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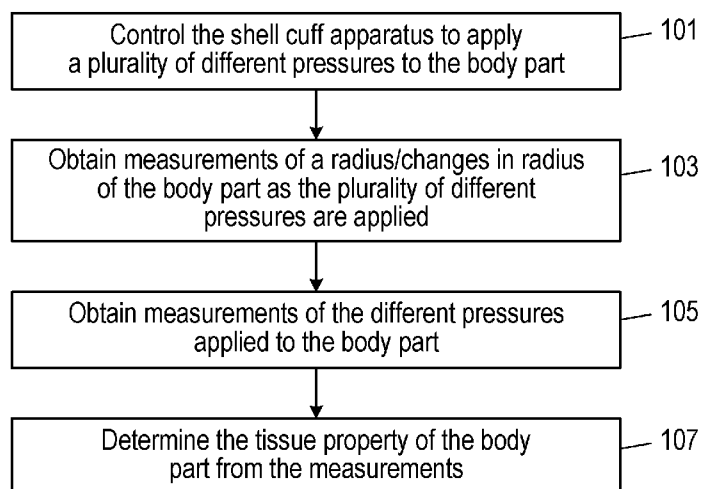


Fig. 6

(57) Abstract: According to an aspect, there is provided a computer-implemented method of determining a tissue property of a body part. The method comprises controlling (101) a shell cuff apparatus to apply a plurality of different pressures to the body part, wherein the shell cuff apparatus at least partially encloses the body part, and wherein the shell cuff apparatus comprises a hard shell portion placed at least partially around the body part and an inflatable cuff surrounding the hard shell portion that is inflated and/or deflated to apply the plurality of different pressures to the body part via the hard shell portion; obtaining (103) radius measurements of a radius of the body part, or changes in the radius of the body part, as the plurality of different pressures are applied to the body part; obtaining (105) pressure measurements of the different pressures applied to the body part; and determining (107) the tissue property of the body part from the radius measurements and the pressure measurements.

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DETERMINING A TISSUE PROPERTY OF A BODY PART

FIELD OF THE INVENTION

This disclosure relates to the use of a shell cuff apparatus to determine a tissue property of a body part, and in particular to a computer-implemented method, a device and a system for determining a tissue property of a body part.

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BACKGROUND OF THE INVENTION

Blood pressure (BP) is an important indicator of health in a person/subject. In the US it is estimated about 30% of the adult population has high blood pressure. Hypertension is a common health problem which has no obvious outward symptoms. Blood pressure generally rises with aging and the risk of becoming hypertensive in later life is considerable. Persistent hypertension is one of the key risk factors for strokes, heart failure and increased mortality. The condition of a subject can be improved by lifestyle changes, healthy dietary choices and medication. Particularly for high-risk patients, continuous 24-hour blood pressure monitoring is very important by means of systems which do not impede ordinary daily life activities. Continuous monitoring of blood pressure can also be useful for patients in a healthcare environment, such as a hospital e.g. in the operating room (OR) or the Intensive Care Unit (ICU).

In some cases absolute measurements of blood pressure can be obtained, and in other cases relative measurements of blood pressure can be obtained, for example a measurement of a change in blood pressure. In particular, blood pressure can change over short time windows, e.g. of the order of a few minutes, and these changes can be relevant for further medical examination and possibly medical intervention.

There are a number of different techniques available for measuring blood pressure, and/or changes in blood pressure. Some of these techniques measure blood pressure itself, while other techniques measure other physiological characteristics of the subject and use these as surrogates for blood pressure, for example by relating changes or values of the physiological characteristic to changes or values of blood pressure. Some of the techniques for directly measuring blood pressure require invasive access to the arteries of the subject, for example via a catheter, or the use of bulky/inconvenient equipment such as inflatable cuffs. However, some of the physiological characteristics used as surrogates for blood pressure can be measured using simple and/or unobtrusive sensors applied to the body of the subject.

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WO 2014/121945 describes a blood pressure measuring system comprising a kinking-proof shell. The blood pressure measurement system is configured to surround a patient's body part and

has pressurization means for applying pressure to the body part and a kinking-proof shell that is located between the pressurization means and the body part. A pressure transducer or pressure sensor pad can be provided on an inner surface of the shell. This arrangement is referred to as a “shell cuff” or “shell cuff apparatus”, and the arrangement essentially prevents damping of the pulsatile signals by components of the device and/or by air outside of the device, which translates into a high signal quality and enables blood pressure to be measured with similar accuracy to invasive techniques that use a catheter to directly access the blood. This is achieved by the hydraulically optimised contact of the pressure transducer or pressure sensor pad with the body part that is relevant for the measurement.

Fig. 1 is a simplified diagram showing a shell cuff apparatus 2 placed at least partially around a body part 4, in this case an arm. The shell cuff apparatus 2 includes a hard shell encased by an inflatable cuff, with a sensor pad 6 arranged on the inside of the hard shell to measure pulse waves 8 passing through the blood vessel 10 (artery) in the body part 4. The pulse wave 8 causes a pressure wave in the tissue in the body part 4 and this is measured by the sensor pad 6.

In more detail, a pulse wave 8 in the brachial artery 10 causes a change in the skin movement. This change in skin movement causes a compression of a liquid in the sensor pad 6. The liquid in the sensor pad 6 can be, for example, a silicone oil such as AK10. The sensor pad 6 is connected via a flexible tube filled with a similar liquid as in the sensor pad 6 to a pressure sensor (not shown in Fig. 1) which measures the pressure signal. In the current application the liquid in the pressure sensor is, for compatibility reasons, a different liquid to the liquid in the sensor pad 6. For example the liquid in the pressure sensor may be glycerol.

The liquid-filled sensor pad 6 is enclosed by a hard shell in the shell cuff apparatus 2 to provide stiffness and cause a sufficient signal to be measured by the pressure sensor. Similar to a standard blood pressure cuff there is an actuator (e.g. an air pump) that can fill a cuff with air, to compress the hard shell and thus the arm tissue, so the brachial artery can be closed.

The blood pressure measurements by the shell cuff apparatus 2 need to be calibrated against blood pressure measurements obtained using an invasive technique (with these measurements being obtained at the same time). This calibration can be done for a large set of patients with different levels of tissue stiffness as the tissue stiffness will influence the transfer of the pressure signal from the artery 10 to the outside of the body part 4. As the shell cuff apparatus 2 includes a hard shell, shell cuffs with different sizes are provided to accommodate patients (subjects) with different arm diameters.

