be determined more accurately. The control systems implemented to control valve openings can enable a rapid, dynamic response to changes in the proportion of iC4 in the sample stream. The instrumentation implemented to measure concentration and density can be simple—mere lengths of pipes—in comparison to other systems that can be used to perform similar operations, e.g., Fourier-Transform, Near-Infrared Spectroscopy (FTNIR) system. Encoding temperature corrections in the control feedback loop can decrease or eliminate the need to measure control sample temperature. When the system is a continuously flowing sample system, the need for a more complex sample handling system with frequently cycling shutoff valves can be negated. Consequently, the need for maintenance and periodic replacement of the valves can also be negated. The system can output diagnostic information—flow rate, pressure, and temperature—periodically.

The details of one or more implementations are set forth in the accompanying drawings and the description below. Other features, aspects, and advantages of the subject matter will become apparent from the description, the drawings, and the claims.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a diagram of an example of an acid catalyst analyzer system.

FIG. 2A is a flowchart of an example process for determining concentrations of components of a volume sampled from a process stream.

FIG. 2B is a flowchart of another example process for determining concentrations of components of a volume sampled from a process stream.

FIG. 3 is a flowchart of an example process for calibrating three flowmeters.

Like reference numbers and designations in the various drawings indicate like elements.

DETAILED DESCRIPTION

Methods, apparatuses, and systems for determining concentrations of components of a multiphase flow, for example,
a mixture of HF mix and iC₄, are described below. In general, one or more flowmeters, one or more separation vessels, and one or more valves can be operated to separate one of the components (for example, iC₄) from the multiphase flow (for example, a mixture of HF mix and iC₄), and to determine the concentration of the components of the multiphase flow (for example, HF, ASO, water, and iC₄). For instance, readings from the flowmeters can be used to determine, rather than estimate, a concentration of each component of the multiphase flow. In some implementations, the position of a valve through which a component passes can be regulated such that noise in the readings taken by the flowmeters can be decreased or eliminated.

For example, in one implementation, a system receives a sample volume that includes a 4-component fluid that includes isobutane (iC₄) and HF mix (HF, water, and ASO). The sample volume flows through a first flowmeter into a separation vessel that separates the iC₄ from the HF mix. The HF mix is flowed through a second flowmeter and other instrumentation, and readings from the flowmeter and other instrumentation are used to determine the concentrations of the components of the HF mix. In particular, percentage concentrations of water, HF, and ASO are determined such that the sum of the percentage concentrations totals 100% for the HF mix. The iC₄ separated by the separation vessel is flowed from the separation vessel through a valve to a third flowmeter. The readings taken using the third flowmeter are used to determine a concentration of iC₄ flowing through the third flowmeter. As described above, the percentage concentration of each component of the HF mix is compared with the actual concentration of iC₄ in the sample volume to determine concentrations of each component in the sample volume. In an alternative or additional implementation, to maintain a balanced separation of the iC₄, a control feedback loop is implemented to control a position of the valve that controls the amount of iC₄, which has been separated by the separation vessel, that can flow through the third flowmeter, so that the third meter is filled only with iC₄, rather than a mixture of HF mix and iC₄. In this manner, the combined mass flowing through the second and third flowmeters may be kept, on average, equal to the mass flowing through the first flowmeter. This may decrease or eliminate noise in the readings taken by the third flowmeter, and further allow the direct measurement of the density of the iC₄.

FIG. 1 is an example of an acid catalyst analyzer system 100. The system 100 is implemented to determine the concentrations of the components of a multiphase flow, for example, HF mix that includes HF, water, and ASO. In general, the system 100 samples a volume of a combination of HF mix and hydrocarbons (for example, isobutane, iC₄) from a process sample point 102 using a valve 104. The system 100 includes a first flowmeter 108, a second flowmeter 106, and a third flowmeter 110. Each of the flowmeters 106, 108, 110 can be a Coriolis flowmeter using a bent tube or a Coriolis flowmeter using a straight flowtube.

The sample volume, which is a combination of iC₄ and HF mix, is flowed through the second flowmeter 106. The iC₄ is in the sample volume because it is an input into the alkylation process. The system 100 includes a 3-way valve 112 that receives the flow from the second flowmeter 106. The valve 112 can be set in a first configuration to flow the HF mix and the iC₄ into a separation vessel 114 and in a second, separate configuration to cross-calibrate the three flowmeters. When the 3-way valve 112 is set in the first configuration, the sampled volume flows from the second flowmeter 106 into the separation vessel 114. The vessel 114 is sized to separate the HF mix from the iC₄ in the sampled volume based on a difference in the densities of the two fluids. Specifically, the HF mix and the iC₄ are immiscible; also, the density of iC₄ (approximately 600 kg/m³) is less than that of the HF mix (approximately 1000 kg/m³). The separation vessel 114 is sized such that the high density HF mix settles to a bottom region of the vessel 114 while the low density iC₄ rises to a top region of the vessel 114 due to the density difference. In some implementations, the separation vessel 114 is large relative to a diameter of the pipe that samples the volume from the process sample point 112. For example, each of a width of the separator vessel 114 and a height of the separator vessel 114 can be at least three times the diameter of the pipe that samples the volume.

