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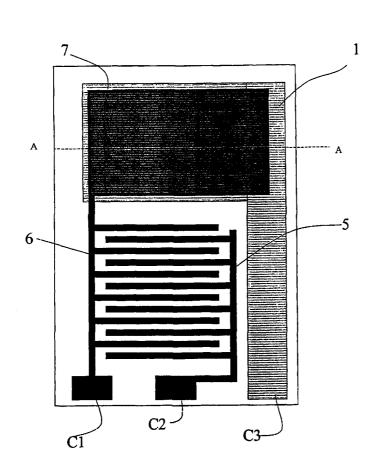
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(54) Title: METHOD AND APPARATUS FOR MEASURING WATER CONTENT



(57) Abstract: The invention relates to a method and apparatus for measuring the water content of a liquid. In the method, a property of the liquid is measured electrically for one parameter. According to the invention, the properties of the liquid are measured at least substantially simultaneously by another electrical method, too, whereby the properties of the liquid are measured using both a relative-value and an absolute-value measurement method.

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METHOD AND APPARATUS FOR MEASURING WATER CONTENT

The invention relates to a method according to the preamble of claim 1 for measuring the water content of a liquid.

The invention also relates to an apparatus for measuring the water content of a liquid.

US Pat. No. 5,331,287 describes a sensor, wherein interdigitated electrodes (finger electrodes) are coated with a conducting polymer. Water contained in the oil hydrates the polymer and thus modifies its conductivity. Also detects possible acids by way of protonation.

US Pat. No. 5,642,098 discloses a ring oscillator circuit, wherein electrical properties of the oil are measured with a number of measurement heads which measure the same parameter.

US Pat. No. 5,644,239 measures the electrical conductivity of a liquid (oil) at two elevated temperatures. The technique may be complemented with a possible optical measurement of oil opacity. A "figure of quality" may then be computed for the oil from these parameters.

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US Pat. No. 5,656,767 describes a sensor system for measuring the change of an electrical parameter value (e.g., capacitance) in oil as a function of time. The same oil at a clean (dry) state may be used as a reference value. The same technique may be varied in multiple ways, e.g., by heating the oil sample.

Conventional techniques are handicapped in many aspects. Common methods for sensing absolute water volume content over the entire range of 0-100 % are the measurement of the dielectric coefficient and measurement of IR absorption. Both of these methods have in common that they require a zeroing step of the measurement system, whereby the reading must be reset to zero water content when the sensor is

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brought to measure an entirely dry (water-free) liquid. This step can be accomplished as a discrete zeroing operation or by using a sample of entirely dry oil in the sensor as a reference.

An additional complication arises therefrom that such a zero setting is typically dependent on the temperature.

Also other factors besides the water content may affect the zero-value with the aging of the liquid.

Methods measuring the absolute water content are favored at high water contents (in the order of several per cent).

At lower water contents, problems generally arise from the marginal detection threshold and offset uncertainty (error of zero setting).

A relative value (aw) measurement method gives information on the water content value in relation to that of a fully saturated situation. However, a conversion to the volume percentage value of absolute water content remains undefined unless a conversion factor for the liquid being measured is known. The aw measurement method is suitable for use at low water content levels (nonsaturated and not emulsified), whereby the measurement has a sufficiently high sensitivity. Moreover, the method is free from zeroing problems.

It is an object of the present invention to overcome the drawbacks of the abovedescribed techniques and to provide an entirely novel type of method and apparatus for measuring the water content of a liquid.

The goal of the invention is achieved by way of measuring the water content of the oil/liquid using two different methods simultaneously, whereby the measurement technique is based on an absolute value measurement method complemented with a relative value measurement method.

More specifically, the method according to the invention is characterized by what is stated in the characterizing part of claim 1.

- Furthermore, the apparatus according to the method is characterized by what is stated in the characterizing part of claim 9.
 - The invention offers significant benefits.
- The combination of an absolute value measurement method with a relative value measurement method (aw-type of measurement) makes it possible to eliminate the need for a zeroing step in the absolute value measurement.
- By way of performing a sequence of rapidly repeated measurements at different temperatures, it is also possible to eliminate errors caused by temperature variations.
 - In the following, the invention will be examined in more detail by means of exemplifying embodiments illustrated in the appended drawings, in which:
- FIG. 1 shows a top view of an embodiment of an electrode structure according to the invention; and
 - FIG. 2 shows a sectional view of the sensor structure of FIG. 1 along line A-A.
- As shown in FIGS. 1 and 2, the embodiment according to the invention can be implemented using a structure, wherein onto the surface of a substrate 4 is formed a combination of three electrodes. The surface of the substrate 4 has directly deposited thereon a pair 3 of bottom electrodes formed by electrodes 5 and 6. In the illustrated case, the electrodes are shaped into finger electrodes, more specifically disposed as interdigitated electrodes, whereby the interelectrode surface is maximized by using an electrode structure having the finger electrodes displaced between each other. The length of the adjacent edges of the interdigitated electrodes 5 and 6 is equivalent to