Currently, the effect of individual variation of tissue properties cannot be taken into account in the measurements obtained using a shell cuff apparatus. Instead, a variation of tissue properties is only taken into account according to on average tissue properties expected for a particular size of shell cuff. There can be, however, for a single size of shell cuff, a significant difference in tissue properties between subjects. For instance, subjects with large obese arms will have different tissue properties compared to subjects with muscular arms of similar size.

Thus, it is desirable to be able to determine a tissue property of the body part of the subject, so that this can, for example, be taken into account in any estimates or measurements of physiological characteristics (e.g. blood pressure) that are derived from the measurements obtained using the shell cuff apparatus.

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SUMMARY OF THE INVENTION

The techniques described herein use a sensor to obtain measurements of a radius of the body part, or changes in the radius of the body part, as a number of different pressures are applied to the body part using the shell cuff apparatus. A tissue property of the body part can be determined from a function or relationship between changes in the radius of the body part and the pressures applied by the shell cuff apparatus.

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According to a first aspect, there is provided a computer-implemented method of determining a tissue property of a body part. The method comprises controlling a shell cuff apparatus to apply a plurality of different pressures to the body part, wherein the shell cuff apparatus at least partially encloses the body part, and wherein the shell cuff apparatus comprises a hard shell portion placed at least partially around the body part and an inflatable cuff surrounding the hard shell portion that is inflated and/or deflated to apply the plurality of different pressures to the body part via the hard shell portion; obtaining radius measurements of a radius of the body part, or changes in the radius of the body part, as the plurality of different pressures are applied to the body part; obtaining pressure measurements of the different pressures applied to the body part; and determining the tissue property of the body part from the radius measurements and the pressure measurements.

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According to a second aspect, there is provided a computer-implemented method of determining a measurement of a physiological characteristic of a subject. The method comprises determining a tissue property of a body part of the subject according to the first aspect or any embodiment thereof, using the determined tissue property to calibrate pressure measurements by a sensor pad positioned inside the hard shell portion; and determining a measurement of the physiological characteristic using the calibrated pressure measurements.

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According to a third aspect, there is provided a device configured to determine a tissue property of a body part. The device is configured to, and/or the device comprises a processing unit configured to: control a shell cuff apparatus to apply a plurality of different pressures to the body part (which can comprise the processing unit outputting a control signal to the shell cuff apparatus), wherein the shell cuff apparatus at least partially encloses the body part, and wherein the shell cuff apparatus comprises a hard shell portion placed around at least a portion of the body part and an inflatable cuff surrounding the hard shell portion that is inflated and/or deflated to apply the plurality of different pressures to the body part via the hard shell portion; obtain (e.g. receive) radius measurements of a radius of the body part, or changes in the radius of the body part, as the plurality of different pressures are applied to the body part; obtain (e.g. receive) pressure measurements of the different pressures applied to

the body part; and determine the tissue property of the body part from the radius measurements and the pressure measurements.

According to a fourth aspect, there is provided a device configured to determine a measurement of a physiological characteristic of a subject. The device is configured according to the fourth aspect, or any embodiment thereof, and is also configured to use the determined tissue property to calibrate pressure measurements by a sensor pad positioned inside the hard shell portion; and determine a measurement of the physiological characteristic using the calibrated pressure measurements.

According to a fifth aspect, there is provided a system for determining a tissue property of a body part. The system comprises a device according to the third or fourth aspects; a shell cuff apparatus configured to at least partially enclose a body part, wherein the shell cuff apparatus comprises a hard shell portion configured to be placed around at least a portion of the body part and an inflatable cuff surrounding the hard shell portion that is configured to be inflated and/or deflated to apply the plurality of different pressures to the body part via the hard shell portion; and one or more sensors for providing (e.g. obtaining) the radius measurements and the pressure measurements.

According to a sixth aspect, there is provided a computer program product comprising a computer readable medium having computer readable code embodied therein, the computer readable code being configured such that, on execution by a computer, or by the processing unit of the device according to the third or fourth aspects, or any embodiment thereof, the computer or device is caused to perform the method according to the first aspect, the second aspect, or any embodiment thereof.

Various embodiments of the device according to the fourth and fifth aspects are also envisaged in which the device is additionally or further configured to operate according to the various embodiments of the methods described herein.

These and other aspects will be apparent from and elucidated with reference to the embodiment(s) described hereinafter.

BRIEF DESCRIPTION OF THE DRAWINGS

Exemplary embodiments will now be described, by way of example only, with reference to the following drawings, in which:

Fig. 1 is a simplified diagram of a shell cuff apparatus around a body part of a subject;

Fig. 2 is a block diagram of a system for determining a tissue property of a body part that comprises a device and a shell cuff apparatus;

Fig. 3 shows an exemplary hard shell portion with a radius sensor in the form of a strain gauge;

Fig. 4 illustrates a cross-section of a hard shell portion around a simplified model of a body part;

Fig. 5 illustrates a sensor pad being pressed into a body part; and

Fig. 6 is a flow chart illustrating a method of determining a tissue property of a body part.

DETAILED DESCRIPTION OF EMBODIMENTS

As noted above, the techniques described herein use a sensor to obtain measurements of a radius of the body part, or changes in the radius of the body part, as a number of different pressures are applied to the body part using the shell cuff apparatus. A tissue property of the body part can be determined from a function or relationship between changes in the radius of the body part and the pressures applied by the shell cuff apparatus. The tissue property can be any of: the elastic modulus, the non-linear E-modulus, the Poisson ratio, ν_{arm} , the fat free mass (FFM), and the adipose fat mass (AFM).

The body part radius or change in body part radius can be measured using a sensor that is referred to herein as a 'radius sensor'. A number of different types of sensors can be used for this purpose. For example, the radius sensor can be or comprise a strain gauge that is either integrated into the hard shell portion of the shell cuff apparatus, or a separate measurement device based on a metal strip, that can be used to measure both the initial body part radius (i.e. before pressure is applied), as well as the change in body part radius as pressure is applied on the body part.

The pressure applied to the body part can be measured directly using a sensor pad associated with the shell cuff apparatus, or indirectly via a measurement of pressure in the inflatable cuff combined with a known friction coefficient between overlapping parts of the hard shell portion, the total length of the hard shell portion and the measured change in body part radius with cuff pressure. The total length of the hard shell portion is a fixed parameter of the hard shell portion, and is length of the shell enclosing the body part (approximately given by $2\pi R$, with R the body part radius) plus the amount of overlap of the opposite sides of the hard shell portion. A mismatch between both methods being too large is an indication that there is an error in the pressure measured by the sensor pad, which will automatically lead to an error in cardiac output or other physiological characteristic derived from the measured pressure. This can especially occur with small adults with stiff arms. With the proposed techniques, the correct pressure on the body part can be determined, and hence the potential error can be avoided or compensated for.

