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SYSTEM AND METHOD FOR MULTIPHASE FLOW METERING ACCOUNTING FOR DISSOLVED GAS

SYSTEM UND VERFAHREN ZUR MEHRPHASENDURCHFLUSSMESSUNG UNTER BERÜCKSICHTIGUNG VON GELÖSTEM GAS

SYSTÈME ET PROCÉDÉ DE MESURE DE DÉBIT MULTIPHASE PRISE EN COMPTE DU GAZ DISSOLU

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FIELD OF INVENTION

[0001] The present invention relates generally to flowmeters and more specifically to Coriolis flowmeters that are adapted to measure multiphase fluids.

BACKGROUND

[0002] Various different flowmeters are used in industry to provide information about the flow rate of multiphase fluids. The fluids that are metered can include mixtures of liquids and gases. This situation is commonly encountered in the oil and gas industry, where the fluids produced are commonly a mixture of oil, water, and gas. However, the need to meter multiphase fluids also occurs in other industries as well.

[0003] One type of flowmeter is a Coriolis flowmeter. A Coriolis flowmeter includes an electronic transmitter and a vibratable flowtube through which fluid to be metered can be passed. The transmitter maintains flowtube vibration by sending a drive signal to one or more drivers and performs measurement calculations based on signals from a pair of sensors that measure movement of the flowtube. The physics of the device dictate that Coriolis forces act along a section of the flowtube between the sensors, resulting in a phase difference between the generally sinusoidal sensor signals. This phase difference is generally proportional to the mass flow rate of the fluid passing through the measurement section of the flowtube. Thus, the phase difference provides a basis for a mass flow measurement of fluid flowing through the flowtube. The frequency of oscillation of the flowtube of a Coriolis meter varies with the density of the process fluid in the flowtube. The frequency value can be extracted from the sensor signals (for example by calculating the time delay between consecutive zero crossings) so that the density of the fluid can also be obtained by analyzing the sensor signals.

[0004] Coriolis meters are widely used throughout various different industries. The direct measurement of mass flow is frequently preferred over volumetric-based metering, for whereas the density and/or volume of a material may vary with temperature and/or pressure, mass is unaffected. This is particularly important in the oil and gas industry, where energy content and hence product value is a function of mass. The term 'Net Oil' is used in the oil and gas industry to describe the oil flow rate within a three-phase or a liquid (oil/water) stream. A common objective in the oil and gas industry is to determine the net oil produced by each well in a plurality of wells because this information can be important when making decisions affecting production from an oil and gas field and/or for optimizing production from an oil and gas field.

[0005] The inclusion of gas in a liquid stream introduces errors in the mass flow and density measurements of a Coriolis meter. Laboratory trials can be used to characterize how mass flow rate and density errors relate to other parameters, such as the observed flow rate and observed reduction in density from that of the pure fluid. These trials can be used to develop empirical models that provide corrections to account for some of the error associated with the presence of multiphase fluids including gas and liquid phases. These empirically-based corrections can result in improved performance of Coriolis meters in field operations. Additional details concerning use of a Coriolis meter to meter multiphase fluids are provided in U.S. Patent Nos. 6,311,136; 6,505,519; 6,950,760; 7,059,199; 7,313,488; 7,617,055; 7,188,534; 7,614,312; 7,660,681 and 8,892,371 as well as document WO 2012/027409 A2.

[0006] The present inventors have made various improvements, which will be described in detail below, applicable to the field of Coriolis flowmeters and applicable to the field of net oil and gas testing.

SUMMARY

[0007] One aspect of the invention is a system for metering flow of a fluid as specified in independent claim 1.
[0008] Another aspect of the invention is a net oil and gas skid as specified in independent claim 9.
[0009] Still another aspect of the invention is a method of metering flow of a multiphase fluid as specified in independent claim 12.

BRIEF DESCRIPTION OF THE DRAWINGS

[0010] FIG. 1 is a perspective of one embodiment of a Coriolis flowmeter;
FIG. 2 is a side elevation of the Coriolis flowmeter shown in Fig. 1;
FIG. 3 is a schematic diagram of one embodiment of a Coriolis flowmeter processor showing its inputs and outputs;
FIG. 4 is a schematic diagram of one embodiment of a system for generating empirical data associated with multiphase flow through the Coriolis meter;
FIG. 5 is a schematic diagram of another embodiment of a system for generating empirical data associated with
multiphase flow through the Coriolis meter;
FIG. 6 is a chart plotting the drive gain vs. nominal gas volume fraction (GVF) for a set of data from a Coriolis meter while metering a mixture of gas and water;
FIG. 7 is a chart plotting density drop vs. nominal GVF for a set of data from a Coriolis meter while metering a mixture of gas and water;
FIG. 8 is a chart plotting mass flow rate error vs. nominal gas volume fraction (GVF) for a set of data from a Coriolis meter while metering a mixture of gas and water;
FIG. 9 is a chart plotting density error vs. nominal GVF for a set of data from a Coriolis meter metering a mixture of gas and water;
FIG. 10 is a chart plotting the drive gain vs. nominal gas volume fraction (GVF) for a set of data from a Coriolis meter while metering a mixture of gas and oil that causes the actual GVF to differ significantly from the nominal GVF;
FIG. 11 is a chart plotting density drop vs. nominal GVF for a set of data from a Coriolis meter while metering a mixture of gas and oil that causes the actual GVF to differ significantly from the nominal GVF;
FIG. 12 is a chart plotting mass flow rate error vs. nominal gas volume fraction (GVF) for a set of data from a Coriolis meter while metering a mixture of gas and oil that causes the actual GVF to differ significantly from the nominal GVF;
FIG. 13 is a chart plotting density error vs. nominal GVF for a set of data from a Coriolis meter metering a mixture of gas and oil that causes the actual GVF to differ significantly from the nominal GVF;
FIG. 14 is a schematic diagram illustrating one embodiment of system for obtaining multiphase flow rate and density corrections for a Coriolis meter;
FIG. 15 is a schematic diagram of one embodiment of a method of training and using a neural network to obtain multiphase flow rate and density corrections for a Coriolis meter;
FIG. 16 is a schematic diagram illustrating another embodiment of a system for obtaining multiphase flow rate and density corrections for a Coriolis meter;
FIG. 17 is a schematic diagram illustrating another method of training and using a neural network to obtain multiphase flow rate and density corrections for a Coriolis meter;
FIG. 18 is a chart showing a relationship between pressure, temperature, and the solution gas - oil ration (Rs);
FIG. 19 is a chart showing a relationship between pressure, temperature, and the formation volume factor (Bo);
FIG. 20 is a front elevation of one embodiment of a net oil and gas skid including the Coriolis flowmeter of Figs. 1 and 2;
FIG. 21 is a side elevation of the net oil and gas skid shown in Fig. 20;
FIG. 22 is a schematic diagram of one embodiment of a flow conditioner in the net oil and gas skid shown in Figs. 20 and 21; and
FIG. 23 is a schematic diagram illustrating one embodiment including the net oil and gas skid illustrated in Figs. 20 and 21 in combination with a compact separation system to facilitate metering fluids with relatively high GVF.

[0011] Corresponding reference numbers indicate corresponding parts through the drawings.

DETAILED DESCRIPTION

[0012] One embodiment of a Coriolis flowmeter, generally designated 215, is illustrated in FIGS. 1 and 2. The flowmeter 215 includes one or more conduits 18, 20 (also referred to as a flowtube), one or more drivers 46a, 46b for driving oscillation of the conduit(s), and a pair of motion sensors 48a, 48b that generate signals indicative of the oscillation of the conduit(s). In the illustrated embodiment, there are two conduits 18, 20 two drivers 46a, 46b and two motion sensors 48a, 48b and the drivers and motions sensors are positioned between the conduits so each driver and motion sensor is operable for both of the conduits. It is understood however that a Coriolis flowmeter may have only a single conduit and/or may have a single driver. It is also understood the conduit(s) may have different configurations than the conduits 18, 20 in the illustrated embodiment.

[0013] As illustrated in FIGS. 1 and 2, the flowmeter 215 is designed to be inserted in a pipeline (not shown) having a small section removed or reserved to make room for the flowmeter. The flowmeter 215 includes mounting flanges 12 for connection to the pipeline, and a central manifold block 16 supporting the two parallel planar loops 18 and 20 which are oriented perpendicularly to the pipeline. The drivers 46a, 46b and sensors 48a, 48b are attached between each end of loops 18 and 20. The drivers 46a, 46b on opposite ends of the loops 18, 20 are energized by a digital controller (not shown) with current of equal magnitude but opposite sign (i.e., currents that are 180° out-of-phase) to cause straight sections 26 of the loops 18, 20 to rotate about their co-planar perpendicular bisector 56 (FIG. 2). Repeatedly reversing (e.g., controlling sinusoidally) the energizing current supplied to the drivers 46a, 46b causes each straight section 26 to undergo oscillatory motion that sweeps out a bow tie shape in the horizontal plane about the axis 56 of symmetry of the loops. The entire lateral excursion of the loops 18, 20 at the lower rounded turns 38 and 40 is small, on the order of 1/16 of an inch for a two foot long straight section 26 of a pipe having a one inch diameter. The frequency of oscillation is typically about 80 to 90 Hertz, although this can vary depending on the size and configuration of the flowtube(s).
[0014] The sensors 48a, 48b are positioned to detect movement of the flowtube at different locations on the flowtube and output sensor signals indicative of the detected movement. As will be understood by those skilled in the art, the Coriolis effect induces a phase difference between the two sensors 48a, 48b that is generally proportional to mass flow rate. Also, the resonant frequency of the loops 18, 20 will vary as a function of density of the fluid flowing therethrough. Thus, the mass flow rate and density can be measured by analyzing the signals from the sensors 48a, 48b. The Coriolis meter 215 has a processor 101 (Fig. 3) configured to receive the sensor signals from the sensors 48a, 48b, determine a phase difference between the sensor signals, and use the determined phase difference to determine a fluid flow rate through the flowtube. The processor 101 is also configured to determine a frequency of one or more of the sensor signals and use the determined frequency to determine a density of the fluid in the flowtube.

[0015] Various corrections can be applied to the basic measurements resulting from the phase difference between the signals from the sensors 48a, 48b and the frequency. For example, multiphase flow introduces highly variable damping on the flowtube, up to three orders of magnitude higher than in single phase conditions. In addition, the mass flow and density measurements generated under multiphase flow conditions are subject to large systematic and random errors, for which correction algorithms can be defined and implemented by the processor 101.

[0016] As will be described in more detail below, the processor 101 is suitably configured to apply corrections that account for the influence of dissolved gas on the measurements from the Coriolis meter. Moreover, the processor 101 is suitably configured to determine the amount of dissolved gas in the fluid flowing through the flowtube. The ability to account for the effects of dissolved gas and/or measure the amount of dissolved gas can be important in many different types of industries.

[0017] One important industry in which dissolved gas is a concern is the oil and gas industry. Typically, high pressure within an oil reservoir results in much or all of the gas being dissolved within the oil phase. Above the bubble point pressure all of the gas is dissolved. As pressure of the petroleum fluid is reduced with each stage of the extraction process, the gas phase is increasingly released from the oil phase to become free gas. However, the oil phase typically remains saturated with dissolved gas, the quantity of which varies with the pressure and temperature conditions as well as the miscibility of the particular oil and gas concerned. Thus, there is a mass exchange between the free gas phase and the dissolved gas phase as temperature and pressure conditions vary throughout the extraction process. The standard in the oil and gas industry is to report flow rates at standard temperature and pressure conditions in order to provide consistent measurements regardless of the operating conditions. At standard conditions, almost all gas is free gas and there is substantially no dissolved gas.

[0018] However, dissolved gas is largely invisible to Coriolis meters, to a first approximation. If a liquid with dissolved gas but no free gas is passed through a Coriolis meter, the following is observed:

(i) The density of the liquid including dissolved gas is close to that of the pure liquid with no dissolved gas. Standard oil industry models exist for predicting the change in density associated with dissolved gas, which is typically a reduction in value, and which is often referred to as the expansion factor. This change in liquid density is typically small compared to the density of the equivalent mixture of oil and free gas;

(ii) The mass flow of the liquid including dissolved gas is the sum of the mass flow rates for the liquid and gas flows, but at the pressures commonly encountered in the oil and gas industry, the mass of the dissolved gas is likely to be very low compared with the mass of liquid; and

(iii) The drive gain (which as used herein means the ratio between the drive current (amp) and the voltage (v) of sensor signals) required to maintain oscillation of the flowtube is an indication of the energy needed to maintain flowtube oscillation. The drive gain is low when pure liquids or pure gases flow through the flowtube, but much higher (typically by two orders of magnitude) when a multiphase fluid is in the flowtube. Even the smallest mixture of free gas with liquid can cause the drive gain to rise rapidly, which makes this a key measurement in detecting of presence of free gas. However, with dissolved gas present but with free gas absent, the drive gain shows no significant rise.