Once the HF mix and the iC₄ have been separated, when the second valve 116 is open, pure HF mix (that includes HF, water, and ASO) flows through a conductivity sensor and transmitter 118, the first flowmeter 108, and a pressure sensor and transmitter 120, which collectively can measure parameters—conductivity, density, pressure, respectively—of the pure HF mix. Using readings from the pressure sensor 120, the conductivity sensor 118, and the first flowmeter 108, a computer system 124 (for example, a flow computer) can determine the concentration of HF, water, and ASO, respectively, in the HF mix. Specifically, the computer system 124 can determine a percentage concentration of water based on the conductivity sensor measurements and a percentage concentration of either HF or ASO from the first flowmeter 108 and pressure sensor 120 measurements. The computer system 124 can infer the concentration of either ASO or HF as a difference, from 100%, of the sum of the percentage concentrations of water and either HF or ASO. Notably, at this stage, the concentrations of HF, water, and ASO are determined as a percentage of the HF mix, and not that of the sample volume.

The separated iC₄ flows through the first valve 122. From the valve, the separated iC₄ stream flows through the third flowmeter 110, which takes density and mass flow rate readings of the iC₄. Based on the readings measured by the third flowmeter 110, the computer system 124 can determine a concentration of iC₄ in the sample volume. Having determined the concentration of iC₄, the computer system 124 can determine a concentration of HF mix in the sample volume, and then determine the concentration of each component of the HF mix in the sample volume. For example, if the percentage concentration of water in the HF mix is 1%, and the concentration of iC₄ in the sample is determined to be 7%, then the percentage concentration of iC₄ in the sample is 93% (i.e., 100%-7%). Similarly, if the percentage concentration of ASO and HF in the HF mix are 4% and 95%, respectively, then the percentage concentration of ASO and HF mix in the sample volume are 3.72% and 88.35%, respectively.

Returning to the flow of the fluids through the system 100, after flowing through the instrumentation that includes the conductivity sensor 118, the first flowmeter 108 and the pressure sensor 120, and the first flowmeter 108, the HF mix and iC₄ stream, respectively, are then mixed again and flowed through valve 128 to the process return point 130. By continuously measuring a mass flow rate of iC₄ through the second flowmeter 106, the computer system 124 can also implement a mass balance calculation to ensure a balanced separation of the HF mix and the iC₄ (i.e., a separation in which the combined mass through the first flowmeter and the second flowmeter is equal, on average, to the mass through the second flowmeter). In addition, the computer system 124 can
cross-calibrate the first, second, and third flowmeters to ensure that density and mass flowrate measurements are accurate.

In some systems, the iC4 may be estimated to account for 6% of the sample volume sampled from the process sample point 102. However, the proportion of iC4 in the sample volume may vary and may not always be 6%. Using the techniques described below, the system 100 can calculate, rather than estimate or assume, the proportion of iC4. Based on the calculation of iC4 concentration, the system 100 can determine the percentage concentrations of the components of the HF mix in the sample volume. Furthermore, by implementing a dynamic response control system, the noise in the readings of the instrumentation can, in some cases, be decreased. Consequently, the system 100 can be implemented to control and optimize concentrations of the components of the HF mix in the sample volume.

The instrumentation shown in FIG. 1 (first, second, and third flowmeters 108, 106, 110, conductivity sensor and transmitter 118, pressure sensor and transmitter 120) are connected to the computer system 124. For example, the computer system 124 can include one or more computers or processors. In some implementations, the instrumentation transmits values of measured parameters to the computer system 124. For example, the second flowmeter 106 transmits density, mass flow rate, and volumetric flow rate values to the computer system 124. The computer system 124 can include a computer storage medium to store the values received from the instrumentation. The computer system 124 also includes a computer-readable medium storing computer software instructions executable by data processing apparatus to perform calculations related to mass balance, volumetric flow balance, and calculations to determine density, conductivity, and concentration.

In addition, the computer system 124 is coupled to the valves (the first valve 122, the second valve 116, the 3-way valve 112) to transmit instructions and control a state of each valve. For example, in response to receiving mass flow rate measurements from the flowmeters 106, 108, and 110, the computer system 124 can determine a level by which the first valve 122 should be opened or closed. The computer system 124 can transmit a signal to open (or close) the first valve 122 by that level. In some implementations, the computer system 124 can transmit a signal to partially open or partially close a valve. Alternatively, the computer system 124 can transmit a signal to partially open or partially close a valve. Strength of the signal that the computer system 124 transmits can correspond to a level to which the valve is to be opened or closed. The computer system 124 may implement a control system such as a proportional-integral-derivative (PID) control loop to control the position of the valve, for example, based on the mass flow rate measurements from the flowmeters 106, 108, and 110 according to the parameters described further below.

The density reading of the second flowmeter 106, which is exposed to the complete sample volume from process sample point 102, varies with the proportion of iC4 in the sample volume. However, the presence of two immiscible fluids (HF mix and iC4) may not induce any bias errors into the readings. On average, the density and mass flow readings generated by the second flowmeter 106 match the true mean values of the sample volume. In particular, if the mass flow and density readings are taken over an averaging period, then the averaged mass flow and density values can be considered as being sufficiently representative of the sample volume. In some implementations, the averaging period can be, for example, 1 minute or 5 minutes or 10 minutes, where the measurement update from each instrument can be of the order of 1 second.

The selection of the averaging period can vary between different systems or within the same system. The averaging period can be selected by, for example, monitoring the standard deviation of the density time series. A short averaging period can provide a good dynamic response to changes in the iC4 content, while providing stable readings for the mass balance calculation described below.