Thus, embodiments of the techniques enable one or more tissue properties to be determined which can be used for individual calibration of measurements by the sensor pad pressure to determine actual cardiac output, or other physiological characteristic. Further, embodiments provide for the detection of potential errors in measurements of cardiac output or other physiological characteristics, especially for small adults with stiff arms, and correct for these errors.

Fig. 2 illustrates an exemplary system 20 for determining a tissue property of a body part. The system 20 comprises a shell cuff apparatus 22 and a device 24. In some embodiments, the system 20 can also be used to determine or measure one or more physiological characteristics of the subject based on the determined tissue property.

The shell cuff apparatus 22 comprises an inflatable cuff 26 and a hard shell portion 28. The hard shell portion 28 is to be placed around at least a portion of a body part, such as an arm or leg, and the inflatable cuff 26 encloses the hard shell portion 28 (i.e. the inflatable cuff 26 is placed around the hard shell portion 28 so that when the inflatable cuff 26 is inflated, it presses the hard shell portion 28 on to the body part). The hard shell portion 28 can be formed from any suitable material, for example a plastic material with an elastic modulus in the range between 0.4 and 5 GPa. Such plastic material can be High Density Polyethylene (HDPE). The length and/or thickness of the hard shell portion 28 can vary with the diameter of the body part that the shell cuff apparatus 22 is to be used on (e.g. the arm or leg). In particular, the larger the diameter of the body part, the larger the length and thickness of the hard shell portion 28. A typical thickness can be in the range between 0.5 and 5 mm, or for example between 1 and 2 mm, with a typical total length can be in the range between 0.25 and 0.7 metres, and for example between 0.3 to 0.5 metres.

The shell cuff apparatus 22 also comprises a pump 30 that is connected to the inflatable cuff 26 (e.g. via a connecting tube 32). The pump 30 is controllable (for example in response to a control signal) to selectively inflate the inflatable cuff 26. The pump 30 may also be able to selectively deflate the inflatable cuff 26, and/or a valve (not shown) may be provided for enabling the inflatable cuff 26 to be selectively deflated.

To enable a tissue property of the body part beneath the inflatable cuff 26 and the hard shell portion 28 to be determined, a radius sensor 34 is provided for measuring the radius of the body part or changes in the radius of the body part as a number of different pressures are applied to the body part by the inflatable cuff 26 and the hard shell portion 28. The radius sensor 34 can take any of a number of different forms, as outlined further below. The radius sensor 34 outputs a measurement signal that comprises measurements of the radius, or changes in the radius, over time.

In some embodiments, for example where the system 20 is to be used for determining a physiological characteristic of the subject that takes into account the determined tissue property, the shell cuff apparatus 22 can further comprise a sensor pad 36 positioned on an inner surface of the hard shell portion 28, i.e. a surface of the hard shell portion 28 that faces the body part when the hard shell portion 28 is placed around the body part. When the inflatable cuff 26 is inflated, the inflatable cuff 26 and the hard shell portion 28 press the sensor pad 36 against the body part to enable the sensor pad 36 to measure the pressure in the body part, for example due to pulse waves passing through the body part. The sensor pad 36 outputs a measurement signal that comprises measurements of the pulse waves or pressure waves over time. The sensor pad 36 may be a liquid-filled pad. The sensor pad 36 can be connected via a flexible tube filled with a similar liquid as in the sensor pad 36 to a pressure sensor outside of the hard shell portion 28 which measures the pressure. The liquid in the sensor pad 36 can be, for example, a silicone oil such as AK10, and the liquid in the pressure sensor can be a different liquid to the liquid in the sensor pad 36, such as glycerol.

In some embodiments, the measurements from the sensor pad 36 can be used to determine the radius of the body part or changes in the radius of the body part, in which case a separate radius sensor 34 is not required.

Measurement signals from the radius sensor 34 and the sensor pad 36 (whichever combination of the radius sensor 34 and the sensor pad 36 are present in the shell cuff apparatus 22) are provided to the device 24 for analysis.

The device 24 operates according to the techniques described herein to determine a tissue property of a body part of a subject. The device 24 is configured to receive the measurement signal from the radius sensor 34 and/or the measurement signal from the sensor pad 36. In some embodiments, the device 24 is configured to control the operation of the pump 30 (for example by sending a control signal to the shell cuff apparatus 22/pump 30), and thereby initiate the inflation of the inflatable cuff 26 at an appropriate time.

The device 24 may be in the form of, or be part of, a computing device, such as a server, desktop computer, laptop, tablet computer, smartphone, smartwatch, etc., or other types of device typically found in clinical environments, such as a patient monitoring device (e.g. a monitoring device located at the bedside of a patient in a clinical environment) that is used to monitor (and optionally display) various physiological characteristics of a subject/patient, including blood pressure, cardiac output (CO), stroke volume (SV), stroke volume variation (SVV), or pulse contour stroke volume (PCSV).

The device 24 includes a processing unit 38 that controls the operation of the device 24 and that can be configured to execute or perform the methods described herein. The processing unit 38 can be implemented in numerous ways, with software and/or hardware, to perform the various functions described herein. The processing unit 38 may comprise one or more microprocessors or digital signal processors (DSPs) that may be programmed using software or computer program code to perform the required functions and/or to control components of the processing unit 38 to effect the required functions. The processing unit 38 may be implemented as a combination of dedicated hardware to perform some functions (e.g. amplifiers, pre-amplifiers, analog-to-digital convertors (ADCs) and/or digital-to-analog convertors (DACs)) and a processor (e.g., one or more programmed microprocessors, controllers, DSPs and associated circuitry) to perform other functions. Examples of components that may be employed in various embodiments of the present disclosure include, but are not limited to, conventional microprocessors, DSPs, application specific integrated circuits (ASICs), field-programmable gate arrays (FPGAs), hardware for implementing a neural network and/or so-called artificial intelligence (AI) hardware accelerators (i.e. a processor(s) or other hardware specifically designed for AI applications that can be used alongside a main processor).