[0019] Thus, the mass flow and density measurements from the Coriolis meter are not significantly affected by the presence of dissolved gas because the errors in these measurements are small or negligible compared to those induced by a mixture of the same liquid with the equivalent volume of free gas. Accordingly, conventional Coriolis meters dismiss the effects of dissolved gas as negligible and do not attempt to determine the amount of dissolved gas in the fluid.

[0020] Although the presence of dissolved gas in and of itself does not influence the behavior of the Coriolis meter, dissolved gas is actually a source of significant error from Coriolis meters during multiphase flow. This is because corrections to account for the presence of multiple phases in a fluid metered by prior art Coriolis meters predominantly use empirical models. While these models make limited use of physical models such as correcting gas density for pressure and temperature, they do not account for dissolved gas. It has been determined that this can introduce substantial errors in the corrected gas flow rates when applied to conditions where there the solubility of gas in the oil phase is different from what is was when the model data was collected.

[0021] The Coriolis meter 215 described herein addresses this problem. The quantity of dissolved gas is modeled or
predicted from the known physical properties of the liquid and gas phases, together with the observed temperature and pressure conditions. A number of standard oil industry models provide estimates of gas solubility and the resulting change in the fluid density. One such example is provided in "Correlations for Fluid Physical Property Prediction", M.E. Vasquez and H.D. Beggs, JPT 968 - 70, June 1980. Another option is to use a black oil model which provides a PVT calculation based on bulk properties of the gas, oil and water phases. This model is practical for oil and gas production applications where the bulk fluid composition can change quickly due to slugging or changes in the operating conditions. A compositional model could be used instead and would be appropriate for applications with constant compositions or on-line compositional measurements, such as medical applications.

[0022] The influence of dissolved gas is taken into account during model development, which is typically done on the basis of data collected during laboratory trials. Here a physical model is developed to predict how much gas has dissolved, while only the residual free gas is used in the development of empirical corrections for multiphase behavior in the Coriolis meter and other meters (e.g. a water cut meter). As will be shown, this can produce multiphase models which yield corrections to mass flow and density that are more consistent. The second area where dissolved gas is taken into account is during real-time operation, where an appropriate model of dissolved gas is included so that the total gas flow (free + dissolved) can be estimated and reported by the meter.

[0023] Thus, one advantage of the Coriolis meter 215 described herein is that it can provide better flow rate and density measurements during multiphase flow including a miscible gas because of the improvements in its ability to provide corrections to the raw flow rate and raw density measurements during multiphase flow. Another advantage of the Coriolis meter 215 is that it can produce a measurement of dissolved gas, which can make it easier to report flow rates at standard conditions in the oil and gas industry and which can be desirable in other applications as well. It is not necessary for any particular meter to achieve both of these advantages. For example, a meter can provide the improved flow rate and density measurements using the technology described herein without determining and/or without outputting a measurement of dissolved gas.

Model for Multiphase Flow Corrections

[0024] A model is derived from empirical data to enable the Coriolis meter 215 to provide corrections to raw mass flow rate and raw density measurements to account for the effects multiphase flow on the meter. The model is developed by subjecting a Coriolis meter to a range of multiphase flow conditions in order to record and model its mass flow and density errors. The model will be described herein in reference to oil, water, and gas to provide a detailed example, but it is understood the same process can be used with different materials.

[0025] A three-phase oil, water, and gas mixture can be specified in a number of ways. One way is to specify the mass flows of the oil, water and gas components. Here the oil and water flows will be stated in kg/s, while the gas flow rate (at the low operating pressures considered here) will be stated in g/s. Another way to specify a three-phase mixture of oil, water, and gas is to specify the total liquid flow rate for oil and water (usually in kg/s), water cut (between 0 percent and 100 percent) and Gas Volume flow Fraction or GVF (from 0 percent up to 50 percent and beyond). When the density of each phase is known along with temperature and pressure conditions and there is no slip between phases it is straightforward to convert from one of these specifications to the other. The first description (mass flows of each phase) can be convenient when specifying the accuracy of final results, while the latter description (liquid flow/water cut/GVF) can be more convenient for describing experimental conditions, particularly during model development. A two-phase flow of gas and pure oil can be considered a specific case of three-phase flow where the water cut is zero. Similarly, a two-phase flow of gas with pure water can be considered as another specific case of three-phase flow where the water cut is 100 percent.

[0026] Figure 4 shows an experimental setup for developing empirical models of Coriolis meter mass flow rate and density errors during multiphase flow. The Coriolis meter under test is integrated into a net oil and gas skid 121, which includes the Coriolis meter 215 illustrated above, along with a pressure sensor positioned to measure a pressure of the fluid in the flowtubing, a temperature sensor positioned to measure a temperature of the fluid in the flowtube, and a water cut meter positioned to measure the water cut of the fluid. A separator system 131 provides separated water and oil streams, while a liquid nitrogen tank 133 is used to supply gas that is mixed with the liquids to make a multiphase flow. Each phase is metered separately by one of a set of single phase meters 141, before being combined and passed through the skid 121. For each experiment performed, steady conditions are established for the desired oil, water and gas flow rates, and then the single phase reference measurements from meters 141 are compared with the outputs of the skid 121, averaged over the duration of the experiment.

[0027] In a simpler alternative arrangement (Fig. 5), there is no separation of oil and water. The water cut is selected in an off-line procedure by replacing a calculated volume of the current oil/water mixture with either pure water or pure oil, depending on whether the water cut is to be increased or decreased. Only if the new water cut is to be 0% or 100% must the entire volume of liquid be replaced. Once the desired water cut has been established, then a series of experiments can be carried out with a range of flowrates and GVF settings, where the water cut is kept constant by continuous mixing.
of the storage vessel. When a new water cut is required, the off-line procedure is repeated.

Effects of Dissolved Gas in Model Development

[0028] In contrast to prior art models developed for Coriolis meters, the calculation of GVF, and of mass flow and density errors, do not rely on the assumption that no gas dissolves into the liquid mixture.

[0029] The interaction between the specific oil, water and gas used may result in a proportion of the gas being dissolved in the liquid mixture. However, the solubility of the gas may vary. For example, petroleum gas is more soluble in oil than it is in water, which means the water cut affects the gas solubility relative to the oil/water mixture. In the case of lower water cuts (e.g., pure oil continuous flow, where the mixture/emulsion consists of water droplets in oil) gas can go into solution more easily than it can in the case of higher water cuts (e.g., pure water continuous flow, where the mixture/emulsion consists of oil droplets in water). Little or no gas may go into solution in the event of very high water cuts.

[0030] The data in Figs. 6-13 contrasts the behavior of the same 80mm diameter Coriolis meter over very similar liquid mass flow and GVF conditions, with and without gas dissolving into the liquid. The pressure for the experiments generating this data was relatively low (300 kPa - 500 kPa absolute), which means the gas density can often be considered negligible because it is relatively small compared to the liquid density. The operating temperature was approximately 40 °C. It is known that the impact of multiphase mixtures on a Coriolis meter can be reduced where the fluid viscosity is high (the relative movement of gas and liquid within the flowtube is reduced). However, in this experiment the viscosity of the oil was too low to dampen the impact of multiphase flow on the Coriolis meter.

[0031] Figures 6-9 show, respectively, the drive gain, density drop, mass flow error and density error from a Coriolis meter during an experiment in which water cut was held steady at about 100 percent (i.e., no oil) and gas void fraction was varied by changing the amount of gas mixed with the water. In this experiment very little gas would be able to dissolve into the liquid because of the high water cut and the fact that this particular gas is substantially insoluble in water. In each case, the x-axis shows the nominal GVF of the multiphase mixture i.e. the percentage of gas by volume at the pressure and temperature conditions of the Coriolis flowtube, assuming that all the gas injected into the liquid stream remains free, without any gas being dissolved. The default assumption inherent in the prior art is that all gas injected into the liquid, as measured by the reference gas meter, remains as free gas in the multiphase mixture and therefore contributes towards the nominal GVF of the multiphase mixture. In Figs. 6-9 no distinction is shown between different liquid flow rates, water cuts, or other conditions under which the meter is operated. This data illustrates the expected behavior of a Coriolis meter when free gas is added to a liquid.

[0032] The drive gain (Fig. 6) is shown on a logarithmic scale, and indicates the amount of energy required to maintain flowtube oscillation. As various definitions of drive gain may be used, the expression applied here is as follows:

\[ \text{drive gain} = \frac{\text{drive current (amps)}}{\text{sensor voltage (v)}} \]

[0033] There is a considerable spread of drive gain values at zero nominal GVF. Some of this may be caused by marginal quantities of gas still present in the liquid after the reference gas supply has been cut off during a series of experiments. Nevertheless there is a clear gap between the drive gain values at zero and non-zero nominal GVF. With the exception of two points, all non-zero GVF points have higher drive gain than all the zero GVF points. The data suggests very roughly an order of magnitude (i.e. a factor of ten) increase in the drive gain with the onset of any free gas. More generally, while the exact values of the drive gain may vary significantly for different flowmeter designs, flowtube orientations and other factors, a substantial rise in drive gain is commonly observed when free gas mixed with liquid passes through the Coriolis meter.

[0034] The density drop is shown in Figure 7. Here the density drop is defined as follows:

\[ \text{density drop} = \left( \rho_L - \rho_g \right) / \rho_L \times 100 \text{ percent} \]

where \( \rho_L \) is the density of the liquid only (note that this may be a mixture of water and oil, depending upon the water cut) and \( \rho_g \) is the apparent or observed density generated by the Coriolis mass flow meter. Note that by convention the density drop is shown as positive when the observed density reading falls below that of the fluid density. As expected, the density drop at zero GVF is also zero, but as the GVF increases from zero the density drop rises quickly, indicating that the presence of free gas results in a drop in the observed density reading.

[0035] Figures 8 and 9 show the mass flow and density errors for the Coriolis meter, defined as follows:
density error = \((\rho_a - \rho_m) / \rho_m \times 100\) percent  
mass flow error = \((\dot{m}_a - \dot{m}_m) / \dot{m}_m \times 100\) percent m

where \(\rho_m\) is the calculated true density of the multiphase mixture, \(\dot{m}_m\) is the true mass flow of the multiphase mixture, and \(\dot{m}_a\) is the apparent mass flow rate from the Coriolis mass flow meter.

The mass flow and density errors show very broadly similar behavior, starting at zero for zero GVF and dropping rapidly for low but non-zero GVF, before spreading out to form a wider range of errors.

Figures 10-13 show the corresponding behavior for drive gain, density drop, mass flow error and density error for the same flowmeter over the same range of GVF and volumetric flow rate, but with low water cut (i.e., oil continuous flow), which results in some of the gas measured by the reference gas meter prior to injection being dissolved into the liquid before reaching the Coriolis meter. In each case the x-axis shows the nominal GVF, assuming all gas measured by the reference gas meter remains free and does not go into solution.

In Fig. 10, in comparison with Fig. 6, there is no longer a clear distinction between the drive gain values for zero GVF and slightly above-zero GVF. When the nominal GVF exceeds about 8 percent, however, the drive gain is consistently above the range of values observed at zero GVF. This indicates some of the gas must be going into solution, as the effect of free gas on the drive gain is always significant for low viscosity fluids. In particular, it appears that almost all of the gas goes into solution when the nominal GVF is about 8 percent or less.

In Fig. 11, (compare with Fig. 7), there is negligible density drop as the nominal GVF increases from 0 percent to about 5 percent. The density drop is not consistently above zero until about 10% nominal GVF. Because the density drop is based on deviation from the pure fluid density, this graph shows that for up to about 5 percent nominal GVF, the reported mixture density is the same as that of the pure liquid. This also indicates at least some of the gas is going into solution.

