

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
24 March 2005 (24.03.2005)

PCT

(10) International Publication Number  
**WO 2005/025399 A2**

(51) International Patent Classification<sup>7</sup>: **A61B**

(21) International Application Number:  
PCT/IL2004/000835

(22) International Filing Date:  
12 September 2004 (12.09.2004)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:  
60/502,212 12 September 2003 (12.09.2003) US  
60/502,210 12 September 2003 (12.09.2003) US  
60/553,142 16 March 2004 (16.03.2004) US  
60/581,376 22 June 2004 (22.06.2004) US

(71) Applicant (for all designated States except US): **OR-NIM MEDICAL LTD.** [IL/IL]; P.O. Box 32372, 16 Bavli Street, 61323 Tel Aviv (IL).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **BALBERG,**

Michal [IL/IL]; 19 Nof Harim, 96190 Jerusalem (IL). **PERY-SCHECHTER, Revital** [IL/IL]; 52 Hatizmoret Street, 75582 Rishon Lezion (IL). **OLSHANSKY, Michal** [IL/IL]; 16 Bavli Street, 62331 Tel Aviv (IL).

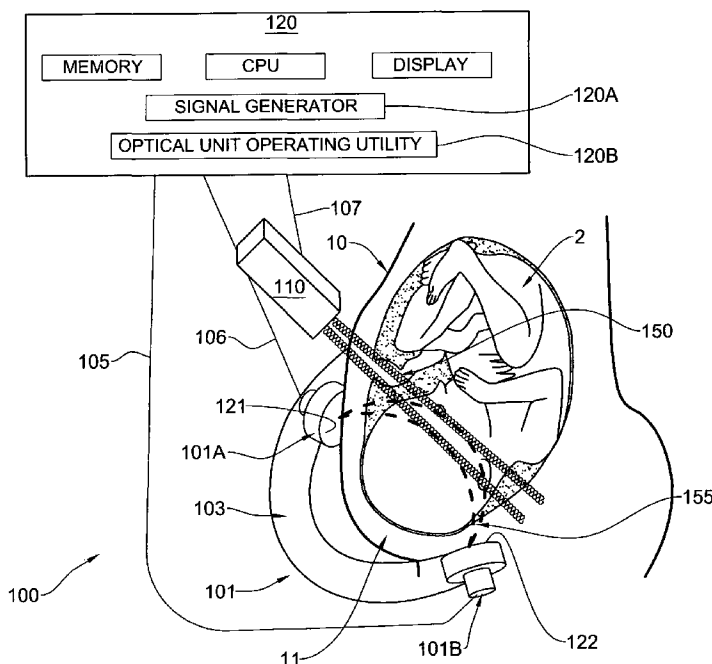
(74) Agent: **REINHOLD COHN AND PARTNERS**; P.O.B. 4060, 61040 Tel-Aviv (IL).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM,

[Continued on next page]

(54) Title: METHOD AND APPARATUS FOR NONINVASIVELY MONITORING PARAMETERS OF A REGION OF INTEREST IN A HUMAN BODY



(57) Abstract: A method and system are presented for use in noninvasive monitoring at least one parameter of a region of interest in a human body. The system comprises a measurement unit and a control unit. The measurement unit comprises an optical unit having an illumination assembly configured to define at least one output port for illuminating light, and a light detection assembly configured to define at least one light input port for collecting light and to generate measured data indicative of the collected light; and an acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range. The measurement unit is configured and operable to provide an operating condition such that the acoustic waves of the predetermined frequency range overlap with an illuminating region within the region of interest and substantially do not overlap with a region outside the region of interest, and that the detection assembly collects light scattered from the region of interest and light scattered from the region outside the region of interest. The measured

data is thus indicative of scattered light having both ultrasound tagged and untagged light portions, thereby enabling to distinguish between light responses of the region of interest and the region outside the region of interest. The control unit is connectable to the optical unit and to the acoustic unit to operate these units, and is responsive to the measured data and preprogrammed to process and analyze the measured data to extract therefrom data indicative of a light response of the region of interest and determine the at least one desired parameter.

WO 2005/025399 A2



ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

**Published:**

- *without international search report and to be republished upon receipt of that report*

## **METHOD AND APPARATUS FOR NONINVASIVELY MONITORING PARAMETERS OF A REGION OF INTEREST IN A HUMAN BODY**

### **FIELD OF THE INVENTION**

This invention relates to a method and apparatus for noninvasive monitoring of parameters of a region of interest in a human body, such as oxygen saturation and/or concentration of analyte(s) in blood.

### **5 BACKGROUND OF THE INVENTION**

Monitoring of the well-being of the fetus inside the uterus is very important and is carried periodically with respect to various parameters of the fetus. One of the important parameters to be monitored is oxygen saturation. Various techniques have been developed to enable noninvasive measurements of  
10 oxygen saturation.

For example, U.S. Patent No. 5,494,032 discloses an oximeter for reliable clinical determination of blood oxygen saturation in a fetus. This technique utilizes a multiple frequency light source which is coupled to an optical fiber. The output of the fiber is used to illuminate blood containing tissue of the fetus.  
15 The reflected light is transmitted back to the apparatus where the light intensities are simultaneously detected at multiple frequencies. The resulting spectrum is then analyzed for determination of oxygen saturation. The analysis method uses multivariate calibration techniques that compensate for nonlinear spectral response, model interfering spectral responses and detect outlier data with high  
20 sensitivity.

A pulse oximetry based technique for determining the fetal arterial blood oxygenation is disclosed in the following article: A. Zourabian et al., "Trans-abdominal monitoring of fetal arterial blood oxygenation using pulse oxymetry", Journal of Biomedical Optics, Vol. 5, No. 4, October 2000, pp. 391-405.

- 2 -

U.S. Patent No. 6,041,248 describes a method and apparatus for frequency encoded ultrasound-modulated optical tomography of dense turbid media. The apparatus includes a function generator producing a frequency sweep signal which is applied to an ultrasonic transducer. The ultrasonic transducer produces  
5 ultrasonic wave in a turbid medium. Coherent light from a laser is passed through turbid medium where it is modulated by the ultrasonic wave. A photomultiplier tube detects the light which passes through the turbid medium. The signal from the photomultiplier tube is fed to an oscilloscope and then to a computer where differences in light intensity at different frequencies can determine the location of  
10 objects in the turbid medium.

The conventionally used techniques for monitoring the well-being of the fetus inside the uterus utilize measuring the fetal-heart-rate (FHR) by placing sensors on the skin of the mother's abdomen proximal to the fetus. These sensors transmit acoustic waves and provide data indicative of the Doppler shift of an  
15 acoustic wave reflected from the fetal heart, enabling calculation of the heart rate based on this shift. A normal fetal-heart-rate (FHR) pattern is usually associated with the delivery of a normal well-oxygenated infant. However, a non-reassuring FHR is not always associated with the delivery of a compromised infant.

In the case of non-reassuring FHR, the fetal blood oxygen saturation level  
20 can be measured only post membrane rupture by either fetal scalp sampling, which measures the pH level of the fetal blood, or by attaching a pulse oximeter to the presenting part of the fetal head during labor. Both of these methods are performed following the rupture of membranes where the fetal scalp and/or cheeks can be reached.

Another important procedure to be done to monitor the well-being of the fetus consists of assessing the maturity of fetal lungs, which is one of the major concerns of pre-term deliveries. If the baby is delivered and the lungs are not mature, the baby may develop Respiratory Distress Syndrome (RDS), which can result either in fetal death or in long-lasting periods of repeated respiratory  
30 difficulty.

- 3 -

In cases where intervention is considered in the course of pregnancy (such as caesarean section or induction of labor) and there is a need to assess the maturity of the lungs, amniotic fluid is drained. Measuring phospholipids in amniotic fluid as the lecithin/sphingomyelin ratio using the thin-layer chromatography method has been the established clinical procedure for predicting fetal lung maturity. Although it is the clinical "gold standard" method, it remains a time-consuming process, has a large intralaboratory and interlaboratory coefficient of variation, and requires expertise. In addition, the procedure of amniotic fluid drainage itself is invasive and suffers a small risk of abortion. Additional techniques that are used for assessing lung maturity levels include measuring the number of lamellar bodies in a volume of amniotic fluid, measuring the prostaglandin level in amniotic fluid and measuring the fluorescence polarization of a sampled amniotic fluid.

When a fetus is acutely distressed, for example as a result of strangulation by the umbilical cord, the bowel content, meconium, may be passed into the amniotic fluid (AF). Assessment of meconial contamination of AF is important in the management of late pregnancy. It appears in nearly one third of all fetuses by 42 weeks of gestation. In cases where the fetus gasps during delivery, inhaling the sticky meconium into the upper respiratory tract results in partial airways obstruction. Meconium aspiration syndrome occurs in 0.2% to 1% of all deliveries and has a mortality rate as high as 18%. The disease is responsible for 2% of all prenatal deaths.

To date, meconium stained amniotic fluid is diagnosed following the rupture of membranes, when the amniotic fluid is drained. However, in cases where the fetus head is tightly fitted in the pelvis, the amniotic fluid is not drained out resulting in misdiagnosis of the potential harmful outcome to the respiratory tract.

## SUMMARY OF THE INVENTION

There is accordingly a need in the art to facilitate noninvasive monitoring of parameters of a region of interest in a human body, by providing a novel noninvasive method and apparatus.

5       The technique of the present invention provides for monitoring blood and/or tissue parameters and/or parameters of fluids of a region of interest in a human body, for example the concentration of an analyte in blood, fluid reservoirs or tissue regions in a human body; as well as fetus condition in utero (e.g., the fetal oxygen saturation level as well as the concentration of analyte in  
10 fetal blood; and the maturity of fetal lungs and the presence of meconium, prior to membrane rupture).

It should be understood that the term "*region of interest*" signifies a tissues region or a fluid contained in a reservoir or cavity inside a body. The region of interest may be a fetus region (e.g., fetus head), amniotic fluid, vicinity  
15 of blood vessels, etc. The term "*fetus-related region of interest*" used herein signifies either one of fetus and amniotic fluid regions.

The main idea of the present invention consists of non-invasively monitoring the optical properties of a region of interest in a human (or animal) body utilizing the principles of ultrasound tagging of light, which in the present  
20 invention is aimed at distinguishing between optical responses of the region of interest in the selected volume (e.g., fetus, amniotic fluid, blood vessel) and the surroundings outside the region of interest; and/or significantly improving pulse oximetry based measurements.

According to one aspect of the present invention, a body portion (e.g., the  
25 abdomen of a pregnant woman) containing a region of interest (fetus) is irradiated with light (e.g., of at least two different wavelengths) and is irradiated with acoustic waves, in a manner to ensure optimal operating condition for measurements. This operating condition is such that the illuminating light and acoustic waves overlap within the region of interest and thus light scattered from  
30 the region of interest is "tagged" by acoustic waves (i.e., the frequency of light is

- 5 -

modulated by the frequency of the acoustic waves) while substantially do not overlap in a region outside the region of interest, and to ensure that detected light includes a portion of light scattered by the region of interest and tagged by acoustic waves and a portion of untagged light scattered by the region outside the  
5 region of interest. This allows for distinguishing between light responses of the region of interest and its surroundings (e.g., fetus and maternal tissues). It should be understood that the term “acoustic wave” refer to acoustic radiation of either one of the following type: continuous wave, pulses, bursts.

It should be understood that for the purposes of the present invention the  
10 term “*maternal tissues*” used herein refers to all the tissues within a region surrounding the fetus-related region of interest (fetus itself or amniotic fluid containing the fetus). Considering the region of interest is fetus, the term “*maternal tissues*” refers to maternal tissues, amniotic fluid, and uterine wall.

According to another aspect of the present invention, the above operating  
15 condition is used in pulse oximetry measurements for determining oxygen saturation level in a region of interest (in mammalian blood and/or blood vessels). Measured data that needs to be analyzed is, for example, in the form of a power spectrum of ultrasound-tagged light response of the region of interest, which is practically insensitive to minor movements of regions outside the region  
20 of interest, while pure pulse oximetric measurements are highly sensitive to such movements.

Preferably, the present invention in either of its aspects utilizes obtaining of measured data in the form of time dependent and/or wavelength dependent variations of ultrasound-tagged light signals for at least two wavelengths of  
25 illuminating light.

The present invention provides for non-invasively determining such parameters as oxygen saturation level in the region of interest (e.g., fetus, blood vessel), concentration of a substance or a structure within the region of interest (e.g., fetus, amniotic fluid), the presence and concentration of lamellar bodies in  
30 amniotic fluid for determining the level of lung maturity of the fetus, the

- 6 -

presence and/or concentration of meconium in the amniotic fluid, presence and/or concentration of blood in the amniotic fluid; as well as for noninvasive monitoring the optical properties of other extravascular fluids such as pleural, pericardial, peritoneal (around the abdominal and pelvis cavities) and synovial  
5 fluids. It is important to note that according to the invention, acoustic (ultrasound) radiation used for measurements needs not be focused, since the measurements utilize ultrasound tagging solely for the purposes of distinguishing between light responses of the region of interest and its surroundings, and/or for increasing signal to noise ratio of ultrasound tagging based measurements.

10 The present invention utilizes the principles of oximetry for processing the measured data. Accordingly, the illumination with at least two different wavelengths is applied. Preferably, the light response signals are collected over a time period larger than a heart beat, and the principles of pulse oximetry are used to determine the oxygen saturation.

15 Preferably, a measurement unit (an illumination assembly, a light detection assembly, and an ultrasound transducer arrangement) is placed in close contact with the respective body portion (e.g., maternal tissues being in contact with amniotic sac containing a fetus). As indicated above, the illumination assembly is configured and operable to illuminate the body portion with at least  
20 two wavelengths. The ultrasound transducer arrangement is configured and operable to transmit acoustic waves into the same volume from which the light detector collects scattered light.

The light detection assembly may be oriented for collecting both back scattered light and forward scattered light.

25 Preferably, the present invention utilizes ultrasound imaging, carried out prior to measurements and aimed at determining optimal positioning of the illumination assembly, light detection assembly and acoustic waves propagation to thereby provide the operating condition for measurements. The ultrasound imaging may and may not utilize the same ultrasound transducer arrangement  
30 that is used for measurements. Preferably, the invention also provides for using



ultrasound radiation for determining such parameters of blood in the region of interest (e.g., fetus) as blood flow, tissue velocity profile, etc. To this end, reflections of ultrasound radiation from the irradiated region are analyzed using any known suitable Doppler-based techniques. The incident ultrasound radiation  
5 may be in the form of continuous waves or pulses (gates).

According to an embodiment of the present invention maternal oxygen saturation level is detected using the same apparatus being used to measure fetal oxygen saturation level.

The present invention can be used for measuring in more than one fetus  
10 presented inside the uterus. In this case, the oxygen saturation level (or other fetal parameters) of each fetus is measured independently using the same apparatus; or several different apparatuses, one for each fetus, all associated with the same control unit (data processing and analyzing utility). Each fetus is located using an ultrasound imaging system, and the optimal arrangement of the light sources,  
15 detectors and ultrasound transducers is determined for monitoring the oxygen saturation level of each fetus.

There is thus provided according to one aspect of the invention a monitoring system for use in non-invasively monitoring at least one parameter of a region of interest in a human body, the system comprising:

- 20 - a measurement unit comprising an optical unit having an illumination assembly configured to define at least one output port for illuminating light, and a light detection assembly configured to define at least one light input port for collecting light scattered from the illuminated body portion and to generate measured data indicative of the collected light; and an  
25 acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range; the measurement unit being configured and operable to provide an operating condition such that the acoustic waves of the predetermined frequency range overlap with an illuminating region within the region of interest and substantially do not overlap with a region  
30 outside the region of interest, and that the detection assembly collects light

- 8 -

scattered from the region of interest and light scattered from the region outside the region of interest, the measured data being thereby indicative of scattered light having both ultrasound tagged and untagged light portions, thereby enabling to distinguish between light responses of the region of interest and the region outside the region of interest;

5

- a control unit, which is connectable to the optical unit and to the acoustic unit to operate these units, the control unit being responsive to the measured data and preprogrammed to process and analyze the measured data to extract therefrom a data portion associated with the light response of the region of interest and determine said at least one parameter of the region of interest.

10

According to another aspect of the invention, there is provided a monitoring system for use in non-invasively monitoring at least one parameter of a region of interest in a human body, the system comprising:

15

- a measurement unit comprising an optical unit configured for generating illuminating light and for collecting light and generating measured data indicative of the collected light; and an acoustic unit configured to generate unfocused acoustic waves of a predetermined ultrasound frequency range; the measurement unit being configured and operable to provide an operating condition such that the acoustic waves of the predetermined frequency range overlap with an illuminating region within the region of interest while substantially not overlapping within a region outside the region of interest, and that the detection assembly collects light scattered from the region of interest and light scattered from the region outside the region of interest, the measured data being thereby indicative of scattered light having both ultrasound tagged and untagged light portions, thereby enabling to distinguish between light responses of the region of interest and the region outside the region of interest;

20

25

- a control unit, which is connectable to the optical unit and to the acoustic unit to operate these units, the control unit being responsive to the

30

- 9 -

measured data and preprogrammed to process and analyze the measured data to extract therefrom a data portion associated with the light response of the region of interest and determine said at least one parameter of the region of interest.

5 According to yet another aspect of the invention, there is provided a system for use in noninvasive monitoring at least one parameter of a region of interest in a human body, the system comprising:

- 10 - a measurement unit comprising an optical unit having an illumination assembly configured to define at least one output port for illuminating light, and a light detection assembly configured to define at least one light input port for collecting light and to generate measured data indicative of the collected light; and an acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range;
- 15 - a control unit, which is connectable to the optical unit and to the acoustic unit to operate these units, the control unit being responsive to the measured data and preprogrammed to process and analyze the measured data to extract therefrom data indicative of a light response of the region of interest and determine the at least one desired parameter, the control unit being operable to provide optimal positioning of the optical unit and  
20 the acoustic unit to satisfy an operating condition for measurements, said operating condition being such that acoustic waves of the predetermined frequency range and illuminating light overlap within the region of interest and substantially do not overlap in a region outside the region of interest, and in that the detection assembly collects the light scattered from  
25 the region of interest and light scattered from the region outside the region of interest; the measured data being thereby indicative of scattered light having both ultrasound tagged and untagged light portions, thereby enabling to analyze the measured data to extract therefrom a data portion associated with the light response of the region of interest and determine  
30 said at least one parameter of the region of interest.

- 10 -

According to yet another aspect of the invention, there is provided a probe device for use in a system for monitoring at least one parameter of a region of interest in a human body, the probe comprising: a support structure configured to contact a body portion, said support structure carrying an array of at least two  
5 light output ports arranged in a spaced-apart relationship and being optically coupled to a light source assembly, an array of light input ports arranged in a spaced-apart relationship and being optically coupled to a light detection assembly, and at least one acoustic output port of an acoustic unit, the arrangement of the light ports and the acoustic unit being such as to allow  
10 selection of at least one of said light output ports, at least one of the light input ports and at least one of the acoustic output ports such that acoustic waves of a predetermined frequency range coming from said at least one selected acoustic output port and illuminating light coming from said at least one selected light output port overlap within a region of interest in the body, and in that said at least  
15 one light input port collects light scattered from the overlapping region and light scattered from outside the region of interest.

According to yet another aspect of the invention, there is provided a probe device for use in a system for monitoring at least one parameter of a region of interest in a human body, the probe comprising: a support structure configured to  
20 contact a body portion, said support structure carrying at least one light output port optically coupled to a light source assembly, at least two light input ports optically coupled to a light detection assembly, and at least one acoustic output port of an acoustic unit, the arrangement of the light and acoustic ports being such as to allow selection of the light and acoustic ports for measurements such  
25 that, with the selected ports, acoustic waves of a predetermined frequency range and illuminating light overlap within a region of interest in the body and that the at least one light input port collects light scattered from the overlapping region and light scattered from outside of the region of interest.

According to yet another aspect of the invention, there is provided a probe  
30 device for use in a system for monitoring at least one parameter of a region of

- 11 -

interest in a human body, the probe comprising: a support structure configured to contact a body portion, said support structure carrying at least two light output port optically coupled to a light source assembly, at least one light input port optically coupled to a light detection assembly, and at least one acoustic output  
5 port of an acoustic unit, the arrangement of the light and acoustic ports being such as to allow selection of the light and acoustic ports for measurements such that, with the selected ports, acoustic waves of a predetermined frequency range and illuminating light overlap within a region of interest in the body and that detected light includes scattered from the overlapping region and light scattered  
10 from outside of the region of interest.

According to yet another aspect of the invention, there is provided a method for use in noninvasive monitoring at least one parameter of a region of interest in a human body, the method comprising: operating an optical unit and an acoustic unit so as to provide that ultrasound waves of a predetermined  
15 frequency range and illuminating light overlap within the region of interest and substantially do not overlap with a region outside the region of interest, thereby producing measured data indicative of collected light including scattered light having ultrasound tagged and untagged light portions, thereby enabling extraction of a light response of the region of interest from all the other light  
20 portions in the collected light.

According to yet another aspect of the invention, there is provided a method for use in noninvasive monitoring oxygen saturation level, the method comprising: applying ultrasound tagging of light in pulse oxymetric measurements, obtaining measured data indicative of time dependent variations  
25 of ultrasound tagged light signals scattered from a region of interest as a function of at least one of time and  $\lambda$ , for at least two different wavelength of illuminating light, and analyzing the measured data to calculate the oxygen saturation level.