According to industrial knowledge, the percentage by volume of iC4, as a proportion of the sample volume, is estimated to be 6%. If the actual percentage by volume of iC4 in the sample volume matches the estimate and that the system 100 is operating in an entirely balanced way, i.e., the volume of iC4 in the separation vessel 114 is also 6%. If the first valve 122 was closed completely and the second valve 116 were open, the proportion of iC4 in the separation vessel 114 will steadily increase because the HF mix will escape through the second valve 116 whereas the iC4 film will not. In this scenario, because the liquids are incompressible and there are no gasses present in the system, the volumetric flow rates are balanced. In other words:

\[ Q_2 = Q_1 \]

The computer system 124 can calculate \( Q_2 \) for each flowmeter by receiving mass flow rate and density readings from the flowmeter. When the first valve 122 is closed:

\[ Q_2 = Q_1 \]

With reference to the mass flow balance in this scenario, because the separation vessel 114 is large, when the first valve 122 is closed, the vessel 114 begins to accumulate more iC4. However, until it is effectively full of iC4, the vessel 114 still manages to filter the iC4 out from the stream passing through the first flowmeter 108. This is achieved by substituting the equivalent volume of high density HF for low density iC4 in the stream sent to the first flowmeter 108, through the action of buoyancy in the separating vessel. Consequently:

\[ M_f (mass flow rate through second flowmeter) = M_i + M \]

The computer system 124 can receive mass flow rate readings from each flowmeter. When valve 1 is shut, \( M_i = 0 \). Thus, although the volumetric flow rate through the second flowmeter 106 is equal to the sum of the volumetric flow rates through the first flowmeter 108 and the third flowmeter 110, a mass flow imbalance exists because the sample volume that flows through the second flowmeter 106 contains low density iC4 while the sample volume that flows through the first flowmeter 108 is pure high density HF mix. In some implementations, the mass flow comparison can be performed over the averaging period described above.

Further, the imbalance is temporary, i.e., until the separation vessel 114 is effectively full of iC4. At that time the first flowmeter 108 will receive a combination of HF mix and iC4, and the mass balance will (on average) be restored. In addition, the imbalance that will be created even when the first valve 122 is completely closed will be small. With 6% iC4 in the sample stream, and assuming densities of 600 kg/m³ and 1000 kg/m³ for iC4 and HF mix, respectively, the mass flow difference between \( M_i \) and \( M_i + M_j \) will be 2.4%.

In another scenario, if the second valve 116 were closed and the first valve 122 were fully open, then:

\[ Q_1 = 0; M_f = 0 \]
Because all of the sample volume will flow through the third flowmeter 110:

\[ Q_3 = Q_2 \]

The top region of the separation vessel 114 includes iC\(_4\) only. Thus, initially, only iC\(_4\) flows through the third flowmeter. Consequently:

\[ M_3 > M_1 + M_3 \]

Mass balance will be restored when the layer of iC\(_4\) at the top region of the separation vessel 114 is exhausted. When the first valve 122 is fully open, if the second valve 116 is assumed to also be open (rather than entirely closed), more iC\(_4\), for example, twice the nominal flowrate of iC\(_4\), will be allowed through the third flowmeter 110. In other words, whereas the volume of iC\(_4\) flowing through the third flowmeter 110 would contain 6% iC\(_4\) by volume when the second valve 116 is fully open, the level by which the second valve 116 is open causes the volume of iC\(_4\) to be 12%. In this scenario also,

\[ M_3 > M_1 + M_3 \]

In sum, if the first valve 122 is maintained at an optimum open level, then a mass flow imbalance between the mass flow rate through the second flowmeter 106 and the first and third flowmeters (108 and 110, respectively) can be decreased or eliminated. If the first valve 122 is less open than the optimum open level, then the second flowmeter 106 measures less mass flow than the first and second flowmeters (108 and 110, respectively) combined. Conversely, if the first valve 122 is more open than the optimum open level, then the second flowmeter 106 measures more mass flow than the first and third flowmeters (108 and 110, respectively) combined.

The first valve 122 can remain open at a level sufficient to ensure that all of the iC\(_4\) in the sample volume is separated to the top region of the separation vessel 114 and that no iC\(_4\) enters the conductivity sensor and transmitter 118, the first flowmeter 108, and pressure sensor and transmitter 120, that measure the concentrations of HF, water, and ASO in the iC\(_4\)-free HF mix. In some situations, the separation vessel 114 can have so large a capacity that changes in the iC\(_4\) mass flow rate (and hence its proportion in the sample volume) are masked. In such situations, a change in the iC\(_4\) may not be observed in the third flowmeter 110 unless the position of the first valve 110 is adjusted. Because the aforementioned mass imbalance is related to a level at which the valve 110 is open, the system 100 can detect and adjust for changes in the proportion of iC\(_4\) using the control features of the computer system 124. To do so, the computer system 124 can execute computer software instructions stored on the computer-readable medium to perform the following function:

If \[ M_2 > M_1 + M_3 \] open valve 1 (122) further;

If \[ M_2 > M_1 + M_3 \] close valve 1 (122) further.

For example, flow control can be implemented as a PID control loop designed to maintain the value of \([M_2 - (M_1 + M_3)]\) at a set point of zero. By performing the aforementioned operations, the system 100 can facilitate a balanced separation of the HF mix and iC\(_4\) stream, so that the third flowmeter 110 will respond rapidly and accurately with changes in the flow rate of iC\(_4\) in the sample stream.