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the area of superimposed electrodes in a planar capacitor. The width and interelectrode gap of the electrodes 5 and 6 typically are in the range of 5 to 500 micrometers. In addition to its interdigitated finger electrode portion, the electrode 6 has a planar area 7. Furthermore, the electrodes 5 and 6 include contact pad areas C1 and C2 for connection of the electrodes to measurement means. Onto the bottom electrode pair 3 is deposited a polymer layer 2 having a thickness of 0.5-5 micrometer typical. The area of the polymer layer 2 may also be extended over the interdigitated finger electrode structure 5 and 6, whereby it also functions as a passivation layer and reduces the effect of conducting particles contained in the liquid sample on the measurement result. Onto the polymer layer 2 is deposited a water-permeable top electrode 1 with a contact pad area C3 for external connections. The top electrode 1 is aligned above the rectangular solid area 7 of the electrode 6 in order to form a planar capacitor structure.

- The structure illustrated in FIGS. 1 and 2 is utilized as follows. The dielectric coefficient of the liquid under measurement is measured over the contacts C1 and C2 of the electrodes 5 and 6. Respectively, the relative value measurement is performed over the electrodes 6 and 1, that is from the contacts C1 and C3.
- Advantageously, the liquid whose water content is measured is oil, but the water content measurement according to the invention may also be performed on other liquids such as a hydraulic fluid, gasoline or a coolant as well.

According to the invention, the measurement of the activity of water can be carried out as follows.

Firstly, it must be noted that the water activity measurement is only an exemplifying embodiment of the relative value measurement technique according to the invention.

As shown in FIGS. 1 and 2, the sensor may comprise a polymer layer 2 deposited between two electrodes as an element, whose absorbency of water is a function of the activity of water in its immediate environment. Typically, such sensors are used

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as relative humidity transducers, for example. This type of measurement method is characterized in that the measurement result indicates the degree of water activity, that is:

$$5 aw = F(ppm/ppm_s(T)), (1)$$

where

ppm = volumetric proportion of water in solution×10⁶

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 $ppm_s = volumetric proportion of water in saturated solution × 10^6$

The function F may for example be assumed to have a linear nature, whereby:

$$aw = ppm/ppm_s(T)$$
 (2)

The value of aw varies from 0 (entirely dry liquid) to 1 (entirely saturated water solution).

Hence, a mere water activity measurement gives information on how close the situation is either to an emulsified state or a water-separation state. However, when the state of emulsification or water separation is reached, the value of water activity becomes very close to 1, whereby no information on the state of the liquid can be obtained. Nevertheless, the method can render at values of aw < 0.9 in a very sensitive manner such oil/liquid-independent information that, for example, is related to the lubrication properties of the measured sample.

At room temperature, the value of ppm_s may vary from 20 ppm for a basic oil having no additives to values above 10,000 ppm for lubrication oils rich with additives.

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Next, the measurement technique of the dielectric coefficient is elucidated in more detail. The measurement of the dielectric coefficient is only one exemplifying

embodiment of the absolute-value concentration measurements possible within the scope of the invention.

In the measurement of the dielectric coefficient of a liquid, the sensor may be implemented either using the interdigitated electrode (finger-electrode) structure according to FIGS. 1 and 2 or, alternatively, formed into a coaxial pair of electrodes brought into contact with the liquid to be measured. The output signal of the sensor is dependent on both the dielectric coefficient and water content of the liquid under measurement:

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$$\varepsilon_{\rm r} = \varepsilon_0 + \rm F(ppm) \tag{3}$$

where

15 ε_0 = dielectric coefficient of entirely dry liquid,

F(ppm) = a function dependent on the water content. Over a limited range of water content, the function may be assumed to be linear, that is:

$$20 \varepsilon_{r} = \varepsilon_{0} + a \times ppm (4)$$

where

a = constant independent of the liquid type.

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An advantage of the dielectric coefficient measurement technique is that it covers the entire possible measurement range from 0 to 100 vol-% water. A disadvantage of the method is that its sensitivity at the low end of water content (where the greatest interest usually lies) is low and that the value of ε_0 must be known. Typically, the measurement system is calibrated using dehydrated oil as a standard.

If both of the above-described measurement techniques are employed simultaneously, a number of different calibration techniques can be used for a given liquid.

If the liquid is known by the value of its $ppm_s(T)$, the following approach is possible:

First, the values of aw and ε_r are measured simultaneously. If aw is less than 1, formulas 2 and 4 may be assumed to be valid.

Then, the combination of formulas 2 and 4 can be solved for ε_0 and ppm. It is also possible to estimate the value of ppm_s(T) if the type of the liquid is known, whereby the result thus obtained, however, remains slightly more inaccurate.