According to yet another aspect of the invention, there is provided a method for use in noninvasive monitoring at least one parameter of a region of  
30 interest in a human body, the method comprising:

- 12 -

- providing an optical unit having an illumination assembly configured to define at least one output port for illuminating light; and a light detection assembly configured to define at least one light input port for collecting light, and to generate measured data indicative of the collected light; and  
5 providing an acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range;
- providing an optimal positioning of the optical and acoustic units with respect to each other and with respect to the region of interest, said optimal positioning satisfying an operating condition resulting in that the  
10 ultrasound waves of the predetermined frequency range overlap with the illuminating light within the region of interest, while substantially not overlapping a region outside the region of interest, and in that the detection assembly collects light scattered from the overlapping region and from the region outside the region of interest;
- 15 - operating the optical and acoustic units, when in the optimal positioning, and generating measured data indicative of collected light including scattered light having ultrasound tagged and untagged light portions, thereby enabling extraction of a light response of the region of interest from all the other light portions in the collected light.

20 According to yet another aspect of the invention, there is provided method for use in noninvasive monitoring at least one parameter of a fetus-related region of interest, the method comprising:

- providing an optical unit having an illumination assembly configured to define at least one output port for the illuminating light; and a light  
25 detection assembly configured to define at least one light input port for collecting light, and to generate measured data indicative of the collected light; and providing an acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range;
- providing an optimal positioning of the optical and acoustic units with  
30 respect to each other and with respect to the fetus-related region of

- 13 -

interest, said optimal positioning satisfying an operating condition resulting in that the ultrasound waves of the predetermined frequency range overlap with the illuminating light within the fetus-related region of interest, while substantially not overlapping in a maternal tissues region outside the fetus-related region of interest, and in that the detection assembly collects light scattered from the fetus-related region of interest and from the maternal tissues region;

- operating the optical and acoustic units, when in the optimal positioning, and generating measured data indicative of collected light including scattered light having ultrasound tagged and untagged light portions, thereby enabling extraction from the measured data a data portion indicative of a light response of the fetus-related region of interest.

According to yet another aspect of the invention, there is provided a method for operating a monitoring system configured for noninvasive monitoring at least one parameter of a region of interest in a human body, which system comprises an optical unit and an acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range, the method comprising:

operating the monitoring system to provide an optimal positioning of the optical and acoustic units with respect to each other and with respect to the region of interest to satisfy an operating condition for measurements, said operating condition resulting in that the ultrasound waves of the predetermined frequency range overlap with illuminating light generated by the optical unit within the region of interest, while substantially not overlapping in a region outside the region of interest, and in that a detection assembly of the optical unit collects light scattered from the region of interest and from the region outside the region of interest, thereby obtaining measured data indicative of scattered light having ultrasound tagged and untagged light portions, and enabling extraction from said measured data a data portion indicative of a light response of the region of interest.

The technique of the present invention may be used for noninvasive monitoring of various parameters of human blood and tissue. More specifically, the present invention is useful for monitoring fetal blood conditions and is therefore described below with respect to this application.

## 5 BRIEF DESCRIPTION OF THE DRAWINGS

In order to understand the invention and to see how it may be carried out in practice, preferred embodiment will now be described, by way of non-limiting examples only, with reference to the accompanying drawings, in which:

**Fig. 1A** schematically illustrates a monitoring apparatus according to one  
10 embodiment of the invention for monitoring oxygen saturation of a fetus or any other region of interest in a human or animal body;

**Fig. 1B** exemplifies an operating method of the apparatus of Fig. 1A;

**Fig. 2** schematically illustrates a monitoring apparatus, according to another embodiment of the present invention;

15 **Fig. 3A** schematically illustrates a monitoring apparatus according to the invention configured to be capable of monitoring the fetal and maternal oxygen saturation;

**Fig. 3B** illustrates a flow diagram of the main steps in a method of the invention using the apparatus of Fig. 3A;

20 **Fig. 3C** shows a flow diagram of a specific example of the method of Fig. 3B;

**Fig. 3D** schematically illustrates yet another example of a monitoring system of the present invention configured for monitoring the amniotic fluid condition;

25 **Fig. 4** schematically illustrates a monitoring apparatus, according to yet another embodiment of the present invention;

**Fig. 5** schematically illustrates a monitoring apparatus, according to yet another embodiment of the present invention;



- 15 -

**Figs. 6A and 6B** show bottom and side views, respectively, of a flexible probe according to one embodiment of the invention;

**Figs. 7A and 7B** show bottom and side views, respectively, of a flexible probe according to another embodiment of the invention.

## 5 DETAILED DESCRIPTION OF THE INVENTION

Referring to **Fig. 1A**, there is schematically illustrated a monitoring apparatus, generally designated **100**, constructed and operated as a fetal oxygen saturation monitor according to the invention. It should, however, be understood that the apparatus configuration is suitable for measuring various other  
10 parameters of a fetus **2** (such as the concentration of an analyte in the fetal blood, or the perfusion of an analyte/metabolite in fetal or maternal tissues). It should be understood that the apparatus of the present invention may be used for monitoring blood or tissue parameters of a human being.

The apparatus **100** includes such main constructional parts as a  
15 measurement unit formed by an optical unit **101** including an illumination assembly **101A** and a light detection assembly **101B**; and an acoustic unit including a transducer arrangement **110**. In the present example of Fig. 1A, the detection assembly includes a single detection unit. In this connection, it should be noted that the term "single detection unit" not necessarily signifies a single  
20 detector, but may refer to an array of detectors, provided they are associated with the same location with respect to the illuminated region.

The optical and acoustic units are connectable to a control unit **120**. The control unit **120** is typically a computer system including *inter alia* a power supply, a control panel with input/output functions, a memory utility, a data  
25 presentation utility (e.g., display), a data acquisition assembly, and a data processing and analyzing utility (e.g. CPU). The control unit **120** includes a signal generator (e.g. function generator) **120A** to control the operation of the transducer arrangement **110**, and an appropriate utility **120B** for operating the optical unit **101**. The CPU is preprogrammed for receiving measured data

- 16 -

coming from the detection assembly **101B** and processing this data to determine the desired parameter, e.g., oxygen saturation of the fetus.

In the present example, the optical unit **101** is configured as a portable probe including a support structure **103** carrying at least a part of the illumination assembly **101A** and at least a part of the detection assembly **101B**. The illumination assembly **101A** is preferably configured for generating light of at least two different wavelengths. To this end, the illumination assembly may include at least two light emitters (e.g., laser diodes), one emitting narrow bandwidth photons of a wavelength within the range of 605nm to 805nm and the other emitting photons of a wavelength within the range of 800nm to 1300nm. The illumination assembly **101A** may for example be preprogrammed to produce the different wavelength components at different times, or simultaneously produce wavelength components with different frequency- and/or phase-modulation. Accordingly, the control unit **120** is preprogrammed to identify, in a signal generated by the detection assembly **101B**, the corresponding wavelength of the irradiating light, using time, phase or frequency analysis.

The illumination assembly **101A** may include light emitter(s) carried by the support structure **103** and communicating with the control unit **120** via an output port **121** of the light emitter(s) using wires **106** or wireless signal transmission. Alternatively, the light emitter(s) may be located outside the support structure **103** (e.g., within the control unit **120**) and a light guiding assembly **106** (e.g., optical fibers) is used for guiding light to the output port **121** located on the support structure **103**.

The detection assembly **101B** includes one or more light detectors. This may be a photomultiplier tube, or preferably an image pixel array, e.g., CCD or an array of photodiodes. It should be noted that, for the purposes of the present invention, an input port **122** of the detection assembly **101B** is larger than that used for imaging by means of diffuse light. In diffuse light imaging, localization is achieved by small input ports, otherwise light from a large volume is collected. According to the invention, light collection from a large volume is desired, since

- 17 -

localization is achieved by the ultrasound tagging. Hence, the input port **122** of the detection assembly **101B** is optimized to collect light from a substantially large volume of tissue and/or blood, for example by using large area detectors or CCD cameras or an array of detectors comprising a single input port.

5 As indicated above, the detection assembly **101B** may include two separate detectors or an array of detectors. Each detector may be coupled to a bandpass filter configured for transmitting light of a corresponding one of the wavelengths produced by the illumination assembly **101A**. The bandpass filters may include high-pass, low-pass and bandpass optical filters. Alternatively  
10 narrow bandwidth detectors can be used.

It should be understood that the detector(s) may be accommodated outside the support structure (probe) **103**, e.g., may be located within the control unit **120**, and returned light (light response) may be guided from the input port **122** of the detection assembly via light guiding means **105** (e.g., optical fibers). It should  
15 also be understood that the connectors **105** and **106** may be electric wires connecting the control unit **120** to the illumination assembly and detection assembly located on the probe **103**, or the connection may be wireless.

Thus, generally, the terms "*illumination assembly*" and "*detection assembly*" or "*detection unit*" as carried by a support structure which is brought  
20 in contact with a human body, are constituted by at least light transmitting and receiving ports. Probes (kits) of the present invention including light transmitting and receiving ports and preferably also acoustic ports, will be described further below with reference to Figs. 6A-6B and 7A-7B.

The control unit **120** (its signal generator **120A** and CPU) is connected to  
25 the transducer arrangement **110** using cables **107** and/or using wireless means.

An example of a monitoring method of the present invention, using the apparatus **100**, will now be described with reference to **Fig. 1B**.

Step 1: Prior to performing the actual measurements, an optimal positioning of the assemblies of the optical unit and of the acoustic unit with  
30 respect to a region of interest (fetus) is provided to satisfy an operating condition

- 18 -

for measurements. The operating condition is such that both light (at least a portion of the illuminating light) and the acoustic radiation irradiate the same region (volume) simultaneously, while substantially not overlapping in outside regions (maternal tissues **11**); and that the detection assembly detects light scattered from both the region of interest and regions outside thereof. Preferably, the region where the ultrasound and light overlap is the region of interest (fetus **2**), but generally a region outside the region of interest may be selected to be overlapping region. Generally speaking, the positioning of the optical unit and transducer arrangement with respect to the fetus **2** is such as to enable distinguishing between scattered photons collected from the maternal tissues **11** and from the fetus **2** using ultrasound tagging of light.

This pre-positioning utilizes an ultrasound imaging of the region of interest. To this end, an imaging system of any known suitable configuration may be used, which may utilize the same transducer arrangement **110** used for the measurement process or another ultrasound transducer(s). Ultrasound images of the maternal tissues **11** (e.g. abdomen, uterus) and of the fetus **2** are acquired and analyzed by the control unit **120** (which in this case is installed with a suitable image processing utility) or another appropriately preprogrammed computer system, to determine the optimal positioning of the optical unit **101** (namely the illumination assembly **101A** and the detection assembly **101B**) relative to the fetus and relative to the acoustic unit **110**.

It is important to note that according to the invention, ultrasound tagging is utilized for the purposes of "tagging" a light response from a selected region of interest (fetus), thus enabling processing of detected tagged and untagged light portions to identify the light response of the selected region of interest. This is contrary to the known techniques where ultrasound tagging is used for imaging purposes to enable two and three-dimensional imaging.

The illumination assembly **101A** is preferably placed at the shortest distance to the fetus **2**, preferably to the fetal head. It should be understood that other organs or tissues of fetus **2** may be chosen for measurements as well.

- 19 -

Preferably, the illumination assembly **101A** is placed such that a light path between the illumination assembly **101A** and the fetus **2** is that suffering the least attenuation at the wavelengths chosen for measurements, as compared to the other paths. The distance between the illumination assembly **101A** and the  
5 detection unit **101B** is preferably determined to be at least equal to and preferably larger than the distance between the illumination assembly **101A** and the head of the fetus **2**.

Preferably, the support structure **103** is configured to define various positions for attaching the detection unit **101B** and/or the illumination assembly  
10 **101A** to be at the correct distance between them. For example, these positions may be determined by using a sliding bar (not shown) that is attached to the light detection unit **101B** and can be secured to the support structure **103** using a small screw or a latch. Alternatively, a plurality of light output ports and/or plurality of light input ports are provided on the support structure **103** and the control unit  
15 **120** operates to select the appropriate light source(s) and detector(s) (light output port and light input port) for measurements. This selection is based on the signals generated by each detector and on the geometry of the maternal tissues and the position of the fetus.

Additionally, the illumination assembly **101A** and the detection unit **101B**  
20 are placed such that the light output port **121** of the illumination assembly and the light input port **122** of the detection assembly are in close contact with an outer skin **10** of the maternal abdomen. Optionally, an index matching oil or adhesive is used to reduce reflection of light from the outer skin **10**. The adhesive may be used to secure the apparatus **100** to a specific location on the maternal  
25 abdomen. Alternatively, or additionally, a belt can be used to prevent movement of the apparatus **100**.

Once the position of the illumination assembly **101A** and the detection assembly **101B** is fixed, the ultrasound transducer arrangement **110** is positioned such that acoustic waves **150** generated by the transducer arrangement **110** are  
30 coupled into the maternal abdomen, propagate through uterus and amniotic fluid,

- 20 -

and reach the fetus **2**. For example, in the case the illumination assembly **101A** and detection assembly **101B** are appropriately placed to illuminate and collect light scattered by the fetus head, the transducer **110** is placed such that the acoustic waves **150** propagate through the same region of the head from which  
5 scattered photons **155** are detected by the detection assembly **101B**. The transducer **110** may be fixed to an appropriate location using an ultrasound transmitting adhesive or using gel for acoustic coupling, and optionally a belt for fixing the transducer to one location. Alternatively, the ultrasound transducer arrangement **110** is configured as a phased array transducer producing a focused  
10 beam that is being scanned over a region of skin **10** overlaying the maternal tissues **11**.

Step 2: Having optimally positioned the illumination assembly **101A**, detection unit **101B** and ultrasound transducer arrangement **110**, measurements are taken by appropriately operating the apparatus **100**. The control unit **120**  
15 actuates the illumination assembly **101A** to generate photons **155** of at least two wavelengths. The photons **155** propagate through maternal tissues, through the uterine wall, and reach the fetus **2**. A portion of photons **155** is absorbed by hemoglobin in the fetus blood, and a portion of photons **155** is scattered by tissues and cells of the fetus **2** and of the mother. A portion of the scattered  
20 photons **155** propagates through the maternal tissues **11** and reaches the detection assembly **101B**. The latter collects at least a part of this portion of the scattered photons **155** and generates measured data indicative thereof, i.e., an electric signal in response to the number of photons that are collected at the input port **122** of the detection unit at a specific point in time for each irradiating  
25 wavelength generated by the illumination assembly **101A**.

It should be noted that, in the case the detection assembly **101B** is spaced from the illumination assembly **101A** a distance equal to or larger than twice the minimal distance between the fetus **2** and the illumination assembly **101A**, the detection unit **101B** collects both back and forward scattered photons. In the case  
30 the illumination assembly **101A** includes a laser with a coherence length larger

- 21 -

than the optical path of scattered photons in the tissue, an interference pattern resulting in a speckle image is generated on the input port **122** of the detection assembly. In order to detect and analyze the speckle image, the detection assembly **101B** may include an array of detectors with an individual size  
5 comparable to that of individual speckle. The illumination assembly **101A** may be configured and operable to produce a continuous stream of photons **155** (CW), or a time modulated stream (at a certain frequency  $W$ ), or a train of pulses.

In the present example of Fig. 1A, a portion of detected photons scattered by the fetus **2** are tagged by ultrasound waves, while detected photons scattered  
10 by the maternal tissues **11** are untagged. As photons **155** illuminate the fetus **2**, the transducer **110** generates acoustic waves **150** that propagate through maternal tissues to irradiate the same volume of the fetus **2** from which scattered photons **155** are detected by the detection assembly **101B**. The interaction of acoustic waves **150** with photons **155** results in that the frequency of photons **155** is  
15 shifted by the frequency of acoustic waves **150** (acousto-optic effect). These frequency-shifted or frequency-modulated photons are thus “tagged” and can be identified. The detection assembly **101B** detects both the frequency shifted photons (“tagged photons”) and the photons at the original frequency (“untagged photons”) at both wavelengths. The detection assembly **101B** generates measured  
20 data (electric signals) in response to both the tagged and untagged photons.

Step 3: The control unit **120** processes the measured data using an appropriate algorithm according to the type of detection used. For example: in the case of a single (large area) detector, heterodyne detection (e.g., as described by [Lev A. and B.G. Sfez Optics Letters (2002) 27 (7) 473-475]) is used to  
25 separate data indicative of the signal of the tagged photons; when a CCD camera is used and a full speckle image is detected, the technique described by [Leveque-Fort et al. in Optics Communication 196 127-131 (2001)] is used to determine the optical signal of photons scattered from the particular volume which is tagged by ultrasound waves.

- 22 -

Using the above, or other suitable, techniques, it is possible to determine the effective attenuation of photons **155** as they propagate through the fetus **2**. To this end, ultrasound radiation may be applied such that acoustic waves **150** propagate through different depths of the fetal tissues (e.g., by displacing the transducer arrangement with respect to the body or by using a phase array transducer). Accordingly, the absorption coefficient and the reduced scattering coefficient can be isolated in the two wavelengths chosen for illumination. For example, using a similar equation to equation 4 of Lev et al. referenced above:

$$10 \quad x = \frac{\gamma_6^O - \frac{\mu_{\text{eff},6}}{\mu_{\text{eff},8}} \gamma_8^O}{\left[ (\gamma_8^H - \gamma_8^O) - \frac{\mu_{\text{eff},6}}{\mu_{\text{eff},8}} (\gamma_6^H - \gamma_6^O) \right]}$$

it is possible to determine the oxygen saturation level of the fetus. Here,  $x$  is the fraction of deoxyhemoglobin,  $\gamma$  are the molar extinction coefficients of oxyhemoglobin(O) and deoxyhemoglobin (H) at both wavelengths (in the referenced paper, 6 stands for 690nm and 8 for 820nm) and  $\mu_{\text{eff},6}$  and  $\mu_{\text{eff},8}$  are the measured attenuation coefficients at 690 and 820nm, respectively.

The ultrasound transducer **110** is kept at a specific location, which is optimal for propagating acoustic waves through the same volume of the fetal body (such as the head) from which scattered photons **155** are detected by the detection assembly **101B**. The beam size of transducer **110** is such that the cross section volume between photons **155** and acoustic waves **150** is as large as possible, whether focused or not, for increasing the signal to noise ratio (SNR) of the detection system, without compromising the sensitivity to detect only the fetal oxygen saturation and not the maternal one.

As indicated above, the present invention utilizes ultrasound tagging for the purposes of distinguishing between light responses of the regions of the fetus **2** and the region of maternal tissues **11**. Preferably, the frequency of acoustic



- 23 -

waves generated by the transducer arrangement **110** is in the range of 50kHz-8MHz, and more preferably - lower than 1MHz. This frequency range provides a better SNR for ultrasound tagged light, as it increases the fraction of photons that are tagged, but results in a lower focusing resolution. This is in contrast to  
5 imaging modalities known in the art, where it is desired to improve the imaging resolution and thus higher frequencies and minimal cross section are conventionally chosen. In addition, the detection assembly **101B** collects forward and back scattered photons according to the preferred geometry of FOSM **100**. Therefore, a number of photons collected by the detection assembly **101B** is  
10 higher than in cases of reflection based imaging disclosed in the above references, thus enabling an improved SNR. Hence, the invention enables using safer light energies for illumination. It should be understood that such a configuration, although rendering high resolution imaging more complicated than the case where primarily back scattered photons are detected, is highly suitable  
15 for fetal oximetry.

The control unit **120** analyzes both back and forward scattered tagged photons to determine the optical attenuation of light propagating through the fetal head. Consequently, the control unit **120** needs not perform high resolution imaging of the fetus, but rather just analyze the collected photons **155** scattered  
20 by a large volume of the fetal tissues.

Step 4: The control unit **120** processes that portion of the measured data, which is associated with tagged photons scattered from the fetus (identified as described above), to determine the desired parameter of the fetus – oxygen saturation in the present example. Two modalities can optionally be used to  
25 determine the oxygen saturation level of a fetus intrauterine, one being based on measuring the average oxygen saturation level (known as oximetry) and the other being based on measuring the oxygen saturation level correlated with changes in the blood volume during the cardiac cycle (known as pulse oximetry).

Oxygen saturation  $S$  is a ratio between the concentration of oxygenated  
30 hemoglobin [HbO] and the total concentration of hemoglobin [HbT] in blood:

- 24 -

$$S = [\text{HbO}]/[\text{HbT}] \quad (*100\%) \quad [1]$$

$$[\text{HbT}] = [\text{HbO}] + [\text{Hb}] \quad [2]$$

5 wherein [Hb] is the concentration of deoxygenated hemoglobin.

The saturation  $S$  can be extracted from the attenuation coefficient measured for at least two wavelengths  $\lambda_1$  and  $\lambda_2$ , where the molar absorption and scattering coefficients for Hb and HbO at each wavelength are known in the literature. It should be noted that more than two wavelengths can be used, to  
10 improve sensitivity of the measurement.