An example of pseudo-code executed by the computer system 124 to determine the concentration of iC\(_4\) in the sample stream is described below. The following variables are defined in the pseudocode:

<table>
<thead>
<tr>
<th>Variable</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>T</td>
<td>number of seconds between samples from Coriolis meters (for example 1 s)</td>
</tr>
<tr>
<td>K</td>
<td>number of samples in averaging period (for example 60) sample counter</td>
</tr>
<tr>
<td>M1</td>
<td>Mass flow rate from Mass Flow Meter 1, updated every T seconds, in kg/s</td>
</tr>
<tr>
<td>D1</td>
<td>Density reading from Mass Flow Meter 1, updated every T seconds, in kg/m^3</td>
</tr>
<tr>
<td>M2, M3, D2, D2</td>
<td>As above for Mass Flow Meter 2 and 3</td>
</tr>
<tr>
<td>Q1</td>
<td>Volumetric Flow from Mass Flow Meter 1 derived from M1 and D1, in liters/s</td>
</tr>
<tr>
<td>Q2, Q3</td>
<td>As above for Mass Flow Meter 2 and 3</td>
</tr>
<tr>
<td>isoB_V</td>
<td>Percentage of isobutene by Volume, %</td>
</tr>
<tr>
<td>M1_sum</td>
<td>M1 accumulated over K samples to collect samples</td>
</tr>
<tr>
<td>M2_sum, M3_sum</td>
<td>As above for Mass Flow Meter 2 and 3</td>
</tr>
<tr>
<td>M1_pv, M2_pv, M3_pv</td>
<td>As above for Mass Flow Meter 2 and 3</td>
</tr>
<tr>
<td>Q1_sum</td>
<td>Q1 averaged over K samples to collect samples</td>
</tr>
<tr>
<td>Q2_sum, Q3_sum</td>
<td>As above for Mass Flow Meter 2 and 3</td>
</tr>
<tr>
<td>Q2_sum/K</td>
<td>Q1 averaged over K samples to collect samples</td>
</tr>
<tr>
<td>MB</td>
<td>Mass Balance, kg/s, based on averaged data</td>
</tr>
<tr>
<td>MBP</td>
<td>Mass Balance Percentage, %</td>
</tr>
<tr>
<td>QB</td>
<td>Volume Balance, ft/s, based on averaged data</td>
</tr>
<tr>
<td>QBP</td>
<td>Volume Balance Percentage, %</td>
</tr>
</tbody>
</table>

Initialization:
\[ i = 0; M_1\_sum = 0; M_2\_sum = 0; M_3\_sum = 0; Q_1\_sum = 0; Q_2\_sum = 0; Q_3\_sum = 0; \]

Operation:

```
While (true) /* loop forever */
  1. i = i + 1;
  2. Collect M1, D1, M2, D2, M3, D3 from Coriolis Mass Flow Meters 1,2,3;
  3. Q1 = 1000 * M1/D1;
  4. Q2 = 1000 * M2/D2;
  5. Q3 = 1000 * M3/D3;
  6. Iso_BV = 100 * Q3/(Q1 + Q3);
  7. M1_sum = M1 + M1;
  8. M2_sum = M2 + M2;
  9. M3_sum = M3 + M3;
  10. Q1_sum = Q1 + Q1;
  11. Q2_sum = Q2 + Q2;
  12. Q3_sum = Q3 + Q3;
  13. Display/Record M1, D1, Q1, M2, D2, Q2, M3, D3, Q3, Iso_BV;
  14. if i = K /* do average calculations and flow control */
     a. M1_ave = M1_sum/K
     b. M2_ave = M2_sum/K
     c. M3_ave = M3_sum/K
     d. MB = M1_ave + M3_ave - M2_ave
     e. MBP = MB/M2_ave;
     f. Adjust valve position using PI controller with target MBP = 0
     g. Q1_ave = Q1_sum/K
     h. Q2_ave = Q2_sum/K
     i. Q3_ave = Q3_sum/K
     j. QBP = Q1_ave + Q3_ave - Q2_ave /* Volumetric Balance as Diagnostic */
     k. QBP = QBP/1000
     l. Display MB, MBP, valve position, QB, QBP
     m. i = 0; M1_sum = 0; M2_sum = 0; M3_sum = 0;
     n. Q1_sum = 0; Q2_sum = 0;
     o. Q3_sum = 0; /* reset totals */
  15. End /* i = K */
```

The proportion of iC\(_4\) in the sample stream is given by mass as follows:

\[ \% \text{iso-butane by mass} = 100\% \times \frac{M_3}{M_2 + M_3} \]

and by volume as follows:

\[ \% \text{iso-butane by volume} = 100\% \times \frac{Q_3/(Q_1 + Q_3)}{Q_2/1000} \]
The readings from the first and third flowmeters (108 and 110, respectively) are used for this calculation as they will have reduced noise compared with the readings from the second flowmeter 106.

Once the mass flow of iC₄ through the third flowmeter 110 is controlled by opening and closing the first valve 122, as described above, the concentrations of the components of the HF mix can be determined from the volume of HF mix that flows from the separation vessel 114 through the second valve 116 through the conductivity sensor and transmitter 118, the first flowmeter 108, and the pressure sensor and transmitter 120. When other trace components of the HF mix are neglected, the HF mix can be approximated to be a ternary solution, which can be modeled as a 3-component system consisting of HF, water, and ASO. The concentrations of the components of a ternary solution can be determined by determining the concentrations of two of its components and inferring the third as a remainder to make up a total of 100%. A two-component solution (i.e., a binary solution) can be characterized by obtaining a density measurement of one component. The density of the component should vary uniquely with solution composition because determination of one component allows the other to be inferred as the remainder to make up a total of 100%. Analogously, a ternary solution can be characterized by two density measurements, as long as each of the two density measurements varies uniquely with the respective concentration of the component.

Electrolytic conductivity sensors can be used to determine a concentration of water in HF. Water dissolves in undergoing the following dissociation reaction:

\[ 2\text{HF} + \text{H}_2\text{O} \rightarrow \text{HF}_2^+ + \text{H}_3\text{O}^+. \]

The dissolution products are ionized species, which makes the solution conductive. In binary HF/water solutions, the concentration of conductive species and therefore the conductivity is determined by the water concentration. The third major component of the HF mix, i.e., ASO, dissolves without the creation of ions. Consequently, a conductivity measurement is an uncompromised predictor of water concentration in the reaction mixture.

