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(54) **Determining concentrations of components of a mixture**

Bestimmung der Konzentrationen von Bestandteilen eines Gemisches

Détermination des concentrations des composants d'un mélange

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Description

TECHNICAL FIELD

5 **[0001]** This document relates to determining concentrations of components in a multiphase flow.

BACKGROUND

10 **[0002]** 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 being consumed in the process. The alkylation process turns low-molecular weight hydrocarbons, which formerly were waste, into a component of gasoline.

15 **[0003]** 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.

20 **[0004]** 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

35 **[0005]** 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.

40 **[0006]** 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.

50 **[0007]** 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.

[0008] US 2010/023275 A1 discloses an apparatus for concentration determination of at least one component in an acid catalyst for hydrocarbon conversion containing an unknown concentration of an acid, an acid-soluble -oil and water. This apparatus has the disadvantage of noise in the readings of the flowmeter.

[0009] Accordingly it is an object of the present invention to provide an acid catalyst analyzer system having lower noise in the readings of the flowmeter.

[0010] This object is achieved by an acid catalyst analyzer system in accordance with claim 1. Preferred embodiments

of the invention are disclosed in the dependent claims.

BRIEF DESCRIPTION OF THE DRAWINGS

[0011]

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.

[0012] Like reference numbers and designations in the various drawings indicate like elements.

DETAILED DESCRIPTION

[0013] Methods, apparatuses, and systems for determining concentrations of components of a multiphase flow, for example, a mixture of HF mix and iC_4 , 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, iC_4) from the multiphase flow (for example, a mixture of HF mix and iC_4), and to determine the concentration of the components of the multiphase flow (for example, HF, ASO, water, and iC_4). 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.

[0014] For example, in one implementation, a system receives a sample volume that includes a 4-component fluid that includes isobutane (iC_4) and HF mix (HF, water, and ASO). The sample volume flows through a first flowmeter into a separation vessel that separates the iC_4 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 iC_4 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 iC_4 flowing through the third flowmeter. As described below, the percentage concentration of each component of the HF mix is compared with the actual concentration of iC_4 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 iC_4 , a control feedback loop is implemented to control a position of the valve that controls the amount of iC_4 , which has been separated by the separation vessel, that can flow through the third flowmeter, so that the third meter is filled only with iC_4 , rather than a mixture of HF mix and iC_4 . 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 iC_4 .

[0015] 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, iC_4) 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.

[0016] The sample volume, which is a combination of iC_4 and HF mix, is flowed through the second flowmeter 106. The iC_4 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 iC_4 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 iC_4 in the sampled volume based on a difference in the densities of the two fluids. Specifically, the HF mix and the iC_4 are immiscible; also, the density of iC_4 (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 iC_4 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.

[0017] Once the HF mix and the iC_4 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.

[0018] The separated iC_4 flows through the first valve 122. From the valve, the separated iC_4 stream flows through the third flowmeter 110, which takes density and mass flow rate readings of the iC_4 . Based on the readings measured by the third flowmeter 110, the computer system 124 can determine a concentration of iC_4 in the sample volume. Having determined the concentration of iC_4 , 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 iC_4 in the sample is determined to be 7%, then the percentage concentration of HF mix in the sample is 93% (i.e., 100% - 7%), and the percentage concentration of water in the sample volume is 0.93% (i.e. 1% of 93%). 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.

[0019] 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 iC_4 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 iC_4 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 iC_4 (i.e., a separation in which the combined mass through the first flowmeter and the third 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.

[0020] In some systems, the iC_4 may be estimated to account for 6% of the sample volume sampled from the process sample point 102. However, the proportion of iC_4 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 iC_4 . Based on the calculation of iC_4 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.

[0021] 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.

[0022] 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 entirely open or entirely 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.

[0023] 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 iC_4 in the sample volume. However, the presence of two immiscible fluids (HF mix and iC_4) 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 iC_4 content, while providing stable readings for the mass balance calculation described below.

[0024] According to industrial knowledge, the percentage by volume of iC_4 , as a proportion of the sample volume, is estimated to be 6%. If the actual percentage by volume of iC_4 in the sample volume matches the estimate and that the system 100 is operating in an entirely balanced way, i.e., the volume of iC_4 in the separation vessel 114 is also 6%. If the first valve 122 were closed completely and the second valve 116 were open, the proportion of iC_4 in the separation vessel 114 will steadily increase because the HF mix will escape through the second valve 116 whereas the iC_4 , that has risen to the top region of the separation vessel 114, 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:

$$Q_2 \text{ (volumetric flow rate through second flowmeter)} = Q_1 + Q_3$$

[0025] The computer system 124 can calculate Q_i for each flowmeter by receiving mass flow rate and density readings from the flowmeter. When the first valve 122 is closed:

$$Q_3 = 0; Q_2 = Q_1$$

[0026] 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 iC_4 . However, until it is effectively full of iC_4 , the vessel 114 still manages to filter the iC_4 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 iC_4 in the stream sent to the first flowmeter 108, through the action of buoyancy in the separating vessel. Consequently:

$$M_2 \text{ (mass flow rate through second flowmeter)} < M_1 + M_3$$

The computer system 124 can receive mass flow rate readings from each flowmeter. When valve 1 is shut, M_3 is 0.

[0027] 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 flow meter 106 contains low density iC_4 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.

[0028] Further, the imbalance is temporary, i.e., until the separation vessel 114 is effectively full of iC_4 . At that time the first flowmeter 108 will receive a combination of HF mix and iC_4 , 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% iC_4 in the sample stream, and assuming densities of 600 kg/m³ and 1000 kg/m³ for iC_4 and HF mix, respectively, the mass flow difference between M_2 and $(M_1 + M_3)$ will be 2.4%.

[0029] In another scenario, if the second valve 116 were closed and the first valve 122 were fully open, then:

$$Q_1 = 0; M_1 = 0$$

Because all of the sample volume will flow through the third flowmeter 110:

$$Q_2 = Q_3$$

[0030] The top region of the separation vessel 114 includes iC_4 only. Thus, initially, only iC_4 flows through the third flowmeter 110. Consequently:

$$M_2 > M_1 + M_3$$

[0031] 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,

$$M_2 > M_1 + M_3$$

[0032] 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.

[0033] 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.