The processing unit 38 is connected to a memory unit 40 that can store data, information and/or signals for use by the processing unit 38 in controlling the operation of the device 24 and/or in executing or performing the methods described herein. For example the memory unit 40 can store any of: radius measurements, pressure measurements, measurement signals from one or both of a radius sensor

34 and a pressure sensor 36, for subsequent use by the processing unit 38. In some implementations the memory unit 40 stores computer-readable code that can be executed by the processing unit 38 so that the processing unit 38 performs one or more functions, including the methods described herein. In particular embodiments, the program code can be in the form of an application for a smartwatch, smartphone, tablet, laptop or computer. The memory unit 40 can comprise any type of non-transitory machine-readable medium, such as cache or system memory including volatile and non-volatile computer memory such as random access memory (RAM), static RAM (SRAM), dynamic RAM (DRAM), read-only memory (ROM), programmable ROM (PROM), erasable PROM (EPROM) and electrically erasable PROM (EEPROM), and the memory unit 40 can be implemented in the form of a memory chip, an optical disk (such as a compact disc (CD), a digital versatile disc (DVD) or a Blu-Ray disc), a hard disk, a tape storage solution, or a solid state device, including a memory stick, a solid state drive (SSD), a memory card, etc.

In some embodiments, the device 24 comprises a user interface 42 that includes one or more components that enables a user of device 24 to input information, data and/or commands into the device 24, and/or enables the device 24 to output information or data to the user of the device 24. Information that can be input via the user interface 42 can include any of: gender, age, height, weight, body mass index (BMI), etc. Information that can be output by the user interface 42 can include any of: information about a tissue property, a value of a tissue property, information about a physiological characteristic of the subject, a physiological characteristic value, etc. The user interface 42 can comprise any suitable input component(s), including but not limited to a keyboard, keypad, one or more buttons, switches or dials, a mouse, a track pad, a touchscreen, a stylus, a camera, a microphone, etc., and/or the user interface 42 can comprise any suitable output component(s), including but not limited to a display screen, one or more lights or light elements, one or more loudspeakers, a vibrating element, etc.

It will be appreciated that a practical implementation of a device 24 and/or shell cuff apparatus 22 may include additional components to those shown in Fig. 2. For example the device 24 may also include a power supply, such as a battery, or components for enabling the device 24 to be connected to a mains power supply. The device 24 may also include interface circuitry for enabling a data connection to and/or data exchange with other devices, including the shell cuff apparatus 22 (for example if the shell cuff apparatus 22 is not directly connected to the device 24), servers, databases and user devices.

In some embodiments, the shell cuff apparatus 22 may comprise one or more further sensors for measuring other properties. For example, in embodiments where the radius sensor 34 is a strain gauge, a temperature sensor can also be provided that can be used to measure the temperature of the environment in which the shell cuff apparatus 22 is being used. The measured temperature can be used to correct the measurement of strain by the strain gauge for changes in temperature. Alternatively, multiple strain gauges can be provided that are coupled in a half or full Wheatstone bridge arrangement to automatically correct for changes in temperature.

In another approach, in implementations where the hard shell portion 28 is tapered to better match the shape of the body part (e.g. arm), multiple radius sensors 34 can be provided on different parts of the hard shell portion 28 to measure the radius or change in radius of those parts.

Fig. 3 shows an exemplary hard shell portion 28 with a radius sensor 34 in the form of a strain gauge. The hard shell portion 28 comprises a sheet of material that is curved or 'rolled up' into a cylindrical shape so that it can be placed around a body part. The leading edges of the material overlap in a region 50 as shown in Fig. 3. As the inflatable cuff 26 is inflated, the hard shell portion 28 is compressed against the body part, the amount of overlap in region 50 increases and the diameter of the cylinder formed by the hard shell portion 28 decreases. A radius sensor 34 in the form of a strain gauge can be attached to the outer surface or inner surface of the hard shell portion 28 to measure the strain caused by deformation of the hard shell portion 28. Alternatively the strain gauge 34 can be integral with the hard shell portion 28. Preferably the hard shell portion 28 is made from a material that has a small coefficient of friction when the material is overlapped. This enables the diameter or size of the hard shell portion 28 change more smoothly as the pressure applied to the outside of the hard shell portion 28 changes.

Fig. 3 also shows an alternative approach which can be used with an existing shell cuff apparatus that does not include a radius sensor 34 to enable the techniques described herein to be used. In this approach, an attachable portion 52 is provided that can be attached to, or placed on, the outer surface or inner surface of a hard shell portion 28, for example at position 54, and that comprises a radius sensor 34 in the form of a strain gauge. Attachment of the attachable portion 52 can be achieved using any suitable attachment means, such as an adhesive, clamp, screw, bolt, etc. Deformation of the hard shell portion 28 when the inflatable cuff 26 is inflated causes corresponding deformation of the attachable portion 52, and the radius sensor 34 measures the strain associated with this deformation. In a particular implementation, the attachable portion 52 is formed from steel that is 0.01mm to 5mm thick. The force (clamping force) with which the attachable portion 52 is attached to the hard shell portion 28 should be small compared to the pressure that can be exerted by the inflatable cuff 26 when it is inflated in order to avoid influencing the pressure in the inflatable cuff 26.

The processing unit 38 or the radius sensor 34 may comprise suitable electronic components or software to enable the measurements of the radius or changes in radius by the radius sensor 34 to be made and/or interpreted. For example, in embodiments where the radius sensor 34 is a strain gauge, the radius sensor 34 or the processing unit 38 can comprise an electrical circuit in the form of a Wheatstone bridge that is used to obtain the measurement from the strain gauge. A low-pass frequency filter may also be provided in the processing unit 38 or the radius sensor 34 for filtering out low frequency noise such as pressure pulses from the pump 30 and/or heart beat pulses. A typical cut-off frequency is preferably lower than 20 Hertz (Hz) and more preferably lower than 0.5 Hz.

As noted above, the radius sensor 34 can take any of a number of different forms, and a strain gauge arranged as shown in Fig. 3 is one possibility. Other types of radius sensor 34 can be a

sensor or sensors for measuring one or more dimensions, or change in dimension, of the hard shell portion 28. Dimensions that could be measured to determine the radius or change in radius include the diameter of the hard shell portion 28, the circumference of the hard shell portion 28, the tangential displacement of the hard shell portion 28 or the amount of overlap of the hard shell portion 28 in region 50. Types of sensors that can be used to measure one or more dimensions include optical sensors (e.g. a camera, imaging unit, or laser-based system, which may or may not require the placement or presence of visual markers on the hard shell portion 28), proximity sensors, pressure (touch) sensors arranged at different positions on the hard shell portion 28 to measure the amount of overlap in region 50, an electrical property sensor (e.g. that measures resistance, conductance, impedance, etc. between two parts of the hard shell portion 28 in the overlap region 50), an accelerometer that measures the movement of the hard shell portion 28 from which the deformation of the hard shell portion 28 can be derived, and a volume sensor that measures a volume of fluid that flows to and/or from the body part beneath the hard shell portion 28 as the inflatable cuff 26 is inflated and deflated (a measured change in volume of the body part is indicative of the change in volume of the hard shell portion 28 surrounding the body part).