In Fig. 12, (compare with Fig. 8), the mass flow rate error is close to zero for low nominal GVFs up to about 5 percent, and only when the nominal GVF is in excess of about 15 percent is the corresponding mass flow error consistently less than zero. Again, this indicates some of the gas is going into solution and is not free gas when it flows through the Coriolis meter.

However, the greatest difference in behavior is shown in the density error (Fig. 13, compare with Fig. 9), where the density error for the liquid/gas mixture is consistently positive for nominal GVFs less than about 8 percent, and density errors are only consistently negative for nominal GVFs above about 20 percent. Clearly, if gas is being dissolved into the liquid this will result in a positive shift in the density measurement compared with a nominal mixture density where it is assumed that all the gas is free. Thus, here again, the results indicate some of the gas is dissolved as it flows through the Coriolis meter.

Overall, the data illustrated in Figs. 6-13 for this particular set of experiments show a clear difference in the response of the Coriolis meter to, on the one hand, a multiphase flow having near 100 percent water cut, where the gas is substantially immiscible with the liquid and any non-zero nominal GVF has an impact on the behavior of the Coriolis meter (such as increased drive gain and density drop, and negative mass flow and density errors), and on the other a multiphase flow near 0 percent water cut, where the gas is miscible, and where for nominal GVFs of up to about 5 percent there is limited or no impact on the density drop and mass flow error. In essence, the Coriolis meter simply fails to detect or respond to the presence of gas dissolved in the liquid. Further, the results indicate that the discrepancies between the nominal GVF and the actual GVF during the development of empirically-based mass flow rate and density corrections can introduce error into empirically-based mass flow rate and density corrections developed using experiments that do not address the issues associated with dissolved gas.

Accordingly, to address this issue the quantity of dissolved gas is modeled from the known physical properties of the liquid and gas phases, together with the observed temperature and pressure conditions. A number of oil industry models provide estimates of gas solubility and the resulting change in the fluid density, for example "Correlations for Fluid Physical Property Prediction", M.E. Vasquez and H.D. Beggs, JPT 968 - 70, June 1980. The results described herein are based on a black oil model which provides a PVT calculation based on bulk properties of the gas, oil and water phases. This model is practical for oil and gas production applications where the bulk fluid composition can change quickly due to slugging or changes in the operating conditions. A compositional model could be used with the same methodologies and would be appropriate for applications with constant compositions or on-line compositional measurements, such as medical applications.

Thus, instead of using nominal GVF when developing empirically-based mass flow rate and density corrections for a Coriolis meter, the level of free gas flowing through the Coriolis meter is calculated based on the reference measurement of total gas flow and the gas solubility model. Then the response of the Coriolis meter to free gas can be modeled better, which results in more accurate corrections for mass flow rate and density errors associated with free gas.

Empirical mass flow rate and density corrections for a Coriolis meter may be implemented in the form of a
mapping between a given set of inputs (e.g., phase difference between sensor signals, frequency of flowtube oscillation, temperature, pressure, drive gain, etc.) to a set of outputs (e.g., corrected mass flow rate and corrected density). For example, a neural network can be used. In general the inputs to a neural network are based on data available to the metering system, either through raw measurements or calculations based on these measurements. The inputs are selected to provide a smooth, continuous input in relation to the outputs. A common neural network design for gas-liquid mixtures is shown in Fig. 14. Note that two separate neural networks are used for the mass flow rate correction and the density correction, which is represented in Fig. 14 by the horizontal line in the box for the neural network. One example of a method for training and using this neural network for use with immiscible fluids is outlined in Fig. 15. The equations used in each step are defined in Table 2, which is provided at the end of the detailed description portion of the specification.

For immiscible fluids (for example where the liquid is predominately water), the phase mass flow rates are the same for standard and actual conditions.

[0046] When gas dissolves in the liquid, the same neural network design can be used in combination with a PVT model. For example, the black oil model uses correlations or tabular data to estimate the amount of dissolved gas and density changes at different temperatures and pressures. The density of the in-situ free gas phase can be determined using the AOA8 model, for example, although other correlations could also be used instead.

[0047] Many correlations are available for the black oil model. One example is the Vazquez model. Key parameters in the black oil model are:

- (i) $\rho_{o,sc}$, $\rho_{w,sc}$, $\rho_{g,sc}$ - standard liquid, gas and water densities used to define the fluids. This is normal input data for the meter. These densities are converted from the meter input standard temperature to 60 °F for use in the correlations.
- (ii) GOR, the gas - oil ratio, is a measure of the gas to oil production rates at standard conditions. This is obtained from surface measurements on the separate gas and liquid phases over a period of time.
- (iii) Rs, the solution gas - oil ratio, accounts for gas that is dissolved in the oil on a volumetric basis. This relationship can be rearranged to predict the bubble point pressure. A free gas phase is present at pressures below the bubble point pressure. At pressures above the bubble point pressure, the gas is under saturated and the amount dissolved is limited by the GOR.
- (iv) Bo, the formation volume factor, adjusts the oil density for the dissolved gas. The oil is lighter due to the presence of dissolved gases, even at high pressures.
- (v) Rw and Bw are similar factors to account for gas dissolved in water. Gas is far less soluble in water than oil and can usually be neglected in the Coriolis application.

[0048] The model suitably includes a gas solubility tuning factor that allows the model to be tuned to account for the fact that actual petroleum fluids do not always behave in the same way. Experiments can be run to improve the tuning of the model for use with a particular set of petroleum fluids. Alternatively, it is sometimes known within the oil and gas industry what tuning factors work well with fluids from a particular oil reservoir or region. The appropriate tuning factors can be input based on this industry knowledge. The tuning factor can be set to a nominal value of 1.0, which matches typical properties of heavy crude oil. When the tuning factor is set to zero, the model becomes an immiscible model.

The tuning factor can be configured as a function of water-cut. A commercial physical property calculation package, such as Schneider Electric SimSci™ PIPEPHASE, can be used to obtain full characterization of multiphase phase flow.

[0049] An example of a miscible fluid neural network model is shown in Fig. 16. One example of a method of training and using this neural network is outlined in Fig. 17. The method generally corresponds to the method outlined in Fig. 15, but there are differences associated with inclusion of calculations for dissolved gas. The equations associated with the miscible fluid neural network are listed in Table 3.

[0050] A simplified parametric model or look-up table based on the correlations can be built into an online multiphase flow meter. For example, Fig. 18 shows curves illustrating the calculated variation with pressure and temperature of the Rs parameter with standard oil and gas densities. These curves show that the amount of dissolved gas increases with increasing pressure, but decreases with increasing temperature. Similarly, as illustrated in Fig. 19, curves can be obtained for Bo as a function of temperature and pressure. After taking into account the tuning factor (CRs) - the gas saturated oil density reduces with increasing temperature and dissolved gas, leading to increasing Bo.

[0051] The dissolved gas model is suitably used during development of the empirical mass flow rate and density corrections for multiphase flow so that the inputs are more accurate than in conventional techniques. For example, one set of calculations are reproduced in Table 4 at the end of the detailed description. Some things stand out in these example calculations. For example, the liquid densities and GVFs calculated with the benefit of the gas solubility model are significantly different from the nominal values. While the nominal value for liquid density was 830.3 kg/m³, the actual liquid density as estimated with the benefit of the gas solubility model was only 825.3 kg/m³. This is corroborated by the scientific understanding that the inclusion of dissolved gas in the oil will reduce the density of the oil (e.g., from 797.5 kg/m³ to about 791.7 kg/m³ per the data from Table 4). The actual GV estimated with the benefit of the gas solubility
model (11.7 percent) was also much lower than the nominal GVF (18.3) percent. This is consistent with the data in Figs. 6-13 indicating that at least some of the gas does in fact dissolve into the oil during three-phase flow trials in the laboratory.

Accordingly, the mass flow rate and density corrections for multiphase flow can be improved using the techniques described herein. Thus, the Coriolis meter 215 is suitably configured to determine mass flow rates for oil, water, and/or gas, and/or density using a mapping (e.g., neural network in combination with gas solubility model) that accounts for the effects of dissolved gas. This can result in a Coriolis meter that provides more accurate mass flow rate and/or density readings during multiphase flow.

The data in Figs. 6-13 was generated using a particular set of multiphase fluids comprised of oil, water and gas from the petroleum industry. It is understood that the effects of dissolved gas can be different when different sets of multiphase fluids are involved in the process. For example, carbon dioxide (CO₂) dissolves easily in water, in contrast to gas from petroleum wells. However, the same process can be used to reduce the error associated with dissolved gas by taking the different materials into account when selecting, adapting, and or developing suitable model to account for the presence of dissolved gas.

Application of dissolved gas model in the field

The preceding sections explain how to improve the accuracy and consistency of a Coriolis meter during multiphase flow. However, it is also possible to use a gas solubility model in the field to provide a measure of the amount of dissolved gas in the fluids flowing through the Coriolis meter 215. The gas solubility model used in the field can be the same model used in the laboratory but it could be a different model. The model used in the field could have a different tuning, or even a different structure, than the model for the laboratory. The dissolved gas model most appropriate for a lower pressure laboratory used to develop the Coriolis meter 215 multiphase flow corrections might not be the most appropriate model for on-line operation in a higher pressure upstream application. However, the Coriolis meter's response to free gas should be the same, even though the quantity of any dissolved gas may be different depending on the pressure. This is because, as noted above, the Coriolis meter 215 is not influenced much by the presence of dissolved gas. If desired, the Coriolis meter 215 can be configured so one or more tuning factors can be input or adjusted by a user. This can facilitate deployment of the same Coriolis meter 215, or same type of Coriolis meter, in different oil fields having fluids that behave differently.

One method of applying a gas solubility model in real-time includes the following steps: (i) determine the amount of free gas (GVF) in the multiphase mixture based upon the response of the Coriolis meter; (ii) use the gas solubility model to calculate the amount of dissolved gas; and (iii) calculate the total gas present from the free gas and the dissolved gas. The processor 101 of the Coriolis meter 215 is suitably configured (e.g., programmed) to perform this method and output the amount of dissolved gas and/or total gas. For example, the processor 101 is suitably configured to report the amount (flow rate and/or totalized over time) of oil, water, and total gas (including both free gas and dissolved gas) at standard conditions.

Skid Design

Multiphase flow regimes of (e.g., oil, water, and gas) can present challenges for accurately measuring the flow rates of the liquid and gas components with a Coriolis mass flow meter and Water-Cut meter (measuring water-liquid volume flow ratio at line conditions) metering system. Specifically, variations in the flow regime can create slip conditions where the gas phase velocity can vary significantly from that of the oil and water liquid phase, rendering accurate metering more difficult.

Water-Cut meters operate best with a well-mixed stream of oil and water. When the flow regime includes significant slugging, it can be desirable to prevent Coriolis meters and water cut meters from being subject to positive and then negative flow rates, as might occur if the meter were positioned on the upward leg of a skid. It is also desirable that Coriolis meters are drained properly in zero flow conditions because a partially filled flowtube can create large flow rate and density errors when there is no fluid flowing through the partially filled flowtube.

Figures 20 and 21 show a net oil and gas skid 121 (which is a multiphase metering system) designed to minimize the potential slip condition between gas and liquid phases. The net oil and gas skid 121 includes the Coriolis meter 215, a water cut meter 145 positioned to measure the water cut of fluid, a temperature sensor 147 positioned to measure the temperature of the fluid, and a pressure sensor 149 positioned to measure the pressure of the fluid. It can also be desirable to measure differential pressure across the flowtube because this can facilitate performing gas calculations using the average pressure in the flowtube rather than just the pressure at the flowtube entrance, at the flowtube exit, or at some other specific location. A net oil computer 151 (broadly, a processor) is mounted on the skid 121. The net oil computer 151 is configured to receive signals from the Coriolis meter 215, water cut meter 145, temperature sensor 147, and pressure sensor 149, along with any other measurement devices that may be included on the skid 121. The net oil computer 151 suitably integrates information from all of the devices 215, 145, 147, 149, etc. and performs any
calculations that have not already been performed by the devices needed to determine flow rate and density of the fluid flowing through the skid 121. The net oil computer 151 suitably outputs current flow rate for oil, water and gas independently of one another. The net oil computer 151 suitably also outputs total flow for oil, water, and gas for one or more time periods. The processing required to operate the Coriolis meter 215 and the net oil computer 151 is suitably performed by one or more processors. For example, some of the calculations may be performed in a transmitter of the Coriolis meter 215 while other calculations are performed in a separate processor. It is understood that there is flexibility in where the various calculations and other processing activities required are carried out and the particular choices described in the embodiment detailed herein should not be considered limiting.