As the arteries expand, a blood volume [HbT] is increased by  $[\Delta\text{HbT}]$ , therefore absorption changes periodically. The optical attenuation at  $\lambda_1$  and  $\lambda_2$  is measured at predetermined points (for example, the maxima and minima of a power spectrum of the tagged signal or the processed tagged signal, as defined  
15 below) generated by the detection assembly **101B** during a cardiac cycle. As indicated above, in the present example tagged signal is that associated with the fetus. The saturation  $S$  can be calculated from differences in attenuation of light ( $\Delta\text{OD}$ ) at each wavelength between maxima and minima.

$$20 \quad \Delta\text{OD}^\lambda = (\mu_{\text{HbO}}^\lambda [\text{HbO}] + \mu_{\text{Hb}}^\lambda [\text{Hb}])d = (\mu_{\text{HbO}}^\lambda S + \mu_{\text{Hb}}^\lambda (1-S))[\text{HbT}]d \quad [3]$$

wherein  $\mu_{\text{HbO}}^\lambda, \mu_{\text{Hb}}^\lambda$  are the molar attenuation coefficient of oxygenated and deoxygenated hemoglobin respectively, at wavelength  $\lambda$  ( $\lambda = \lambda_1, \lambda_2$ ) and  $d$  is the distance from the source to the target tissue (fetal or maternal).

25 Defining the ratio  $R$  between  $\Delta\text{OD}^\lambda$  at each wavelength  $\lambda_1$  and  $\lambda_2$ :

$$R = \frac{\Delta\text{OD}^{\lambda_1}}{\Delta\text{OD}^{\lambda_2}} = \frac{[\mu_{\text{HbO}}^{\lambda_1} S + \mu_{\text{Hb}}^{\lambda_1} (1-S)]}{[\mu_{\text{HbO}}^{\lambda_2} S + \mu_{\text{Hb}}^{\lambda_2} (1-S)]} \quad [4]$$

- 25 -

saturation  $S$  is extracted from equation [4] when  $\Delta OD^{\lambda_1}$  and  $\Delta OD^{\lambda_2}$  are measured and the molar attenuation coefficients are known.

According to an embodiment of the present invention, the control unit **120** analyzes signals generated by the detection assembly **101B** in response to each wavelength  $\lambda_1, \lambda_2$  generated by the illumination assembly **101A**. The signals corresponding to tagged photons **155** are selected by the detection assembly **101B** using heterodyne detection, or by the control unit **120** using frequency analysis and/or speckle imaging. These signals are termed "tagged signals". The time dependent amplitude and/or phase of the tagged signals for each wavelength  $\lambda_1, \lambda_2$  is stored in the memory of the control unit **120**, over a specified period of time of at least one fetal heart cycle. To determine the oxygen saturation level of the fetus, the control unit **120** determines the changes in attenuation of tagged signals at each wavelength.

Considering the determination of oxygen saturation of fetus **2** based on oximetry, the time averaged signals generated by the detection assembly **101B** in response to the tagged photons **155** of at least two illuminating wavelengths reaching the input port **122**, are used to determine the oxygen saturation level. Time averaging can be performed over longer time scales than the duration of a fetal heart cycle.

Considering pulse oximetry used for determining oxygen saturation of a fetus, the temporal changes (due to the fetal cardiac cycle) in the blood volume of the fetus are monitored by the control unit **120** by monitoring the low-frequency changes (1-2.5Hz) in the signals generated by the detection assembly **101B** in response to the tagged photons **155** of at least two illuminating wavelengths reaching the input port **122** of the detection assembly. Since the ultrasound frequency is orders of magnitude higher than the fetal heart rate, it is possible to average the signals responsive to tagged photons over a fraction of the fetal heart cycle to improve the SNR of the measurement. Using methods of pulse oximetry, both the oxygen saturation and the pulse rate are determined simultaneously.

- 26 -

The control unit **120** displays the determined fetal oxygen saturation level, along with fetal heart rate, as a function of time. Fetal heart rate is determined by low-frequency analysis of the tagged signals. The control unit **120** optionally alerts using a suitable indication utility (e.g. sound and/or light signal), when  
5 oxygen saturation level drops below a certain threshold (for example 30% or 40%), or when fetal heart rate changes abnormally.

Preferably, monitoring apparatus **100** provides for calibrating for movements of fetus **2** during the measurement. To this end, the control unit **120** operates to determine the position of fetal head relative to the apparatus **100**. This  
10 is carried out either periodically, or upon detection of signals not corresponding to a normal heart rate or oxygen saturation level. The control unit **120** sends a control signal to the transducer arrangement **110** initiating an ultrasound echo measurement. In an echo measurement, the transducer arrangement **110** transmits acoustic waves **150** into maternal tissues, and collects acoustic waves **150**  
15 reflected by fetal and maternal tissues. The reflected signals are analyzed by the control unit **120** (using any conventional ultrasound imaging technique) to determine a position of the fetal head. If a substantial movement is detected, the control unit **120** sends a signal to the transducer arrangement **110** to optionally change a direction of acoustic waves **150** in a new direction corresponding to a  
20 new position of the fetus **2**. Additionally or alternatively, the control unit **120** alerts the operator of apparatus **100** to readjust the position of the apparatus accordingly.

Although the above description refers to a single fetus, it should be understood that the technique of the present invention can easily be adapted for  
25 monitoring several fetuses intrauterio. The location of each fetus is determined using an ultrasound imaging system, and different monitoring apparatuses (i.e., optical and acoustic units) or an integrated multi-fetuses apparatus are used. All the monitoring apparatuses can be hooked to the common control system that controls each apparatus separately, and processes the signals using the same or

- 27 -

different processing utilities. A display shows the oxygen saturation level of each fetus separately along with its heart rate and other parameters.

The present invention also provides for advantageously utilizing the principles of ultrasound tagging of light in pulse oximetry for monitoring oxygen saturation in a localized region of interest in a human or animal body (without a fetus). Turning back to **Fig. 1A**, the optical unit **101** is configured as a pulse oximeter, namely includes an illumination assembly **101A** configured to generate light of at least two different wavelengths and a light detector **101B**; and is used in combination with the transducer arrangement to significantly improve the pulse oximetric measurements. The monitoring apparatus **100** may be configured to operate in a transmission mode (light transmission based detection), such as the conventional pulse oximeter placed on a finger or earlobe. In this case, the support structure **103** is located such that the illumination assembly **101A** is co-linear with the detection assembly **101B**: the illumination assembly **101A** is placed at one side of the tissue and the detection assembly **101B** is placed at the opposite side of the tissue, therefore ballistic and scattered light emitted from illumination assembly **101A** are detected by detection assembly **101B**. The transducer arrangement **110** is placed such that acoustic waves overlap with an illuminated region in the region of interest from which scattered light reaches the detection assembly **101B**, which is preferably the region encompassing a blood vessel (e.g. an artery) or a collection of arterial vessels. In other applications, requiring reflection based detection from a region of interest (“reflection mode”), the apparatus **100** is located as described for the fetus-related application, where the region of interest preferably encompasses a blood vessel (e.g. an artery) or a collection of arterial vessels. Such an arrangement is superior to conventional pulse oximeter as it is not affected by incoherent ambient light, and more importantly is less affected by motion of the tissue relative to illumination and detection assemblies, as long as the region of interest is kept illuminated and the acoustic waves propagate through it.

It should be understood that using the ultrasound tagging of light in the pulse oximetry based measurements significantly improves the measurements, since the measured power spectrum of an ultrasound-tagged light signal is practically insensitive to movements of the region of interest under  
5 measurements, which is the common problem of the typical pure pulse oximetry measurements.

**Fig. 2** exemplifies another configuration of a fetal oxygen saturation monitor, generally designated **200**, of the present invention. To facilitate understanding, the same reference numbers are used for identifying components  
10 that are common in all the examples of the invention. In the apparatus **200**, an illumination assembly **101A** includes a light source **201A** mounted within a control unit **120** and an optical fiber **201B** guiding light from the light source to the region of interest (fetus). The optical fiber **201B** is inserted into the vaginal track of the pregnant woman (using suitable means, for example a flexible  
15 support structure which is not specifically shown). The optical fiber **201B** is positioned such that it is in close contact with the cervix or to amniotic membranes prior to rupture. Optionally, following membrane rupture optical fiber **201B** is attached to the presenting part of the fetal head with the aid of a support structure. A light detection assembly **101B** is placed on the maternal  
20 abdomen, such that it collects photons scattered by the fetal blood. The detection assembly **101B** is connected to a suitable utility of the control unit **120** via wires (as shown in the figure) or wireless. An ultrasound transducer arrangement **110** is also placed transabdominally such that acoustic waves **150** can propagate through the part of the head of a fetal **2** being closest to the optical fiber **201B**  
25 and detection assembly **101B**. In some cases, it may be advantageous to introduce the transducer arrangement **110** also through the vaginal track.

Reference is made to **Fig. 3A** exemplifying a preferred embodiment of a monitoring apparatus **300** according to the invention. The apparatus **300** is configured generally similar to the above-described apparatus **100**, namely  
30 includes an acoustic transducer arrangement **110**, and an optical unit **101** having

- 29 -

a probe **103** carrying at least a part of an illumination assembly **101A** and at least a part of a detection assembly. Here, the detection assembly is formed by two detection units **101B** and **101C** associated with different locations with respect to the illuminated region defined by a location of the illumination assembly **101A**.

5 The additional detection unit **101C** is also attached to the support structure (probe) **103** and is connected to a control unit **120** via electrical cable (not shown) or wireless means.

One of the detection units – detection **101C** in the present example, is located in the proximity of the illumination assembly **101A**, and the other

10 detection unit **101B** is located at a larger distance from the illumination assembly. In the present example, the detection unit **101C** is located between the illumination assembly **101A** and the detection unit **101B**. Generally, the arrangement of the illumination assembly and detection units is such that one of the detection units (detection unit **101C**) is located close to the illumination

15 assembly to therefore detect photons **165** scattered from regions outside the fetus **2** (i.e., light reaches the detector **101C** prior to reaching the fetus); and the other detector **101B**) is more distant from the illumination assembly and thus detects photons **155** scattered from the fetus and propagating through the maternal tissues region and being thereby affected by the maternal tissues.

20 The transducer arrangement **110** is aligned and/or scanned, such that it transmits acoustic waves **150** to the volume within the illuminated region of the fetus from which photons **155** are detected by the detection unit **101B**, and substantially does not irradiate the maternal tissue region from which photons **165** are collected by the detection unit **101C**.

25 Signals (measured data) generated by the detection unit **101C** may be used by the control unit **120** to determine the maternal oxygen saturation level and heart rate simultaneously, which may thus be displayed. Generally, the use of the additional detection unit **101C** located close to the illumination assembly **101A** assists in separating a light response of the fetus region **2** from that of the

30 maternal tissues' region **11**, since the detection unit **101C** so-positioned will

- 30 -

detect scattering effect of light that traveled through the maternal tissues and did not reach the fetus, and which is thus indicative of the maternal region response only. The other detection unit **101B** practically detects photons **155** including the tagged response of the fetus and the tagged response of the maternal tissues.

5           It should be understood that generally the detection unit detects the tagged light response of the fetus affected by the maternal tissues. Hence, the expression “tagged response of the maternal tissues” means photons tagged (by ultrasound) inside the volume of the fetus being scattered by maternal tissue.

Both the tagged and untagged responses of the maternal tissues’ region are  
10   identically frequency modulated by the mother’s heart rate. Hence, the first measured data from the detection unit **101C**, which is mainly indicative of the untagged light response of the maternal tissues, can be used to analyze the second measured data from the detection unit **101B** to separate a signal indicative of a light response of the fetus from that of the maternal tissues.

15           **Fig. 3B** illustrates a flow chart of the main operational steps of a method of the invention utilizing the monitoring system **300**.

Step 1: First, optimal positioning of the illumination assembly, detection assembly and acoustic transducer arrangement is provided as described above. This positioning ensures that acoustic waves interact with the region of interest  
20 (fetus volume) from which photons **155** are detected at the detector **101C** and substantially do not interact with the region outside the region of interest (maternal tissues) from which photons **165** are detected by the detector **101C**.

Step 2: Actual measurements are performed when at the optimal positions of the illumination, detection and acoustic assemblies. Measured data includes:  
25 (1) a first data portion generated by the detection unit **101C** and indicative of the untagged photons coming from the maternal tissues; and (2) a second data portion generated by the detection unit **101B** and indicative of the photons including tagged and untagged photons coming from the fetus, and untagged photons coming from the maternal tissues.



- 31 -

Step 3: The measured data is processed to filter out, the contribution of tagged and untagged photons scattered by regions outside the region of interest, to the measured signal, wherein this contribution is identified as that having frequency modulation by the mother's heart rate, as previously identified from the data portion (1). Hence, the so-separated light response of the fetus can be processed to determine the desired parameter of the fetus. The control unit **120** may use signals generated by the detection units **101B** and **101C** to determine fetal and optionally maternal oxygen saturation levels.

More specifically, the apparatus **300** operates as follows: The illumination assembly **101A** simultaneously generates photons of two different wavelengths (generally, at least two wavelengths). Photons denoted **155** are photons scattered from maternal and fetal tissues and reaching an input port **122** of the detection unit **101B**, i.e., photons scattered from a tagged volume of tissue, that is intermittently or continuously radiated by acoustic waves **150** generated by transducer **110**.

The transducer arrangement **110** may, for example, be operated to generate a burst of acoustic waves, with a delay of at least  $t_{on}$  between the end of one burst and the onset of another burst. Time  $t_0$  is the time it takes the acoustic burst to reach the target fetal tissues (e.g. head). The duration of the burst  $\Delta t_0$  is determined such that at time  $t_f$ , bounded by a condition  $t_0 \leq t_f \leq (t_0 + \Delta t_0)$ , the acoustic pulse propagates primarily through target fetal tissues (i.e., through a volume  $\Delta V$  of fetal tissues). Therefore, during this time  $t_f$  the acoustic burst reaches the target fetal tissues, and acoustic waves are hardly propagating through maternal tissues. A portion of photons **155** propagating through the same volume of fetal tissues during time  $t_f$  is tagged. Whereas, photons **165** are those propagating only through maternal tissues at the same time  $t_f$  and are therefore untagged (since they did not interact with the ultrasound irradiated region).

The detection unit **101C** is placed at a distance  $Q$  from the illumination assembly **101A**, such that its input port collects primarily photons **165** that are not scattered from the tagged volume. The detection unit **101C** is optionally

- 32 -

moved until it does not collect tagged photons, and is then fixed in the appropriate position. It should be noted that, alternatively, the detection units **101B** and **101C** are fixed in place, and a position of the ultrasound transducer arrangement **110** is adjusted to be such that the tagged photons **155**, scattered from fetal tissues, primarily reach the detection unit **101B** and not the detection unit **101C**.

**Fig. 3C** more specifically exemplifies the data processing procedure for processing measured data from the apparatus **300**. The detection unit **101B** receives photons **155** including tagged and untagged photons scattered by the uterus and tagged photons scattered by the fetus, while the detection unit **101C** receives primarily only untagged photons **165** scattered by the maternal tissues. A signal that is generated by the detection assembly **101B** in response to collected photons **155** is referred to as "signal *A*". A signal generated by the detection unit **101C** in response to photons **165** is referred to as "signal *B*".

According to this example, two models are used to describe the propagation of light in a multi layer tissue system. Such models are described for example by Keinle *et al.* in *Physics in Medicine and Biology* 44: 2689–2702 (1999). One model (Model A) includes the parameters representing some of the tissues through which photons **165** propagate from the illumination assembly **101A** through a medium until they reach the detection unit **101C**, and the other model (Model B) includes the parameters representing some of the tissues in the medium through which tagged photons **155** propagate until they reach the detection unit **101B**. The models include known parameters, such as the molar absorption and scattering coefficients of blood cells, and of oxygenated hemoglobin and deoxygenated hemoglobin at each of the wavelengths of illuminating photons. In addition, the models may include the thickness of the layers (maternal and fetal), presence and volume of amniotic fluid in the light path and other parameters that are measured during the operation of apparatus **300** (as described above with reference to apparatus **100** of Fig. 1). Some tissue

- 33 -

parameters in the model may be averaged or other manipulations of the known or measured parameters of the real tissues in models A and B may be carried out.

Given a certain source amplitude, and the known separation between the illumination assembly **101A** and the detection unit **101C**, model A is used to  
5 calculate the expected time dependent photon flux, or light intensity at the input port of the detection unit **101C**. The expected time dependent photon flux or light intensity is used to calculate the expected signal (termed "signal *C*") that can be generated by the detection unit **101C** in response to such a photon flux. Signal *C* actually presents theoretical data for untagged photons at the location of  
10 detection unit **101C**, while signal *B* presents real measured data for untagged photons collected by the detection unit **101C**. The parameters of model A are adjusted such that signal *C* is made equal to or closely resembles signal *B* (best fitting). Signal processing techniques based on optimization algorithms, such as neural network, can be used to optimally determine the parameters of model A.  
15 The parameters are used to calculate the optical properties of some of the tissues through which photons **165** propagate.

Additionally or alternatively, certain parameters of models A and B (such as thicknesses of maternal tissues, in particular uterine wall, and/or tensions of muscles) may be unknown, and be determined during operation. During  
20 contractions, the thickness of the uterine wall and the tension of the muscles change. Controller **120** determines the thickness of the uterine wall as a function of time by optimizing primarily this parameter of model A. Once determined, these parameters are used to determine contractions' duration and amplitude. Alternatively, tissue velocity measurements are performed by ultrasound  
25 assembly **110**, using techniques known in the art for echocardiography. Transducer arrangement **110** emits acoustic pulses (not shown) that are reflected back by uterine muscles. The reflected pulses are Doppler shifted with respect to the emitted acoustic pulses. Controller **120** analyzes the reflected signals to determine the thickness and velocity of the muscles. During contractions the  
30 thickness and velocity change, therefore controller **120** monitors these changes as

- 34 -

a function of time. Consequently, controller **120** displays the amplitude and duration of the contractions. The apparatus **300** thus provides information needed to monitor the progression of labor (contractions' duration and amplitude) in addition to fetal well being (heart rate and oxygen saturation).

5 In addition, signal *B* is optionally used to extract maternal oxygen saturation level, by using the time dependent amplitudes of the signals generated by the detection unit **101C** in response to photons **165** of at least two wavelengths.

It may generally be assumed that the optical properties of tissues outside  
10 the region of interest (outside fetus) through which both photons **155** and **165** propagate are similar (for example maternal abdominal tissues). Alternatively, it may be assumed that by determining the parameters and optical properties of the tissues through which photons **165** propagate, one can deduce, within a reasonable error, the optical properties of corresponding tissues (e.g., other areas  
15 of maternal abdominal tissues) through which photons **155** propagate. The parameters calibrated by signal *B* and the optical properties of the tissues through which photons **165** propagate are then used to calibrate model B that describes the propagation of photons **155** through maternal and fetal tissues.

The time dependent amplitude of signal *A* at all wavelengths of photons  
20 **155** is processed by the control unit **120** using techniques known in the art, such as digital Fourier transformations and analog or digital filtering, to extract, from the entire signal *A*, a signal portion corresponding to the tagged photons **155**. This signal portion is termed "tagged signal *A*". Tagged signal *A* is that modulated at the ultrasound frequency generated by the transducer arrangement  
25 **110**. The amplitude of the power spectra of the tagged signal *A* at the ultrasound frequency (or related to the ultrasound frequency), the modulation width of its power spectra or other features of tagged signal *A*, such as its phase, are termed together as "processed tagged signal *A*". This processed tagged signal *A* is actually indicative of both the maternal tissues response and the fetus response

- 35 -

tagged by ultrasound. In addition, the signal *A* contains information which is not modulated at the ultrasound frequency, termed “untagged signal *A*”.

According to this specific embodiment, untagged signal *A* may also be used in the data processing and analyzing procedure, for example to determine  
5 some of unknown parameters of model B and further optimize this model. For example, untagged signal *A* may contain signals which are modulated by maternal cardiac cycle, and have a modulation frequency of 0.5-2Hz corresponding to maternal heart rate  $F_m$ . Signal *B* is also modulated at the same frequency, as photons 165 propagate through maternal tissues containing the  
10 same pulsating blood. Consequently, untagged signal *A* and signal *B* may be used to calibrate model B relative to model A, where differences and similarities between untagged signal *A* and signal *B* are used to optimize the parameters of model B.

In addition, tagged signal *A* and/or processed tagged signal *A* are also  
15 modulated at maternal heart rate, as tagged photons 155 pass through maternal tissues before and after they pass through the tagged volume. Consequently, tagged signal *A* and/or processed tagged signal *A* modulated at this low frequency may be used in conjunction with untagged signal *A* and/or signal *B* to extract the portion of tagged signal *A* that is affected by absorption by fetal  
20 blood. This portion calculated for all wavelengths of photons 155, is used to extract the fetal oxygen saturation level.

According to another embodiment of the invention, only tagged signal *A* and untagged signal *A* at all the wavelengths of photons 155 are used to extract fetal oxygen saturation levels. According to yet another embodiment, tagged  
25 signal *A* and/or processed tagged signal *A* are used to determine fetal oxygen saturation at all fetal heart rates  $F_f$ , where  $F_f \neq F_m$  (or more precisely  $F_f > F_m + BW$ , where  $BW$  is the bandwidth of the detection system, as fetal heart rate is usually faster than maternal heart rate). First, tagged signal *A* is extracted (separated) by the control unit 120 as described above. Then, the modulation amplitudes of  
30 tagged signal *A* and processed tagged signal *A* at  $F_f$  and  $F_m$  are determined.