[0034] 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:

Variable	Definition
T	number of seconds between samples from Coriolis meters (for example 1s)
K	number of samples in averaging period (for example 60)
I	sample counter
M1	Mass flow rate from Mass Flow Meter 1, updated every T seconds, in kg/s.
D1	Density reading from Mass Flow Meter 1, updated every T seconds, in kg/m ³ .
M2, M3, D2, D2	As above for Mass Flow Meter 2 and 3
Q1	Volumetric Flow from Mass Flow Meter 1 derived from M1 and D1, in litres/s.
Q2, Q3	As above for Mass Flow Meter 2 and 3

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(continued)

Variable	Definition
isoB_V	Percentage of isobutene by Volume, %
M1_sum	M1 accumulated over K samples to collect samples
M2_sum, M3_sum	As above for Mass Flow Meter 2 and 3
M1_av	M1 averaged over K samples
M2_av, M3_av	As above for Mass Flow Meter 2 and 3
Q1_sum	Q1 accumulated over K samples to collect samples
Q2_sum, O3_sum	As above for Mass Flow Meter 2 and 3
Q1_av	Q1 averaged over K samples
Q2_av, Q3_av	As above for Mass Flow Meter 2 and 3
MB	Mass Balance, kg/s, based on averaged data
MBP	Mass Balance Percentage, %
QB	Volume Balance, l/s, based on averaged data
QBP	Volume Balance Percentage, %
Initialisation: I = 0; M1_sum = 0; M2_sum = 0; M3_sum = 0; Q1_sum = 0; Q2_sum = 0; O3_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. $M1_sum = M1_sum + M1$

8. $M2_sum = M2_sum + M2$

9. $M3_sum = M3_sum + M3$

10. $Q1_sum = Q1_sum + Q1$

11. $Q2_sum = Q2_sum + Q2$

12. $O3_sum = Q3_sum + Q3$

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. $QB = Q1_ave + Q3_ave - Q2_ave$ /* Volumetric Balance as Diagnostic */
 k. $QBP = QB/Q2ave$
 l. Display MB, MBP, valve position, QB, QBP
 m. $I = 0; M1_sum = 0; M2_sum = 0; M3_sum = 0; Q1_sum = 0; Q2_sum = 0; Q3_sum = 0;$ /* reset totals */

15. End /* I == K */

[0035] The proportion of iC_4 in the sample stream is given by mass as follows:

$$\% \text{ iso-butane by mass} = 100\% * M_3 / (M_1 + M_3),$$

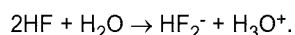
and by volume as follows:

$$\% \text{ iso-butane by volume} = 100\% * Q_3 / (Q_1 + Q_3)$$

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.

[0036] Once the mass flow of iC_4 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.

[0037] Electrolytic conductivity sensors can be used to determine a concentration of water in HF. Water dissolves in undergoing the following dissociation reaction:



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.

[0038] 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.

[0039] 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 alkylation mixture while water does not. Consequently, density can be used to determine the composition of either HF or ASO in the HF mix.

[0040] 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.

[0041] 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:

$$\%HF + \%ASO + \%Water = 100\%$$

[0042] 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:

$$\% \text{ water} = f_1(\text{cond}) + f_1(p) + f_1(T)$$

$$\% \text{ HF} = f_2(\text{cond}) + f_2(p) + f_2(T)$$

[0043] Then,

$$\% \text{ ASO} = 100 - \% \text{ HF} - \% \text{ water}$$

[0044] 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 iC_4 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 iC_4 in the sample volume may cause some HF to flow through the third flowmeter 110.

[0045] 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 iC_4 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 110 receive a mixture of HF mix and iC_4 . 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 iC_4 and the first flowmeter 108 receives no iC_4 , density readings by the flowmeters will be different from each other. Using the techniques described below, this difference can be used to determine the proportion of iC_4 by mass and volume.

[0046] The computer system 124 can receive readings measured by the flowmeters and determine average readings over averaging periods. The averaging periods can be selected to ensure that short-term variations in the HF mix to iC_4 proportions 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 .

[0047] 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:

Variable	Definition
T	number of seconds between samples from Coriolis meters (for example 1s)
K	number of samples in averaging period (for example, 60)
I	sample counter
M1	Mass flow rate from Mass Flow Meter 1, updated every T seconds, in kg/s.
D1	Density reading from Mass Flow Meter 1, updated every T seconds, in kg/m ³ .
M2, M3, D2, D3	As above for Mass Flow Meter 2 and 3
Q1	Volumetric Flow from Mass Flow Meter 1 derived from M1 and D1, in litres/s.
Q2, Q3	As above for Mass Flow Meter 2 and 3
isoB_V	Percentage of isobutene by Volume, %
M1_sum	M1 accumulated over K samples to collect samples
M2_sum, M3_sum	As above for Mass Flow Meter 2 and 3
M1_av	M1 averaged over K samples
M2_av, M3_av	As above for Mass Flow Meter 2 and 3
Q1_sum	Q1 accumulated over K samples to collect samples
Q2_sum, Q3_sum	As above for Mass Flow Meter 2 and 3
Q1_av	Q1 averaged over K samples
Q2_av, Q3_av	As above for Mass Flow Meter 2 and 3
D_iC ₄	Density of iC ₄ (assumed 593.4 kg/m ³)
D_HF	Calculated density of HF mix
V_iC ₄	Calculated volumetric flow rate of iC ₄
V_HF	Calculated volumetric flow rate of HF mix

[0048] Variables M1, D1, M2, D2, M3, and D3 are measured by the first, second, and third flowmeters, respectively. The computer system 124 determines corresponding time averages for the measurements, for example, by executing the pseudo-code described above. The computer system 124 then calculates the density of the HF mix and volumetric flow rates of iC₄ and HF mix as follows.

[0049] The volumetric and mass flow rates through the flowmeters should balance. In other words:

$$V2_{av} = V1_{av} + V3_{av} \text{ (expected volumetric balance)}$$

$$M2_{av} = M1_{av} + M3_{av} \text{ (expected mass flow balance)}$$

[0050] 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 iC₄ and HF mix (for example, at low velocity flows) can cause density offsets.