If the value of ppm_s(T) of the liquid is unknown, the following procedure can be carried out:

The sensor output is measured at two (unknown) water content values. If the value of aw is smaller than 1 in both cases, an equation group of four equations and four unknown variables is obtained, which means the equations can be solved in unique manner. Assigning subindices 1 and 2 for the measurement results of the two sessions, respectively, the following formula can be written:

$$\varepsilon_0 = (aw_2 \times \varepsilon_{r1} - aw_1 \times \varepsilon_{r2})/(aw_2 - aw_1)$$
 (5)

This procedure may also be arranged to take place automatically during the continuous function of the measurement apparatus if the water content of the liquid under measurement varies.

If the measurement data is collected by more than two pairs of values, the unknown terms can be fitted to the data using, e.g., the least squares method.

The benefit of the latter technique is that possible changes in the values of ε_0 or $ppm_s(T)$ due to temperature variations or aging/soiling of the liquid can be compensated for. In fact, the changes detected in ε_0 serve as an indicator of aging in oil.

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If the temperature of an oil sample or, respectively, of a sample flow, is changed so rapidly that the water content of the sample can be assumed to stay substantially unchanged, it is possible to determine the temperature dependence of the dielectric coefficient of an entirely dry liquid by way of measuring essentially simultaneously the ε_r and temperature of the liquid at at least two temperatures.

For example it can be assumed assuming that ε_0 is a linear function with temperature:

$$\varepsilon_{\rm r} = b0 + b1 \times T + a \times ppm \tag{6}$$

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whereby the following formula can be formed

$$\varepsilon_{r}(T2) - \varepsilon_{r}(T2) = b1 \times (T2 - T1) \tag{7}$$

wherefrom the coefficient b1 can be solved. Also in this case, it is possible to collect values at a greater number of temperatures and then fit the measurement results with the help of the least squares method. This technique gives a continuously reliable parameter estimate value for ε_0 as the temperature varies.

Also the temperature dependence of ppm_s(T) can be determined simultaneously.

Over a limited temperature range, a general assumption may be made as:

$$ppm_s(T) = c0 \times e^{(C1 \times T)}$$
(8)

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 $c0 = value \text{ of ppm}_s \text{ at } T = 0 \text{ }^{\circ}\text{C}$

c1 = coefficient of temperature dependence.

Then, the value of c1 can be solved by way of measuring aw at at least two temperatures with the assumption that the water content of the liquid remains unchanged during the measurements:

$$c1 = LN(aw1/aw2)/(T2-T1)$$
 (9)

Advantageously, these two determinations may be carried out simultaneously.

According to a preferred embodiment of the invention, the zeroing operation is performed automatically each time the measurement of aw gives a sufficiently low value. The lower the value of aw the more accurate the zeroing operation. The uncertainty of the method is associated with the value of ppm_s(T) that is dependent on the type of liquid under measurement and thus can be obtained by an "intelligent guess".

According to another preferred embodiment of the invention, two samples of the liquid/oil having different water contents are taken. Then, the measurement system may intentionally be set to measure two samples of different water contents or, alternatively, gradual accumulation of data from the measured process is utilized, whereby the natural variation of water content in the monitored process is availed of. This approach also gives a value for ppm_s(T) on the basis of which it is possible according to the invention to compute from the measured value of aw the correct value of ppm without resorting to an "intelligent guess".

If a continuous data collection from the monitored process is performed using simultaneously a sliding-window technique for "dumping" obsolete data, it is also possible to compensate for changes in both the value of ppm_s and ϵ_0 due to aging of the liquid/oil.

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In the case that the process also is subjected to temperature changes, the temperature dependencies of ppm_s and ϵ_0 can also be resolved.

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What is claimed is:

1. Method for measurement of water content of a liquid, in which method a property of the liquid is measured electrically for one parameter,

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characterized in that

- the properties of the liquid are measured at least substantially simultaneously also by another electrical method, whereby the properties of the liquid are measured using both a relative-value measurement method and an absolute-value measurement method resulting in the measurement of the liquid for its dielectric coefficient and relative water content.
- 2. Method according to claim 1, c h a r a c t e r i z e d in that said measurement is carried out using a capacitive sensor.
 - 3. Method according to any one of foregoing claims, characterized in that said measurement is repeated at two different temperatures in a so rapid succession that the water content of the liquid may be assumed to stay at least substantially constant.
 - 4. Method according to any one of foregoing claims, c h a r a c t e r i z e d in that changes in the results of water content measurement due to aging of the liquid are compensated for by virtue of using only the most recent data of the measurement history for the compensation for changes in the response of the measurement system.
 - 5. Method according to any one of foregoing claims, c h a r a c t e r i z e d in that the aging of said liquid, advantageously oil, is indicated on the basis of changes in the value of ε_0 .