- 36 -

Tagged photons **155** are modulated at  $F_f$ , however a modulation at  $F_m$  may also exist, as tagged photons **155** also propagate through maternal tissues. When this modulation is small, its contribution at higher harmonics (i.e.,  $2F_m$ ,  $3F_m$ ) is negligible. The amplitudes of tagged signal  $A$ , processed tagged signal  $A$  and  
5 untagged signal  $A$  modulated at frequency  $F_m$  are optionally used to determine certain tissue parameters in model B. Using these parameters, tagged signal  $A$  is calibrated to correspond primarily to fetal contributions. Fetal oxygen saturation is extracted from features (such as the modulation amplitude, the bandwidth of the modulation, autocorrelation etc.) of the calibrated tagged signal  $A$  and/or  
10 processed tagged signal  $A$  modulated at  $F_f$  at all wavelengths of photons **155**.

In some cases where the modulation of tagged signal  $A$  at  $F_f$  in the range of  $2F_m - BW < F_f < 2F_m + BW$  can not be neglected, signal  $B$  and untagged signal  $A$  are used to determine fractions of tagged photons **155** that are modulated by maternal blood, by fitting the parameters of models A and B as described above.  
15 Once the parameters are determined, fractions of tagged photons **155** that are modulated by maternal blood and fetal blood can be determined using known methods, for example such as Monte Carlo simulations. Using the results of the simulations, tagged signal  $A$  is calibrated to correspond primarily to fetal contributions. The calibrated signal is then used to extract fetal oxygen saturation  
20 levels as described above.

Turning back to **Fig. 3A**, it should be noted that the acoustic transducer arrangement may be accommodated such that acoustic waves propagate towards the fetus along an axis passing between the illumination assembly and the detection assembly. For example, the transducer arrangement or its associated  
25 ultrasound port is located on the same support structure **103**. The optical unit is preferably operated to start illumination/detection a certain predetermined time after the generation of the acoustic radiation, which is the time needed for the acoustic radiation of a given frequency to arrive at the region of interest (fetus). This ensures that light detected by the detection unit **101C** is not affected  
30 (tagged) by the acoustic radiation.

- 37 -

Alternatively or additionally, the apparatus of the present invention, for example configured as the above-described apparatus **300**, can be used to monitor the optical properties of the amniotic fluid surrounding the fetus. In this case, a region within the amniotic fluid presents a region of interest, and as indicated above the term “maternal tissues” refers to regions outside the region of interest. Optical properties of the amniotic fluid may include, for example, the absorption coefficient, the scattering coefficient, the reduced scattering coefficient and the refractive index of the fluid. Such optical properties are used to calculate the concentration of lamellar bodies, blood or meconium dispersed within the amniotic fluid. The calculated concentration is optionally compared to a threshold level as described below.

As illustrated schematically in **Fig. 3D**, to monitor amniotic fluid, apparatus **300A** is configured and positioned similar to the above-described apparatus **300**, whereas the region of interest **2A** is defined by a substantial volume of amniotic fluid, being at the shortest optical path to light output and/or input ports of illumination and detections assemblies. The substantial volume is that which allows for an overlap of illuminating light and an ultrasound beam within that volume. For example, in the case the illumination assembly **101A** and detection unit **101B** are appropriately placed to illuminate and collect light scattered by the amniotic fluid **2A**, a transducer arrangement **110** is placed such that acoustic waves **150** propagate through the same region of the amniotic fluid **2A** from which scattered photons **155** are detected by a detection unit **101B**. Preferably, the detection assembly is configured such that a detection unit **101C** collects untagged photons propagating through maternal tissues **11A** (a region outside the region of interest), and the detection unit **101B** collects tagged and untagged photons propagating through a substantial volume of the amniotic fluid (region of interest).

The apparatus **300A** may be used for determining the optimal positioning of illumination/detection and ultrasound assemblies, having a plurality of input and output ports, such that the ultrasound beam is scanned over different

locations inside the body and the autocorrelation or power spectrum of signals generated by each detection unit are determined by a controller **120** in response to photons scattered from different volumes within the body overlapping with the ultrasound beam. As the line-width of the autocorrelation or power spectrum of the tagged signals, around the frequency of the ultrasound radiation, is different when tagging is performed inside a fluid volume than when performed in a tissue or bone volume, the controller **120** can determine, by monitoring the line width, when the ultrasound beam is used to optimally tag a volume of the amniotic fluid.

10 For monitoring amniotic fluid, the illumination assembly **101A** includes one or more light sources generating a plurality of wavelengths (either simultaneously or sequentially) from 300nm to 12 $\mu$ m. For example, a plurality of wavelengths that are absorbed and/or scattered by lamellar bodies contained in amniotic fluid is chosen for determining the concentration of lamellar bodies.

15 Preferably, the plurality of wavelengths is less absorbed by water. Such wavelengths may be chosen in the range of near infrared, i.e., 600nm-1300nm.

At each wavelength, the control unit **120** optionally determines models A and B (as described above) for the overlaying maternal tissues **11A** (region outside the region of interest) and the amniotic fluid (region of interest), and

20 determines tagged and untagged signal *A* and untagged signal *B* as described above. Processed tagged signal *A* and calibrated tagged signal *A* are used to determine the reduced scattering coefficient and the absorption coefficient of the tagged volume of amniotic fluid as explained below.

The control unit **120** then determines the concentration of lamellar bodies, blood or meconium in the amniotic fluid. The output of the control unit **120** is

25 displayed on the integrated display or communicated via wireless means or cables to another display or electronic processor. Possible outputs include but are not limited to a light signal indicating higher or lower concentration of lamellar bodies relative to a predetermined threshold, a number shown on the display

30 corresponding to the concentration of lamellar bodies in addition to a display of



- 39 -

the threshold number for that value, a sound indicating high or low concentration of lamellar bodies relative to a threshold. The control unit 120 optionally displays a “mature” or “premature” signal, without quantitative information about the concentration of lamellar bodies, or displays “stained” or “clear” signal for the case of meconium staining.

Lamellar bodies are produced by type II alveolar cells in increasing quantities as fetal lungs mature. They are composed almost entirely of phospholipid and represent the storage form of the surfactant. Their diameter is about 0.5-2  $\mu\text{m}$ , and their index of refraction is about 1.475. Consequently, when using the above wavelengths range, it is clear that Mie scattering dominates the scattering process of light from lamellar bodies. Choice of specific wavelengths depends on the optimal signal to noise ratio (SNR) of the apparatus used. The difference in wavelengths used for illumination has to provide a sufficient change in the scattering coefficient that can be detected by the system. The relationship between the wavelength and the reduced scattering coefficient  $\mu_{s'}$  of monodisperse scattering dielectric spheres is known in the literature to be:

$$\mu_{s'} = 3.28\pi a^2 \rho \left( \frac{2\pi a}{\lambda} \right)^{0.37} (m-1)^{2.09} \quad [5]$$

wherein  $a$  is the radius of the dielectric spheres,  $\rho$  is their volume density,  $\lambda$  is the wavelengths in vacuum,  $m = \frac{n_s}{n_0}$  where  $n_s$  and  $n_0$  are the refractive indices of the spheres and the surrounding material respectively.

The reduced scattering coefficient  $\mu_{s'}$  is related to the scattering coefficient  $\mu_s$  using the following equation:

$$\mu_{s'} = (1-g)\mu_s \quad [6]$$

wherein  $g$  is the anisotropy factor related to the size and geometry of the scattering centers.

Both the reduced scattering coefficient and the scattering coefficient can be determined according to an embodiment of the present invention.

- 40 -

As an example, light at a plurality of wavelengths is generated by the illuminating assembly **101A**. Wavelength selection is based on the absorption and scattering coefficients of the amniotic fluid. For example, amniotic fluid with no blood or meconium absorbs almost equally light at around 735nm and 780nm.

5 Therefore, light distributions inside amniotic fluid at these two wavelengths will differ due to wavelength dependant changes of the reduced scattering coefficient of the fluid. This reduced scattering coefficient depends on the size, volume concentration and relative index of refraction of the lamellar bodies (as evident from equation 5 above). Therefore, by determining the reduced scattering

10 coefficients at different wavelengths, the control unit **120** determines the concentration of lamellar bodies in the fluid using equation 5 (or a modified equation 5 which includes physiological parameters of lamellar bodies instead of monodisperse spheres).

In order to determine the presence of meconium in the fluid, a different

15 selection of wavelengths is used. It is known that meconium primarily contains blood and billirubin, therefore at least two wavelengths are used: one that is absorbed by blood or billirubin (for example 660nm) and the other that is weakly absorbed by either one of them (for example 1064nm). These wavelengths may for example be used in addition to the above mentioned wavelengths 735 and

20 780nm. Irradiating the amniotic fluid with these two wavelengths (e.g., 660nm and 1064nm) will result in a different light distribution at the shorter wavelengths when meconium is present than in the case of no meconium. The absorption coefficient of the fluid at these wavelengths is also used to determine the concentration of meconium in the fluid.

25 The control unit **120** determines the processed tagged signal *A* at the plurality of wavelengths used to illuminate the tissues, and stores them in memory. Since the time dependant variations in the optical properties of the amniotic fluid are very slow, tagged signal *A* at each wavelength can be integrated over long periods (much longer than the fetal or maternal heart beats).

30 At those wavelengths, which are absorbed almost equally in the fluid, the

variations between the signals obtained at different wavelengths are proportional to the reduced scattering coefficient. From these variations of the measured parameters (listed below), the control unit **120** determines the reduced scattering coefficient.

5           The control unit **120** determines the reduced scattering coefficient of the amniotic fluid by determining, for each illuminating wavelength, at least one of the following parameters:

(a) The amplitude of the power spectra or autocorrelation of processed tagged signal *A* corresponding to the frequency of the ultrasound waves;

10           (b) The line width of the power spectra or autocorrelation of processed tagged signal *A* (for example, the full width at half max around the frequency of the ultrasound waves);

(c) The spatial attenuation of the amplitude of the power spectrum of tagged signal *A* corresponding to the ultrasound frequency.

15           In the latter case, the ultrasound beam scans the illuminated region such that the overlapping volume, within the amniotic fluid, is varied along the optical path between the illumination assembly **101A** and the detection unit **101B**. For each location, tagged signal *A* or processed tagged signal *A*, is stored in memory, and the attenuation of the amplitude of the power spectrum of tagged signal *A*  
20 corresponding to the ultrasound frequency is determined.

In order to determine the absorption coefficient, following the determination of the reduced scattering coefficient at the above wavelengths, different wavelengths which are differently absorbed in the fluid (such as 1064nm or 660nm) are used and the above parameters are determined according  
25 to a specific method from those listed above. The absorption coefficient depends on the product of the concentration of the chromophores or structures within the fluid and the molar absorption coefficients (which are known in the literature). In order to determine the absorption coefficient, following the determination of the reduced scattering coefficient at the above wavelengths (735nm and 780nm),  
30 different wavelengths which are differently absorbed in the fluid (such as

- 42 -

1064nm or 660nm) are used to illuminate the body region. From the determined parameters of the signals at each wavelengths, the optical attenuation is obtained. The optical attenuation  $\mu_{\text{eff}}$  is known to depend on both the reduced scattering ( $\mu_{s'}$ ) and ( $\mu_a$ ) absorption coefficients where  $\mu_{\text{eff}} = \sqrt{3\mu_a(\mu_a + \mu_{s'})}$ , since the 5 reduced scattering coefficients were determined before, the absorption coefficients can be extracted.

Therefore, the concentration of absorbing centers (like meconium) is determined similar to the concentration of hemoglobin, as described above.

In some embodiments of the present invention, a threshold value for 10 mature lungs is an input parameter to the control unit **120** prior to its operation. In some embodiments of the present invention, several parameters are being input into the control unit **120** prior to operation. Some parameters are measured by an ultrasound imager, such as, thickness of uterine wall or thickness of abdominal wall. Additional parameters may include weight of gravida and 15 duration of gestation (relative to last menstrual period or based on other indicators). Some of these parameters are used to calculate the threshold value to which the optical properties of the amniotic fluid, or the concentration of lamellar bodies, are compared. These parameters can optionally be used to calculate the concentration of the lamellar bodies from the optical signals according to an 20 algorithm that uses the optical properties of maternal tissues **11A** to extract the optical attenuation in amniotic fluid **2A** as explained above.

It should be noted that similar to the configuration of **Fig. 2**, illumination assembly **101A** can be inserted transvaginally to illuminate a volume of amniotic fluid through the cervix. Choice of wavelengths for transcervical illumination 25 may be different than choice of wavelengths for abdominal illumination, since the composition of the different tissue layers in between the illumination assembly **101A** and the amniotic fluid **2A** is different for the two configurations. For example, skin (epidermis) contains melanin that highly absorbs in the ultraviolet region, whereas the cervix has little or no melanin.

While it is preferred that the control unit **120** determines the concentration of lamellar bodies or meconium, it is not always necessary to determine these concentrations. A database of signals can be defined by recording signals collected by the controller's memory or features of these signals (e.g. amplitude, phase, frequency, time dependence, wavelets or principal components). The database may contain data from measurement obtained for premature fetal lungs and mature fetal lungs. For lung maturity classification by the database, a similarity metric is defined. The obtained signals are classified according to the best fit to mature or immature lungs using clustering, neural networks and/or other classification algorithms. For meconium stained analyses, the database can contain signals from stained and clear amniotic fluid. Features from the ultrasound image taken prior or during the assay are used to categorize the measured signals in the database.

Whereas the above examples relate to measuring the optical properties of amniotic fluid, similar apparatus can be designed for noninvasive measuring the optical properties of other extravascular fluids such as pleural (around the lungs), pericardial (around the heart), peritoneal (around the abdominal and pelvis cavities) and synovial (around the joints) fluids.

Reference is made to **Fig. 4** exemplifying a monitoring apparatus **400** according to yet another embodiment of the invention. The apparatus is constructed and operable to enable determining properties of both maternal and fetal regions, for example detecting maternal and fetal oxygen saturation levels along with other tissue components. The apparatus **400** includes a measurement unit (optical and acoustic units) carried by a flexible probe **403**; and is connectable to a control unit **120**. The control unit **120** is similar to the above described control unit, namely is a computer system including *inter alia* a power supply, a control panel with input/output functions, a display unit, a function generator, electronic circuits, filters and processors, etc. Additionally, the control unit **120** may contain light sources, light detectors and acoustic sources. Electric wires, optical fibers and/or wireless means connect the control unit **120** to the

- 44 -

elements of the measurement unit carried by the flexible probe **403**. The flexible probe **403** includes light output ports **121A** and **121B** associated with an illumination assembly **101A**, and light input ports **122A** and **122B** associated with a detection assembly **101B**. It should be understood that the elements **121A** and **121B** carried by the probe **403** may be light sources (such as lasers, laser diodes or LEDs) or the output faces of optical fibers (or fiber bundles) whose input faces are coupled to light sources contained outside the probe, e.g., in the control unit **120**. Similarly, elements **122A** and **122B** may be light detectors (such as for example diodes or CCD cameras) or the input faces of optical fibers or fiber bundles whose exit faces are coupled to light detectors located outside the probe, e.g., contained in the control unit **120**. In the case actual detectors are placed on the flexible probe **403**, the detectors are preferably mechanically and electronically isolated such that acoustic waves propagating from an acoustic output port **245** minimally affect the collection of photons by the detectors and the transduction of light signals into electronic signals. If the ultrasound transducer arrangement **110** is also placed on the probe **403**, then the configuration is such as to prevent RF and other electronic signals generated by the transducer arrangement from interfering with the collection of photons by the detectors and with the transduction of light signals into electronic signals. Such a shielding of the detectors may for example include electrical isolation by appropriate materials that are poor conductors or create a Faraday cage around a detector, mechanical isolation by using appropriate material that attenuates the propagation of sheer acoustic waves through the probe itself or through maternal tissues. Detectors may be connected to the probe **403** using connecting parts (possibly detachable) that isolate mechanical and electrical signals at the frequencies generated by the ultrasound transducer arrangement and at other frequencies.

The detection assembly generates electronic signals in response to the amplitude and phase of photons reaching the input ports **122A** and **122B**. These electronic signals may be filtered by analog or digital filters, for example

- 45 -

bandpass filters, that are appropriately provided being connected to the data processing utility of the control unit **120** or being a part of this processing utility. The bandwidth of these filters can be fixed or changed by the control unit **120**. Tuning of the bandwidth can be performed optically by heterodyne detection or  
5 by a plurality of filters having different bandwidths, or by tunable filters which are coupled to each detector. Alternatively, filters can be electronic.

It should be noted that more than two input and output light ports may be provided in the monitoring apparatus **400**, only two pairs of such ports being shown in the present example for the purposes of simplifying the illustration. In  
10 addition, each port may serve as a dual input and output light port by using a fiber combiner/splitter that couples light into and out of one optical fiber. The ports may be arranged in a one-dimensional array or a two-dimensional array to improve flexibility of use.

Preferably, the input and output ports are arranged such that an equal  
15 optical path through body tissues exists between pairs of output and input ports. For example, light input and output ports are arranged such that they are not coplanar, but rather form sets of isosceles triangles between output and input ports, having equal light paths between one output port and two input ports (or *vice versa*).

20 The flexible probe **403** also contains the ultrasound output port **245** from which acoustic waves **255** (shown as semicircles **255**) are directed towards the region of interest. Acoustic waves **255** are generated by the ultrasound transducer arrangement **110** that may be located on the flexible probe **403** and connected to a function generator in the control unit **120** using electric wires **236** or using  
25 wireless means. Alternatively, the ultrasound transducer **110** may be located outside the probe **403**, e.g., in the control unit **120**, and acoustic waves be transmitted to the output port **245** using suitable acoustic waveguides **236**. In the example of Fig. 4, the output port **245** is located between the light input and output ports. The output port **245** may be placed at any location of the flexible  
30 probe **403** (i.e. to the right of output port **121A** or the left of input port **122A**).

- 46 -

Several ultrasound output ports, at different locations along the flexible probe **403**, may be used being coupled to the same ultrasound transducer arrangement or to different ultrasound transducer arrangements.

When different ultrasound transducer arrangements are used, the transducer arrangements may generate acoustic waves of the same frequency modulation, or each may generate a different frequency modulation. When different frequencies are being generated, the control unit **120** controls the modulation at each transducer arrangement according to the spatial locations of each output port associated with each transducer arrangement, such that light propagating through the same volume as the acoustic waves and collected through one or several light input ports is analyzed based on the correct frequency modulation of the corresponding transducer arrangement (as will be described below for one such transducer arrangement). Different transducer arrangements can generate acoustic waves at the same time intervals, or during different time intervals.

The flexible probe **403** is placed in contact with a maternal skin **10** in a region overlaying a fetal tissue **2** contained in maternal tissues. The positioning of the flexible probe **403** may be performed with the aid of an ultrasound imaging system that is used to determine the optimal location of light ports and acoustic output port relative to the fetus (as described above). Different fetal organs or tissues can be used to monitor fetal oxygen saturation including the placenta.

The flexible probe **403** is placed in contact with the maternal skin **10** using adhesive pads or by applying pressure to maternal abdomen using a belt. The acoustic output port **245** is coupled to the skin **10** using an acoustic coupling material such as gel or a hydrogel adhesive.

Upon attachment of the probe **403**, the control unit **120** is turned on in a calibration mode. During the calibration mode, the control unit **120** sends an electronic signal to the function generator connected to the ultrasound transducer arrangement causing it to generate a series of acoustic pulses. The acoustic pulses



- 47 -

are transmitted through the ultrasound output port **245** and propagate through maternal tissues **11** and fetal tissues **2**. A portion of the acoustic pulses is reflected from each surface or impedance mismatched layer that the pulses encounter during their propagation. The reflected portion is collected through the acoustic output port **245** and transmitted to an ultrasound transducer of the imaging system (which may be the same transducer arrangement **110** used for measurements). The transducer arrangement used for imaging may be a single acoustic transmitter, a single acoustic transceiver, an array of transmitters and/or receivers, an ultrasound Doppler imager, a phased array, or a complete imaging system capable of transmitting and receiving acoustic signals. The control unit **120** analyzes the signals generated by the transducer in response to the portion of reflected pulses and determines a distance between the ultrasound output port **245** and the fetal tissues **2** (by multiplying the speed of sound in the tissue by half the time difference  $\Delta\tau$  between the emission of the acoustic pulses from the output port **245** and the collection of the reflected pulses in the said port **245**).

The control unit **120** then determines an appropriate distance to be provided between the light output port **121A** or **121B** and the light input port **122A** or **122B**, such that photons **250** emitted from the output port **121A** will propagate through fetal tissues **2** before reaching the input port **122A** or **122B**. The control unit **120** can select which output and input ports are used from a plurality of light ports arranged at different spatial locations, such that at least one input port collects photons, emitted from at least one output port, that propagate through the same fetal tissue **2** through which acoustic waves **255** propagate. The control unit **120** also determines which of the other light input ports collect photons that propagate through maternal tissues **11** and not through fetal tissues **2**. Additionally, during the calibration mode, the control unit **120** determines a desired frequency bandwidth  $\Delta f_1$  that is to be used during the monitoring. For example, the control unit **120** selects the frequency bandwidth  $\Delta f_1$  corresponding to that optimally filtered by analog or digital electronic filters connected to the light detectors.