[0051] 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:

$$D_{HF} = D1_{av}$$

[0052] The second flowmeter 106 and the third flowmeter 110 see different proportions of HF mix and iC₄. The computer

system 124 can calculate density differences for these two flowmeters as follows:

$$D2_drop = [(D_HF - D2_av)/D_HF] \times 100\% \text{ (\% drop in density at 2}^{nd}\text{ flowmeter)}$$

$$D3_drop = [(D_HF - D3_av)/D_HF] \times 100\% \text{ (\% drop in density at 3}^{rd}\text{ flowmeter)}$$

[0053] A K factor can be used to convert density drop into volumetric ratio for iC₄.

[0054] $K_factor = (D_HF - D_iC_4)/D_HF$ (density drop to volumetric ratio for iC₄) The computer system 124 can determine the volumetric proportion of iC₄ for the entire flow, based on the readings of the second flowmeter 106:

$$iC_4_V = D2_drop/K_factor \text{ (\% by volume of iC}_4\text{ in fluid received by 3}^{rd}\text{ flowmeter)}$$

$$V_iC_4 = V2_ave \times iC_4_V/100 \text{ (volumetric flowrate of iC}_4\text{ through 2}^{nd}\text{ flowmeter)}$$

[0055] The computer system 124 can implement a consistency check using the readings of the third flowmeter 110.

$$iC_4_V_check = D3_drop/K_factor \text{ (\% volume of iC}_4\text{ in fluid received by 3}^{rd}\text{ flowmeter)}$$

$$V_iC_4_check = V3_ave \times iC_4_V_check/100 \text{ (volumetric flowrate of iC}_4\text{ through 3}^{rd}\text{ flowmeter)}$$

[0056] The computer system 124 can check that $V_iC_4 = V_iC_4_check$.

[0057] Having determined the concentration proportion of iC₄ in the sample volume, the computer system 124 can determine the concentration proportion of each of HF, water, and ASO in the HF mix, and also the concentration proportion of each of HF, water, and ASO in the sample volume using techniques described above with reference to FIG. 2A.

[0058] FIG. 2A is a flowchart of an example process 200 for determining concentrations of components of a volume sampled from a process stream. The sample volume that includes HF mix and iC₄ can be received through a first flowmeter (202). The HF mix can be separated from the iC₄ (204). The separation yields pure HF mix, which can be flowed through the conductivity sensor 118, the first flowmeter 108, and the pressure sensor 120 (206). The iC₄ can be flowed through the third flowmeter 110 (208). The mass flow rate through the second flowmeter 106 (M₁) can be compared with the mass flow rate through the first flowmeter 108 and the third flowmeter 110 (M₂ + M₃) (210). It can be checked if M₁ > M₂ + M₃ or if M₁ < M₂ + M₃ (212). If lesser, then the opening of the valve through which iC₄ flows through the third flowmeter (110) can be increased further (214). If greater, then the valve opening can be decreased further (216) to maintain a balance in the mass of iC₄ through the third flowmeter 110. The conductivity of the pure HF mix can be measured (step 218), for example, by the conductivity sensor. The conductivity sensor can then transmit the measured conductivity to the computer system 124. The concentration proportion of water in the HF mix (i.e., percentage of water in the HF mix) can be determined based on the conductivity (2220), for example, by the computer system 124. Mass flow and density of the fluid can be measured (222), for example, by the first flowmeter 108. The concentration proportion of either HF or ASO (i.e., percentage of either HF or ASO in the HF mix) can be determined based on the measured density (224), for example, by the computer system 124 that receives the mass flow and density measurements from the first flowmeter 108. The concentration of the remaining component (i.e., percentage of either ASO or HF in the HF mix), can be determined as a difference, from 100%, of the sum of the concentration of water and either HF or ASO (226). The 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.

[0059] 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 valve control is not employed. Similarly to FIG. 2A, a sample volume that includes HF mix and iC_4 can be received through a first flowmeter. The HF mix and the iC_4 can be separated using the separation vessel. The pure HF mix can be flowed through the second flowmeter. The partially separated iC_4 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.

[0060] 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 iC_4 (238). A volumetric proportion of iC_4 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 iC_4 as a product of average volumetric flow rate through the second flowmeter and the volumetric proportion of iC_4 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 iC_4 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 iC_4 through the second flowmeter can be compared with the volumetric proportion of iC_4 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 iC_4 in the sample volume and a concentration of HF, ASO, and water in the HF mix (250).

[0061] FIG. 3 is a flowchart of an example process 300 for calibrating three flowmeters. In some implementations, the system 100 can cross-calibrate the first, second, and third flowmeters (106, 108, and 110, respectively) to ensure an accurate determination of the flow rate of iC_4 . 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 (described above) 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.

[0062] 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 and 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.

[0063] Examples of flowmeters that can be implemented in system 100, are described below. Types of flowmeters include digital flowmeters. For example, U.S. Patent No. 6,311,136 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. Patent 6,505,519 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).

[0064] 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.

[0065] 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 storage substrate, a random or serial access memory array or device, or a combination of one or more of them. Moreover, while a computer storage medium is not a propagated signal, a computer storage medium can be a source or destination of computer program instructions encoded in an artificially-generated propagated signal. The computer storage medium can also be, or be included in, one or more separate physical components or media (for example, multiple CDs, disks, or other storage devices).

[0066] 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.

[0067] A computer program (also known as a program, software, software application, script, or code) 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 in a portion of a file 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.

[0068] 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).

[0069] 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), to name just a few. 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. The processor and the memory can be supplemented by, or incorporated in, special purpose logic circuitry.

[0070] To provide for interaction with a user, implementations of the computer system 124 can include a display device, 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.

[0071] While this specification contains many specific implementation details, these should not be construed as limitations on the scope of any inventions 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.

[0072] 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.

[0073] 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.

[0074] Embodiments of the invention are disclosed in claim 1. Preferred embodiments are disclosed in the dependent claims.

[0075] According to the invention the operations of a system according to the invention may 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 mixture.

[0076] According to the invention the acid can be Hydrofluoric acid.

[0077] According to the invention the first flowmeter can be a Coriolis flowmeter using a bent tube.

[0078] According to the invention the second flowmeter can be a Coriolis flowmeter using a straight tube.

[0079] According to the invention, the hydrocarbon fluid can include a first component, a second component, and a third component, and the operations of the system can further comprise determining a concentration of each of the first component, the second component, and the third component in the hydrocarbon fluid. Preferably the operations of the system can 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 hydrocarbon fluid and the concentration of the acid catalyst mix in the fluid flow.