6. Method according to any one of foregoing claims, c h a r a c t e r i z e d in that, in the measurement of relative water content, an auxiliary medium is used for absorbing thereto the water contained in the liquid under measurement.

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- 7. Method according to claim 6, c h a r a c t e r i z e d in that said auxiliary medium is a thin-film polymer layer.
 - 8. Method according to claim 6 or 7, c h a r a c t e r i z e d in that the water content of said auxiliary medium is determined by way of measuring its dielectric coefficient.
 - 9. Apparatus for measurement of the water content of a liquid, said apparatus comprising one electrical sensor means (5, 6 or 1, 6) for measuring the water content of a liquid, c h a r a c t e r i z e d in that

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the apparatus also includes a second electrical sensor means (1, 6 or 5, 6) for measuring the water content of a liquid, said second sensor means (1, 6 or 5, 6) measuring a different parameter than that measured by said first electrical sensor means (5, 6 or 1, 6), said sensor means being such that one of them measures the properties of the liquid by a relative-value measurement method and the other by an absolute-value measurement method, whereby one sensor means (5, 6) is sensitive to changes in the dielectric coefficient and the other sensor means is sensitive to the relative water content.

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- 10. Apparatus according to claim 9, c h a r a c t e r i z e d in that said sensor means (5, 6) sensitive to changes in dielectric coefficient is formed by two interdigitated finger electrodes (5, 6).
- 11. Apparatus according to claim 9 or 10, c h a r a c t e r i z e d in that one electrode (6) of the sensor pair adapted to perform the measurement of dielectric

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coefficient also forms a part of the measurement electrode pair (1, 6) adapted to perform the measurement of the relative water content.

- 12. Apparatus according to claim 9, c h a r a c t e r i z e d in that the sensor means sensitive to changes in the dielectric coefficient is formed by a coaxial structure, wherein one electrode is formed by a center pin and the jacket has a net-like structure and is permeable to water.
- 13. Apparatus according to claim 9, c h a r a c t e r i z e d in that the sensor means adapted for measuring the relative water content contain an auxiliary medium capable of absorbing water contained in the liquid under measurement.
 - 14. Apparatus according to claim 13, characterized in that said auxiliary medium is a thin-film polymer layer.
 - 15. Apparatus according to claim 13 or 14, characterized in that the apparatus contains means adapted to measure the dielectric coefficient of said auxiliary medium whereupon the relative water content of said auxiliary medium can be determined.

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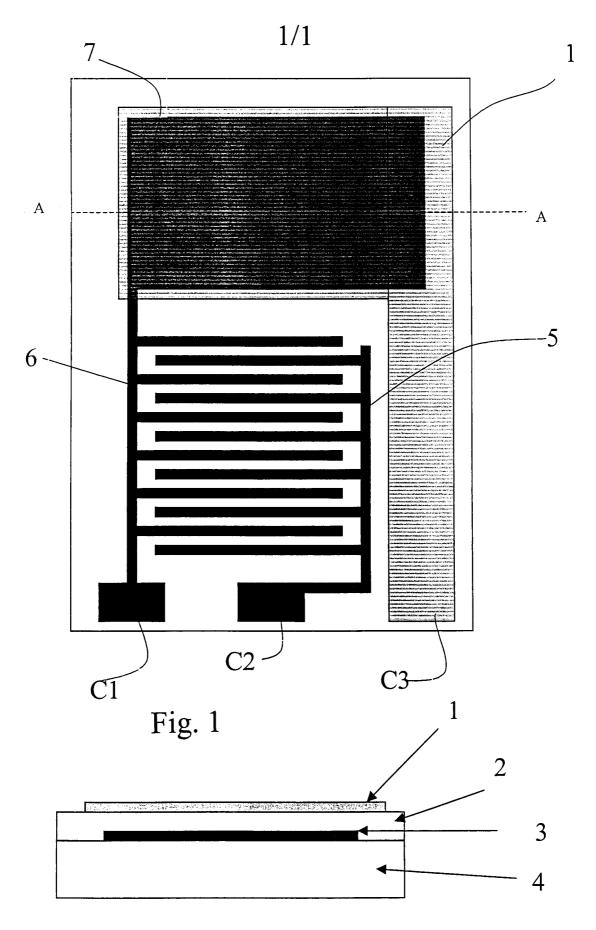


Fig. 2

INTERNATIONAL SEARCH REPORT

International application No.

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A. CLASSIFICATION OF SUBJECT MATTER

IPC7: G01N 27/02, G01N 27/22, G01N 33/28
According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPODOC, WPI

C. DOCUMENTS CONSIDERED TO BE RELEVANT

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See patent family annex.

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Date of the actual completion of the international search

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/FI 00/00542

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No. PCT/FI 00/00542

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