[0059] The net oil and gas skid 121 is configured to maintain a conditioned flow profile as a multiphase fluid flows through the skid. A flow conditioner 153 (Fig. 22) is positioned along the top section of the skid 121 to receive the multiphase fluid flowing through the skid to mix the fluid before it reaches the flowtube of the Coriolis meter 215 and before it reaches the water cut meter 145. As illustrated in Fig. 22, the flow conditioner 153 suitably includes a plurality of tabs 155 projecting inwardly from the wall 157 of a conduit 159 positioned to convey the multiphase fluid through the skid 121. The tabs 155 generate turbulence in the flowing fluid, which results in mixing of the fluid. It is understood that other types of flow conditioners could be used instead of the flow conditioner 153 illustrated in Fig. 22.

[0060] The conduit 159 is also configured to have a rise and fall in the skid 121, which tends to disrupt any slip condition developed in the flow. For example, one leg 161 of the conduit 159 extends upwardly and another leg 163 of the conduit extends downwardly. A horizontal section 165 of the conduit connects the upward and downward legs 161, 163 to another. The rise and fall of fluid as it flows through these segments 161, 163, 165 helps facilitate good mixing of the gas a liquid phases.

[0061] The water cut meter 145 is downstream of the Coriolis meter 215. The fluid is also mixed as it flows through the flowtube of the Coriolis meter 215 due to the bends therein. Thus, the Coriolis meter 215 also helps ensure the fluid is sufficiently mixed when it reaches the water cut meter 215.

[0062] In the net oil and gas skid 121 illustrated in Figs. 20 and 21, the Coriolis meter 215 and water cut meter 145 are installed in the downward leg 163 of the conduit 159, which can be desirable to limit the amount of reverse flow through the Coriolis meter 215 and water cut meter 145 (e.g., in cases where slugging conditions are frequently encountered). However, it is understood that it may be desirable to install the Coriolis meter 215 elsewhere in a net oil and gas skid, such as on the upward leg or horizontal leg. Likewise, the water cut meter 145, temperature sensor 147, and pressure sensor 149 can be installed at other locations than those illustrated in Figs. 21 and 22 if desired.

[0063] The operational range of the net oil and gas skid 121 may be increased for high gas flow, high GVF flows by the inclusion of a compact partial separation system 171 upstream of the skid, as illustrated in Fig. 23. The purpose of the separator is to reduce the quantity of free gas in the liquid stream provided to the Coriolis meter 215 and the other components of the 121. The separator 161 has a gas output 173 and a liquid output 175. The liquid output 175 is directed from the separator 171 to the flowtube of the Coriolis meter 215 and the gas output is diverted from the flowtube to reduce the amount of free gas in the multiphase fluid before it flows through the flowtube. The separator 171 may lack the capacity to achieve complete separation of the gas and liquids, but this is not a problem because the capability of the net oil and gas skid 121 to handle multiphase flow. A separator 171 designed to achieve only partial separation can be advantageous because of its reduced size, cost, and complexity. Thus, in some instances it may be desirable to use a separator 171 that lacks capacity to achieve complete separation.

[0064] Another Coriolis meter 215', which may be identical to the Coriolis meter 215 described above is positioned to meter the gas output from the separator 171. It is understood that the gas output 175 can be metered by a different type of meter, including a meter that is not a Coriolis meter if desired. However, it may be desirable in some cases to use a Coriolis meter to meter the gas output 175 from the separator 171 because a control system (not shown) of the separator can monitor the density readings and drive gain of the Coriolis meter to detect any liquid carry-over (i.e., liquid that is undesirably included in the gas output). Short-term occurrences of liquid carry-over are to be expected from time to time, during which the measurement from the system would become invalid if undetected. When these liquid carry-over events are detected (e.g., by monitoring the density readings and drive gain of the Coriolis meter 215'), a SEVA approach, as described in U.S. Patent No. 5,570,300 can be adopted where the measurement state can be temporarily set to "DAZ-ZLED", and the system output may be replaced by the current test average, or a recent average of known reliable data. The cumulative duration of liquid carry-over events is suitably recorded and reported to the operator so that its impact on the overall test can be taken into consideration.

[0065] When introducing elements of aspects of the invention or the embodiments thereof, the articles "a," "an," "the," and "said" are intended to mean that there are one or more of the elements. The terms "comprising," "including," and "having" are intended to be inclusive and mean that there may be additional elements other than the listed elements.

[0066] Not all of the depicted components illustrated or described may be required. In addition, some implementations and embodiments may include additional components. Variations in the arrangement and type of the components may be made without departing from the scope of the claims as set forth herein. Additional, different or fewer components may be provided and components may be combined. Alternatively or in addition, a component may be implemented by
several components.

[0067] The above description illustrates the aspects of the invention by way of example and not by way of limitation. This description enables one skilled in the art to make and use the aspects of the invention, and describes several embodiments, adaptations, variations, alternatives and uses of the aspects of the invention, including what is presently believed to be the best mode of carrying out the aspects of the invention. Additionally, it is to be understood that the aspects of the invention are not limited in application to the details of construction and the arrangement of components set forth in the following description or illustrated in the drawings. The aspects of the invention are capable of other embodiments and of being practiced or carried out in various ways. Also, it will be understood that the phraseology and terminology used herein is for the purpose of description and should not be regarded as limiting.

[0068] Having described aspects of the invention in detail, it will be apparent that modifications and variations are possible without departing from the scope of aspects of the invention as defined in the appended claims. It is contemplated that various changes could be made in the above constructions, products, and process without departing from the scope of aspects of the invention. In the preceding specification, various preferred embodiments have been described with reference to the accompanying drawings. It will, however, be evident that various modifications and changes may be made thereto, and additional embodiments may be implemented, without departing from the broader scope of the aspects of the invention as set forth in the claims that follow. The specification and drawings are accordingly to be regarded in an illustrative rather than restrictive sense.

[0069] The Abstract is provided to help the reader quickly ascertain the nature of the technical disclosure. It is submitted with the understanding that it will not be used to interpret or limit the scope or meaning of the claims.
<table>
<thead>
<tr>
<th>Variable</th>
<th>Description</th>
<th>Units</th>
</tr>
</thead>
<tbody>
<tr>
<td>$E_o$</td>
<td>Oil Formation Volume Factor</td>
<td>[dimensionless]</td>
</tr>
<tr>
<td>$E_w$</td>
<td>Water Formation Volume Factor</td>
<td>[dimensionless]</td>
</tr>
</tbody>
</table>

**Corrections**

- $C_{de}$: Coriolis NN correction for total fluid density
- $C_{mf}$: Coriolis NN correction for total mass flow
- $C_{Bo}$: Black oil tuning for dissolved gas in oil
- $C_{Bo_w}$: Black oil tuning for dissolved gas in water

*Laboratory fluids with approx. 20% solubility compared to hydrocarbon gas in Vazquez model, are accommodated by setting $C_{Bo} = 0.2$. This tuning factor may be defined as a function of the measured water cut.*

<table>
<thead>
<tr>
<th>$\text{GVF}$</th>
<th>Gas Volume Flow Fraction</th>
<th>Volume %</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\text{LVF}$</td>
<td>Liquid Volume Flow Fraction</td>
<td>Volume %</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>$\dot{m}_{\text{phase,conditions}}$</th>
<th>Mass flow rate</th>
<th>kg/s</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phase</td>
<td></td>
<td></td>
</tr>
<tr>
<td>o oil</td>
<td></td>
<td></td>
</tr>
<tr>
<td>w water</td>
<td></td>
<td></td>
</tr>
<tr>
<td>g gas</td>
<td></td>
<td></td>
</tr>
<tr>
<td>gf gas free</td>
<td></td>
<td></td>
</tr>
<tr>
<td>go gas in oil</td>
<td></td>
<td></td>
</tr>
<tr>
<td>gw gas in water</td>
<td></td>
<td></td>
</tr>
<tr>
<td>l liquid</td>
<td></td>
<td></td>
</tr>
<tr>
<td>m mixture</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Conditions:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>sc standard conditions</td>
<td></td>
<td></td>
</tr>
<tr>
<td>ac actual conditions</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>$\rho_{raw}$</th>
<th>Mixture raw measured density</th>
<th>kg/m$^3$</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>$\rho_{drop}$</th>
<th>Density drop</th>
<th>%</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>$\rho_{\text{phase,conditions}}$</th>
<th>Phase density</th>
<th>kg/m$^3$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phase</td>
<td></td>
<td></td>
</tr>
<tr>
<td>o oil</td>
<td></td>
<td></td>
</tr>
<tr>
<td>w water</td>
<td></td>
<td></td>
</tr>
<tr>
<td>g gas</td>
<td></td>
<td></td>
</tr>
<tr>
<td>gf gas free</td>
<td></td>
<td></td>
</tr>
<tr>
<td>go gas in oil</td>
<td></td>
<td></td>
</tr>
<tr>
<td>gw gas in water</td>
<td></td>
<td></td>
</tr>
<tr>
<td>l liquid</td>
<td></td>
<td></td>
</tr>
<tr>
<td>m mixture</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Conditions:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>sc standard conditions</td>
<td></td>
<td></td>
</tr>
<tr>
<td>ac actual conditions</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>$P$</th>
<th>Absolute Pressure of fluid</th>
<th>kPa</th>
</tr>
</thead>
</table>

| $P_o$ | Pressure at standard conditions | |

<table>
<thead>
<tr>
<th>$Q_{\text{phase,conditions}}$</th>
<th>Volume flow rate</th>
<th>kg/s</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phase</td>
<td></td>
<td></td>
</tr>
<tr>
<td>o oil</td>
<td></td>
<td></td>
</tr>
<tr>
<td>w water</td>
<td></td>
<td></td>
</tr>
<tr>
<td>g gas</td>
<td></td>
<td></td>
</tr>
<tr>
<td>gf gas free</td>
<td></td>
<td></td>
</tr>
<tr>
<td>go gas in oil</td>
<td></td>
<td></td>
</tr>
<tr>
<td>gw gas in water</td>
<td></td>
<td></td>
</tr>
<tr>
<td>l liquid</td>
<td></td>
<td></td>
</tr>
<tr>
<td>m mixture</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Conditions:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>sc standard conditions</td>
<td></td>
<td></td>
</tr>
<tr>
<td>ac actual conditions</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Symbol</td>
<td>Description</td>
<td>Unit</td>
</tr>
<tr>
<td>--------</td>
<td>-------------</td>
<td>------</td>
</tr>
<tr>
<td>$R_g$</td>
<td>Solution gas oil ratio (standard conditions)</td>
<td>m³ / m³</td>
</tr>
<tr>
<td>$R_w$</td>
<td>Solution gas water ratio (standard conditions)</td>
<td>m³ / m³</td>
</tr>
<tr>
<td>$T$</td>
<td>Temperature of fluid</td>
<td>°C or K</td>
</tr>
<tr>
<td>$T_o$</td>
<td>Temperature at standard conditions</td>
<td></td>
</tr>
<tr>
<td>$T_{time}$</td>
<td>Time</td>
<td>seconds</td>
</tr>
<tr>
<td>$W_{cut}$</td>
<td>Water cut – Instrument measurement</td>
<td>Vol/Vol %</td>
</tr>
<tr>
<td>$WT_{phase}$</td>
<td>Measured weight of phase over time</td>
<td>kg</td>
</tr>
<tr>
<td>$Z$</td>
<td>Gas non ideal compressibility factor</td>
<td></td>
</tr>
</tbody>
</table>
Table 2: Steps in neural network calculations – Immiscible Fluids