Claims

1. Acid catalyst analyzer system comprising:

a first flowmeter configured to receive a sample volume that includes an acid catalyst mix and a hydrocarbon fluid, the acid catalyst mix including an acid, water, and acid soluble oils (ASO);

a separation vessel configured to receive the sample volume from the first flow meter, and to separate the acid catalyst mix from the hydrocarbon gas;

a second flowmeter adapted to receive the hydrocarbon fluid that has been separated from the acid catalyst mix, through a first valve;

a third flowmeter adapted to receive the acid catalyst mix that has been separated from the hydrocarbon fluid through a second valve; and

data processing apparatus operatively coupled to the first flowmeter 108, the second flowmeter, and the third flow meter, the data processing apparatus configured to execute computer software instructions stored on a computer-readable medium to perform operations comprising:

receiving a first set of mass flow and density readings, a second set of mass flow and density readings, and a third set of mass flow and density readings from the first flowmeter, the second flowmeter, and the third flowmeter, respectively,
determining a concentration of the hydrocarbon fluid in the sample volume, and
transmitting a signal to control a level by which the first valve is open to maintain a mass balance between the first flowmeter, and the second and the third flowmeters.

2. System according to claim 1, the operations further comprising:

determining a first mass flow through the first flowmeter based on the first set of mass flow and density readings;
determining a second mass flow through the second flowmeter based on the second set of mass flow and density readings;
determining a third mass flow through the third flowmeter based on the third set of mass flow and density readings;
comparing the first mass flow to a sum of the second mass flow and the third mass flow;
upon determining that the first mass flow is less than the sum, increasing the level by which the first valve is open, or
upon determining that the first mass flow is greater than the sum, decreasing the level by which the first valve is open.

3. System according to claim 1 or 2, further comprising a conductivity sensor coupled to the second flowmeter to receive the acid catalyst mix that has been separated from the hydrocarbon fluid, the conductivity sensor configured to measure a conductivity of the acid catalyst mix, wherein the conductivity sensor is operatively coupled to the data processing apparatus to transmit the measured conductivity to the data processing apparatus.

4. System according to any of claims 1 to 3, further comprising a pressure sensor coupled to the second flowmeter to receive the acid catalyst mix that has been separated from the hydrocarbon fluid, the pressure sensor configured to measure a pressure of the acid catalyst mix, wherein the pressure sensor is operatively coupled to the data processing apparatus to transmit the measured pressure to the data processing apparatus.

5. System according to claim 4, the operations further comprising:

determining a concentration of the water in the acid catalyst mix based on the received conductivity, wherein the concentration of the water is proportional to the measured conductivity;
determining a concentration of the acid in the acid catalyst mix based on the received third set of mass flow and density readings from the third flowmeter, wherein the concentration of the acid is proportional to the density readings in the third set; and
determining a concentration of the ASO as a remainder of a sum of the concentration of the water and the concentration of the acid from 100%.

6. System according to any of the preceding claims, wherein the operations further comprise determining the concentration of each of the water, the acid, and the ASO in the sample volume based on the concentration of the hydrocarbon fluid in the sample volume and the concentration of each of the water, the acid, and the ASO in the acid catalyst mix.

7. System according to any of the preceding claims, 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.

8. System according to any of the preceding claims, wherein the separation vessel includes a first inlet to receive the sample volume, the separation vessel sized to separate the acid catalyst mix and the hydrocarbon fluid based on a difference in a density of the acid catalyst mix and a density of the hydrocarbon fluid.

9. System according to any of the preceding claims, wherein the acid is Hydrofluoric acid.

10. System according to any of the preceding claims, wherein the first flowmeter is a Coriolis flowmeter using a bent tube.

11. System according to any of the preceding claims, wherein the second flowmeter is a Coriolis flowmeter using a straight tube.

Patentansprüche**1. System zur Analyse eines sauren Katalysators, umfassend:**

einen ersten Durchflussmesser, der so konfiguriert ist, dass er ein Probenvolumen aufnimmt, das ein saures Katalysatorgemisch und ein Kohlenwasserstofffluid enthält, wobei das saure Katalysatorgemisch eine Säure, Wasser und säurelösliche Öle (ASO) enthält;
 ein Trenngefäß, das so konfiguriert ist, dass es das Probenvolumen vom ersten Durchflussmesser aufnimmt und das saure Katalysatorgemisch vom Kohlenwasserstoffgas trennt;
 einen zweiten Durchflussmesser, der dazu geeignet ist, das Kohlenwasserstofffluid, das vom sauren Katalysatorgemisch getrennt wurde, über ein erstes Ventil aufzunehmen;
 einen dritten Durchflussmesser, der dazu geeignet ist, das saure Katalysatorgemisch, das vom Kohlenwasserstofffluid getrennt wurde, über ein zweites Ventil aufzunehmen; und
 eine Datenverarbeitungsvorrichtung, die funktionsfähig mit dem ersten Durchflussmesser 108, dem zweiten Durchflussmesser und dem dritten Durchflussmesser gekoppelt ist, wobei die Datenverarbeitungsvorrichtung so konfiguriert ist, dass sie Computersoftwarebefehle ausführt, die auf einem computerlesbaren Medium gespeichert sind, um Vorgänge auszuführen, umfassend:

Empfangen eines ersten Satzes von Massenstrom- und Dichtemesswerten, eines zweiten Satzes von Massenstrom- und Dichtemesswerten und eines dritten Satzes von Massenstrom- und Dichtemesswerten von dem ersten Durchflussmesser, dem zweiten Durchflussmesser bzw. dem dritten Durchflussmesser, Bestimmen einer Konzentration des Kohlenwasserstofffluids im Probenvolumen und Übertragen eines Signals zum Steuern eines Ausmaßes, mit dem das erste Ventil geöffnet ist, um einen Massenausgleich zwischen dem ersten Durchflussmesser und dem zweiten und dem dritten Durchflussmesser aufrechtzuerhalten.