In embodiments where a separate radius sensor 34 is not provided and the measurements from the sensor pad 36 are used to determine the radius or changes in radius of the body part, a measurement of the total air (or other fluid) volume in the inflatable cuff 26 and a measurement of the pressure in the inflatable cuff 26 can be combined to give a relationship between the change in radius of the body part and the pressure on the body part. The volume or air (or other fluid) in the inflatable cuff 26 can be measured using a flow meter, with the measurements of air (fluid) flow being integrated over time to give the volume of air (fluid) in the inflatable cuff 26.

An alternative method to determine the body part radius without a radius sensor 34 is in the case of an overlapping hard shell portion 28. It can be shown that a relation for the body part radius, $R(\Delta p_i)$, can be derived as:

$$R(\Delta p_i) = \frac{\frac{1}{2}\mu(1+\varepsilon)L_t}{1-\varepsilon+\mu\pi(1+\varepsilon)} \quad (1)$$

where ε is the efficacy (a ratio between sensor pad pressure and cuff pressure, which is described further below), μ is a friction factor between the overlapping shell parts and L_t is the total length of the hard shell portion 28, which is different for different shell cuff sizes. Thus, to determine the body part radius, the efficacy ε and the friction factor μ are required to be known or measured. Although this approach is simple and does not require a separate radius sensor, it is likely to be less accurate as the friction factor μ may vary between overlapping shell parts, and the model that is used to calculate the body part radius is not fully applicable due to the effect described below with reference to Fig. 5.

As described above, it is useful to determine one or more tissue properties of the subject as the response of a pressure sensor (the sensor pad 36) in the hard shell portion 28 is strongly influenced by the tissue properties of a subject. In principle, it is possible to determine, from the change in radius of

the body part with the applied pressure on the outside of the body part, a non-linear elastic modulus (E-modulus) of the body part. The details of these relations will be discussed further below, but it can be understood that a relatively large decrease in body part radius with increasing pressure is an indication of a subject with different tissue properties compared to a subject that has a smaller decrease in radius for the same changes in pressure.

The diagram in Fig. 4 illustrates a cross-section of a hard shell portion 28 around a simplified model of a body part 60. The body part 60 is assumed to have a circular cross-section and includes a bone 62 at the centre of the body part 60 with radius R_b , an artery 64 and other soft tissue 66. In this model the soft tissue 66 is considered to be uniform (i.e. uniform density and other material properties), but it will be appreciated that more complicated models can be used that includes different tissue sections representing fat, muscle, etc. A radius sensor 34 is provided on the hard shell portion 28, and can be, for example, a strain gauge. A sensor pad 36 is positioned on the inside surface of the hard shell portion 28 and is pressed into the body part 60. The hard shell portion 28 is wrapped around the body part 60 and there is an overlap region 50 where leading edges of the hard shell portion 28 overlap with each other. The amount of overlap (i.e. the length of the shell overlap, L_o) is given by:

$$L_o = L_t - 2\pi R_i(\Delta p_i) \quad (2)$$

where L_t is the total length of the hard shell portion 28 in the direction around the body part 60, $R_i(\Delta p_i)$ is the radius of the body part as a function of pressure applied on the inside of the body part 60 (which is denoted Δp_i). The total length L_t of the hard shell portion 28 is a fixed parameter of the hard shell portion 28, measured from one side of the hard shell portion 28 to the other side of the hard shell portion 28. With reference to Fig. 4, the total length L_t is the full length of the hard shell portion 28 around the body part 60, including the length of the hard shell portion 28 in the overlap region 50. The pressure applied on the inside of the body part 60 is measured by the sensor pad 36. Although not shown in Fig. 4, an inflatable cuff 26 is arranged around the hard shell portion 28 so that when the inflatable cuff 26 is inflated, pressure (as indicated by the arrows directed in towards the hard shell portion 28) is applied to the hard shell portion 28.

The thickness of the hard shell portion 28, t is given by:

$$t = R_o - R_i \quad (3)$$

where R_o is the distance from the centre of the body part 60 to the outer surface of the hard shell portion 28.

An initial radius of the body part, denoted R_{i0} , can be measured or determined using the radius sensor 34 when no or a very small pressure is being applied to the body part 60 by the inflatable cuff 26.

The radius sensor 34 is used to measure the radius or changes in radius of the body part 60 as the pressure in the inflatable cuff 26 (Δp_o) is varied. The sensor pad 36 is used to measure the pressure applied to the body part 60 (Δp_i) by the hard shell portion 28. The following equation (4) sets out

how one or more tissue properties (namely the elastic modulus, non-linear E-modulus, the Poisson ratio, v_{arm} , the fat free mass (FFM), and the adipose fat mass (AFM)) can be derived from the changes in radius ΔR_i and the pressure applied to the body part 60:

$$\Delta R_i = \frac{-1}{E_{arm}(\Delta p_i)} \frac{R_{i0}(1-v_{arm}^2)(R_{i0}^2-R_b^2)\Delta p_i}{(R_{i0}^2(1+v_{arm})+R_b^2(1-v_{arm}))} \quad (4)$$

5 with

$$E_{arm}(\Delta p_i) = E_0 + k \cdot \Delta p_i \quad (5)$$

where E_0 is the elastic modulus of the tissue 66, k is a parameter describing a non-linear E-modulus of the tissue 66, v_{arm} is the Poisson ratio of the tissue 66 (assuming the tissue 66 to be a uniform material). The Poisson ratio will be of the order of 0.4 – 0.5, with 0.5 for an incompressible isotropic material deformed elastically at small strains.

It will be appreciated that the relationship shown in equation (4) is based on the simplified body part model in Fig. 4, and different relationships can be derived for more complicated models, e.g. where the body part is considered to be made up of different types of tissue such as muscle and fat.

15 In the above approach, the pressure on the body part 60 (Δp_i) is measured using the sensor pad 36. However, in an alternative approach, the pressure on the body part 60 can be determined by measuring the pressure in the inflatable cuff 26 (the “cuff pressure”), which is denoted Δp_o . Due to the properties of the hard shell portion 28, the cuff pressure Δp_o is different (larger) than the pressure Δp_i on the inside of the hard shell portion 28 which is compressing the body part 60.