Following the calibration mode, the control unit **120** operates an array of fixed output and input light ports by modulating (including time gating) the light sources connected only to the chosen output ports, or modulating the output ports themselves, and analyzing the signals generated by the detectors coupled to the  
5 chosen input ports.

As indicated above, the control unit **120** determines the desired frequency bandwidth  $\Delta f_1$  to be used during the monitoring as that corresponding to a frequency bandwidth optimally filtered by analog or digital electronic filters connected to the light detectors. The bandwidth that these filters optimally  
10 transmit is fixed or varied by the control unit **120** during the operation of the apparatus **400**. The control unit **120** controls a portion of the frequency bandwidth generated by the function generator to correspond to the frequency bandwidth that is optimally transmittable by the electronic filters connected to the light detectors. Alternatively, the bandwidth of the filters is varied by the  
15 control unit **120** to correspond to a portion of the frequency bandwidth generated by the function generator. The control unit **120** also controls the frequency modulation of acoustic waves **255** such that waves having a frequency outside the frequency bandwidth  $\Delta f_1$  are for example generated during different portions of the acoustic pulse.

20 The control unit **120** controls the time dependent generation of the frequency modulated acoustic waves. The control unit **120** determines a time period  $\Delta t_1$  needed for signal acquisition such that optimal signal-to-noise ratio (SNR) for determining fetal oxygen saturation is obtained during the measurements. The time period  $\Delta t_1$  is shorter than a time difference between the  
25 fetal heart beats, when pulse oximetry is used for data analysis. The control unit **120** also determines the frequency modulation parameters such that the desired frequency bandwidth  $\Delta f_1$ , (or phase) propagates through the fetal tissue **2** during the time period  $\Delta t_1$ . The onset of time period is at time  $t_1$  equal to about  $\Delta\tau/2$  where the acoustic pulse reaches the fetal tissues **2** and time  $t_2$  is determined by  
30  $t_2=t_1+\Delta t_1$ .

- 49 -

In the present example of **Fig. 4**, the operation of apparatus **400** in a “monitor mode” or actual measurement mode is shown. During the monitor mode, the control unit **120** activates the light source(s) associated with the output port **121A** to emit photons **250** and **251**, and actuates the light source(s) associated with the output port **121B** to emit photons **252**. The light ports may be associated with different light sources or with one or more common light sources. A single light source and preferably two light sources, emitting light of at least two different wavelengths simultaneously, are connected to the output ports, whereas one light source may be connected to more than one output port. Light sources connected to different output ports, or output ports themselves, may be activated during different time periods or with different characteristics (such as different modulation frequency or phase), such that the control unit **120** can distinguish between photons **251** and **252** reaching input port **122B** of the detection unit.

The control unit **120** also activates a function generator that, in turn, activates the ultrasound transducer arrangement **110** to generate acoustic waves **255** transmitted through the output port **245**. The acoustic wave frequency (or phase) generated by the function generator is modulated by the control unit **120** such that the acoustic waves reaching a region of the fetal tissue **2** illuminated by photons **250** will have a predetermined frequency bandwidth  $\Delta f_1$ . If the bandwidth  $\Delta f_1$  is fixed, then the control unit **120** determines the frequency modulation of the function generator controlling the generation of ultrasound waves **255**, such that acoustic waves **255** modulated at a frequency within  $\Delta f_1$  reach the fetal tissues **2** at time  $t_1$ . In addition, acoustic waves with a frequency within  $\Delta f_1$  substantially do not propagate through other tissues during the time period  $\Delta t_1$  following time  $t_1$ . Accordingly, the control unit operates the detection assembly such that the light detectors associated with the input ports **122A** and **122B** start collection of photons at time  $t_1$  and end the collection process during  $t_2$ . Alternatively or additionally, the control unit **120** controls the activation of light sources associated with the output ports **121A** and/or **121B** at time  $t_1$  and

ends the activation at time  $t_2$ . During the time period  $\Delta t_1$ , the input port **122A** and/or input port **122B** collect photons propagating through the same fetal tissues through which acoustic waves **255** propagate (a time delay in photon propagation through the tissue is neglected).

5           **Fig. 5** exemplifies a monitoring apparatus **500** of a somewhat different configuration. Apparatus **500** differs from the above-described apparatus **400** in the arrangement of input and output ports (or light sources and detectors) within a flexible probe **403**. Here, light port **121A'** functions as an output port (associated with the illumination assembly), light ports **121B'** and **122A'**  
10 function as both output and input ports (associated with the illumination assembly and detection unit), and light port **122B'** functions as an input port (associated with the detection unit).

In addition, all input ports may be time gated to collect light propagating through specific tissues during a certain time period. Additionally or  
15 alternatively, the output ports may be activated during different time intervals. For example, the input ports **121B'** or **122B'** are activated during a time interval  $\Delta t_g$ , following the introduction of light from the output ports **121A'** or **122A'**, respectively, such that only photons **256** propagating from the output port **121A'** to the input port **121B'** or photons **257** propagating from the output port **122A'** to  
20 the input port **122B'** are collected during that time interval. The output ports **121A'** and **122A'** are therefore activated at different time intervals, such that the input port **122B'** does not collect photons **251** and **257** simultaneously. During a time interval  $\Delta t_k$ , different from  $\Delta t_g$ , the input port **122A'** is activated, following the introduction of light from the output ports **121A'**, such that only photons **250**  
25 propagating through fetal tissues **2**, are detected. As will be explained below, during the time interval  $\Delta t_g$  the light input ports collect light propagating primarily through maternal tissues, being untagged. During time interval  $\Delta t_g$  acoustic waves **255** do not propagate through the same volume from which light is collected by the input ports **121B'** and **122B'**. During time interval  $\Delta t_k$  the light

- 51 -

input ports collect light propagating primarily through maternal and fetal tissues, being tagged or untagged.

It is preferred that the time interval  $\Delta t_k$  corresponds to the time interval  $\Delta t_1$  defined above, where both intervals start at  $t_1$ . It is preferred that the intervals  $\Delta t_k$  and  $\Delta t_g$  are closely spaced in time such that no substantial modulation in blood or tissue characteristics occurs in between these time intervals.

Alternatively, several light input ports may be activated during different time intervals corresponding to light propagation through either maternal tissues alone or maternal and fetal tissues, and different acoustic waves can optionally propagate through all volumes from which light is collected allowing both tagged and untagged light to be analyzed through every input port.

Tagged and untagged signals collected by all the input ports are used to determine the fetal oxygen saturation level based on the tissue models A and B described above. The amplitudes of the tagged and untagged signals are determined during the time intervals  $\Delta t_k$  and  $\Delta t_g$  respectively.

The filtered electronic signal at a selected bandwidth  $\Delta f_1$ , during the time period  $\Delta t_1$ , starting at  $t_1$ , corresponds to the amplitude of the tagged signal propagating through the fetal tissue 2. Photons propagating through other tissue regions, during the time period  $\Delta t_1$ , where the frequency of the ultrasound waves is outside the frequency bandwidth  $\Delta f_1$ , will have a different modulation frequency and will be attenuated by the filters having the bandwidth of  $\Delta f_1$ . The electronic filters connected to the light detectors can have several bandwidths of optimal transmission ( $\Delta f_2 \dots \Delta f_n$ ) different from  $\Delta f_1$ . Detectors are configured such that during each time period  $\Delta t_i$  ( $i=1..n$ ) starting at time  $t_i$ , where  $t_i$  is the time where portion of acoustic pulse having a frequency modulation with a bandwidth  $\Delta f_i$  reaches the fetal tissue 2, the detectors are tuned to optimally convert photons, modulated by frequency  $\Delta f_i$ , to electrical signals.

For example, as the modulated ultrasound waves propagate through maternal and fetal tissues, at different time periods ( $\Delta t_2 \dots \Delta t_n$ ) different from  $\Delta t_1$ ,

- 52 -

acoustic waves at different frequency modulations propagate through the fetal tissues. The control unit **120** then operates each filter according to the order and duration of the different frequency modulations within the ultrasound pulse, such that each electronic filter having a frequency bandwidth  $\Delta f_i$  is activated at time  
5 periods  $\Delta t_i$  starting at  $t_i$  in accordance with the corresponding frequency bandwidth of the ultrasound wave. Additionally, the electronic filter can have a variable bandwidth that is controlled by the control unit **120** in accordance with the generation of the frequency modulation of the ultrasound wave. The control unit **120** then integrates the electronic signals generated by the light detectors and  
10 filtered by the electronic filters at all frequency bandwidths during the corresponding time periods. The control unit **120** then analyzes signals received from the electronic filters to determine the fetal and/or maternal oxygen saturation levels.

Referring to either one to **Figs. 4 and 5**, the control unit **120** analyzes  
15 tagged signals, reaching at least one of the input ports of the detection units and being modulated at frequency bandwidth  $\Delta f_m$  different from  $\Delta f_i$  ( $i=1..n$ ), during a time period  $\Delta t_i$  starting at  $t_i$ , to determine the properties of maternal tissues through which photons **251** and/or **252** (in Fig. 4) propagate. These signals may optionally be used to determine maternal oxygen saturation levels. The control  
20 unit **120** then uses the tagged signals having a modulation corresponding to frequency bandwidth  $\Delta f_m$  in model B described above to determine characteristics of maternal tissues. The control unit **120** then calibrates model A as described above. The control unit **120** then uses tagged signals, reaching the input ports of the detection units and being modulated at the frequency  
25 bandwidth corresponding to  $\Delta f_i$  during the same time period  $\Delta t_i$  starting at  $t_i$  in accordance with model A to determine fetal oxygen saturation levels.

It should be noted with respect to the above-described examples, that the transducer arrangement **110** may include an ultrasound imaging system and/or and ultrasound Doppler imager, such that ultrasound images or Doppler signals  
30 are acquired during operation of the monitoring apparatus. Prior to and during the

- 53 -

operation, an ultrasound image of tissues including the region of interest is first acquired. The control unit **120** (or the operator) identifies the region of interest by marking this region on the first acquired ultrasound image. Transducer arrangement **110** is then fixed at the same location (by the operator or by using an adhesive), for duration necessary to obtain efficient tagged signals from the region of interest. The control unit **120** determines the corresponding distance, angle and size of the region of interest relative to the transducer arrangement **110**. Then, the control unit **120** operates to select input and output light ports to emit light and collect light propagating through the region of interest and the outside region, as described above (i.e., to ensure optimal positioning during the measurements). The control unit **120** synchronizes the activation of the light output ports and/or light input ports such that emission of ultrasound pulses to the region of interest corresponds to the propagation of light pulses through the region of interest. Alternatively, the control unit **120** synchronizes the emission of acoustic pulses by the transducer arrangement **110** such that the onset, duration and direction of these pulses correspond to the onset and duration of the activation of light input and/or output ports. The control unit **120** analyzes tagged and untagged signals generated by the detection assembly and determines the concentration of an analyte (for example, oxygenated hemoglobin) in the region of interest, as described above. More than one analyte can be monitored simultaneously by using different wavelengths of light, each having a characteristic absorption or scattering by the selected analytes. The control unit **120** then displays simultaneously (using, for example, color coded images) the local concentration of the analyte on the region of interest of the ultrasound image, in addition to any other form of display (e.g. numerical) that is understood by the operator and/or by a machine connected to the control unit **120**. In cases where more than one ultrasound pulse is needed for obtaining efficient tagged light signals, the transducer arrangement is operated without scanning the ultrasound beam, and a single beam is emitted in the direction of the region of interest for the duration needed for an efficient tagged signal acquisition. Yet

- 54 -

another option suitable for cases when the transducer arrangement is moved during operation, such that distances and angles between the transducer arrangement **110** and the region of interest are changed, the control unit **120** tracks the position of the region of interest using techniques and algorithms such as of three-dimensional ultrasound imaging (F. Rousseau, P. Hellier, C. Barillot, 5 *“Calibration Method for 3D Freehand Ultrasound”*, In *Medical Image Computing and Computer Assisted Intervention, MICCAI'03*, Montreal, Canada, November 2003). The control unit **120** then determines the changed distance and angle to the region of interest, and dynamically selects and activates 10 corresponding input and output light ports accordingly during the movement of the transducer arrangement **110**.

In another embodiment of the invention, an imaging apparatus may be used to monitor changes in the concentration of analyte(s) in a region of interest during therapeutic or surgical procedures (such as during the application of high 15 power ultrasound pulses or wave, laser ablation, or chemical procedures). For example, the transducer arrangement **110** may be used for ablation of tumors or malformations in a tissue. During the application of high power ultrasound pulses, light is emitted and collected by illumination and detection assemblies, respectively, to determine the concentration of an analyte indicative of the 20 treatment in the region being ablated. Alternatively, low power ultrasound pulses (that do not cause ablation) intermittently irradiate the region of interest, while low-power light pulses are emitted and collected by illumination and detection assemblies to determine the concentration of an analyte indicative of the treatment. For example, oxygenation of the region of interest is monitored. Such 25 information is used for controlling and monitoring the treatment during application of ultrasound radiation.

Reference is now made to **Figs. 6A and 6B** exemplify yet another configuration of a flexible probe, generally designated **603**, suitable to be used in either one of the above described monitoring apparatuses. The flexible probe **603** 30 includes a flexible support **301**, for example made of electrically isolating



- 55 -

material(s), carrying light ports **303-316** (fiber-ends in appropriate housing, or light sources and/or light detectors as described above) and an acoustic output port **302** (or acoustic transducer arrangement). **Fig. 6A** presents a bottom view of the flexible probe **603** viewed from the side by which it is attachable to a skin, and **Fig. 6B** shows a side view diagram of the flexible probe **603**. Optical fibers or electric wires **330-336** and **340-346** connect the light ports **310-316** and **303-309**, respectively, to a common connector **320**. An isolated electric cable (or acoustic waveguides) connects the acoustic output port **302** to the same connector **320** associated with a control unit. The acoustic port **302** is preferably coupled to an ultrasound transducer arrangement **327** connected to the flexible support **301** using vibration controlling elements. The connector **320** couples the optical fibers and cables attached to the flexible support **301** with optical fibers and electric cables coupled to the control unit (not shown here). The connector **320** may be composed of several connector elements. An adhesive **325** is attached to the bottom side of the support **301**, such that the probe **603** can be fixed to the skin using this adhesive **325**. Adhesive **325** is preferably transparent and produces minimal scattering in a wavelength range used for measurements (i.e., emitted by light sources). Alternatively or additionally, the adhesive **325** may form an optical index matching layer between the light ports and the skin. Alternatively, the adhesive **325** may not cover the light ports at all, or may partially cover them. The adhesive **325** may contain pigments, chromophores or other materials for controlling the transmission of different wavelengths of light. An adhesive gel **326** is located below the acoustic port **302**. The adhesive gel **326** is made from the same or different material as the adhesive **325** and is designed for optimal acoustic coupling between the acoustic port **302** and the skin. Possible materials for adhesives **325** and **326** include hydrogel based adhesives.

The different elements of the flexible probe **603** may be assembled in different ways. For example, the complete probe **603** is assembled prior to operation, and a user only needs to remove a thin layer covering the bottom side of adhesives **325** and **326**. In yet another example, the adhesive **326** is attached to

- 56 -

the acoustic output port **302** (preferably including the acoustic transducer arrangement itself) which is not attached to the probe **603** prior to the device operation. The user first attaches the flexible support **301** to the skin using the adhesive **325**, and then inserts the acoustic port **302** through an appropriately provided opening in the support **301**, where the transducer **327** is optionally  
5 connected to the support **301** using conventional means and is attached to the skin using the adhesive **326**. The latter may be part of adhesive **325**, and only the acoustic output port **302** is inserted and attached to the upper part of the adhesive **326** (being a double sided adhesive). The user first attaches the adhesives **325**  
10 and **326** to skin, then attaches the acoustic port **302** to the adhesive **326**, and then connects the support **301** to the upper side of the adhesive **325** (being a double sided adhesive). Finally, the connector **320** is connected to the cables and fibers from the control unit to allow the operation of the probe. Each element of the flexible probe **603** and the complete probe **603** as a unit may be used only once  
15 and then discarded (i.e., is disposable), or used multiple times.

**Figs. 7A and 7B** show yet another example of a flexible probe **703** according to the present invention. Here, a support **301** carries light ports (or light sources) and several acoustic ports **302**, **319** and **319A** (or acoustic transducer arrangements). Each of the acoustic ports **302**, **319** and **319A** is  
20 coupled to a connector **320** using cables **338**, **339** and **339A**, respectively. Adhesive gels **326**, **329** and **329A** are used to couple the acoustic ports **302**, **319** and **319A**, respectively, to the skin. Similarly, each of the acoustic ports may be separated from the probe **703** when not in use, and inserted by user for as preparation for operation.

25 Those skilled in the art will readily appreciate that various modifications and changes may be applied to the embodiments of the invention as hereinbefore described without departing from its scope defined in and by the appended claims.

**CLAIMS:**

1. A monitoring system for use in non-invasively monitoring at least one parameter of a region of interest in a human body, the system comprising:
  - 5 - a measurement unit comprising an optical unit having an illumination assembly configured to define at least one output port for illuminating light, and a light detection assembly configured to define at least one light input port for collecting light scattered from the illuminated body portion and to generate measured data indicative of the collected light; and an  
10 acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range; the measurement unit being configured and operable to provide an operating condition such that the acoustic waves of the predetermined frequency range overlap with an illuminating region within the region of interest and substantially do not overlap with a region outside the region of interest, and that the detection assembly collects light  
15 scattered from the region of interest and light scattered from the region outside the region of interest, the measured data being thereby indicative of scattered light having both ultrasound tagged and untagged light portions, thereby enabling to distinguish between light responses of the region of interest and the region outside the region of interest;
  - 20 - a control unit, which is connectable to the optical unit and to the acoustic unit to operate these units, the control unit being responsive to the measured data and preprogrammed to process and analyze the measured data to extract therefrom a data portion associated with the light response of the region of interest and determine said at least one parameter of the  
25 region of interest.
2. The system of Claim 1, wherein the acoustic unit is configured for generating unfocused ultrasound radiation.
3. The system of Claim 1, configured to determine oxygen saturation of the region of interest.

4. The system of Claim 1, configured to utilize principles of oximetry to determine the desired parameter of the region of interest.
5. The system of Claim 1, configured to utilize principles of pulse oximetry to determine the desired parameter of the region of interest.
- 5 6. The system of Claim 1, wherein the measurement unit is configured and operable to generate the measured data indicative of variations of the ultrasound tagged light signal as a function of at least one of time and wavelength of the illuminating light, the control unit being configured to analyze the variations of the ultrasound tagged light signal to determine maxima/minima of the signal and  
10 calculate oxygen saturation.
7. The system of Claim 1, wherein the control unit is preprogrammed and operable to provide optimal positioning of the optical unit and the acoustic unit, said optimal positioning satisfying said operating condition.
8. The system of Claim 7, wherein the control unit is operable to provide a  
15 relative displacement between a direction of propagation of the acoustic radiation and at least one of the illumination and detection assemblies so as to provide said optimal positioning.
9. The system of Claim 1, wherein the illumination assembly is configured to define at least two of the spaced-apart light output ports.
- 20 10. The system of Claim 9, wherein the control unit is operable to select at least one of the light output ports for measurements, to provide said operating condition.
11. The system of Claim 1, wherein the detection assembly is configured to define at least two spaced-apart light input ports.
- 25 12. The system of Claim 11, wherein the control unit is operable to select at least one of the light input ports for measurements, to provide said operating condition.
13. The system of Claim 10, wherein the detection assembly is configured to define at least two spaced-apart light input ports.

14. The system of Claim 13, wherein the control unit is operable to select at least one of the light input ports for measurements, to provide said operating condition.
15. The system of Claim 8, wherein the control unit is operable to displace at least one of the light input or output ports with respect to the direction of the acoustic waves propagation.
16. The system of Claim 1, wherein the illumination assembly is configured and operable to generate the illuminating light of at least two different wavelengths.
- 10 17. The system of Claim 16, where the two different wavelengths are selected as those differently absorbable by oxygenated and deoxygenated hemoglobin in the body.
18. The system of Claim 16, where the two different wavelengths are selected as those characterized by the same absorption by tissue or fluid components in the body.
- 15 19. The system of Claim 18, where the two different wavelengths are selected to be differently scattered by tissue or fluid components in the body.
- 20 20. The system of Claim 16, wherein the illumination assembly is operable to generate said at least two different wavelengths at different times, respectively.
21. The system of Claim 16, wherein the at least two wavelengths are differently modulated by at least one of frequency and phase characteristics.
22. The system of Claim 20, wherein the at least two wavelengths are differently modulated by at least one of frequency and phase characteristics.
23. The system of Claim 1, wherein the detection assembly is configured to define the light input port which, when in the operating condition, detects both the light scattered from the region of interest and the light scattered from the outside of the region of interest.
- 25 24. The system of Claim 1, wherein the detection assembly is configured to define an array of at least two spaced-apart light input ports, such that when in the operating condition, at least one of the light input ports collects the tagged
- 30

- 60 -

light coming from the region of interest, and at least one other of the light input ports collects the untagged light scattered from the region outside the region of interest.

25. The system of Claim 1, wherein the control unit is preprogrammed for  
5 operating the optical unit a predetermined time after the operation of the acoustic unit, thereby providing said operating condition.

26. The system of Claim 25, wherein said predetermined time corresponds to a time needed for the acoustic waves of the predetermined frequency to reach the region where the overlapping with the illuminating light is to be provided.