2. System nach Anspruch 1, wobei die Vorgänge weiter umfassen:

Bestimmen eines ersten Massenstroms durch den ersten Durchflussmesser auf der Grundlage des ersten Satzes von Massenstrom- und Dichtemesswerten;
 Bestimmen eines zweiten Massenstroms durch den zweiten Durchflussmesser auf der Grundlage des zweiten Satzes von Massenstrom- und Dichtemesswerten;
 Bestimmen eines dritten Massenstroms durch den dritten Durchflussmesser auf der Grundlage des dritten Satzes von Massenstrom- und Dichtemesswerten;
 Vergleichen des ersten Massenstroms mit einer Summe aus dem zweiten Massenstrom und dem dritten Massenstrom;
 wenn festgestellt wird, dass der erste Massenstrom kleiner als die Summe ist, Erhöhen des Ausmaßes, mit dem das erste Ventil geöffnet ist, oder
 wenn festgestellt wird, dass der erste Massenstrom größer als die Summe ist, Reduzieren des Ausmaßes, mit dem das erste Ventil geöffnet ist.

3. System nach Anspruch 1 oder 2, weiter umfassend einen Leitfähigkeitssensor, der mit dem zweiten Durchflussmesser gekoppelt ist, um das saure Katalysatorgemisch aufzunehmen, das vom Kohlenwasserstofffluid getrennt wurde, wobei der Leitfähigkeitssensor so konfiguriert ist, dass er eine Leitfähigkeit des sauren Katalysatorgemischs misst, wobei der Leitfähigkeitssensor funktionsfähig mit der Datenverarbeitungsvorrichtung gekoppelt ist, um die gemessene Leitfähigkeit an die Datenverarbeitungsvorrichtung zu übertragen.

4. System nach einem der Ansprüche 1 bis 3, weiter umfassend einen Drucksensor, der mit dem zweiten Durchflussmesser gekoppelt ist, um das saure Katalysatorgemisch aufzunehmen, das vom Kohlenwasserstofffluid getrennt wurde, wobei der Drucksensor so konfiguriert ist, dass er einen Druck des sauren Katalysatorgemischs misst, wobei der Drucksensor mit der Datenverarbeitungsvorrichtung funktionsfähig gekoppelt ist, um den gemessenen Druck an die Datenverarbeitungsvorrichtung zu übertragen.

5. System nach Anspruch 4, wobei die Vorgänge weiterhin umfassen:

Bestimmen einer Konzentration des Wassers im sauren Katalysatorgemisch auf der Grundlage der empfangenen Leitfähigkeit, wobei die Konzentration des Wassers proportional zur gemessenen Leitfähigkeit ist;
 Bestimmen einer Konzentration der Säure im sauren Katalysatorgemisch auf der Grundlage des empfangenen

ritten Satzes von Massenstrom- und Dichtemesswerten vom dritten Durchflussmesser, wobei die Konzentration der Säure proportional zu den Dichtemesswerten im dritten Satz ist; und
Bestimmen einer Konzentration der ASO als Rest einer Summe aus der Konzentration des Wassers und der Konzentration der Säure aus 100 %.

6. System nach einem der vorhergehenden Ansprüche, wobei die Vorgänge weiter das Bestimmen der Konzentration des Wassers, der Säure und der ASO im Probenvolumen auf der Grundlage der Konzentration des Kohlenwasserstofffluids im Probenvolumen und der Konzentration des Wassers, der Säure und der ASO im sauren Katalysatorgemisch umfassen.
7. System nach einem der vorhergehenden Ansprüche, weiter umfassend ein Ventil, das mit dem ersten Durchflussmesser, dem zweiten Durchflussmesser und dem dritten Durchflussmesser gekoppelt und so konfiguriert ist, dass es den Durchfluss des vom ersten Durchflussmesser aufgenommenen Probenvolumens nur zum zweiten Durchflussmesser oder nur zum dritten Durchflussmesser oder zu beiden Durchflussmessern gleichzeitig ermöglicht.
8. System nach einem der vorhergehenden Ansprüche, wobei das Trenngefäß einen ersten Einlass zum Aufnehmen des Probenvolumens aufweist, wobei das Trenngefäß so bemessen ist, dass es das saure Katalysatorgemisch und das Kohlenwasserstofffluid auf der Grundlage einer Differenz hinsichtlich einer Dichte des sauren Katalysatorgemischs und einer Dichte des Kohlenwasserstofffluids trennt.
9. System nach einem der vorhergehenden Ansprüche, wobei die Säure Fluorwasserstoffsäure ist.
10. System nach einem der vorhergehenden Ansprüche, wobei der erste Durchflussmesser ein Coriolis-Durchflussmesser mit einem gebogenen Rohr ist.
11. System nach einem der vorhergehenden Ansprüche, wobei der zweite Durchflussmesser ein Coriolis-Durchflussmesser mit einem geraden Rohr ist.

Revendications

1. Système analyseur à catalyseur acide comprenant :

un premier débitmètre configuré pour recevoir un volume échantillon qui inclut un mélange catalyseur acide et un fluide hydrocarbure, le mélange catalyseur acide incluant un acide, de l'eau, et des huiles solubles dans l'acide (HSA) ;

un récipient de séparation configuré pour recevoir le volume échantillon depuis le premier débitmètre, et pour séparer le mélange catalyseur acide depuis le gaz hydrocarbure ;

un second débitmètre adapté pour recevoir le fluide hydrocarbure qui a été séparé du mélange catalyseur acide, via une première valve ;

un troisième débitmètre adapté pour recevoir le mélange catalyseur acide qui a été séparé du fluide hydrocarbure via une seconde valve ; et

un appareil de traitement de données fonctionnellement couplé au premier débitmètre, au second débitmètre, et au troisième débitmètre, l'appareil de traitement de données étant configuré pour exécuter des instructions d'un logiciel informatique stocké sur un support lisible à l'ordinateur afin d'effectuer des opérations comprenant :

la réception d'un premier groupe de lectures de débit massique et de densité, d'un second groupe de lectures de débit massique et de densité, et d'un troisième groupe de lectures de débit massique et de densité depuis le premier débitmètre, le second débitmètre, et le troisième débitmètre, respectivement,

la détermination d'une concentration du fluide hydrocarbure dans le volume échantillon, et la transmission d'un signal pour commander un niveau duquel la première valve est ouverte afin de maintenir un équilibre de masse entre le premier débitmètre, et le second et le troisième débitmètre.

2. Système selon la revendication 1, dans lequel le fonctionnement comprend en outre :

la détermination d'un premier débit massique à travers le premier débitmètre sur la base du premier groupe de lectures de débit massique et de densité ;

la détermination d'un second débit massique à travers le second débitmètre sur la base du second groupe de

lectures de débit massique et de densité ;

la détermination d'un troisième débit massique à travers le troisième débitmètre sur la base du troisième groupe de lectures de débit massique et de densité ;

la comparaison du premier débit massique à une somme du second débit massique du troisième débit massique ;
suite à la détermination que le premier débit massique est inférieur à la somme, l'augmentation du niveau duquel la première valve est ouverte, ou suite à la détermination que le premier débit massique est supérieur à la somme, la diminution du niveau duquel la première valve est ouverte.