20 A simple analytical relation has been found between the pressure as measured by the sensor pad 36 and the cuff pressure. The ratio between these pressures is called the efficacy ε . The efficacy ε depends on the friction factor, the total length of the hard shell portion (L_t), and the actual body part radius (which is a function of the applied pressure). The efficacy ε is given by:

$$\varepsilon = \frac{1 - \frac{1}{2}\mu \left(\frac{L_t}{R(\varepsilon \Delta p_o)} - 2\pi \right)}{1 + \frac{1}{2}\mu \left(\frac{L_t}{R(\varepsilon \Delta p_o)} - 2\pi \right)} \quad (6)$$

25 Hence with the measured body part radius and the known friction factor and the total length of the hard shell portion 28, the pressure on the inside of the body part can be determined via the cuff pressure. It has been found that a more compressible body part will give lower efficacy. It is possible to obtain an efficacy as low as approximately 0.4 with a friction coefficient (of the hard shell portion 28 material of less than 0.15.

30 The derivation of the efficacy ε also enables the measurement of the pressure by the sensor pad 36 to be verified. In particular, for smaller sized body parts that produce a relatively small change in the radius on the application of pressure, there is a risk that the pressure measurement by the

sensor pad 36 is not the actual pressure acting on the body part. This is illustrated in Fig. 5. For a small, stiff body part, the sensor pad 36 may not be pushed completely into the body part and the pressure in the sensor pad 36 may be larger than the pressure acting on the rest of the body part. Therefore, the use of the efficacy ϵ enables the measurement of pressure by the sensor pad 36 to be verified based on the measured cuff pressure.

The flow chart in Fig. 6 illustrates a method of determining a tissue property of a body part according to the techniques described herein. The method can be performed by the device 24, and in particular by the processing unit 38. The method can be performed as a result of the device 24 or processing unit 38 executing suitable computer-readable code, that may, for example, be stored in the memory unit 40.

During the measurement stage of the method, represented by steps 101, 103 and 105, a shell cuff apparatus 22 is placed partially or fully around a body part for which one or more tissue properties is to be determined. The shell cuff apparatus 22 comprises a hard shell portion 28 that is placed partially or fully around the body part, and there is an inflatable cuff 26 surrounding the hard shell portion 28. The inflatable cuff 26 can be inflated and/or deflated via a pump 30 to apply multiple different pressures to the body part over time via the hard shell portion 28.

Thus, in a first step, the shell cuff apparatus 22 is controlled to apply a plurality of different pressures to the body part. The shell cuff apparatus 22 can be controlled by the device 24 or processing unit 38, and in particular a pump 30 can be controlled to inflate and/or deflate the inflatable cuff 26. In step 101 the inflatable cuff 26 may be continuously inflated or deflated, or the inflatable cuff 26 may be inflated or deflated in a step-wise manner (i.e. the inflatable cuff 26 is inflated/deflated to a first pressure, the pressure is held for a short period of time, and the inflatable cuff 26 is inflated/deflated to another pressure, and so on). The processing unit 38 can control the application of the plurality of different pressures by the shell cuff apparatus 22 by outputting a suitable control signal to the shell cuff apparatus 22.

In step 103, measurements of a radius of the body part, or measurements of changes in the radius of the body part, are obtained. These measurements are referred to as 'radius measurements'. Radius measurements are obtained as the plurality of different pressures are applied to the body part (i.e. as the inflatable cuff 26 is inflated and/or deflated). That is, multiple radius measurements are obtained, each for a different applied pressure.

In some embodiments, the radius measurements are obtained using one or more radius sensors 34. The radius sensor or sensors 34 can be sensors configured to measure deformation of the hard shell portion 28. Since the hard shell portion 28 is placed around the body part, deformation of the hard shell portion 28 results in a change in radius of the body part beneath the hard shell portion 28. The radius sensor 34 can be configured or arranged such that deformation of the hard shell portion 28 changes one or more electrical properties of the radius sensor 34, such as the resistance, capacitance, inductance. The

radius measurements can be derived from changes in these electrical properties. In some embodiments, the radius sensor 34 is a strain gauge.

In alternative embodiments, the radius measurements can be obtained from measurements of the pressure in the inflatable cuff 26, and measurements of the rate of fluid flow into and out of the inflatable cuff 26. In these embodiments, a cuff pressure sensor can be provided for measuring the pressure in the inflatable cuff 26, and a flow meter can be provided for measuring the rate of fluid flow into and/or out of the inflatable cuff 26. Thus, in these alternative embodiments, the radius measurements are obtained by processing the measurements of the pressure in the inflatable cuff 26 and the measurements of the rate of fluid flow into and out of the inflatable cuff 26.

In step 105, measurements of the different pressures applied to the body part are obtained. These measurements are referred to as 'pressure measurements'. The pressure measurements are obtained at the same time as the radius measurements, so that, for each radius measurement, there is a corresponding measurement of the pressure being applied to the body part at that time. This enables the change in radius of the body part to be observed for changes in the applied pressure.

In some embodiments, the pressure measurements are obtained using a sensor pad 36 positioned on the inside of the hard shell portion 28. In alternative embodiments, a cuff pressure sensor can be used to obtain measurements of the pressure in the inflatable cuff 26 as the inflatable cuff 26 is inflated and/or deflated, and pressure measurements can be determined from a respective cuff pressure sensor measurement and a corresponding radius measurement (i.e. a radius measurement obtained at generally the same time as the cuff pressure sensor measurement). In particular, a pressure measurement can be determined from the cuff pressure sensor measurement, the corresponding radius measurement, a friction factor (friction coefficient) of the hard shell portion 28, and the total length of the hard shell portion 28.

Once the measurements have been obtained in steps 103 and 105, in step 107 a tissue property of the body part can be determined from the radius measurements and the pressure measurements. In some embodiments, step 107 comprises using a function that relates changes in radius of the body part to one or more tissue properties and the pressure applied to the body part. The function can be based on a model of the structure of the body part. For example, the model can assume that the tissue property is uniform throughout the body part, or that certain parts of the body part comprise fat and other comprise muscle, each with different tissue properties.

In some embodiments, step 107 comprises evaluating equation (4) to determine any of the tissue properties E_0 , k , and v_{arm} .

In some embodiments, once the tissue property has been determined, the tissue property can be used to calibrate pressure measurements obtained using the shell cuff apparatus 22 for the purpose of determining a physiological characteristic of the subject. The physiological characteristic can be any of blood pressure, cardiac output, stroke volume (SV), stroke volume variation, or pulse contour stroke volume (PCSV). Those skilled in the art will be aware of techniques for calibrating pressure

measurements according to a determined tissue property. For example, WO 2019/211210 describes a calibration for SV and PCSV based on the tissue properties FFM and AFM, and the techniques described herein can be used to determine those tissue properties and thereby improve the calibration.