10 27. The system of Claim 11, wherein at least one of the light input ports is located proximate of the light output port thereby collecting the light scattered from the region outside of the region of interest, and the at least one other light input port is located at a larger distance from the light output port thereby collecting the light coming from the region of interest and being tagged by the  
15 ultrasound radiation.

28. The system of Claim 11, wherein at least one of the light input ports is located proximate of the light output port thereby collecting untagged light scattered from the region outside of the region of interest; and the at least one other light input port is located at a larger distance from the light output port  
20 thereby collecting ultrasound tagged light coming from the region of interest through the outside region.

29. The system of Claim 28, wherein the control unit is preprogrammed to process and analyze the first measured data generated by said at least one light input port to determine data indicative of a characteristic of a light response of  
25 the region outside of the region of interest to the illuminating light; and to utilize said data indicative of the light response characteristic in the processing and analyzing of the second measured data generated by said at least one other input port to thereby extract the light response of the region of interest.

30. The system of Claim 11, wherein the control unit is preprogrammed for operating at least one of the light input ports a predetermined time after the operation of the acoustic unit, thereby providing said operating condition.

31. The system of Claim 30, wherein said predetermined time corresponds to  
5 a time needed for the acoustic waves of the predetermined frequency to reach the region where the overlapping with the illuminating light is to be provided.

32. The system of Claim 31, wherein said at least one of the light input ports is located proximate of the light output port thereby collecting untagged light scattered from the region outside of the region of interest; and said at least one  
10 other light input port is located at a larger distance from the light output port as compared to a location of at least one other light input port thereby collecting tagged light coming from the region of interest through the region outside of the region of interest and being therefore affected by the outside region.

33. The system of Claim 32, wherein the control unit is preprogrammed to  
15 process and analyze the first measured data generated by said at least one light input port to determine data indicative of a characteristic of a light response of the region outside of the region of interest to the illuminating light; and to utilize said data indicative of the light response characteristic in the processing and analyzing of the second measured data generated by said at least one other input  
20 port to thereby extract the light response of the region of interest.

34. The system of Claim 1, comprising a support structure configured to be brought in contact with a body, said support structure carrying at least the light output and light input ports of the illumination and detection assemblies, respectively.

25 35. The system of Claim 34, wherein said support structure carries the illumination and detection assemblies.

36. The system of Claim 34, wherein the support structure carries the acoustic unit.

37. The system of Claim 34, wherein the support structure carries an  
30 ultrasound output port of the acoustic unit.

38. The system of Claim 34, wherein the support structure is flexible to wrap the body portion.
39. The system of Claim 1, wherein the acoustic unit is configured for generating ultrasound radiation in the form of one of the following: continuous  
5 wave, pulse, burst.
40. The system of Claim 1, wherein the acoustic unit is configured for generating focused ultrasound radiation, where a focal length corresponding to a distance from the acoustic unit to the region of interest
41. The system of Claim 1, wherein the acoustic unit comprises a phase  
10 array of ultrasound transducers.
42. The system of Claim 1, wherein the control unit is configured for receiving data indicative of reflections of the ultrasound waves from an irradiated volume inside of the body for creating data indicative of an image of the irradiated volume, and analyzing said image data to provide said operating  
15 condition for measurements.
43. The system of Claim 1, wherein the control unit is configured for receiving data indicative of reflections of the ultrasound waves from an irradiated volume inside of the body for determining data indicative of a Doppler shift in the ultrasound radiation reflections.
- 20 44. The system of Claim 1, wherein said predetermined frequency range of the ultrasound radiation is from 50kHz to 8MHz.
45. The system of Claim 1, wherein the ultrasound radiation is of a frequency lower than 1MHz.
46. The system of Claim 1, configured for use in monitoring at least one of  
25 the following parameters: oxygen saturation level of the region of interest, and concentration of a substance or structure in the region of interest.
47. A monitoring system for use in non-invasively monitoring at least one parameter of a region of interest in a human body, the system comprising:
- a measurement unit comprising an optical unit configured for generating  
30 illuminating light and for collecting light and generating measured data



- 63 -

indicative of the collected light; and an acoustic unit configured to generate unfocused acoustic waves of a predetermined ultrasound frequency range; the measurement unit being configured and operable to provide an operating condition such that the acoustic waves of the predetermined frequency range overlap with an illuminating region within the region of interest while substantially not overlapping within a region outside the region of interest, and that the detection assembly collects light scattered from the region of interest and light scattered from the region outside the region of interest, the measured data being thereby indicative of scattered light having both ultrasound tagged and untagged light portions, thereby enabling to distinguish between light responses of the region of interest and the region outside the region of interest;

- a control unit, which is connectable to the optical unit and to the acoustic unit to operate these units, the control unit being responsive to the measured data and preprogrammed to process and analyze the measured data to extract therefrom a data portion associated with the light response of the region of interest and determine said at least one parameter of the region of interest.

**48.** A system for use in noninvasive monitoring at least one parameter of a region of interest in a human body, the system comprising:

- a measurement unit comprising an optical unit having an illumination assembly configured to define at least one output port for illuminating light, and a light detection assembly configured to define at least one light input port for collecting light and to generate measured data indicative of the collected light; and an acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range;

- a control unit, which is connectable to the optical unit and to the acoustic unit to operate these units, the control unit being responsive to the measured data and preprogrammed to process and analyze the measured data to extract therefrom data indicative of a light response of the region

- 64 -

of interest and determine the at least one desired parameter, the control unit being operable to provide optimal positioning of the optical unit and the acoustic unit to satisfy an operating condition for measurements, said operating condition being such that acoustic waves of the predetermined frequency range and illuminating light overlap within the region of interest and substantially do not overlap in a region outside the region of interest, and in that the detection assembly collects the light scattered from the region of interest and light scattered from the region outside the region of interest; the measured data being thereby indicative of scattered light having both ultrasound tagged and untagged light portions, thereby enabling to analyze the measured data to extract therefrom a data portion associated with the light response of the region of interest and determine said at least one parameter of the region of interest.

49. A system for use in non-invasively monitoring at least one parameter of a fetus-related region of interest, the system comprising:

- a measurement unit comprising an optical unit having an illumination assembly configured to define at least one output port for illuminating light, and a light detection assembly configured to define at least one light input port for collecting light and to generate measured data indicative of the collected light; and an acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range;
- a control unit, which is connectable to the optical unit and to the acoustic unit to operate these units, the control unit being responsive to the measured data and preprogrammed to process and analyze the measured data to extract therefrom data indicative of a light response of the region of interest and determine the at least one desired parameter of the region of interest, the control unit being operable to provide optimal positioning of the optical unit and the acoustic unit to satisfy an operating condition for measurements, said operating condition being such that acoustic waves of the predetermined frequency range and the illuminating light overlap

- 65 -

within the region of interest and substantially do not overlap in a maternal tissues region outside the region of interest, and in that the detection assembly collects the light scattered from the fetus-related region and light scattered from maternal tissues region; the measured data being indicative of the tagged light scattered by the fetus-related region and tagged and untagged light scattered by the maternal tissues region, thereby enabling to analyze the measured data to extract the data indicative of the light response of the fetus-related region and determine the at least one desired parameter of the fetus-related region.

5  
10 **50.** A system for use in noninvasive monitoring oxygen saturation level of a region of interest in a human or animal body, the system comprising:

- a measurement unit comprising a pulse oxymeter unit having an illumination assembly configured to define at least one output port for illuminating light of at least two different wavelengths, and a light detection assembly configured to define at least one light input port for collecting light and to generate measured data indicative of the collected light; and an acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range;
- a control unit, which is connectable to the pulse oxymeter unit and to the acoustic unit to operate these units, the control unit being responsive to the measured data and preprogrammed to process and analyze the measured data to determine time dependent variations of ultrasound-tagged light signals for said at least two wavelengths scattered from the region of interest and analyze these time variations to calculate the oxygen saturation of the region of interest.

15  
20  
25  
30 **51.** A probe device for use in a system for monitoring at least one parameter of a region of interest in a human body, the probe comprising: a support structure configured to contact a body portion, said support structure carrying an array of at least two light output ports arranged in a spaced-apart relationship and being optically coupled to a light source assembly, an array of light input ports

arranged in a spaced-apart relationship and being optically coupled to a light detection assembly, and at least one acoustic output port of an acoustic unit, the arrangement of the light ports and the acoustic unit being such as to allow selection of at least one of said light output ports, at least one of the light input  
5 ports and at least one of the acoustic output ports such that acoustic waves of a predetermined frequency range coming from said at least one selected acoustic output port and illuminating light coming from said at least one selected light output port overlap within a region of interest in the body, and in that said at least one light input port collects light scattered from the overlapping region and light  
10 scattered from outside the region of interest.

**52.** A probe device for use in a system for monitoring at least one parameter of a region of interest in a human body, the probe comprising: a support structure configured to contact a body portion, said support structure carrying at least one light output port optically coupled to a light source assembly, at least two light  
15 input ports optically coupled to a light detection assembly, and at least one acoustic output port of an acoustic unit, the arrangement of the light and acoustic ports being such as to allow selection of the light and acoustic ports for measurements such that, with the selected ports, acoustic waves of a predetermined frequency range and illuminating light overlap within a region of  
20 interest in the body and that the at least one light input port collects light scattered from the overlapping region and light scattered from outside of the region of interest.

**53.** A probe device for use in a system for monitoring at least one parameter of a region of interest in a human body, the probe comprising: a support structure  
25 configured to contact a body portion, said support structure carrying at least two light output port optically coupled to a light source assembly, at least one light input port optically coupled to a light detection assembly, and at least one acoustic output port of an acoustic unit, the arrangement of the light and acoustic ports being such as to allow selection of the light and acoustic ports for  
30 measurements such that, with the selected ports, acoustic waves of a

predetermined frequency range and illuminating light overlap within a region of interest in the body and that detected light includes scattered from the overlapping region and light scattered from outside of the region of interest.

**54.** A method for use in noninvasive monitoring at least one parameter of a region of interest in a human body, the method comprising: operating an optical unit and an acoustic unit so as to provide that ultrasound waves of a predetermined frequency range and illuminating light overlap within the region of interest and substantially do not overlap with a region outside the region of interest, thereby producing measured data indicative of collected light including scattered light having ultrasound tagged and untagged light portions, thereby enabling extraction of a light response of the region of interest from all the other light portions in the collected light

**55.** The method of Claim 54, comprising irradiating the region of interest with unfocused ultrasound radiation.

**56.** The method of Claim 54, comprising irradiating the region of interest with focused ultrasound radiation.

**57.** The method of Claim 54, for determining oxygen saturation of the region of interest.

**58.** The method of Claim 54, utilizing principles of oximetry to determine the desired parameter of the region of interest.

**59.** The method of Claim 54, utilizing principles of pulse oximetry to determine the desired parameter of the region of interest.

**60.** The method of Claim 54, comprising applying the illumination of at least two different wavelengths, and generating the measured data indicative of variations of the ultrasound tagged light signal as a function of at least one of time and wavelength of the illuminating light.

**61.** The method of Claim 60, comprising analyzing the time variations to determine maxima/minima of the signal and calculate oxygen saturation of the region of interest.

- 62.** The method of Claim 54, comprising providing a relative displacement between a direction of propagation of the acoustic radiation and at least one of illumination and detection assemblies of the optical unit.
- 63.** The method of Claim 54, comprising selecting for measurements at least  
5 one light output port, from a plurality of spaced-apart light output ports, of the optical unit.
- 64.** The method of Claim 54, comprising selecting for measurements at least one light input port from a plurality of spaced-apart light input ports of the optical unit.
- 10 **65.** The method of Claim 63 comprising selecting for measurements at least one light input port from a plurality of spaced-apart light input ports of the optical unit.
- 66.** The method of Claim 62, comprising displacing at least one of light input or light output ports of the optical unit with respect to the direction of the  
15 acoustic waves propagation.
- 67.** The method of Claim 54, comprising applying ultrasound imaging to the region of interest surrounded by the outside region.
- 68.** The method of Claim 67, wherein the imaging is carried out using said acoustic unit.
- 20 **69.** The method of Claim 54, comprising detecting reflections of the ultrasound radiation from the irradiated body portion and determining a Doppler shift in the detected reflections.
- 70.** The method of Claim 54, comprising operating an illumination assembly of the optical unit to generate the illuminating light of at least two different  
25 wavelengths.
- 71.** The method of Claim 70, wherein said at least two different wavelengths are differently absorbed by oxygenated and deoxygenated hemoglobin.
- 72.** The method of Claim 70, wherein said at least two different wavelengths are equally absorbed by tissue or fluid components.

73. The method of Claim 72, wherein said at least two different wavelengths are differently scattered by tissue or fluid components.
74. The method of Claim 70, comprising operating the illumination assembly to generate the illuminating light of said at least two different  
5 wavelengths at different times, respectively.
75. The method of Claim 70, comprising operating the illumination assembly to generate the illuminating light of said at least two different wavelengths differently modulated by at least one of frequency and phase characteristics.
- 10 76. The method of Claim 74, comprising operating the illumination assembly to generate the illuminating light of said at least two different wavelengths differently modulated by at least one of frequency and phase characteristics.
77. The method of Claim 54, comprising detecting both the light scattered  
15 from the region of interest and the light scattered from the outside of the region of interest.
78. The method of Claim 54, comprising operating at least two spaced-apart light input ports of the optical unit, such that at least one of the light input ports collects the light scattered from the region of interest, and at least one other of  
20 the light input ports collects the light scattered from the outside region.
79. The method of Claim 54, comprising operating the optical unit a predetermined time after the operation of the acoustic unit, said predetermined time corresponding to a time needed for the acoustic waves of the predetermined frequency to reach the region of interest where the overlapping with the  
25 illumination light is to be provided.
80. The method of Claim 54, comprising operating at least two spaced-apart light input ports of the optical unit, such that at least one of the light input ports collects the untagged light scattered from the region outside the region of interest, and at least one other of the light input ports collects the tagged light

coming from the region of interest through the region outside the region of interest.

**81.** The method of Claim 80, comprising collecting the untagged light scattered from the region outside of the region of interest by said at least one  
5 light input port located proximate of the light output port, and collecting the tagged light coming from the region of interest through the region outside the region of interest by said at least one other light input port located at a larger distance from the light output port.

**82.** The method of Claim 81, comprising processing and analyzing the first  
10 measured data generated by said at least one light input port to determine data indicative of a characteristic of a light response of the region outside the region of interest to the illuminating light; and utilizing said data indicative of the light response characteristic in the processing and analyzing of the second measured data generated by said at least one other light input port, to thereby extract from  
15 the second measured data the data indicative of the light response of the region of interest.

**83.** The method of Claim 54, wherein said at least one desired parameter includes at least one of the following: oxygen saturation level; and concentration of a substrate or structure in the region of interest.

20 **84.** The method of Claim 54, for use in noninvasive monitoring a fetus condition.

**85.** The method of Claim 84, wherein said at least one desired parameter is that of fetus blood.

**86.** The method of Claim 84, wherein said at least one desired parameter  
25 includes oxygen saturation level.

**87.** The method of Claim 84, wherein said at least one desired parameter includes concentration of a substance or a structure within the fetus.

**88.** The method of Claim 84, wherein said at least one desired parameter includes concentration of a substance or a structure within amniotic fluid.



89. The method of Claim 88, wherein said at least one desired parameter includes a level of lung maturity of the fetus.

90. The method of Claim 89, comprising determining presence and concentration of lamellar bodies in the amniotic fluid.

5 91. The method of Claim 88, wherein said at least one desired parameter includes at least one of the following: presence of meconium in the amniotic fluid and concentration thereof.

92. The method of Claim 88, wherein said at least one desired parameter includes at least one of the following: presence of blood in the amniotic fluid and  
10 concentration thereof.

93. The method of Claim 54, for noninvasive monitoring optical properties of an extravascular fluid, such as pleural, pericardial, peritoneal and synovial fluids.

94. A method for use in noninvasive monitoring oxygen saturation level, the  
15 method comprising: applying ultrasound tagging of light in pulse oxymetric measurements, obtaining measured data indicative of variations of ultrasound tagged light signals scattered from a region of interest as a function of at least one of time and wavelength of illuminating light, and analyzing the measured data to calculate the oxygen saturation level.

20 95. A method for use in noninvasive monitoring at least one parameter of a region of interest in a human body, the method comprising:

- providing an optical unit having an illumination assembly configured to define at least one output port for illuminating light; and a light detection assembly configured to define at least one light input port for collecting  
25 light, and to generate measured data indicative of the collected light; and providing an acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range;
- providing an optimal positioning of the optical and acoustic units with respect to each other and with respect to the region of interest, said  
30 optimal positioning satisfying an operating condition resulting in that the

ultrasound waves of the predetermined frequency range overlap with the illuminating light within the region of interest, while substantially not overlapping a region outside the region of interest, and in that the detection assembly collects light scattered from the overlapping region and from the region outside the region of interest;

5

- operating the optical and acoustic units, when in the optimal positioning, and generating measured data indicative of collected light including scattered light having ultrasound tagged and untagged light portions, thereby enabling extraction of a light response of the region of interest from all the other light portions in the collected light.

10

**96.** A method for use in noninvasive monitoring at least one parameter of a fetus-related region of interest, the method comprising:

- providing an optical unit having an illumination assembly configured to define at least one output port for the illuminating light; and a light detection assembly configured to define at least one light input port for collecting light, and to generate measured data indicative of the collected light; and providing an acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range;

15

- providing an optimal positioning of the optical and acoustic units with respect to each other and with respect to the fetus-related region of interest, said optimal positioning satisfying an operating condition resulting in that the ultrasound waves of the predetermined frequency range overlap with the illuminating light within the fetus-related region of interest, while substantially not overlapping in a maternal tissues region outside the fetus-related region of interest, and in that the detection assembly collects light scattered from the fetus-related region of interest and from the maternal tissues region;

20

25

- operating the optical and acoustic units, when in the optimal positioning, and generating measured data indicative of collected light including scattered light having ultrasound tagged and untagged light portions,

30

- 73 -

thereby enabling extraction from the measured data a data portion indicative of a light response of the fetus-related region of interest.

**97.** The method of Claim 96, wherein the fetus-related region of interest includes the fetus only.

5 **98.** The method of Claim 97, wherein said at least one parameter includes at least one of oxygen saturation level and concentration of a substance or structure in the fetus region.

**99.** The method of Claim 96, wherein the fetus-related region of interest includes an amniotic fluid region only.

10 **100.** The method of Claim 99, wherein said at least one parameter includes at least one of the following: a level of lung maturity of the fetus; presence of lamellar bodies; concentration of lamellar bodies; presence of meconium; concentration of meconium; presence of blood; and concentration of blood.

**101.** A method for operating a monitoring system configured for noninvasive  
15 monitoring at least one parameter of a region of interest in a human body, which system comprises an optical unit and an acoustic unit configured to generate acoustic waves of a predetermined ultrasound frequency range, the method comprising:

operating the monitoring system to provide an optimal positioning of the  
20 optical and acoustic units with respect to each other and with respect to the region of interest to satisfy an operating condition for measurements, said operating condition resulting in that the ultrasound waves of the predetermined frequency range overlap with illuminating light generated by the optical unit within the region of interest, while substantially not  
25 overlapping in a region outside the region of interest, and in that a detection assembly of the optical unit collects light scattered from the region of interest and from the region outside the region of interest, thereby obtaining measured data indicative of scattered light having ultrasound tagged and untagged light portions, and enabling extraction from said measured data a data portion  
30 indicative of a light response of the region of interest.