In some implementations, the conductivity sensor and transmitter 118 can be an electrodeless, non-contacting conductivity sensor. The sensor 118 can include a bore of inert material, for example, a pipe, through which the HF mix can pass. External to the bore, a coil called a “drive toroid” can generate an oscillating magnetic field, which induces an alternating current in the HF mix. The current, in turn, can generate an oscillating signal in a second external coil called a “sense toroid.” The signal in the sense toroid, which is proportional to HF mix fluid conductivity, can be transmitted to the computer system 124. Because the fluid conductivity is proportional to the water concentration, the computer system 124 can be executed to calculate water concentration based on the received signal.

Once the concentration of water in the HF mix has been measured, a second measurement, i.e., of density, can be used to determine the HF and ASO concentrations, as described below. Light hydrocarbons have densities as low as 0.6 g/cc, whereas the density of pure HF is nearly 1.0 g/cc (almost the same density as water). Thus, the relative concentrations of HF and ASO affect the density of the alklyation mixture while water does not. Consequently, density can be used to determine the composition of either HF or ASO in the HF mix.

In some implementations, the first flowmeter 108 and the pressure sensor and transmitter 120 can be implemented as a Coriolis sensor which measures both density and mass flow rate. A Coriolis sensor is a coiled tube on which a resonant vibration is induced. The system 100 can determine mass flow and density by analysis of phase shift and frequency of the tube vibration, respectively. As with conductivity, the HF mix passes through a simple pipe shaped as a coil. With two measurements—conductivity and density—two components of the HF mix are determined, and the third is the remainder.

Knowing the concentration of water from the conductivity measurement and the concentration of HF or ASO from the density measurement, the concentration of the third component is simply the remainder because:

% HF + % ASO + % Water = 100%

In some alternative situations, temperature and a second-order influence of interactions that may occur between the ASO and water may affect the conductivity and density measurements. All species in a solution affect each other directly or indirectly. In such situations, corrections for temperature and interactions between the components of the HF mix can be built into the system 100. To do so, a chemometric model can be developed for recording conductivity (cond), flow, pressure (P), density (p), and temperature (T) of HF mix. The data gathered from the system can be time-correlated to % HF, % water, and % ASO over a time period, for example, four months. By implementing statistical techniques (for example, multivariate regression), the data can be fit to equations that predict % water and % HF as functions of conductivity, density, and temperature. In equation form:

\[ \% \text{water} = f_1(\text{cond}) + f_2(p) + f_3(T) \]
\[ \% \text{HF} = f_1(\text{cond}) + f_2(p) + f_3(T) \]

Then,

\[ \% \text{ASO} = 100 - \% \text{HF} - \% \text{water} \]

In an alternative situation, the system 100 can be implemented to determine concentrations of components of the sample volume without implementing a control feedback loop, as described previously. In such situations, the fluid that exits the separation vessel 114 and flow through the third flowmeter 110 through the first valve 122 may not be pure iC₄ but may include some HF mix. Such a situation may arise because of one or more of several reasons. For example, it may not be practical to implement control of the first valve 122. The separation process in the separation vessel 114 may only be partial. The slugging of HF and iC₄ in the sample volume may cause some HF to flow through the third flowmeter 110.

In such situations, the concentrations of the components of the sample volume can be determined by placing the first valve 122 in a fixed position. For example, the first valve 122 is fixed such that approximately half of the total flow that flows through the second flowmeter 106 passes through the first flowmeter 108. Notably, the fluid that flows through the first flowmeter 108 is pure HF mix because the iC₄ has been separated from the HF mix in the separation vessel 114. Thus, the first flowmeter 108 receives pure HF mix, and both the second flowmeter 106 and the third flowmeter 108 receive a mixture of HF mix and iC₄. Because each of the first and third flowmeters (108, 110) receive half the sample volume relative to the second flowmeter 106, and because the third flowmeter 110 receives all the iC₄ and the first flowmeter 108 receives no iC₄, density readings by the flowmeters will be different from each other. Using the techniques described above, this difference can be used to determine the proportion of iC₄ by mass and volume.

The computer system 124 can receive readings measured by the flowmeters and determine average readings over aver-
The concentrations of the components of the sample volume average out. The computer system 124 can determine the concentrations of the components of the sample volume based on an assumption that a density of $iC_4$ is 593.4 kg/m$^3$.

An example of calculations executed by the computer system 124 to determine the concentration of $iC_4$ in the sample stream in the alternative implementation is described below. The following variables are defined in the pseudocode:

<table>
<thead>
<tr>
<th>Variable</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>T</td>
<td>number of seconds between samples from Coriolis meter (for example, 60)</td>
</tr>
<tr>
<td>K</td>
<td>number of samples in averaging period (for example, 60)</td>
</tr>
<tr>
<td>I</td>
<td>sample counter</td>
</tr>
<tr>
<td>M1</td>
<td>mass flow rate from Mass Flow Meter 1, updated every T seconds, in kg/s.</td>
</tr>
<tr>
<td>D1</td>
<td>density reading from Mass Flow Meter 1, updated every T seconds, in kg/m$^3$.</td>
</tr>
<tr>
<td>M2, M3, D2</td>
<td>Volumetric Flow from Mass Flow Meter 1 derived from M1 and D1, in liters/s.</td>
</tr>
<tr>
<td>Q1</td>
<td>as above for Mass Flow Meter 2 and 3</td>
</tr>
<tr>
<td>M2_av, M3_av, Q1_av</td>
<td>as above for Mass Flow Meter 2 and 3</td>
</tr>
<tr>
<td>M2_sum, M3_sum, Q1_sum</td>
<td>as above for Mass Flow Meter 2 and 3</td>
</tr>
</tbody>
</table>

The volumetric and mass flow rates through the flowmeters should balance. In other words:

\[ V_{2_{av}} = \frac{V_{1_{av}} + V_{3_{av}}}{2} \] (expected volumetric balance)

\[ M_{2_{av}} = M_{1_{av}} + M_{3_{av}} \] (expected mass flow balance)

If the mass balance readings are noisy, but the mean is near zero (i.e., less than a threshold), then the averaging time can be increased to decrease the noise. If, however, the mean is greater than the threshold, then the flowmeters can be calibrated as described with reference to FIG. 3. Further, slip between the $iC_4$ and HF mix (for example, at low velocity flows) can cause density offsets.