3. Système selon la revendication 1 ou 2, comprenant en outre un capteur de conductivité couplé au second débitmètre pour recevoir le mélange catalyseur acide qui a été séparé du fluide hydrocarbure, le capteur de conductivité étant configuré pour mesurer une conductivité du mélange catalyseur acide, dans lequel le capteur de conductivité est fonctionnellement couplé à l'appareil de traitement de données pour transmettre la conductivité mesurée à l'appareil de traitement de données.

4. Système selon l'une quelconque des revendications 1 à 3, comprenant en outre un capteur de pression couplé au second débitmètre pour recevoir le mélange catalyseur acide qui a été séparé du fluide hydrocarbure, le capteur de pression étant configuré pour mesurer une pression du mélange catalyseur acide, et le capteur de pression étant fonctionnellement couplé à l'appareil de traitement de données pour transmettre la pression mesurée à l'appareil de traitement de données.

5. Système selon la revendication 4, dont le fonctionnement comprend en outre :

la détermination d'une concentration de l'eau dans le mélange catalyseur acide sur la base de la conductivité reçue, dans lequel la concentration de l'eau est proportionnelle à la conductivité mesurée ;

la détermination d'une concentration de l'acide dans le mélange catalyseur acide sur la base du troisième groupe reçu de lectures de débit massique et de densité depuis le troisième débitmètre, dans lequel la concentration de l'acide est proportionnelle aux lectures de densité dans le troisième groupe ; et

la détermination d'une concentration des HSA à titre de reste d'une somme de la concentration de l'eau et de la concentration de l'acide par rapport à 100 %.

6. Système selon l'une quelconque des revendications précédentes, dans lequel le fonctionnement comprend en outre la détermination de la concentration de chacun de l'eau, de l'acide et des HSA dans le volume échantillon sur la base de la concentration du fluide hydrocarbure dans le volume échantillon et de la concentration de chacun de l'eau, de l'acide et des HSA dans le mélange catalyseur acide.

7. Système selon l'une quelconque des revendications précédentes, comprenant en outre une valve couplée au premier débitmètre, au second débitmètre, et au troisième débitmètre, et configurée pour permettre l'écoulement du volume échantillon reçu par le premier débitmètre vers le second débitmètre uniquement ou vers le troisième débitmètre uniquement ou encore vers les deux débitmètres simultanément.

8. Système selon l'une quelconque des revendications précédentes, dans lequel le récipient de séparation inclut une première entrée pour recevoir le volume échantillon, le récipient de séparation ayant une taille propre à séparer le mélange catalyseur acide et le fluide hydrocarbure sur la base d'une différence d'une densité du mélange catalyseur acide et d'une densité du fluide hydrocarbure.

9. Système selon l'une quelconque des revendications précédentes, dans lequel l'acide est de l'acide fluorhydrique.

10. Système selon l'une quelconque des revendications précédentes, dans lequel le premier débitmètre est un débitmètre de Coriolis utilisant un tube cintré.

11. Système selon l'une quelconque des revendications précédentes, dans lequel le second débitmètre est un débitmètre de Coriolis utilisant un tube droit.