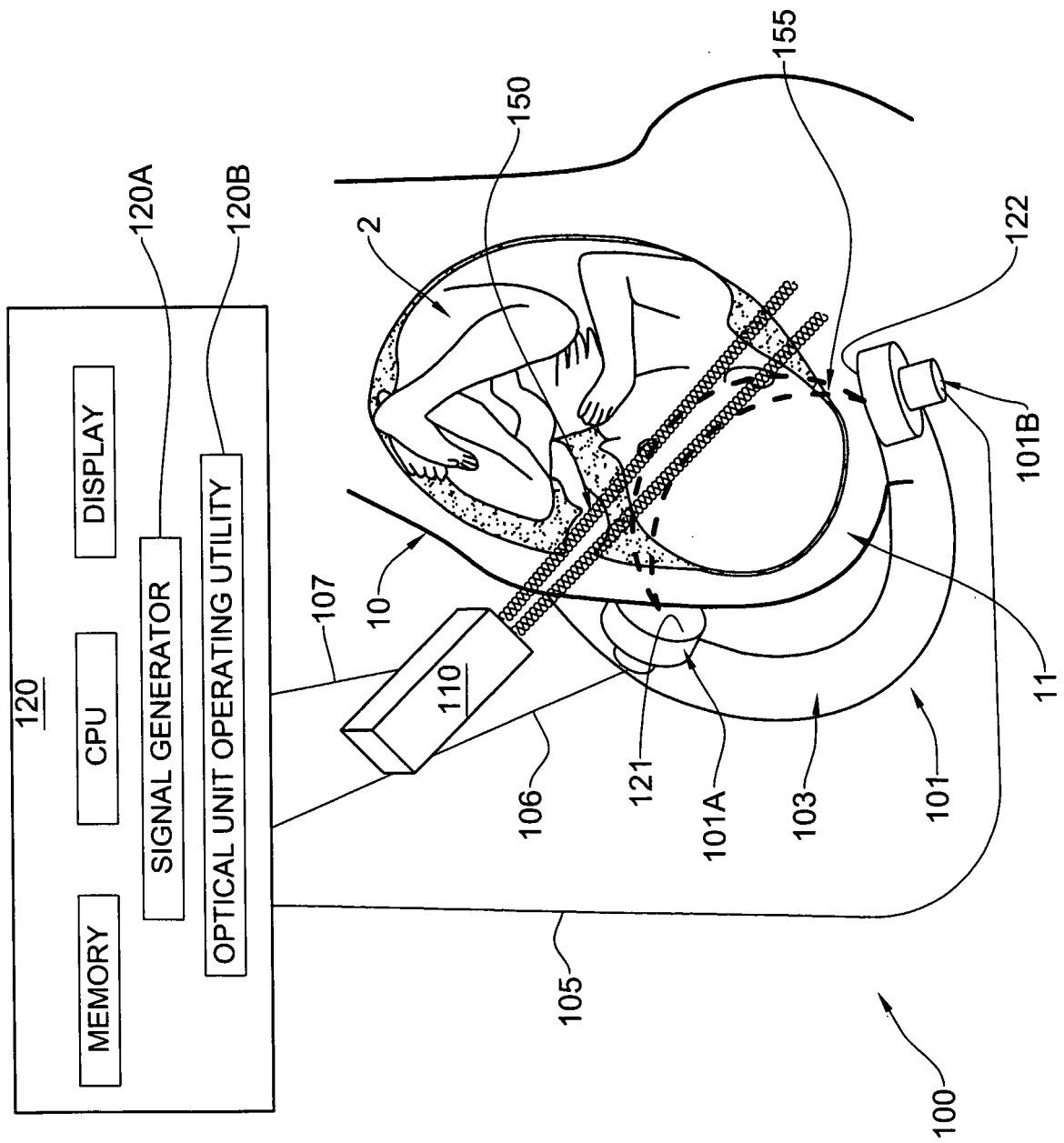


FIG. 1A

2/11

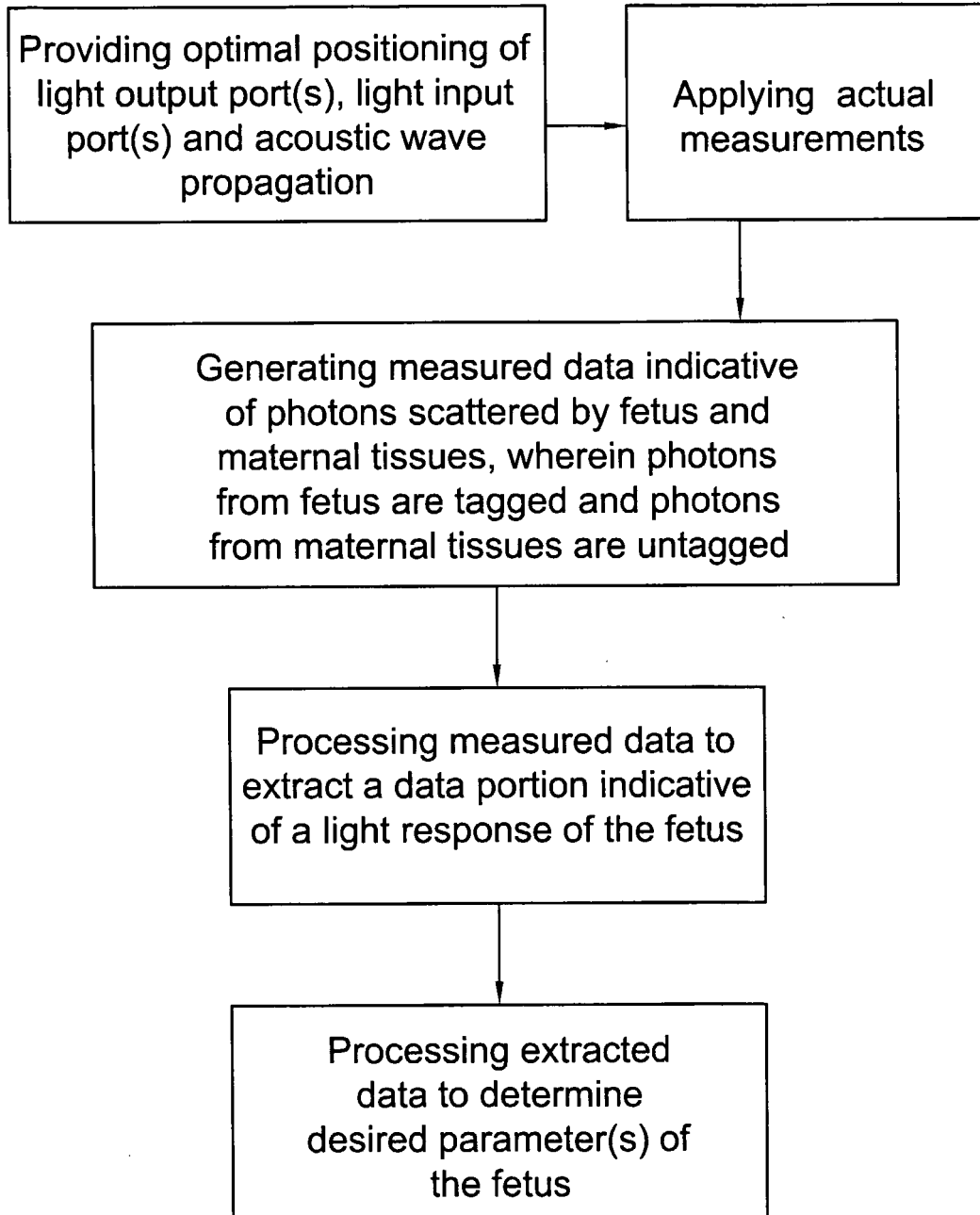


FIG. 1B

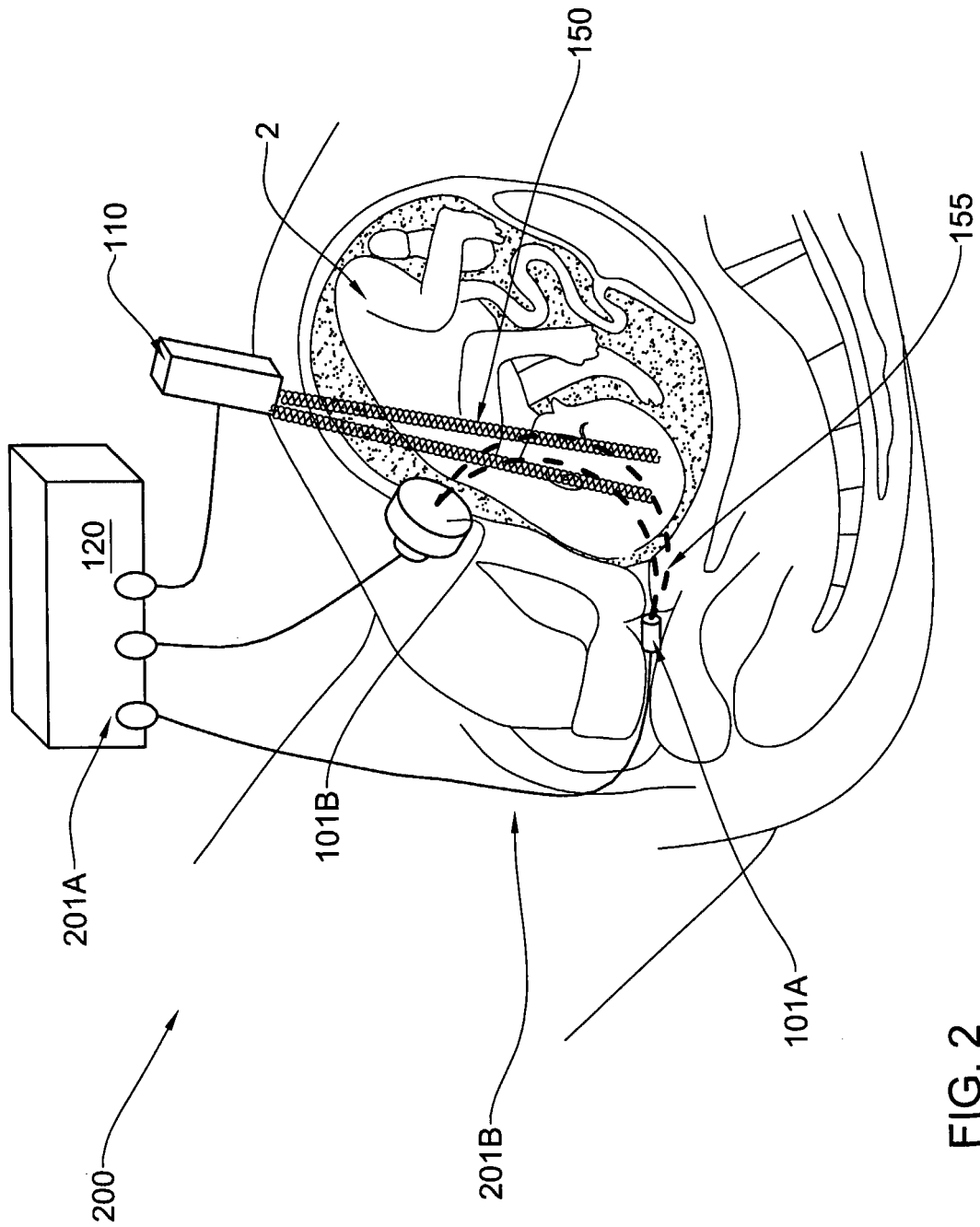


FIG. 2

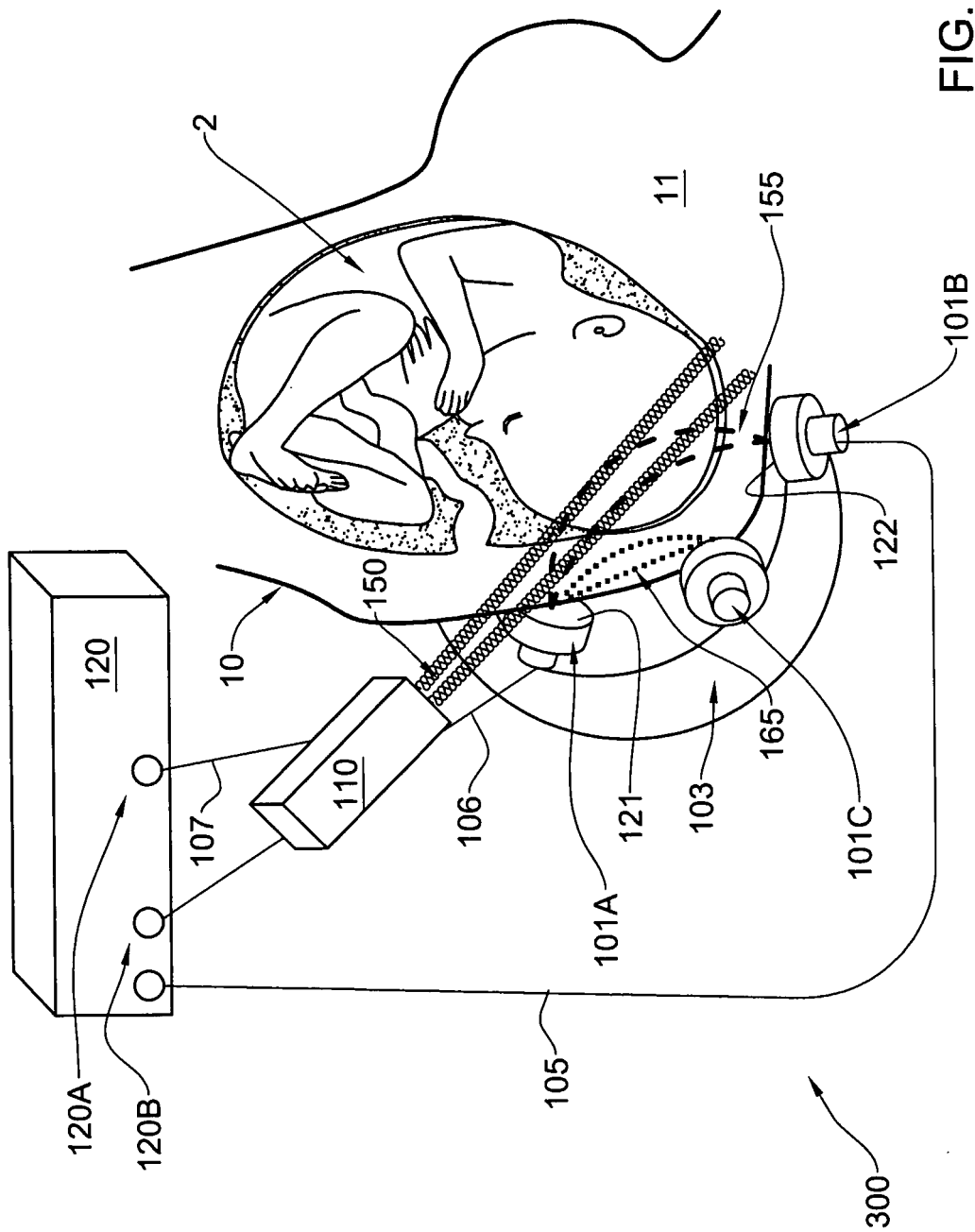


FIG. 3A

5/11

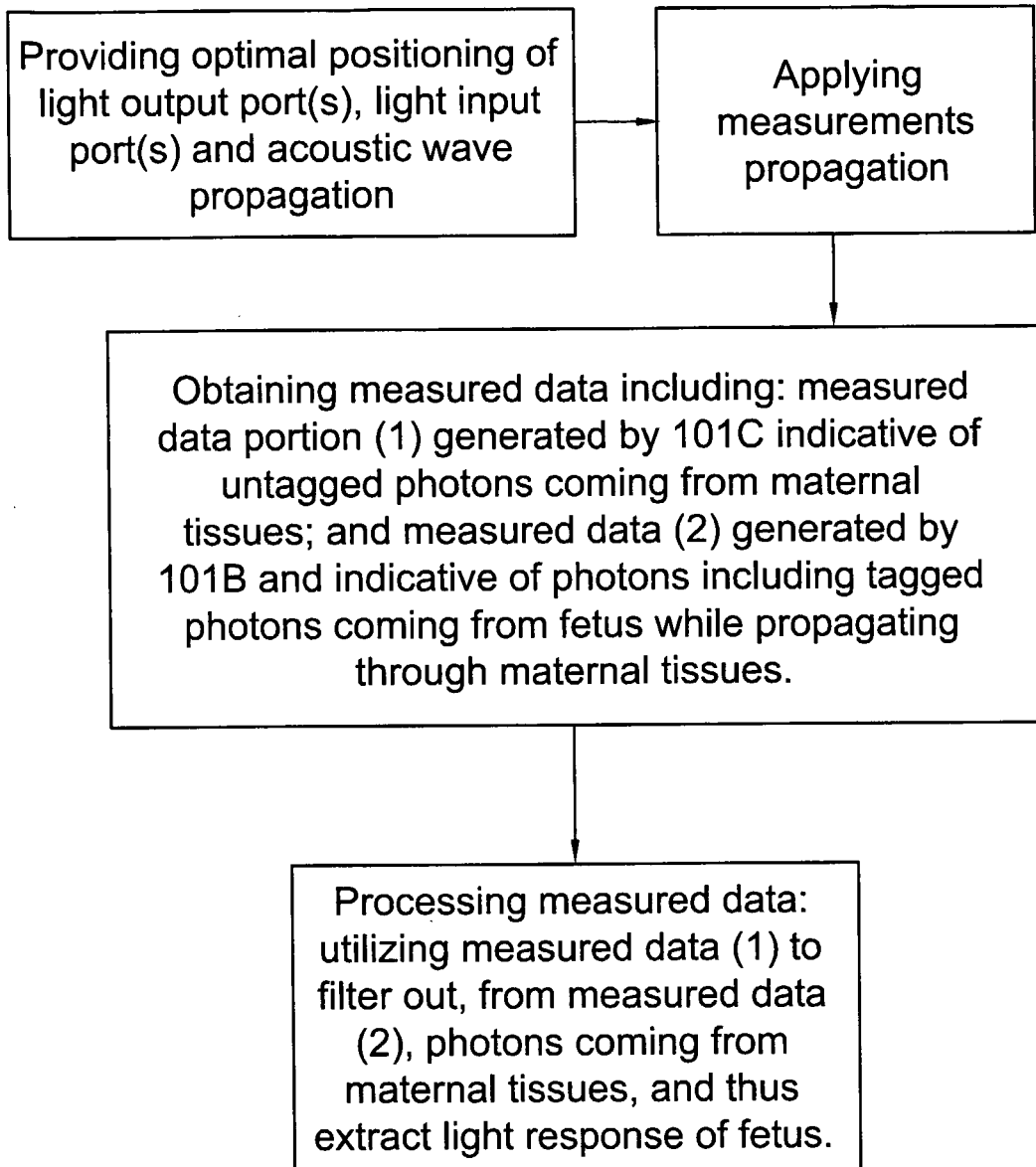


FIG. 3B



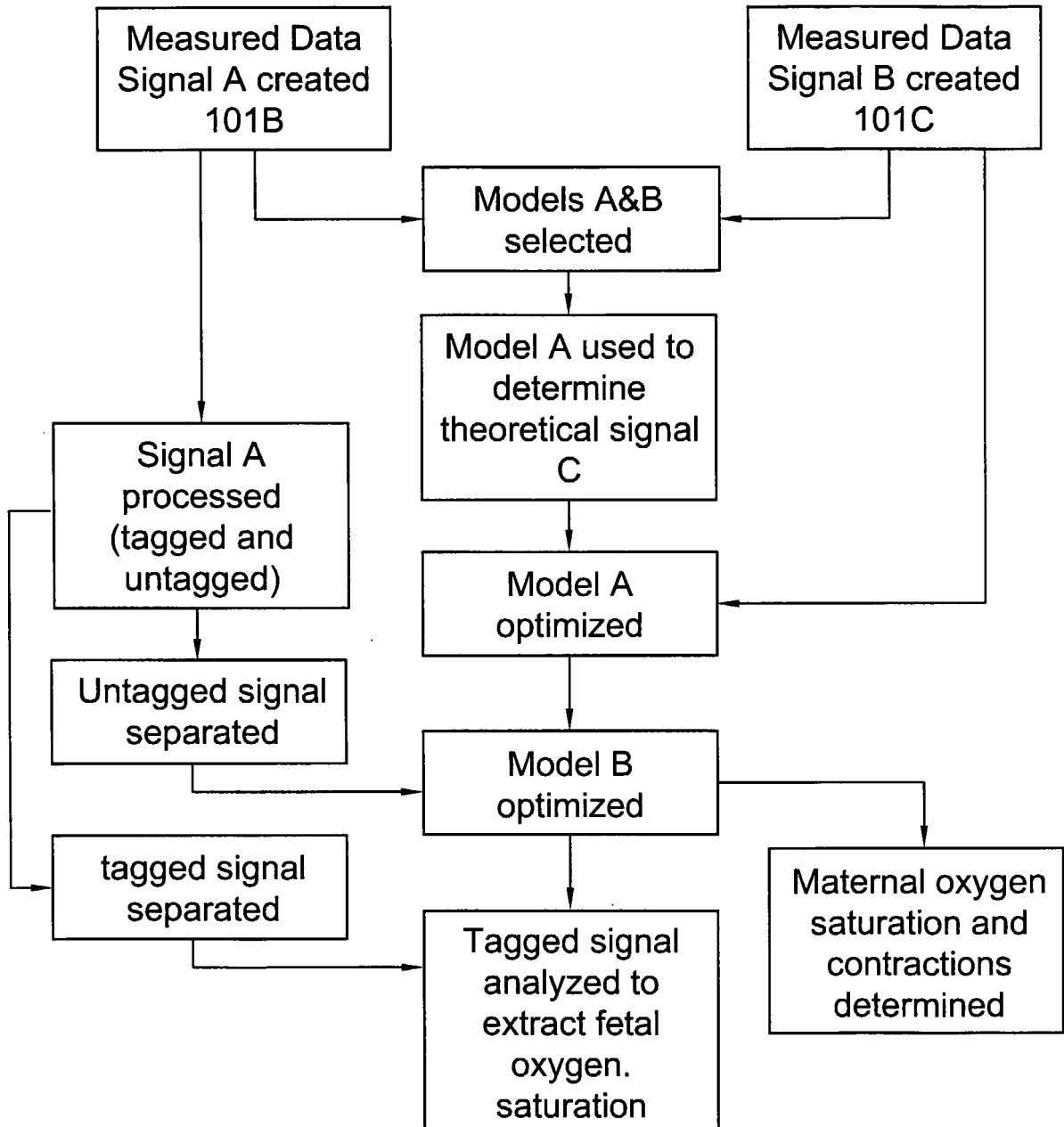


FIG. 3C

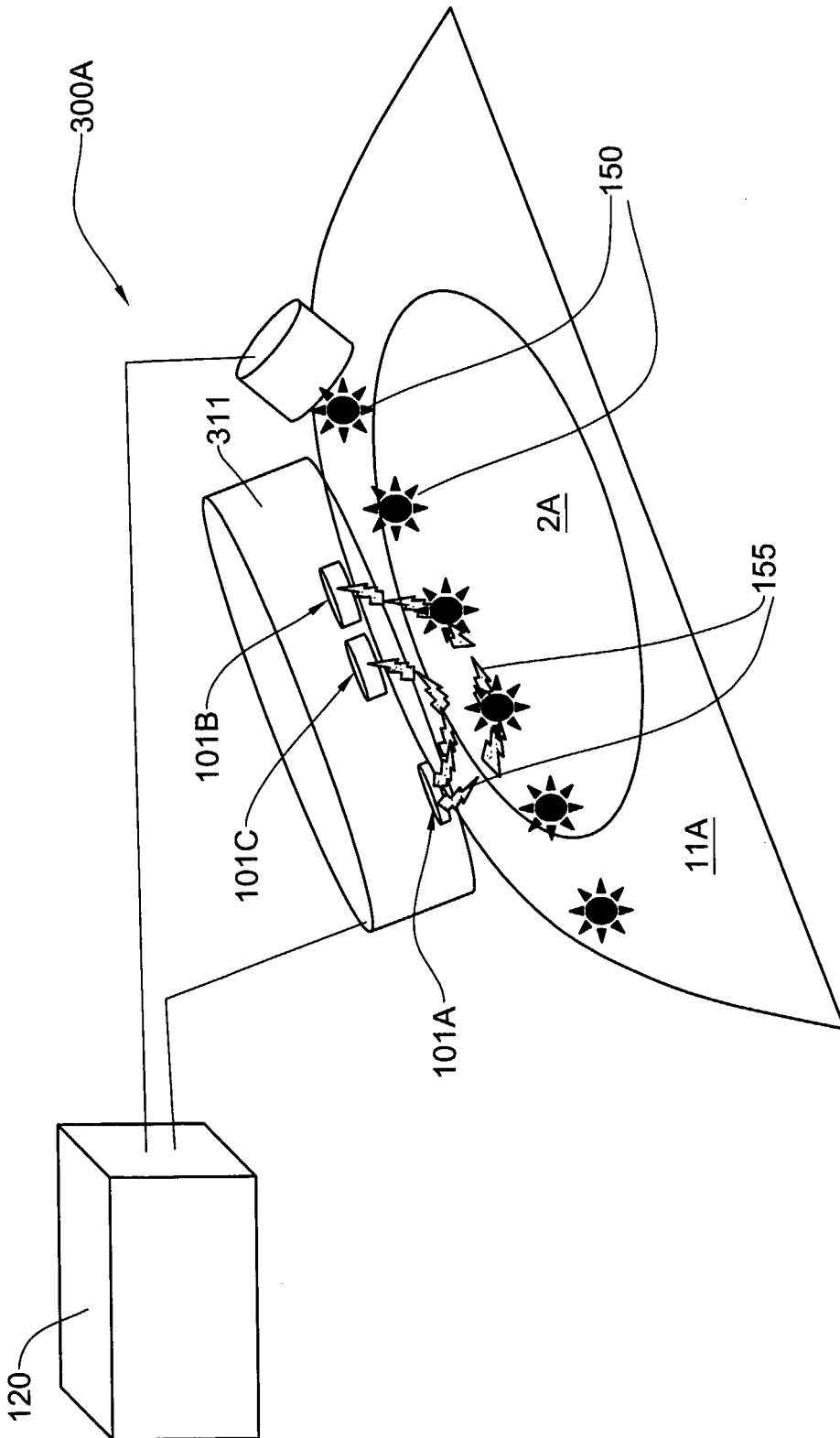


FIG. 3D

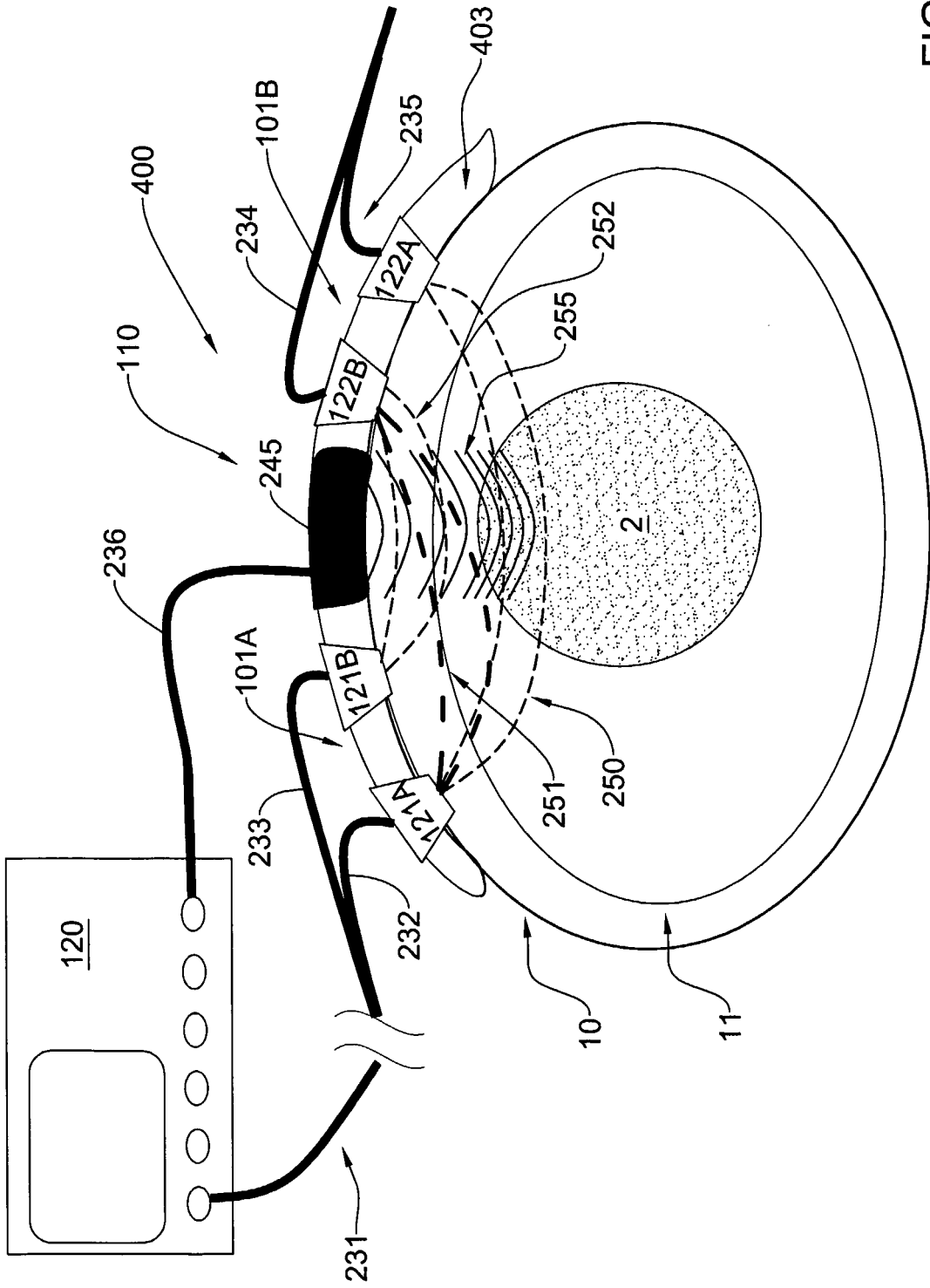


FIG. 4

9/11

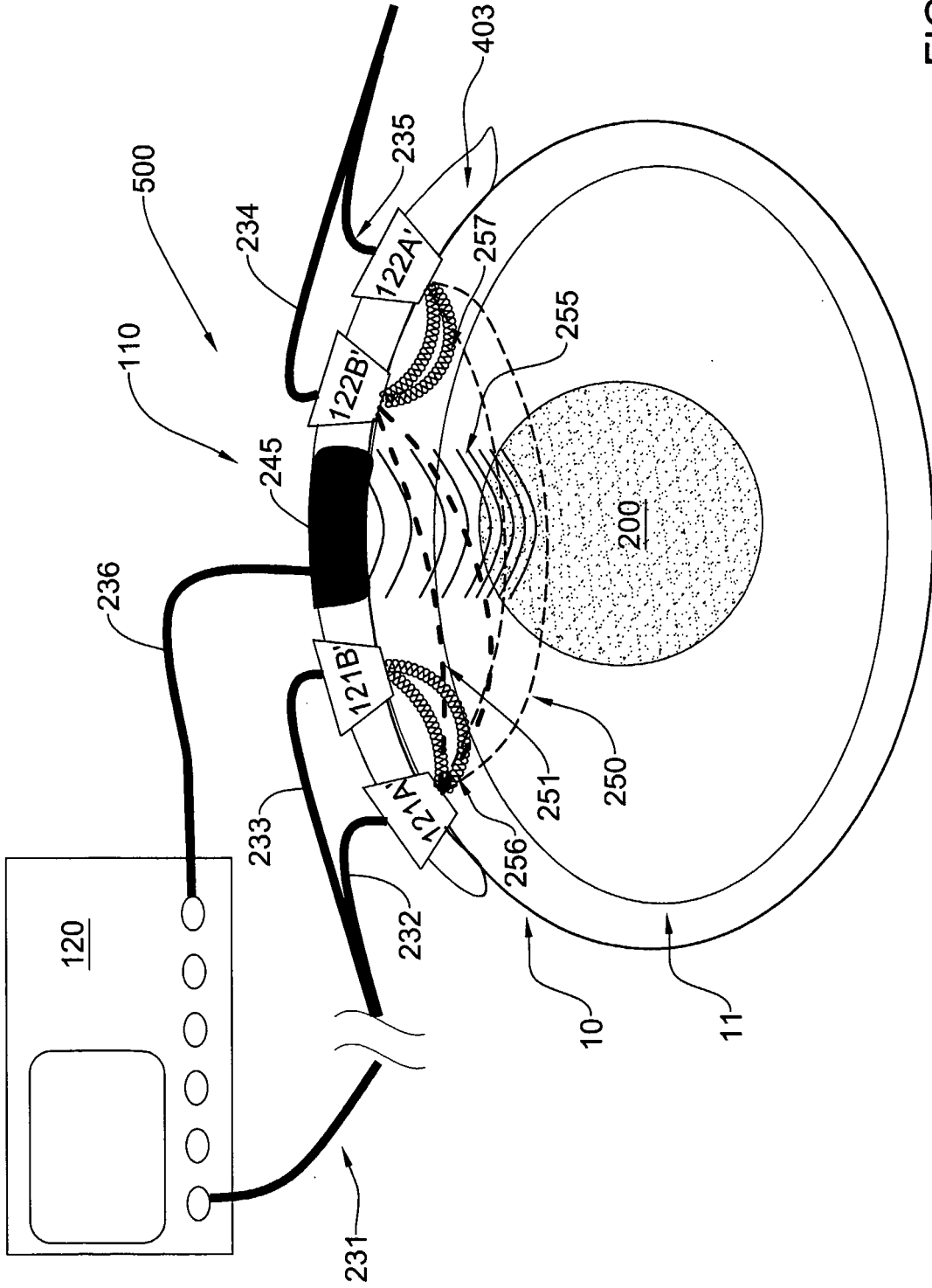


FIG. 5

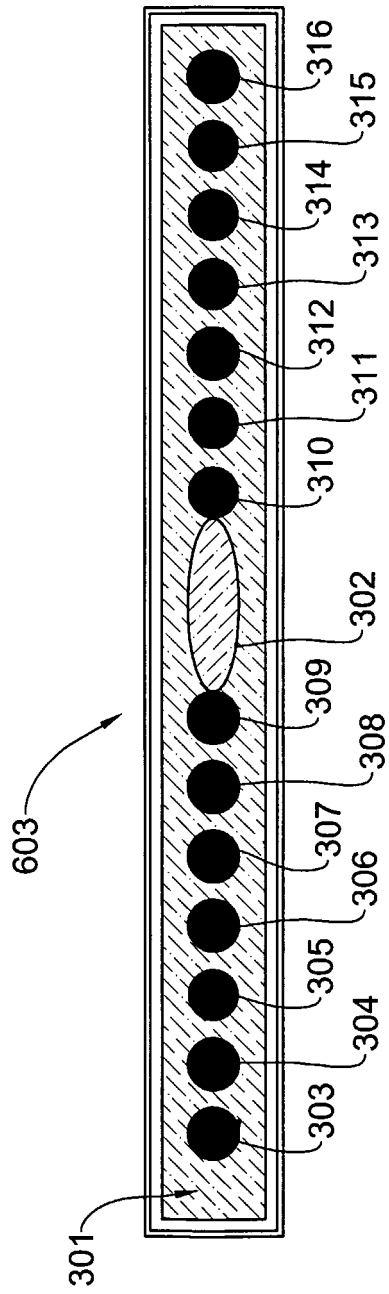


FIG. 6A

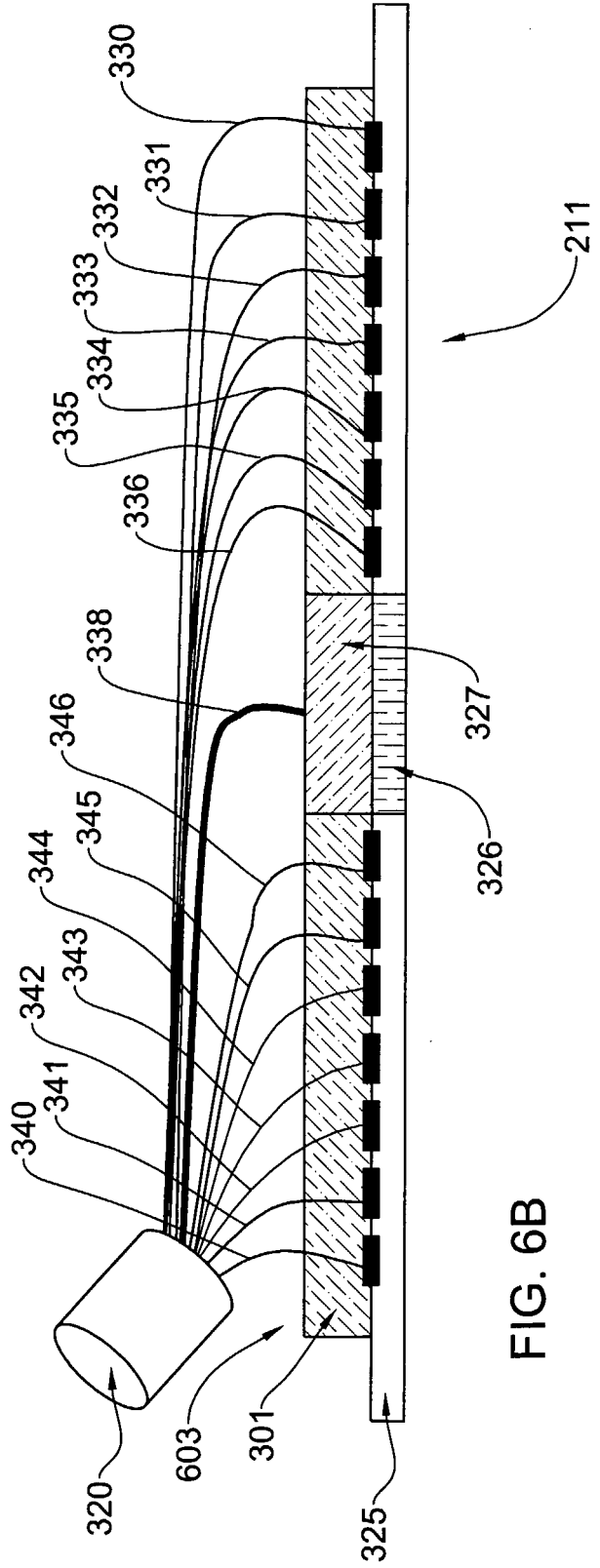


FIG. 6B

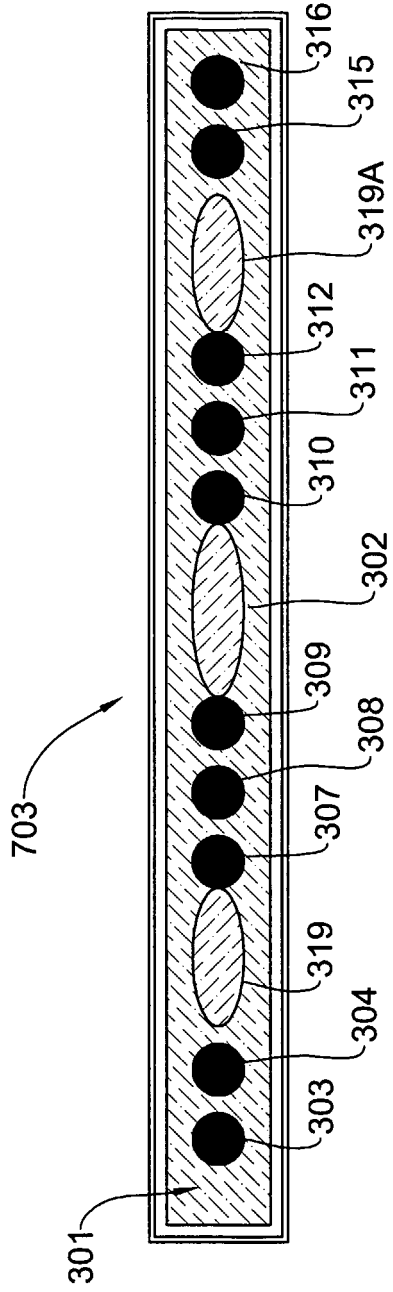


FIG. 7A

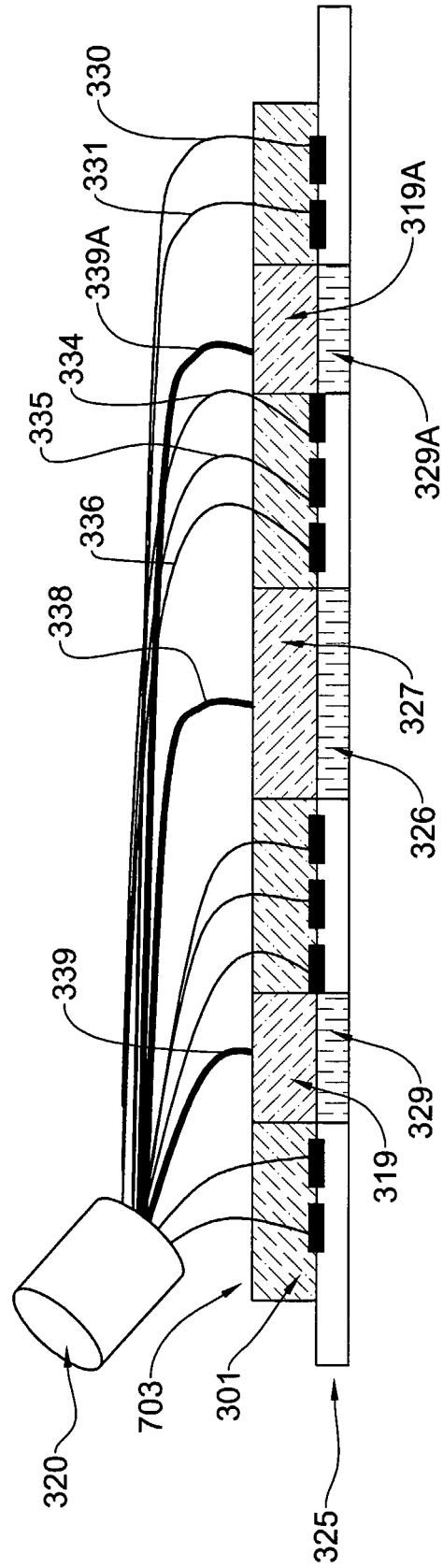


FIG. 7B

专利名称(译)	无创光学监测感兴趣区域		
公开(公告)号	<a href="#">EP1675501A2</a>	公开(公告)日	2006-07-05
申请号	EP2004770506	申请日	2004-09-12