The first flowmeter 108 sees pure HF mix. Consequently, the computer system 124 can update the density readings of the first flowmeter 108 each averaging period:

\[ D_{HF} = D_{1_{av}} \]

The second flowmeter 106 and the third flowmeter 110 see different proportions of HF mix and $iC_4$. The computer system 124 can calculate density differences for these two flowmeters as follows:

\[ D_{2_{drop}} = \left( \frac{(D_{HF} - D_{2_{av}})}{D_{HF}} \right) \times 100\% \] (% drop in density at 2nd flowmeter)

\[ D_{3_{drop}} = \left( \frac{(D_{HF} - D_{3_{av}})}{D_{HF}} \right) \times 100\% \] (% drop in density at 3rd flowmeter)

A $K$ factor can be used to convert density drop into volumetric ratio for $iC_4$.

\[ K_{factor} = \frac{(D_{HF} - D_{iC4})}{D_{HF}} \] (density drop to volumetric ratio for $iC_4$

The computer system 124 can determine the volumetric proportion of $iC_4$ for the entire flow, based on the readings of the second flowmeter 106:

\[ iC_4_{check} = \frac{iC_4_{check}}{V_{100}} \] (volumetric flowrate of $iC_4$ through 2nd flowmeter)

The computer system 124 can implement a consistency check using the readings of the third flowmeter 110.

\[ iC_4_{check} = \frac{iC_4_{check}}{V_{100}} \] (volumetric flowrate of $iC_4$ through 3rd flowmeter)

The computer system 124 can check that $V_iC_4 = V_iC_4_{check}$.
concentration proportion of iC₄ (for example, percentage of iC₄ in the sample volume) can be determined based on second and third flowmeter readings \((228)\). The concentration proportion of each component of the sample volume can be determined based on the previously determined concentration proportions \((230)\). For example, if the percentage of iC₄ in the sample volume was determined to be 7%, then the proportion of HF mix in the sample volume is 93% (100% - 7%). If the percentage of water, HF, and ASO in the HF mix is 1%, 95%, and 4%, respectively, then the percentage of water, HF, and ASO in the sample volume is 0.93% (1% of 93%), 88.35% (95% of 93%), and 3.72% (4% of 93%), respectively.

FIG. 2B is a flowchart of another example process for determining concentrations of components of a volume sampled from a process stream. The example process can be implemented, for example, by the computer system 124 and may be employed when a consistency check, the volumetric flowrate calculated based on the HF mix can be separated using the separation vessel. The pure iC₄ can be flowed through the second flowmeter. The partially separated iC₄ that may include some HF mix can be flowed through the third flowmeter. Each of the first, second, and third flowmeters can take density and flow measurements on the received fluids, and transmit the measurements to the computer system 124.

Average density readings of first, second, and third flowmeters can be determined \((232)\). A percentage drop in density at second flowmeter can be calculated as a percentage of a difference between the density of HF and the average density reading of the second flowmeter, to an average density reading of the first flowmeter \((234)\). A percentage drop in density at third flowmeter can be determined as a percentage of a difference between the density of HF and the average density reading of the third flowmeter, to an average density reading of the first flowmeter \((236)\). A K_factor can be determined to convert density drop into volumetric ratio for iC₄ \((238)\). A volumetric proportion of iC₄ for the entire flow can be calculated as a ratio of the percentage drop in density at the second flowmeter to the K_factor \((240)\). A volumetric flowrate of iC₄ as a product of average volumetric flow rate through the second flowmeter and the volumetric proportion of iC₄ for the entire flow can be determined \((242)\). A ratio of the percentage drop in density at the third flowmeter to the K_factor can be calculated \((244)\). A volumetric flowrate of iC₄ as a product of an average volumetric flow rate through the third flowmeter and the ratio of the percentage drop in density at the third flowmeter to the K_factor can be determined \((246)\). To implement a consistency check, the volumetric flowrate calculated based on the volumetric proportion of iC₄ through the second flowmeter can be compared with the volumetric proportion of iC₄ through the third flowmeter. Similarly to the process described with reference to FIG. 2A, a concentration of HF, ASO, and water in the HF mix can be determined \((248)\). A concentration of HF, ASO, and water in the sample volume can be determined based on the concentration of iC₄ in the sample volume and a concentration of HF, ASO, and water in the HF mix \((250)\).

FIG. 3 is a flowchart of an example process \((300)\) for calibrating three flowmeters. In some implementations, the system \((300)\) can cross-calibrate the first, second, and third flowmeters \((106, 108, 110, 302)\) respectively to ensure an accurate determination of the flow rate of iC₄. Cross-calibration can decrease or eliminate offsets between the readings of the three flowmeters to ensure that a correct mass balance calculation is achieved. To cross-calibrate the flowmeters, the 3-way valve 112 can be set in a second configuration in which no sample volume flows to the separation vessel 114; instead, the sample volumes flow through calibration bypass portion 1 and calibration bypass portion 2 to the first flowmeter 108 and the second flowmeter 110, respectively \((302)\). The 3-way valve 112 is an example of a valve arrangement to enable fluid flow in the first configuration \((300)\) and the second configuration. Another type of valve or multiple valves can be arranged to achieve a function that is equivalent to that of the 3-way valve 112.