<table>
<thead>
<tr>
<th>Calculation</th>
<th>Description</th>
<th>Equation</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. <strong>NN Input</strong></td>
<td>Oil density, ac</td>
<td>$\rho_{o,ac} = f(T_r, \rho_{o,ac})$</td>
</tr>
<tr>
<td>2.</td>
<td>Water density, ac</td>
<td>$\rho_{w,ac} = f(T_r, \rho_{w,ac})$</td>
</tr>
<tr>
<td>3.</td>
<td>Gas density, ac <em>(For this NN, used later)</em></td>
<td>$\rho_g = \rho_{g,ac} \times Z \times \left( \frac{P_{so}}{P_o} \right) \times \left( \frac{T_u}{T_{iso}} \right)$</td>
</tr>
<tr>
<td></td>
<td></td>
<td>$Z$ calculated from AGAB or similar PVT models</td>
</tr>
<tr>
<td>4.</td>
<td>Water cut adjustment</td>
<td>$W_{cut} = f(T_r)$</td>
</tr>
<tr>
<td>5.</td>
<td>Liquid density, ac</td>
<td>$\rho_{l,ac} = \rho_{l,ac} \times \frac{W_{w,ac}}{100} + \rho_{g,ac} \times \frac{(100-W_{cut})}{100}$</td>
</tr>
<tr>
<td>6.</td>
<td>Density drop</td>
<td>$\rho_{drop} = 100 \times \frac{(\rho_{l,ac} - \rho_{w,ac})}{\rho_l}$</td>
</tr>
<tr>
<td>7. <strong>NN Target Prep</strong></td>
<td>Reference phase mass flow rates <em>(from total reference measurements)</em></td>
<td>$\dot{m}<em>{phase,ac} = \frac{W</em>{T,phase}}{\Delta \text{Time}}$</td>
</tr>
<tr>
<td>8.</td>
<td>Reference total mass flow rate</td>
<td>$\dot{m}<em>m = \dot{m}</em>{o,ac} + \dot{m}<em>{w,ac} + \dot{m}</em>{g,ac}$</td>
</tr>
<tr>
<td>9.</td>
<td>Reference phase volume rates</td>
<td>$Q_{phase,ac} = \frac{\dot{m}<em>{phase,ac}}{\rho</em>{phase,ac}}$</td>
</tr>
<tr>
<td>10.</td>
<td>Reference mixture density, ac</td>
<td>$\rho_{m,ac} = \frac{\dot{m}<em>{m,ac}}{(Q</em>{w,ac} + Q_{w,ac} + Q_{g,ac})}$</td>
</tr>
<tr>
<td>Calculation</td>
<td>Description</td>
<td>Equation</td>
</tr>
<tr>
<td>-------------</td>
<td>-------------</td>
<td>----------</td>
</tr>
<tr>
<td>1.</td>
<td><strong>NN Input</strong>&lt;br&gt;Oil density, ac (no dissolved gas)</td>
<td>( \rho_{\text{ac}} = f(\eta, \rho_{\text{ac}}) ) &lt;br&gt;Lab polynomial fit</td>
</tr>
<tr>
<td>2.</td>
<td><strong>Water density, ac (no dissolved gas)</strong></td>
<td>( \rho_{\text{w,ac}} = f(\eta, \rho_{\text{w,ac}}) ) &lt;br&gt;Lab polynomial fit</td>
</tr>
<tr>
<td>3.</td>
<td><strong>Gas density, ac</strong>&lt;br&gt;(For this NN, used later)</td>
<td>( \bar{\rho}<em>g = \rho</em>{\text{gas}} \times Z \times \left( \frac{P_{\text{gas}}}{P_c} \right) \times \left( \frac{T_{\text{gas}}}{T_c} \right) )&lt;br&gt;Z calculated from AGA or similar PVT model</td>
</tr>
<tr>
<td>4.</td>
<td><strong>Water cut adjustment</strong></td>
<td>( W_{\text{cut}} = f(\eta, ...) ) &lt;br&gt;Lab tuning NN</td>
</tr>
<tr>
<td>5.</td>
<td><strong>Liquid density, ac (no dissolved gas)</strong></td>
<td>( \rho_{\text{liq}} = \rho_{\text{w,liq}} \times \frac{W_{\text{liq}}}{100} + \rho_{\text{w,liq}} \times \frac{(100 - W_{\text{cut}})}{100} )</td>
</tr>
<tr>
<td>6.</td>
<td><strong>Density drop</strong></td>
<td>( \rho_{\text{drop}} = 100 \times \frac{(\rho_{\text{liq}} - \rho_{\text{raw}})}{\rho_{\text{liq}}} )</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>---</td>
<td>---</td>
<td>---</td>
</tr>
<tr>
<td>7.</td>
<td>NN Target Prep</td>
<td>Reference phase mass flow rates (from total reference measurements) [ \dot{m}<em>{\text{phase,ref}} = \frac{W \dot{T}</em>{\text{phase}}}{\Delta T \text{ime}} ]</td>
</tr>
<tr>
<td>8.</td>
<td></td>
<td>Reference total mass flow rate [ \dot{m}<em>{\text{t}} = \dot{m}</em>{\text{daw}} + \dot{m}<em>{\text{sw}} + \dot{m}</em>{\text{g}} ]</td>
</tr>
<tr>
<td>9.</td>
<td></td>
<td>Reference phase volume rates at sc [ Q_{\text{phase,sc}} = \frac{\dot{m}<em>{\text{phase,sc}}}{\rho</em>{\text{phase,sc}}} ]</td>
</tr>
<tr>
<td>10.</td>
<td></td>
<td>Reference Gas oil ratio [ GOR = \frac{Q_g}{Q_{\text{daw}}} ]</td>
</tr>
</tbody>
</table>
| 11. |   | Black oil Parameters \[ \begin{align*}
R_o &= C_{\text{ro}} \ast f(\rho_{\text{gas}}, \rho_{\text{o,cor}}, P, T) \\
B_o &= f(\rho_{\text{gas}}, \rho_{\text{o,cor}}, P, T) \\
R_w &= C_{\text{rw}} \ast f(\rho_{\text{gas}}, \rho_{\text{w,cor}}, P, T) \\
B_w &= f(\rho_{\text{gas}}, \rho_{\text{w,cor}}, P, T)
\end{align*} \] |
| 12. |   | Gas rates in each phase \[ \begin{align*}
Q_{\text{daw}} &= R_o \ast Q_{\text{gas}} \\
Q_{\text{sw}} &= R_w \ast Q_{\text{sw}} \\
Q_{\text{g}} &= Q_{\text{gas}} - Q_{\text{daw}} - Q_{\text{sw}} \\
\dot{m}_{\text{daw}} &= Q_{\text{daw}} \ast \rho_{\text{daw}} \\
\dot{m}_{\text{sw}} &= Q_{\text{sw}} \ast \rho_{\text{sw}} \\
\dot{m}_{\text{g}} &= Q_{\text{gas}} \ast \rho_{\text{gas}} \\
\dot{m}_{\text{free,ac}} &= \dot{m}_{\text{g}} - \dot{m}_{\text{daw}} - \dot{m}_{\text{sw}}
\end{align*} \] |
| 13. |   | Free gas training check If \( \dot{m}_{\text{free,ac}} < 0 \) - Undersaturated If under saturated... Gas flow rate range from 0-saturation Gas Apply model of degree of under-saturation or eliminate point from NN modelling process. |
| 14. |   | Phase volume flow rate, ac \[ \begin{align*}
Q_{\text{daw}} &= B_o \ast Q_{\text{daw}} \\
Q_{\text{sw}} &= B_w \ast Q_{\text{sw}} \\
Q_{\text{g}} &= \dot{m}_{\text{free,ac}} \frac{\rho_{\text{g,ac}}}{\rho_{\text{gas}}}
\end{align*} \] |
| 15. |   | Reference Mixture density, ac \[ \rho_{\text{m,ac}} = \frac{\dot{m}_{\text{m,ac}}}{Q_{\text{daw}} + Q_{\text{sw}} + Q_{\text{g}}}
\] \[ C_{\text{dm,target}} = 100 \ast \frac{\rho_{\text{m,ac}}}{\rho_{\text{ref}}} \] |
<p>| 17. | <strong>NN Target</strong> | NN mass flow correction target | $C_{mfs \text{, target}} = \frac{\dot{m}<em>{\text{gas}}}{\dot{m}</em>{\text{raw}}}$ |
| 18. | <strong>NN Corrections</strong> | NN density correction | $C_{\rho} = f(\dot{m}<em>{\text{raw}}, \rho</em>{\text{raw}}, \rho_{\text{gas}})$ |
| 19. | | NN mass flow correction | $C_{mfs} = f(\dot{m}<em>{\text{raw}}, \rho</em>{\text{raw}}, \rho_{\text{gas}})$ |
| 20. | | Mixture density, ac | $\rho_{\text{m,ac}} = \frac{C_{\rho} \cdot \dot{m}<em>{\text{raw}}}{100}$ |
| 21. | | Mixture mass flow rate | $\dot{m}</em>{\text{m}} = \frac{C_{mfs} \cdot \dot{m}<em>{\text{raw}}}{100}$ |
| 22. | | Mixture volume flow rate, ac | $Q</em>{\text{m,ac}} = \frac{\dot{m}<em>{\text{m}}}{\rho</em>{\text{m,ac}}}$ |
| 23. | | Black oil Parameters $R_e, B_o, R_w, B_w$ | Refer to equations in step 11 |
| 24. | | Phase density, ac | $\rho_{\text{ph,ac}} = \frac{\rho_{0,ac}}{B_o}$, $\rho_{\text{w,ac}} = \frac{\rho_{w,ac}}{B_w}$, $\rho_{\text{gas,ac}} = \rho_{\text{gas,ac}} \cdot \frac{W_{\text{gas}}}{100} + \rho_{\text{gas,ac}} \cdot \frac{(100 - W_{\text{gas}})}{100}$, $\rho_{\phi} = \rho_{\phi,ac} \cdot \frac{P_{\text{res}}}{P_{\phi,ac}}$ |
| 25. | | Liquid volume fraction, ac | $LVF = \frac{(\rho_{\text{m,ac}} - \rho_{\text{gas,ac}})}{\rho_{\text{ph,ac}} - \rho_{\text{gas,ac}}}$ |
| 26. | | Phase volume flow rate, ac | $Q_{\phi,ac} = \frac{(100 - W_{\text{gas}})}{100} \cdot LVF \cdot Q_{\text{m,ac}}$, $Q_{\text{w,ac}} = W_{\text{w,ac}} \cdot L VF \cdot Q_{\text{m,ac}}$, $Q_{\text{gas,ac}} = (1 - LVF) \cdot Q_{\text{m,ac}}$ |
| 27. | | Gas rates in each phase | See Step 12 |</p>
<table>
<thead>
<tr>
<th>Page</th>
<th>Free gas measurement check</th>
<th>If $\text{Gain} &lt; \text{Gain Tol} = \text{Undersaturated}$</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Recommended Gain Tol 2X water default for tube</td>
</tr>
<tr>
<td></td>
<td></td>
<td>If under saturated...</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Gas flow rate range from 0 - saturation gas</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Liquid density based on user GOR if given. If</td>
</tr>
<tr>
<td></td>
<td></td>
<td>not given, single phase correlation used</td>
</tr>
<tr>
<td></td>
<td></td>
<td>No comparison to lab data for gas rates</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Report options for gas rate...</td>
</tr>
<tr>
<td></td>
<td></td>
<td>User GOR for gas rate at sc</td>
</tr>
<tr>
<td></td>
<td></td>
<td>N/A or &lt; Saturated value or</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2-4 region estimates based on gain</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Page</th>
<th>Phase mass flow rate, sc</th>
<th>$m_{o,sc} = m_{o,ac} - m_{go,ac}$</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>$m_{w,sc} = m_{w,ac} - m_{gw,ac}$</td>
<td></td>
</tr>
<tr>
<td></td>
<td>$m_{g,sc} = m_{g,ac} + m_{go,ac} + m_{w,sc}$</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Page</th>
<th>Phase volume flow rate, sc</th>
<th>For reporting</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>$Q_{phase,sc} = \rho_{phase,sc}$</td>
</tr>
</tbody>
</table>

Table 4 - Example calculations for laboratory trials:

<table>
<thead>
<tr>
<th>Std. Oil density at 60 °F</th>
<th>814.6</th>
<th>kg/m³</th>
</tr>
</thead>
<tbody>
<tr>
<td>Std. Gas density at 60 °F, 101.325 kPa abs.</td>
<td>1.223</td>
<td>kg/m³</td>
</tr>
<tr>
<td>Actual Pressure</td>
<td>400.0</td>
<td>kPa</td>
</tr>
<tr>
<td>Actual Temperature</td>
<td>40.0</td>
<td>°C</td>
</tr>
<tr>
<td>Tuning constant for gas in oil CRs</td>
<td>0.20</td>
<td></td>
</tr>
<tr>
<td>Calculated actual Oil Rs</td>
<td>0.408</td>
<td></td>
</tr>
<tr>
<td>Calculated gas saturated Oil Bo</td>
<td>1.029</td>
<td></td>
</tr>
<tr>
<td>Calculated actual gas saturated Oil density</td>
<td>791.7</td>
<td>kg/m³</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Assume known true Oil mass flow rate</th>
<th>4.0</th>
<th>kg/s</th>
</tr>
</thead>
<tbody>
<tr>
<td>Oil std. volume flow rate</td>
<td>0.004911</td>
<td>m³/s</td>
</tr>
<tr>
<td>Calc. saturated gas std. volume flow</td>
<td>0.002002</td>
<td>m³/s</td>
</tr>
<tr>
<td>Calc. saturated gas mass flow</td>
<td>2.448</td>
<td>g/s</td>
</tr>
</tbody>
</table>

Now, ignoring negligible dissolved gas in water

<table>
<thead>
<tr>
<th>Assume known true Water mass flow rate</th>
<th>1.0</th>
<th>kg/s</th>
</tr>
</thead>
<tbody>
<tr>
<td>Actual water density, calc. at line temperature</td>
<td>994.2</td>
<td>kg/m³</td>
</tr>
<tr>
<td>Calc. Liquid volume flow rate</td>
<td>0.006059</td>
<td>m³/s</td>
</tr>
</tbody>
</table>
Calc. Water Cut (actual volumetric flow ratio) | 16.6 | %
Calc. Liquid density | 825.3 | kg/m³
Assume known true Gas mass flow rate | 6.0 | g/s
Free Gas mass flow rate | 3.552 | g/s
Free Gas density, calc. at line pressure and temperature | 4.4 | kg/m³
Free Gas volume flow rate | 0.000798 | m³/s
Calc. Free Gas GVF | 11.7 | %
Calc. mixture density | 732.0 | kg/m³
Calc. density drop | 11.6 | %

For comparison, assuming no dissolved gas, the following would have been calculated

Oil density | 797.5 | kg/m³
Liquid volume flow | 0.006022 | m³/s
Water Cut | 16.7 | %
Liquid density | 830.3 | kg/m³
Gas volume flow | 0.001348 | m³/s
'nominal GVF' | 18.3 | %
Mixture density | 679.2 | kg/m³
Density drop | 18.2 | %

Claims

1. System adapted for metering flow of a fluid, the system comprising:
   a vibratable flowtube for receiving a multiphase fluid flow including water and a petroleum fluid;
   a driver (46a, 46b) configured to vibrate the flowtube;
   a pair of sensors (48a, 48b) positioned to detect movement of the flowtube at different locations on the flowtube
   and output sensor signals indicative of the detected movement;
   a pressure sensor (149) positioned to measure a pressure of the multiphase fluid;
   a temperature sensor (147) positioned to measure a temperature of the fluid; and
   one or more processors (101) configured to receive the sensor signals, determine a phase difference between
   the sensor signals, and use the determined phase difference to determine a fluid flow rate through the flowtube,
   wherein said one or more processors (101) are further configured to:

   receive a signal from the pressure sensor (149) indicative of the pressure of the multiphase fluid in the
   flowtube;
   receive a signal from the temperature sensor (147) indicative of the temperature of the multiphase fluid in
   the flowtube; and
   receive a signal from a meter (145) indicative of relative amounts of multiple different liquids in the multiphase
   fluid;

characterized in that said one or more processors (101) are further configured to:

determine an amount of dissolved gas in the multiphase fluid using the pressure, the temperature, and the relative
amounts the multiple liquids in the multiphase fluid, according to a gas solubility model, wherein the gas solubility
model includes an adjustable tuning factor that allows the gas solubility model to be tuned to account for the fact
that actual petroleum fluids do not always behave in the same way, and determine the fluid flow rate by using a mapping between a plurality of inputs and at least one output related to the fluid flow rate, wherein the inputs comprise a raw mass fluid flow rate, pressure of the fluid, temperature of the fluid, and the relative amounts of the multiple liquids in the multiphase fluid.

2. System as set forth in claim 2, wherein one or more processors (101) are configured to determine an amount of free gas in the multiphase fluid, wherein said one or more processors (101) preferably configured to determine an amount of total gas in the multiphase fluid using the amounts determined for dissolved gas and free gas.

3. System as set forth in any one of the preceding claims, further comprising a flow conditioner positioned and structured to mix the multiphase fluid before it enters the flowtube, wherein the flow conditioner preferably comprises a plurality of tabs (155) projecting inwardly from a wall (157) of a conduit (159) positioned to convey the multiphase fluid to the flowtube.

4. System as set forth in any one of the preceding claims, further comprising a conduit upstream of the flow tube and connected to the flowtube for conveying the multiphase fluid to the flowtube, the conduit (159) being configured to have two substantially vertical segments (161, 163) and a substantially horizontal segment (165) between the substantially vertical segments (161, 163), wherein one of the substantially vertical segments (161, 163) is preferably between the substantially horizontal segment (165) and the flowtube, wherein the flowtube is preferably at a lower elevation than the substantially horizontal segment (165), and/or wherein the substantially horizontal segment (165) is preferably connected to each of the substantially vertical segments (161, 163) at upper ends of the substantially vertical segments (161, 163).

5. System as set forth in any one of the preceding claims, further comprising a separator (171) upstream of the flowtube for separating at least some free gas in the multiphase fluid from the liquid phase, the separator (171) having a gas output (173) and a liquid output (175), wherein the liquid output (175) is directed from the separator (171) to the flowtube and the gas output (173) is diverted from the flowtube to reduce the amount of free gas in the multiphase fluid before it flows through the flowtube.

6. System as set forth in any of the preceding claims, wherein the multiple different liquids include water and oil and wherein said meter (145) comprises a water cut meter.

7. System as set forth in any of the preceding claims, wherein the one or more processors (101) are configured to:
determine fluid flow rates for each of said multiple different liquids using a mapping between a plurality of inputs and said fluid flow rates of the different liquids, said inputs comprising a phase difference between the motion sensor signals, a frequency of at least one of the motion sensors signals, the signal from the temperature sensor (147), the signal from the pressure sensor (149), and the signal from said meter (145) indicative of the relative amounts of the multiple different liquids, wherein the mapping accounts for solubility of the gas in the multiple different liquids.

8. System as set forth in any one of the preceding claims, wherein the mapping comprises a neural network, and/or wherein the mapping is based in part on the gas solubility model.

9. Net oil and gas skid (121) adapted for characterizing the output from one or more petroleum wells, the net oil and gas skid (121) comprising a system as set forth in any of the preceding claims, wherein the vibratable flowtube receives a multiphase fluid from said one or more petroleum wells; wherein the net oil and gas skid (121) further comprises a water cut meter (145) adapted to measure relative amounts of oil and water in the multiphase fluid; and wherein said one or more processors (101) are further configured to:
receive a signal from the water cut meter (145) indicative of relative amounts of oil and water in the multiphase fluid;
determine a density of the multiphase fluid in the flowtube using the sensor signals.

10. Net oil and gas skid (121) as set forth in claim 9, wherein the mapping comprises a neural network, wherein the neural network preferably has been trained using empirical data that accounts for the effects of dissolved gas using the gas solubility model and/or wherein the gas solubility model is preferably a first gas solubility model and wherein the neural network has been trained using empirical data that accounts for the effects of dissolved gas using a
second gas solubility model that is different from the first gas solubility model.

11. Net oil and gas skid as set forth in any one of claims 9 to 10, wherein said one or more processors (101) are configured to determine an amount of free gas in the multiphase fluid, wherein said one or more processors (101) are preferably configured to determine an amount of total gas in the multiphase fluid using the amounts determined for dissolved gas and free gas, and/or further comprising a flow conditioner positioned and structured to mix the multiphase fluid before it enters the flowtube, and/or further comprising a separator (171) upstream of the flowtube, the separator (171) having a gas output (173) and a liquid output (175), wherein the liquid output (175) is directed from the separator (171) to the flowtube and the gas output (173) is diverted from the flowtube to reduce the amount of free gas in the multiphase fluid before it flows through the flowtube.

12. Method of metering flow of a multiphase fluid using a system as set forth in any one of claims 1 to 8, the method comprising:

- vibrating a flowtube while the multiphase fluid flows through the flowtube;
- using a pair of sensors (46a, 46b) to detect movement of the flowtube at different locations on the flowtube and outputting a sensor signal from each sensor (46a, 46b);
- measuring a pressure of the multiphase fluid;
- measuring a temperature of the multiphase fluid;
- measuring relative amounts of multiple different liquids in the multiphase flow; and
- determining a flow rate based on a phase difference between the sensor signals, characterized by determining an amount of dissolved gas in the multiphase fluid using the temperature and pressure of the fluid and the relative amounts of the multiple liquids in the multiphase fluid, wherein the amount of dissolved gas is determined according to a gas solubility model, wherein the solubility model includes an adjustable tuning factor that allows the gas solubility model to be tuned to account for the fact that actual petroleum fluids do not always behave in the same way.

13. Method as set forth in claim 12 using a system as set forth in any one of claims 1 to 8 and/or a net oil and gas skid as set forth in any of claims 9 to 11.

Patentansprüche

1. System, das zum Messen der Strömung eines Fluids geeignet ist, wobei das System umfasst:

- ein Strömungsrohr, das zum Vibrieren gebracht werden kann, zum Aufnehmen eines mehrphasigen Fluidstroms, der Wasser und eine Erdölflüssigkeit aufweist;
- einen Treiber (46a, 46b), der konfiguriert ist, um das Strömungsrohr zum Vibrieren zu bringen;
- ein Paar Sensoren (48a, 48b), die so positioniert sind, dass sie die Bewegung des Strömungsrohrs an unterschiedlichen Stellen auf dem Strömungsrohr erfassen, und Sensorsignale ausgeben, die die erfasste Bewegung anzeigen;
- einen Drucksensor (149), der positioniert ist, um einen Druck des mehrphasigen Fluids zu messen;
- einen Temperatursensor (147), der so angeordnet ist, dass er die Temperatur des Fluids misst; und
- einen oder mehrere Prozessoren (101), die konfiguriert sind, um die Sensorsignale zu empfangen, eine Phasendifferenz zwischen den Sensorsignalen zu bestimmen und die bestimmte Phasendifferenz zu verwenden, um eine Fluidströmungsrate durch das Strömungsrohr zu bestimmen, wobei der eine oder die mehreren Prozessoren (101) weiter konfiguriert sind, um:

- ein Signal vom Drucksensor (149) zu empfangen, das den Druck des mehrphasigen Fluids im Strömungsrohr anzeigen;
- ein Signal vom Temperatursensor (147) zu empfangen, das die Temperatur des mehrphasigen Fluids im Strömungsrohr anzeigen; und
- ein Signal von einem Messgerät (145) zu empfangen, das die relativen Mengen oder der relativen Flüssigkeiten in dem mehrphasigen Fluid anzeigen;

dadurch gekennzeichnet, dass der eine oder die mehreren Prozessoren (101) weiter konfiguriert sind, um:

- eine Menge an gelöstem Gas im mehrphasigen Fluid unter Verwendung des Drucks, der Temperatur und der relativen Mengen der mehreren Flüssigkeiten in dem mehrphasigen Fluid gemäß einem Gaslöslich-
keitsmodell zu bestimmen, wobei das Gaslöslichkeitsmodell einen einstellbaren Abstimmfaktor aufweist, der es ermöglicht, das Gaslöslichkeitsmodell dahingehend abzustimmen, dass sich tatsächliche Erdölfüssigkeiten nicht immer in gleicher Weise verhalten, und die Fluidstromungsrate unter Verwendung einer Zuordnung zwischen einer Vielzahl von Eingängen und mindestens einem Abfluss in Bezug auf die Fluidstromungsrate zu bestimmen, wobei die Eingänge eine Rohmassenfluidstromungsrate, den Druck des Fluids, die Temperatur des Fluids und die relativen Mengen der mehreren Flüssigkeiten im mehrphasigen Fluid umfassen.

2. System nach Anspruch 2, wobei die eine oder die mehreren Prozessoren (101) konfiguriert sind, um eine Menge an freiem Gas in dem mehrphasigen Fluid zu bestimmen, wobei die eine oder die mehreren Prozessoren (101) vorzugsweise konfiguriert sind, um eine Menge an Gesamtgas in dem mehrphasigen Fluid unter Verwendung der für gelöstes Gas und freies Gas bestimmten Mengen zu bestimmen.