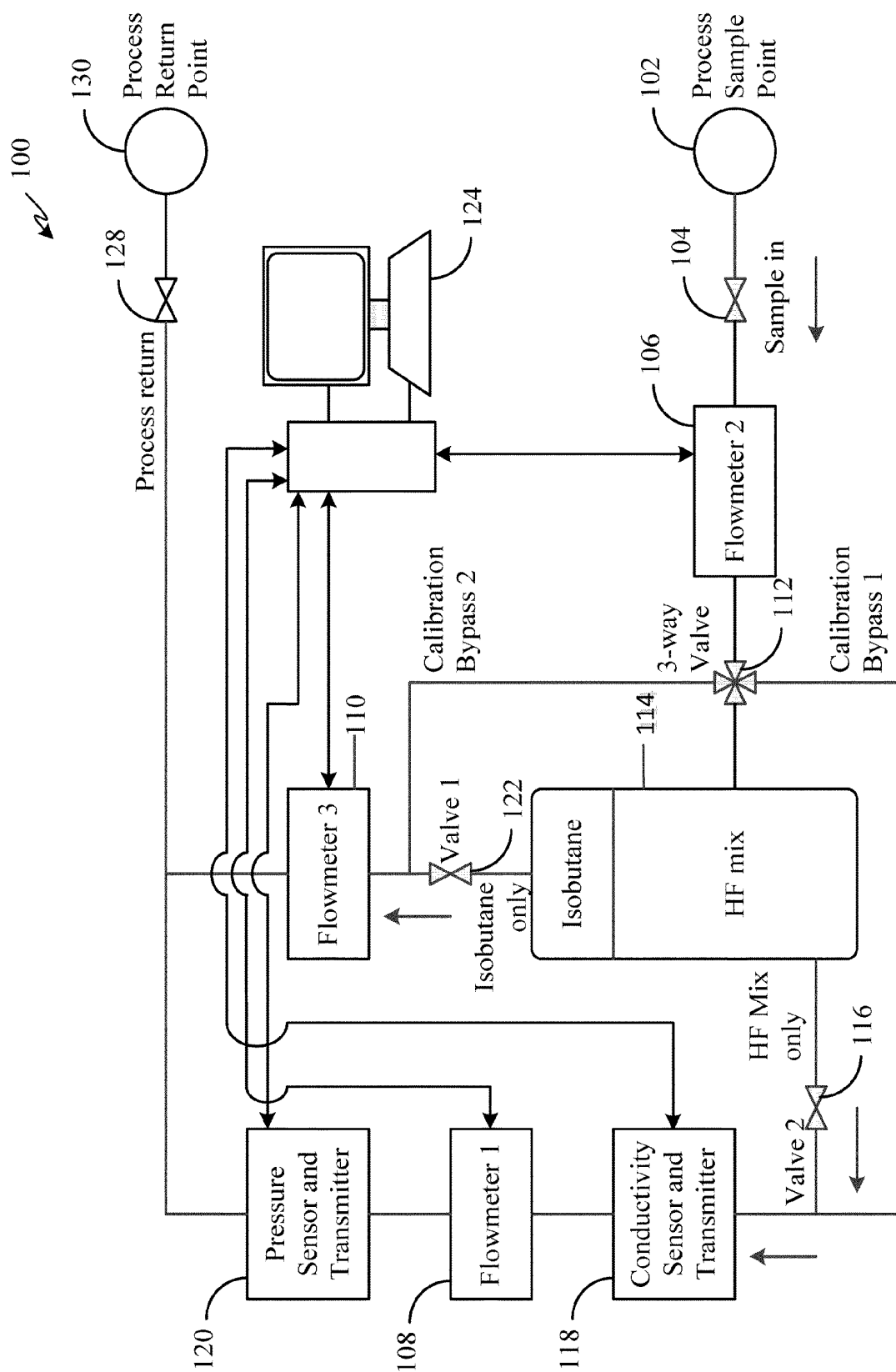


FIG. 1

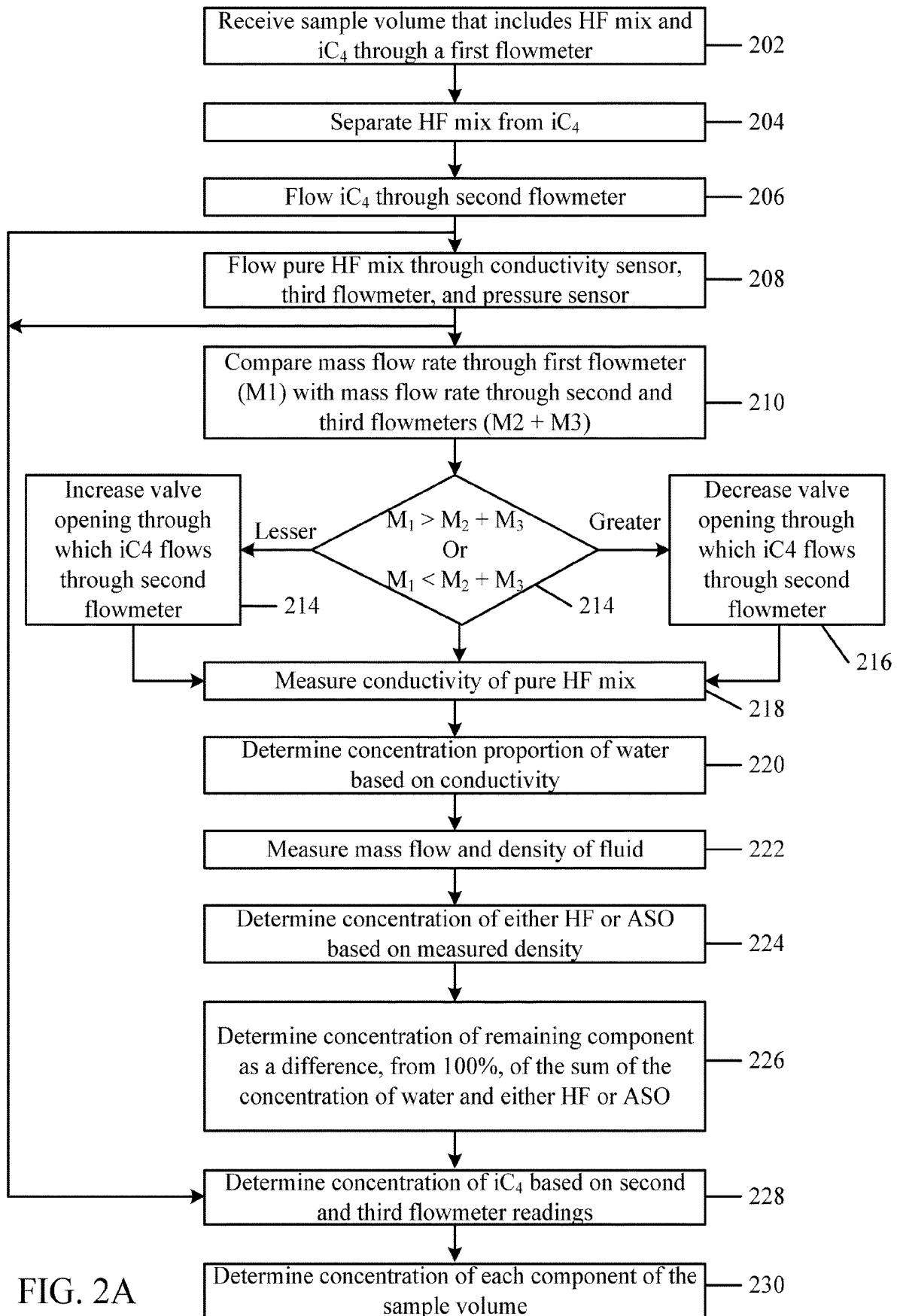


FIG. 2A

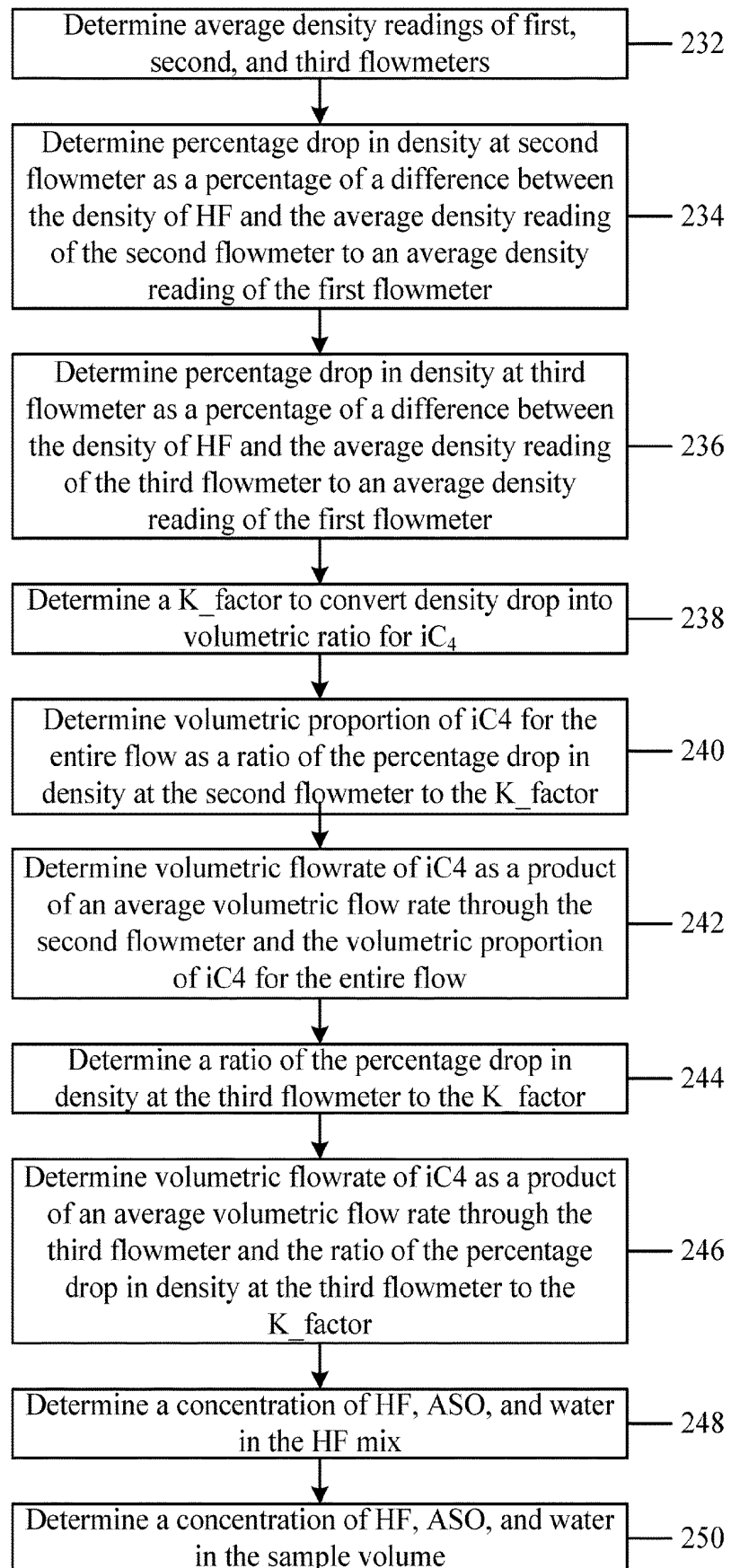