With the 3-way valve 112 set in the second configuration, the second valve 116 can be closed and the first valve 122 can be opened such that all of the sample volume flows first through the second flowmeter 106 and then through the third flowmeter 110 \((304)\). The two flowmeters can transmit mass flow and density readings to the computer system 124. The computer system 124 can average the readings over the averaging period, and compare \((306)\) the readings. It can be checked if the two readings are substantially equal to each other. For example, two readings are substantially equal to each other if the difference between the two readings satisfies a threshold, such as within 0.2% of the reading. If so, then the process \((300)\) ends \((308)\) as the two flowmeters are calibrated. If not, then this indicates that the flowmeters are not calibrated. The flowmeters can then be adjusted \((310)\) and the comparison can be repeated. Similarly, the first and second flowmeters \((108, 106)\), respectively can be cross-calibrated by setting the 3-way valve 112 in the second configuration, closing the first valve 122 and opening the second valve 116. In addition to calibrating the flowmeters by comparing mass flow and density readings, calibration based on volumetric flow rates can also be performed. Specifically, when the 3-way valve 112 is in the second configuration and both the first valve 116 and the second valve 122 are open, then the volumetric flow rate through the second flowmeter 106 should be equal to the sum of the volumetric flow rates through the first flowmeter 108 and through the third flowmeter 110.

Examples of flowmeters that can be implemented in system \((100)\) are described below. Types of flowmeters include digital flowmeters. For example, U.S. Pat. No. 6,311,136, which is hereby incorporated herein by reference, discloses the use of a digital flowmeter and related technology including signal processing and measurement techniques. Such digital flowmeters may be very precise in their measurements, with little or negligible noise, and may be capable of enabling a wide range of positive and negative gains at the driver circuitry for driving the conduit. Such digital flowmeters are thus advantageous in a variety of settings. For example, commonly-assigned U.S. Pat. No. 6,505,519, which is incorporated herein by reference, discloses the use of a wide gain range, and/or the use of negative gain, to prevent stalling and to more accurately exercise control of the flowtube, even during difficult conditions such as two-phase flow (for example, a flow containing a mixture of liquid and gas).

Although digital flowmeters are specifically discussed below, it should be understood that analog flowmeters also exist. Although such analog flowmeters may be prone to typical shortcomings of analog circuitry, for example, low precision and high noise measurements relative to digital flowmeters, they also may be compatible with the various techniques and implementations discussed herein.

Implementations of the computer system 124 and the operations described with reference to the computer system 124 can be implemented in digital electronic circuitry, or in computer software, firmware, or hardware, including the structures disclosed in this specification and their structural
equivalents, or in combinations of one or more of them. Implementations of the computer system 124 can be implemented as one or more computer programs, i.e., one or more modules of computer program instructions, encoded on computer storage medium for execution by, or to control the operation of, data processing apparatus. Alternatively or in addition, the program instructions can be encoded on an artificially-generated propagated signal, for example, a machine-generated electrical, optical, or electromagnetic signal, that is generated to encode information for transmission to suitable receiver apparatus for execution by a data processing apparatus. The computer storage medium can be, or be included in, a computer-readable storage device, a computer-readable medium, a computer readable medium, a memory, a computer storage medium, a computer storage device, a storage device, a computer readable medium, or an optical or magnetic disk. Moreover, while a computer storage medium is not a propagated signal, a computer storage medium can be a source or destination for program instructions encoded in an artificially-generated propagated signal. The computer storage medium can also be, or be included in, on or more separate physical components or media (for example, multiple CDs, disks, or other storage devices).

The operations described in this specification can be implemented as operations performed by the data processing apparatus 124 on data stored on one or more computer-readable storage devices or received from other sources. The term “data processing apparatus” encompasses all kinds of apparatus, devices, and machines for processing data, including by way of example a programmable processor, a computer, a system on a chip, or multiple ones, or combinations of the foregoing. The apparatus can include special purpose logic circuitry, for example, an FPGA (field programmable gate array) or an ASIC (application-specific integrated circuit). The apparatus can also include, in addition to hardware, code that creates an execution environment for the computer program in question, for example, code that constitutes processor firmware, a protocol stack, a database management system, an operating system, a cross-platform runtime environment, a virtual machine, or a combination of one or more of them. The apparatus and execution environment can realize various different computing model infrastructures, such as web services, distributed computing and grid computing infrastructures.

A computer program (also known as a program, software, application, code, or script) can be written in any form of programming language, including compiled or interpreted languages, declarative or procedural languages, and it can be deployed in any form, including as a stand-alone program or as a module, component, subroutine, object, or other unit suitable for use in a computing environment. A computer program may, but need not, correspond to a file in a file system. A program can be stored as a portion of data that holds other programs or data (for example, one or more scripts stored in a markup language document), in a single file dedicated to the program in question, or in multiple coordinated files (for example, files that store one or more modules, sub-programs, or portions of code). A computer program can be deployed to be executed on one computer or on multiple computers that are located at one site or distributed across multiple sites and interconnected by a communication network.

The processes and logic flows described in this specification can be performed by one or more programmable processors executing one or more computer programs to perform actions by operating on input data and generating output. The processes and logic flows can also be performed by, and apparatus can also be implemented as, special purpose logic circuitry, for example, an FPGA (field programmable gate array) or an ASIC (application-specific integrated circuit).