3. System, nach einem der vorhergehenden Ansprüche, weiter umfassend eine Strömungskonditioniervorrichtung, die so positioniert und strukturiert ist, dass sie das mehrphasige Fluid mischt, bevor es in das Strömungsröhr gelangt, wobei die Strömungskonditioniervorrichtung vorzugsweise eine Vielzahl von Laschen (155) umfasst, die von einer Wand (157) einer Leitung (159) nach innen ragen, die so positioniert ist, dass sie das mehrphasige Fluid zum Strömungsröhr transportiert.

4. System nach einem der vorhergehenden Ansprüche, weiter umfassend eine dem Strömungsröhr vorgeschaltete Leitung, die mit dem Strömungsröhr verbunden ist, um das mehrphasige Fluid zum Strömungsröhr zu transportieren, wobei die Leitung (159) so konfiguriert ist, dass sie zwei im Wesentlichen vertikale Segmente (161, 163) und ein im Wesentlichen horizontales Segment (165) zwischen den im Wesentlichen vertikalen Segmenten (161, 163) aufweist, wobei eines der im Wesentlichen vertikalen Segmente (161, 163) vorzugsweise zwischen dem im Wesentlichen horizontalen Segment (165) und dem Strömungsröhr liegt, wobei das Strömungsröhr vorzugsweise in einer niedrigeren Höhe als das im Wesentlichen horizontale Segment (165) liegt, und/oder wobei das im Wesentlichen horizontale Segment (165) vorzugsweise mit jedem der im Wesentlichen vertikalen Segmente (161, 163) an den oberen Enden der im Wesentlichen vertikalen Segment (161, 163) verbunden ist.

5. System nach einem der vorhergehenden Ansprüche, weiter umfassend einen dem Strömungsröhr vorgeschalteten Separator (171) zum Trennen von mindestens etwas freiem Gas in dem mehrphasigen Fluid aus der flüssigen Phase, wobei der Separator (171) einen Gasabfluss (173) und einen Flüssigkeitsabfluss (175) aufweist, wobei der Flüssigkeitsabfluss (175) vom Separator (171) zum Strömungsröhr geführt wird und der Gasabfluss (173) vom Strömungsröhr abgeleitet wird, um die Menge an freiem Gas im mehrphasigen Fluid zu reduzieren, bevor es durch das Strömungsröhr strömt.

6. System nach einem der vorstehenden Ansprüche, wobei die mehreren unterschiedlichen Flüssigkeiten Wasser und Öl aufweisen und wobei das Messgerät (145) ein Wasserableitmessgerät umfasst.

7. System nach einem der vorhergehenden Ansprüche, wobei die eine oder die mehreren Prozessoren (101) konfiguriert sind, um Fluidstromungsrate für jede der mehreren unterschiedlichen Flüssigkeiten unter Verwendung einer Zuordnung zwischen einer Vielzahl von Eingängen und den Fluidstromungsrate der unterschiedlichen Flüssigkeiten zu bestimmen, wobei die Eingänge eine Phasedifferenz zwischen den Bewegungssensorsignalen, eine Frequenz von mindestens einem der Bewegungssensorsignale, das Signal vom Temperatursensor (147), das Signal vom Drucksensor (149) und das Signal vom Messgerät (145) umfassen, das die relativen Mengen der mehreren unterschiedlichen Flüssigkeiten anzeigt, wobei die Zuordnung die Löschlichkeit des Gases in den mehreren unterschiedlichen Flüssigkeiten berücksichtigt.

8. System nach einem der vorhergehenden Ansprüche, wobei die Zuordnung ein neuronales Netzwerk umfasst, und/oder wobei die Zuordnung teilweise auf dem Gaslöslichkeitsmodell basiert.

9. Netto-Öl- und Gaselement (121), das zum Charakterisieren des Abflusses aus einer oder mehreren Erdölohrungen ausgelegt ist, wobei das Netto-Öl- und Gaselement (121) ein System nach einem der vorhergehenden Ansprüche umfasst, wobei das Strömungsröhr, das zum Vibrieren gebracht werden kann, eine mehrphasige Flüssigkeit aus einer oder mehreren Erdölohrungen empfängt; wobei das Netto-Öl- und Gaselement (121) weiter ein Wasserableitmessgerät (145) umfasst, das angepasst ist, um relative Mengen an Öl und Wasser in dem mehrphasigen Fluid zu messen; und wobei der eine oder die mehreren
Prozessoren (101) weiter konfiguriert sind, um:

ein Signal vom Wasserabtrennungsgerät (145) zu empfangen, das die relativen Mengen an Öl und Wasser in dem mehrphasigen Fluid anzeigt;
eine Dichte des mehrphasigen Fluids im Strömungsrohr unter Verwendung der Sensorsignale zu bestimmen.


11. Netto-Öl- und Gaselement nach einem der Ansprüche 9 bis 10, wobei die eine oder die mehreren Prozessoren (101) konfiguriert sind, um eine Menge an freiem Gas in dem mehrphasigen Fluid zu bestimmen, wobei die eine oder die mehreren Prozessoren (101) vorzugsweise konfiguriert sind, um eine Menge oder ein Gasgemisch in dem mehrphasigen Fluid unter Verwendung der für gelösten Gas und freies Gas bestimmten Mengen zu bestimmen, und/oder weiter umfassend eine Strömungskonditioniervorrichtung, die eine Strömungsrichtung und/oder wobei das mehrphasige Fluid mischt, bevor es in das Strömungsrohr gelangt, und/oder weiter umfassend einen dem Strömungsrohr vorgeschalteten Separator (171), wobei der Separator (171) einen Gasabfluss (173) und einen Flüssigkeitsabfluss (175) aufweist, wobei der Flüssigkeitsabfluss (175) vom Separator (171) zum Strömungsrohr gelenkt und der Gasabfluss (173) aus dem Strömungsrohr abgeleitet wird, um die Menge an freiem Gas im mehrphasigen Fluid zu reduzieren, bevor es durch das Strömungsrohr fließt.

12. Verfahren zum Messen der Strömung eines mehrphasigen Fluids unter Verwendung eines Systems nach einem der Ansprüche 1 bis 8, wobei das Verfahren umfasst:

Vibrierenlassen eines Strömungsrohrs, während das mehrphasige Fluid durch das Strömungsrohr strömt;
Verwenden eines Paars von Sensoren (46a, 46b) zum Erfassen einer Bewegung des Strömungsrohrs an unterschiedlichen Stellen auf dem Strömungsrohr und Ausgeben eines Sensorsignals von jedem Sensor (46a, 46b);
Messen eines Drucks des mehrphasigen Fluids;
Messen einer Temperatur des mehrphasigen Fluids;
Messen von relativen Mengen mehrerer unterschiedlicher Flüssigkeiten in der mehrphasigen Strömung; und
Bestimmen einer Strömungsgeschwindigkeit auf der Grundlage einer Phasendifferenz zwischen den Sensorsignalen;
gekennzeichnet durch das Bestimmen einer Menge an gelöstem Gas in dem mehrphasigen Fluid unter Verwendung der Temperatur des Drucks des Fluids und der relativen Mengen der mehreren Flüssigkeiten in dem mehrphasigen Fluid, wobei die Menge an gelöstem Gas gemäß einem Gaslöslichkeitsmodell bestimmt wird, wobei das Löslichkeitsmodell einen einstellbaren Abstimmfaktor aufweist, der es ermöglicht, das Gaslöslichkeitsmodell auf die Menge einzustellen, um zu berücksichtigen, dass sich tatsächliche Erdölfluide nicht immer gleich verhalten.


Revendications

1. Système adapté pour doser l’écoulement d’un fluide, le système comprenant :

un tube d’écoulement susceptible d’être mis en vibration pour recevoir un écoulement de fluide multiphasé incluant de l’eau et un fluide dérivé du pétrole ;
un actionneur (46a, 46b) configuré pour faire vibrer le tube d’écoulement ;
eine Paare von Sensoren (48a, 48b) positionnés pour détecter un mouvement du tube d’écoulement à différents emplacements du tube d’écoulement et délivrer des signaux de capteur indicatifs du mouvement détecté ;
un capteur de pression (149) positionné pour mesurer une pression du fluide multiphasé ;
un capteur de température (147) positionné pour mesurer une température du fluide ; et
un ou plusieurs capteurs (101) configurés pour recevoir les signaux de capteur, pour déterminer une diffé-
RENDEZ DU PHASE ENTRE LES SIGNAUX DE CAPTEUR, ET POUR UTILISER LA DIFFÉRENCE DE PHASE DÉTERMINÉE AFIN DE DÉTERMINER UN DÉBIT DU FLUIDE À TRAVERS LE TUBE D’ÉCOULEMENT, DANS LEQUEL LESDITS UN OU PLUSIEURS PROCESSEURS (101) SONT EN OUTRE CONFIGURÉS POUR :

- RECEVOIR UN SIGNAL DEPUIS LE CAPTEUR DE PRESSE (149) INDICATIF DE LA PRESSION DU FLUIDE MULTIPHASE DANS LE TUBE D’ÉCOULEMENT ;
- RECEVOIR UN SIGNAL DEPUIS LE CAPTEUR DE TEMPÉRATURE (147) INDICATIF DE LA TEMPÉRATURE DU FLUIDE MULTIPHASE DANS LE TUBE D’ÉCOULEMENT ;
- RECEVOIR UN SIGNAL DEPUIS UN DISPOSITIF DE MESURE (145) INDICATIF DES QUANTITÉS RELATIVES DE LA MULTIPlicité DE LIQUIDES DIFFÉRENTS DANS LE FLUIDE MULTIPHASE ;

CARACTÉRISÉ EN CE QUE LESDITS UN OU PLUSIEURS PROCESSEURS (101) SONT EN OUTRE CONFIGURÉS POUR :

- DÉTERMINER UNE QUANTITÉ DE GAZ DISSOUS DANS LE FLUIDE MULTIPHASE EN UTILISANT LA PRESSION, LA TEMPÉRATURE, ET LES QUANTITÉS RELATIVES DE LA MULTIPlicité DE LIQUIDES DANS LE FLUIDE MULTIPHASE, EN ACCORD AVEC UN MODÈLE DE SOLUBILITÉ DE GAZ, DANS LEQUEL LE MODÈLE DE SOLUBILITÉ DE GAZ INCLUT UN FACTEUR D’ADJUSTABLE QUI PERMET D’AJUSTER LE MODÈLE DE SOLUBILITÉ DE GAZ POUR TENTER COMPTER LE FAIT QUE LES FLUIDES DÉRIVÉS DU PÉTROLE RÉELS NE SE COMPORTENT PAS TOUJOURS DE LA MÊME MANIERE, ET
- DÉTERMINER LE DÉBIT DU FLUIDE EN UTILISANT UNE CARTOGRAPHIE ENTRE UNE PLURALITÉ D’ENTRÉES ET AU MOINS UNE SORTIE EN RELATION AVEC LE DÉBIT DU FLUIDE, DANS LEQUEL LES ENTREES COMPRENNENT UN DÉBIT MASSIQUE BRUT DU FLUIDE, LA PRESSION DU FLUIDE, LA TEMPÉRATURE DU FLUIDE, ET LES QUANTITÉS RELATIVES DE LA MULTIPlicité DE LIQUIDES DANS LE FLUIDE MULTIPHASE.

2. SYSTÈME SELON LA REVENDICATION 2, DANS LEQUEL LESDITS UN OU PLUSIEURS PROCESSEURS (101) SONT CONFIGURÉS POUR DÉTERMINER UNE QUANTITÉ DE GAZ LIBRE DANS LE FLUIDE MULTIPHASE, LESDITS UN OU PLUSIEURS PROCESSEURS (101) ÉTANT DE PRÉFÉRENCE CONFIGURÉS POUR DÉTERMINER UNE QUANTITÉ DE GAZ TOTALE DANS LE FLUIDE MULTIPHASE EN UTILISANT LES QUANTITÉS DÉTERMINÉES POUR LE GAZ DISSOUS ET LE GAZ LIBRE.