[标]申请(专利权)人(译)	OR-NIM医疗有限公司		
申请(专利权)人(译)	OR-NIM MEDICAL LTD.		
当前申请(专利权)人(译)	OR-NIM MEDICAL LTD.		
[标]发明人	BALBERG MICHAL PERY SCHECHTER REVITAL OLSHANSKY MICHAL		
发明人	BALBERG, MICHAL PERY-SCHECHTER, REVITAL OLSHANSKY, MICHAL		
IPC分类号	A61B5/00 A61B A61B5/024 A61B5/08		
CPC分类号	A61B5/0097 A61B5/02411 A61B5/08 A61B5/14542 A61B5/1464 A61B5/7264 A61B2562/0233 A61B2562/043 G01S15/8968		
优先权	60/502212 2003-09-12 US 60/502210 2003-09-12 US 60/553142 2004-03-16 US 60/581376 2004-06-22 US		
其他公开文献	EP1675501A4 EP1675501B1		
外部链接	<a href="#">Espacenet</a>		

#### 摘要(译)

一种用于非侵入性地监测人体中感兴趣区域的至少一个参数的方法和系统。该系统包括测量单元和控制单元。测量单元包括光学单元，该光学单元具有照明组件（101A）和光检测组件（101B），以产生指示所收集的光的测量数据；声学单元（110），被配置为产生预定超声频率范围的声波。测量单元提供操作条件，使得声波与感兴趣区域内的照明区域重叠并且基本上不在感兴趣区域外重叠。测量数据指示具有超声标记和未标记光部分的散射光，使得能够区分感兴趣区域的光响应和感兴趣区域之外的区域。