Variations to the disclosed embodiments can be understood and effected by those skilled in the art in practicing the principles and techniques described herein, from a study of the drawings, the disclosure and the appended claims. In the claims, the word "comprising" does not exclude other elements or steps, and the indefinite article "a" or "an" does not exclude a plurality. A single processor or other unit may fulfil the functions of several items recited in the claims. The mere fact that certain measures are recited in mutually different dependent claims does not indicate that a combination of these measures cannot be used to advantage. A computer program may be stored or distributed on a suitable medium, such as an optical storage medium or a solid-state medium supplied together with or as part of other hardware, but may also be distributed in other forms, such as via the Internet or other wired or wireless telecommunication systems. Any reference signs in the claims should not be construed as limiting the scope.

CLAIMS:

1. A computer-implemented method of determining a tissue property of a body part, the method comprising:

controlling (101) a shell cuff apparatus to apply a plurality of different pressures to the body part, wherein the shell cuff apparatus at least partially encloses the body part, and wherein the shell
5 cuff apparatus comprises a hard shell portion placed at least partially around the body part and an inflatable cuff surrounding the hard shell portion that is inflated and/or deflated to apply the plurality of different pressures to the body part via the hard shell portion;

obtaining (103) radius measurements of a radius of the body part, or changes in the radius of the body part, as the plurality of different pressures are applied to the body part;

10 obtaining (105) pressure measurements of the different pressures applied to the body part; and

determining (107) the tissue property of the body part from the radius measurements and the pressure measurements.

15 2. A method as claimed in claim 1, wherein the pressure measurements are obtained using a sensor pad positioned on the inside of the hard shell portion.

3. A method as claimed in claim 1, wherein the step of obtaining (105) pressure measurements comprises:

20 using a cuff pressure sensor to obtain measurements of the pressure in the inflatable cuff as the inflatable cuff is inflated and/or deflated;

determining the pressure measurements by, for a first cuff pressure sensor measurement, determining the pressure measurement from the first cuff pressure sensor measurement and a corresponding radius measurement.

25 4. A method as claimed in claim 3, wherein the step of determining pressure measurements of the different pressures applied to the body part comprises:

determining a pressure measurement as a function of a cuff pressure sensor measurement, a corresponding radius measurement, a coefficient of friction of the hard shell portion and a total length
30 of the hard shell portion.

5. A method as claimed in any of claims 1-4, wherein the step of obtaining (103) the radius measurement comprises using one or more radius sensors.

6. A method as claimed in claim 5, wherein the radius sensor is a sensor configured to measure deformation of the hard shell portion.

7. A method as claimed in claim 5 or 6, wherein the radius sensor is configured or arranged such that deformation of the hard shell portion changes one or more electrical properties of the radius sensor.

8. A method as claimed in any of claims 5-7, wherein the radius sensor is a strain gauge.

9. A method as claimed in claim 1-4, wherein the radius measurements are obtained from measurements of the pressure in the inflatable cuff, and measurements of the rate of fluid flow into and out of the inflatable cuff.

10. A method as claimed in claim 2, or claims 3 or 4 when dependent on claim 2, wherein the radius measurements are obtained from measurements of the pressure in the inflatable cuff, pressure measurements obtained using the sensor pad, a friction factor for the hard shell portion, and the length of the hard shell portion.

11. A method as claimed in any of claims 1-10, wherein the step of determining (107) the tissue property of the body part from the radius measurements and the pressure measurements comprises using a function that relates changes in radius of the body part to one or more tissue properties and the pressure applied to the body part.

12. A method as claimed in any of claims 1-11, wherein the step of determining (107) the tissue property comprises evaluating, using the obtained radius measurements, ΔR_i , and the obtained pressure measurements, Δp_i :

$$\Delta R_i = \frac{-1}{E_{arm}(\Delta p_i)} \frac{R_{i0}(1 - v_{arm}^2)(R_{i0}^2 - R_b^2)\Delta p_i}{(R_{i0}^2(1 + v_{arm}) + R_b^2(1 - v_{arm}))}$$

with

$$E_{arm}(\Delta p_i) = E_0 + k \cdot \Delta p_i$$

where E_0 is the elastic modulus of tissue in the body part, k is a parameter describing a non-linear E-modulus of the tissue, v_{arm} is the Poisson ratio of the tissue, R_{i0} is an initial radius of the

body part when no pressure is applied to the body part by the shell cuff apparatus, and R_b is a radius of a bone in the body part.

13. A computer-implemented method of determining a measurement of a physiological characteristic of a subject, the method comprising:

determining a tissue property of a body part of the subject according to the method of any of claims 1-12;

using the determined tissue property to calibrate pressure measurements by a sensor pad positioned inside the hard shell portion; and

determining a measurement of the physiological characteristic using the calibrated pressure measurements.

14. A device (24) for determining a tissue property of a body part, the device (24) comprising a processing unit (38) configured to:

output a control signal to a shell cuff apparatus (22) to control the shell cuff apparatus (22) to apply a plurality of different pressures to the body part, wherein the shell cuff apparatus (22) at least partially encloses the body part, and wherein the shell cuff apparatus (22) comprises a hard shell portion (28) placed around at least a portion of the body part and an inflatable cuff (26) surrounding the hard shell portion (28) that is inflated and/or deflated to apply the plurality of different pressures to the body part via the hard shell portion (28);

obtain radius measurements of a radius of the body part, or changes in the radius of the body part, as the plurality of different pressures are applied to the body part;

obtain pressure measurements of the different pressures applied to the body part; and
determine the tissue property of the body part from the radius measurements and the pressure measurements.

15. A system (20) for determining a tissue property of a body part, the system comprising:
a device as claimed in claim 14;

a shell cuff apparatus (22) configured to at least partially enclose a body part, wherein the shell cuff apparatus (22) comprises a hard shell portion (28) configured to be placed around at least a portion of the body part and an inflatable cuff (26) surrounding the hard shell portion (28) that is configured to be inflated and/or deflated to apply the plurality of different pressures to the body part via the hard shell portion (28); and

one or more sensors (34, 36) for providing the radius measurements and the pressure measurements.

16. A computer program product comprising a computer readable medium having computer readable code embodied therein, the computer readable code being configured such that, on execution by a suitable computer or the processing unit (38) of the device (24) claimed in claim 14, the computer or device (24) is caused to perform the method of any of claims 1-13.