3. SYSTÈME SELON L’UNE QUELCONQUE DES REVENDICATIONS PRÉCEDANTES, COMPRENANT EN OUTRE UN DISPOSITIF DE CONDITIONNEMENT D’ÉCOULEMENT Positionné ET STRUCTURÉ POUR MÉLANGER LE FLUIDE MULTIPHASE AVANT QU’IL ENTRE DANS LE TUBE D’ÉCOULEMENT, DANS LEQUEL LE DISPOSITIF DE CONDITIONNEMENT D’ÉCOULEMENT COMPREND DE PRÉFÉRENCE UNE PLURALITÉ DE LANGUETTES (155) QUI SE PROJETTENT VERS L’INTÉRIEUR DEPUIS UNE PAROI (157) D’UN CONDUIT (159) POSITIONNÉ POUR CONVOYER LE FLUIDE MULTIPHASE VERS LE TUBE D’ÉCOULEMENT.

4. SYSTÈME SELON L’UNE QUELCONQUE DES REVENDICATIONS PRÉCEDANTES, COMPRENANT EN OUTRE UN CONDUIT EN AMONT DU TUBE D’ÉCOULEMENT ET CONNECTÉ AU TUBE D’ÉCOULEMENT POUR CONVOYER LE FLUIDE MULTIPHASE VERS LE TUBE D’ÉCOULEMENT, LE CONDUIT (159) Étant CONFIGURé POUR PRÉSENTER DEUX SEGMENTS SENSIBLEMENT VERTICAUX (161, 163) ET UN SEGMENT SENSIBLEMENT HORIZONTAL (165) ENTRE LES SEGMENTS SENSIBLEMENT VERTICAUX (161, 163), DANS LEQUEL UN DES SEGMENTS SENSIBLEMENT VERTICAUX (163) SE TROUVE DE PRÉFÉRENCE ENTRE LE SEGMENT SENSIBLEMENT HORIZONTAL (165) ET LE TUBE D’ÉCOULEMENT, DANS LEQUEL LE TUBE D’ÉCOULEMENT SE TROUVE DE PRÉFÉRENCE À UNE HAUTEUR PLUS Basse QUE LE SEGMENT SENSIBLEMENT HORIZONTAL (165), ET/OU DANS LEQUEL LE SEGMENT SENSIBLEMENT HORIZONTAL (165) EST DE PRÉFÉRENCE CONNECTé À CHACUN DES SEGMENTS SENSIBLEMENT VERTICAUX (161, 163) À DES EXTÉRIMITÉS SUPÉRIEURES DES SEGMENTS SENSIBLEMENT VERTICAUX (161, 163).

5. SYSTÈME SELON L’UNE QUELCONQUE DES REVENDICATIONS PRÉCEDANTES, COMPRENANT EN OUTRE UN SÉPARATEUR (171) EN AMONT DU TUBE D’ÉCOULEMENT POUR SÉPARER AU MOINS UNE PARTIE DU GAZ LIBRE DANS LE FLUIDE MULTIPHASE HORS DE LA PHASE LIQUIDE, LE SÉPARATEUR (171) AYANT UNE SORTIE DE GAZ (173) ET UNE SORTIE DE LIQUIDE (175), DANS LEQUEL LA SORTIE DE LIQUIDE (175) EST DIRIGéE DEPUIS LE SÉPARATEUR (171) VERS LE TUBE D’ÉCOULEMENT ET LA SORTIE DE GAZ (173) EST DÉRIVÉE DEPUIS LE TUBE D’ÉCOULEMENT POUR RÉDUIRE LA QUANTITÉ DE GAZ LIBRE DANS LE FLUIDE MULTIPHASE AVANT QU’IL S’ÉCOUELA TRAVERS LE TUBE D’ÉCOULEMENT.

6. SYSTÈME SELON L’UNE QUELCONQUE DES REVENDICATIONS PRÉCEDANTES, DANS LEQUEL LA MULTIPlicité DE LIQUIDES DIFFéRENTS INCLUT DE L’EAU ET DE L’HUile, ET DANS LEQUEL LEDIT DISPOSITIF DE MESURE (145) COMPREND UN DISPOSITIF DE MESURE DE TENUE EN EAU.

7. SYSTÈME SELON L’UNE QUELCONQUE DES REVENDICATIONS PRÉCEDANTES, DANS LEQUEL LESDITS UN OU PLUSIEURS PROCESSEURS (101) SONT CONFIGURÉS POUR : DÉTERMINER LES DÉBITS DE FLUIDE POUR CHACUN DE LADITE MULTIPlicité DE LIQUIDES DIFFéRENTS EN UTILISANT UNE CARTOGRAPHIE ENTRE UNE PLURALITÉ D’ENTRÉES ET LESDITS
débits de fluide des liquides différents, lesdites entrées comprenant une différence de phase entre les signaux des capteurs de mouvement, une fréquence de l’un au moins des signaux de capteur de mouvement, le signal provenant du capteur de température (147), le signal provenant du capteur de pression (149), et le signal provenant dudit dispositif de mesure (145) indicatif des quantités relatives de la multiplicité de liquides différents, dans lequel la cartographie tient compte de la solubilité du gaz dans la multiplicité de liquides différents.

8. Système selon l’une quelconque des revendications précédentes, dans lequel la cartographie comprend un réseau neuronal, et/ou dans lequel la cartographie est basée en partie sur le modèle de solubilité de gaz.

9. Groupe de mesure "skid" (121) de quantité nette de pétrole et de gaz, adapté pour caractériser la production d’un ou plusieurs puits de pétrole, le groupe de mesure (121) comprenant un système selon l’une quelconque des revendications précédentes, dans lequel le tube d’écoulement susceptible d’être mis en vibration reçoit un fluide multiphasé depuis lesdits un ou plusieurs puits de pétrole ; dans lequel le groupe de mesure (121) comprend en outre un dispositif de mesure de teneur en eau (145) adapté pour mesurer des quantités relatives d’huile et d’eau dans le fluide multiphasé ; et dans lesdits un ou plusieurs processeurs (101) sont en outre configurés pour :

recevoir un signal depuis le dispositif de mesure de teneur en eau (145) indicatif des quantités relatives d’huile et d’eau dans le fluide multiphasé ;

déterminer une densité du fluide multiphasé dans le tube d’écoulement en utilisant les signaux des capteurs.

10. Groupe de mesure "skid" (121) selon la revendication 9, dans lequel la cartographie comprend un réseau neuronal, dit réseau neuronal ayant de préférence été soumis à un entraînement en utilisant des données empiriques qui tiennent compte des effets des gaz dissous en utilisant le modèle de solubilité de gaz et/ou dans lequel le modèle de solubilité de gaz est de préférence un premier modèle de solubilité de gaz, et dans lequel le réseau neuronal a été soumis à un entraînement en utilisant des données empiriques qui tiennent compte des effets des gaz dissous en utilisant un second modèle de solubilité de gaz qui est différent du premier modèle de solubilité de gaz.

11. Groupe de mesure "skid" selon l’une quelconque des revendications 9 et 10, dans lequel lesdits un ou plusieurs processeurs (101) sont configurés pour déterminer une quantité de gaz libre dans le fluide multiphasé, dans lequel lesdits un ou plusieurs processeurs (101) sont de préférence configurés pour déterminer une quantité de gaz totale dans le fluide multiphasé en utilisant les quantités déterminées pour le gaz dissous et le gaz libre, et/ou comprenant en outre un dispositif de conditionnement d’écoulement positionné et structuré pour mélanger le fluide multiphasé avant qu’il entre dans le tube d’écoulement, et/ou comprenant en outre un séparateur (171) en amont du tube d’écoulement, le séparateur (171) ayant une sortie de gaz (173) et une sortie de liquide (175), dans lequel le sorti de liquide (175) est dirigée depuis le séparateur (171) vers le tube d’écoulement et la sortie de gaz (173) est dérivée depuis le tube d’écoulement afin de réduire la quantité de gaz libre dans le fluide multiphasé avant qui s’écoule à travers le tube d’écoulement.

12. Procédé pour doser l’écoulement d’un fluide multiphasé utilisant un système selon l’une quelconque des revendications 1 à 8, le procédé comprenant les étapes consistant à :

mettre en vibration un tube d’écoulement alors que le fluide multiphasé s’écoule à travers le tube d’écoulement ;

utiliser une paire de capteurs (46a, 46b) pour déterminer un mouvement du tube d’écoulement à différents emplacements sur le tube d’écoulement et délivrer un signal de capteur depuis chaque capteur (46a, 46b) ;

mesurer une pression du fluide multiphasé ;

mesurer une température du fluide multiphasé ;

mesurer des quantités relatives d’une multiplicité de liquides différents dans le fluide multiphasé ; et déterminer un débit sur la base d’une différence de phase entre les signaux de capteur ;

caractérisé par l’étape consistant à déterminer une quantité de gaz dissous dans le fluide multiphasé en utilisant la température et la pression du fluide et les quantités relatives de la multiplicité de liquides dans le fluide multiphasé, dans lequel la quantité de gaz dissous est déterminée en accord avec un modèle de solubilité de gaz, le dit modèle de solubilité incluant un facteur d’accord ajustable qui permet d’accorder le modèle de solubilité de gaz pour tenir compte du fait que les fluides dérivés du pétrole réels ne se comportent pas toujours de la même manière.
13. Procédé selon la revendication 12 utilisant un système selon l’une quelconque des revendications 1 à 8, et/ou un groupe de mesure "skid" de quantité nette de pétrole et de gaz selon l’une quelconque des revendications 9 à 11.
FIG. 3

- Flow Rate
- Density
- Dissolved Gas
FIG. 7

Density Drop against Nominal GVF, assuming no dissolved gas - Water Continuous

Density Drop (%)  
0  10  20  30  40  50  60  70
0  10  20  30  40  50  60  70
Nominal Gas Volume Fraction (%)
FIG. 9

Density Error against Nominal GVVF assuming no dissolved gas - Water Continuous
FIG. 11

Density Drop against Nominal GVF, assuming no dissolved gas - Oil Continuous
FIG. 12

Mass Flow Error against Nominal GVF, assuming no dissolved gas - Oil Continuous
FIG. 13
FIG. 14

"Raw Mass Flow Liquid" \( \dot{m}_{m,raw} \)
Raw measurement

"Density Drop" \( \rho_{\text{drop}} \)
Involves \( \rho_{m,raw}, \rho_w, \rho_o \) \( W_{\text{cut}} \)

"Water Cut" \( W_{\text{cut}} \)
Measured, possibly corrected

Mass flow

Density

"Mass Flow Correction" \( c_{mf} \)
For \( \dot{m}_{em} \)

"Density Correction" \( c_{dc} \)
Then \( \rho_{m,ac}, \dot{m}_{o,ac}, \dot{m}_{w,ac}, \dot{m}_{g,ac} \)

FIG. 15

Train Neural Network
- Get Raw measurements
- Align with reference data
- Calculate inputs
  - Steps 1-6
- Calculate targets
  - Steps 7-12
- Train NN / Get Results
  - Steps 13-14
- Calculate Phase mass flow rates
  - Steps 15-21

Use Neural Network
- Get Raw measurements
- Calculate inputs
  - Steps 1-6
- Apply
  - Steps 13-14
- Calculate Phase mass flow rates
  - Steps 15-21
**FIG. 16**

"Raw Mass Flow Liquid"  \( \dot{m}_{m,\text{raw}} \)

"Density Drop" \( \rho_{\text{drop}} \)
Involves
\( \rho_{m,\text{raw}} \rho_w \rho_o \ W_{\text{cut}} \)

"Water Cut" \( W_{\text{cut}} \)
Measured, possibly corrected

Mass flow

"Density Correction" \( C_{dc} \)
For \( \rho_{m,sc} \), incorporated gas split, then
\( \rho_{m,sc} \ m_{o,sc} \ m_{w,sc} \ m_{g,sc} \)

"Mass Flow Correction" \( C_{mf} \)
For \( \dot{m}_{tm} \)

**FIG. 17**

**Train NN**

- Get Raw measurements
- Align with reference data
- Calculate NN inputs
  Steps 1-6
- Calculate NN targets
  Steps 7-17
- Train NN / Get Results
  Steps 18-19
- Calculate Phase mass flow rates
  Steps 20-30

**Use NN**

- Get Raw measurements
- Calculate NN inputs
  Steps 2-6
- Apply NN
  Steps 18-19
- Calculate Phase mass flow rates
  Steps 20-30
REFERENCES CITED IN THE DESCRIPTION

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