FIG. 2B

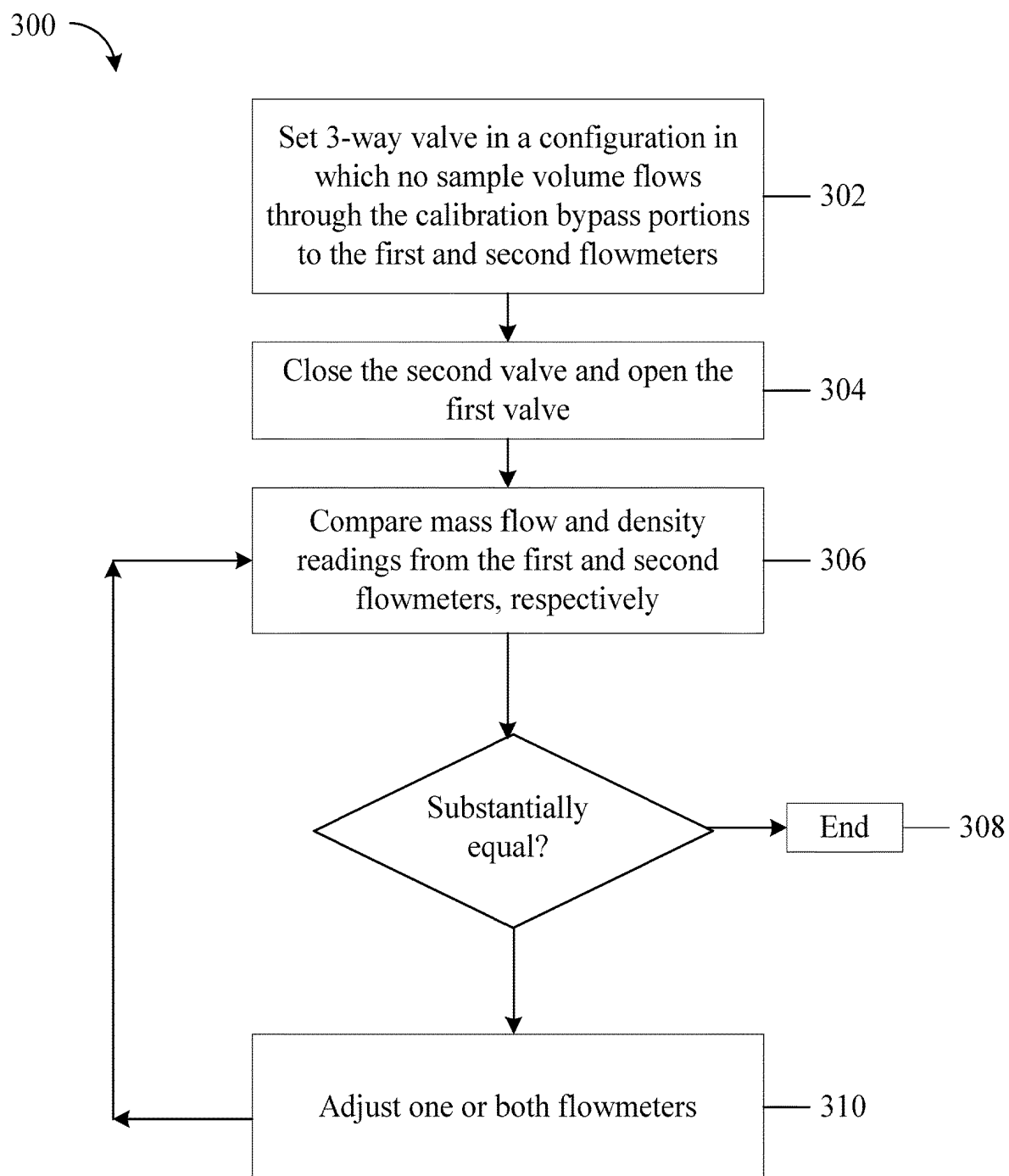


FIG. 3

REFERENCES CITED IN THE DESCRIPTION

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