Processors suitable for the execution of a computer program include, by way of example, both general and special purpose microprocessors, and any one or more processors of any kind of digital computer. Generally, a processor will receive instructions and data from a read-only memory or a random access memory or both. The essential elements of a computer are a processor for performing actions in accordance with instructions and one or more memory devices for storing instructions and data. Generally, a computer will also include, or be operatively coupled to receive data from or transfer data to, or both, one or more mass storage devices for storing data, for example, magnetic, magneto-optical disks, or optical disks. However, a computer need not have such devices. Moreover, a computer can be embedded in another device, for example, a mobile telephone, a personal digital assistant (PDA), a mobile audio or video player, a game console, a Global Positioning System (GPS) receiver, or a portable storage device (for example, a universal serial bus (USB) flash drive). Devices suitable for storing computer program instructions and data include all forms of non-volatile memory, media and memory devices, including by way of example semiconductor memory devices, for example, EPROM, EEPROM, and flash memory devices; magnetic disks, for example, internal hard disks or removable disks; magneto-optical disks; and CD-ROM and DVD-ROM disks.

To provide for interaction with a user, implementations of the computer system 124 can include a display, for example, a CRT (cathode ray tube) or LCD (liquid crystal display) monitor, for displaying information to the user and a keyboard and a pointing device, for example, a mouse or a trackball, by which the user can provide input to the computer. Other kinds of devices can be used to provide for interaction with a user as well; for example, feedback provided to the user can be any form of sensory feedback, for example, visual feedback, auditory feedback, or tactile feedback; and input from the user can be received in any form, including acoustic, speech, or tactile input. In addition, a computer can interact with a user by sending documents to and receiving documents from a device that is used by the user; for example, by sending web pages to a web browser on a user’s client device in response to requests received from the web browser. While this specification contains many specific implementation details, these should not be construed as limitations on the scope of any invention or of what may be claimed, but rather as descriptions of features specific to particular implementations of particular inventions. Certain features that are described in this specification in the context of separate implementations can also be implemented in combination in a single implementation. Conversely, various features that are described in the context of a single implementation can also be implemented in multiple implementations separately or in any suitable subcombination. Moreover, although features may be described above as acting in certain combinations and even initially claimed as such, one or more features from a claimed combination can in some cases be excised from the combination, and the claimed combination may be directed to a subcombination or variation of a subcombination.

Similarly, while operations are depicted in the drawings in a particular order, this should not be understood as requiring that such operations be performed in the particular order shown or in sequential order, or that all illustrated operations
be performed, to achieve desirable results. In certain circumstances, multitasking and parallel processing may be advantageous. Moreover, the separation of various system components in the implementations described above should not be understood as requiring such separation in all implementations, and it should be understood that the described program components and systems can generally be integrated together in a single software product or packaged into multiple software products.

Thus, particular implementations of the subject matter have been described. Other implementations are within the scope of the following claims. In some cases, the actions recited in the claims can be performed in a different order and still achieve desirable results. In addition, the processes depicted in the accompanying figures do not necessarily require the particular order shown, or sequential order, to achieve desirable results. In certain implementations, multitasking and parallel processing may be advantageous.

What is claimed is:

1. A system to determine concentrations of components of a four-component multiphase fluid, the system comprising:
   - a first flowmeter configured to receive a fluid flow and to generate one or more measurements of the fluid flow, the fluid flow including a first component, a second component, a third component, and a fourth component;
   - a separation vessel configured to receive the fluid flow, and to separate the fluid flow into a first constituent that includes the first, second, and third components and a second constituent;
   - a second flowmeter configured to receive the second constituent through a valve and to generate one or more measurements of the second constituent;
   - a third flowmeter configured to receive the first constituent and to generate one or more measurements of the first constituent; and
   - a data processing apparatus operatively coupled to the first flowmeter, the second flowmeter, and the third flowmeter, the data processing apparatus configured to perform operations comprising:
     - receiving the one or more measurements of the fluid flow from the first flowmeter;
     - receiving the one or more measurements of the second constituent from the second flowmeter;
     - receiving the one or more measurements of the first constituent from the third flowmeter; and
     - determining a concentration of the fourth component in the fluid flow.

2. The system of claim 1, wherein the second constituent consists of the fourth component.

3. The system of claim 2, wherein the operations further comprise determining, based on the measurements received from the first, second, and third flowmeters, a level by which the first valve is open to maintain a mass balance between the first, second, and third flowmeters.

4. The system of claim 2, wherein the operations further comprise determining a concentration of each of the first component, the second component, and the third component in the fluid.

5. The system of claim 4, wherein the operations further comprise determining a concentration of each of the first component, the second component, and the third component in the fluid based on the concentration of the fourth component in the fluid and the concentration of each of the first component, the second component, and the third component in the first constituent.

6. The system of claim 1, further comprising a valve coupled to the first flowmeter, the second flowmeter, and the third flowmeter, and configured to enable flow of the sample volume received by the first flowmeter to the second flowmeter only or the third flowmeter only or both flowmeters simultaneously.

7. The system of claim 6, wherein the valve is a 3-way valve.

8. The system of claim 1, wherein the second constituent includes a fourth component in a mixture with the first, second, and third components, wherein the fourth component constitutes a greater proportion in the second constituent than in the fluid flow.

9. The system of claim 8, wherein the operations further comprise determining a concentration of each of the first component, the second component, and the third component in the fluid flow based on the concentration of each of the first component, the second component, and the third component in the first constituent and the concentration of the fourth component in the fluid flow.

10. The system of claim 1, wherein the received one or more measurements of the fluid flow from the first flowmeter, the received one or more measurements of the second constituent from the second flowmeter, and the received one or more measurements of the first constituent from the third flowmeter are used to determine the concentration of the fourth component in the fluid flow.

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