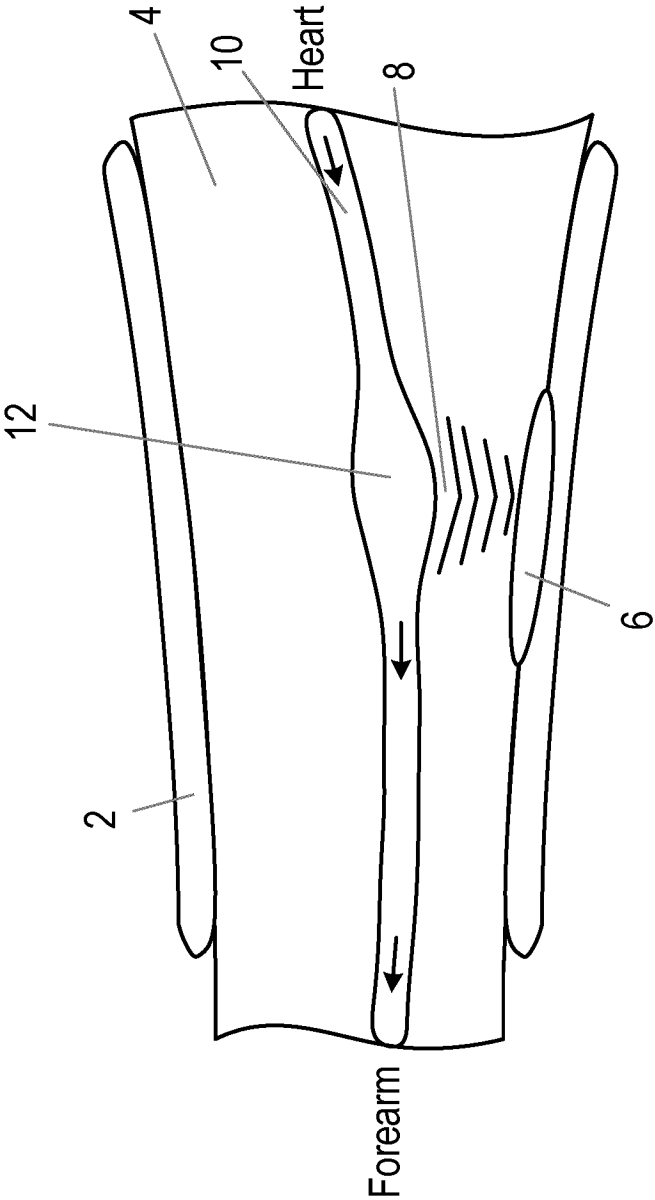


Fig. 1

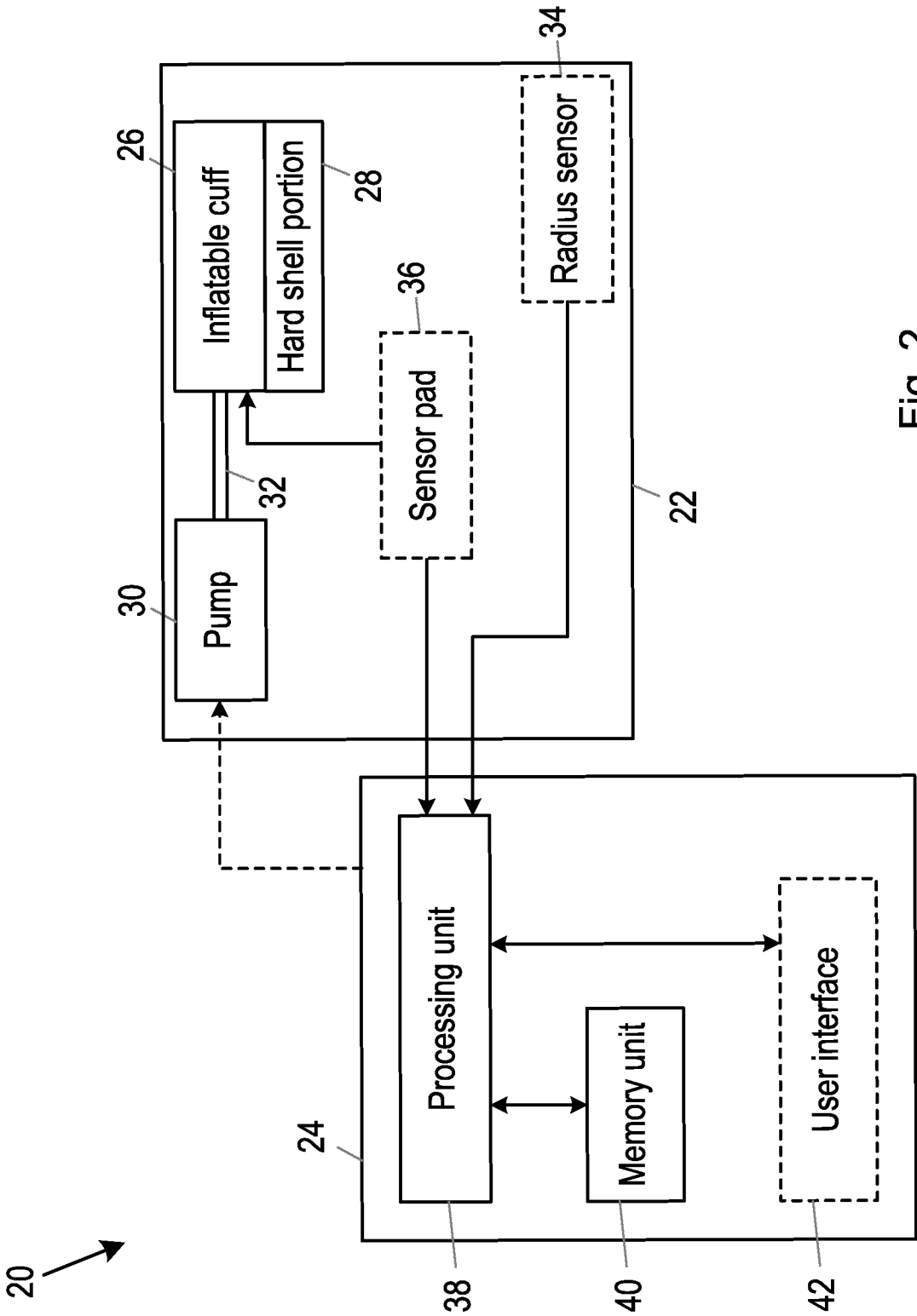


Fig. 2

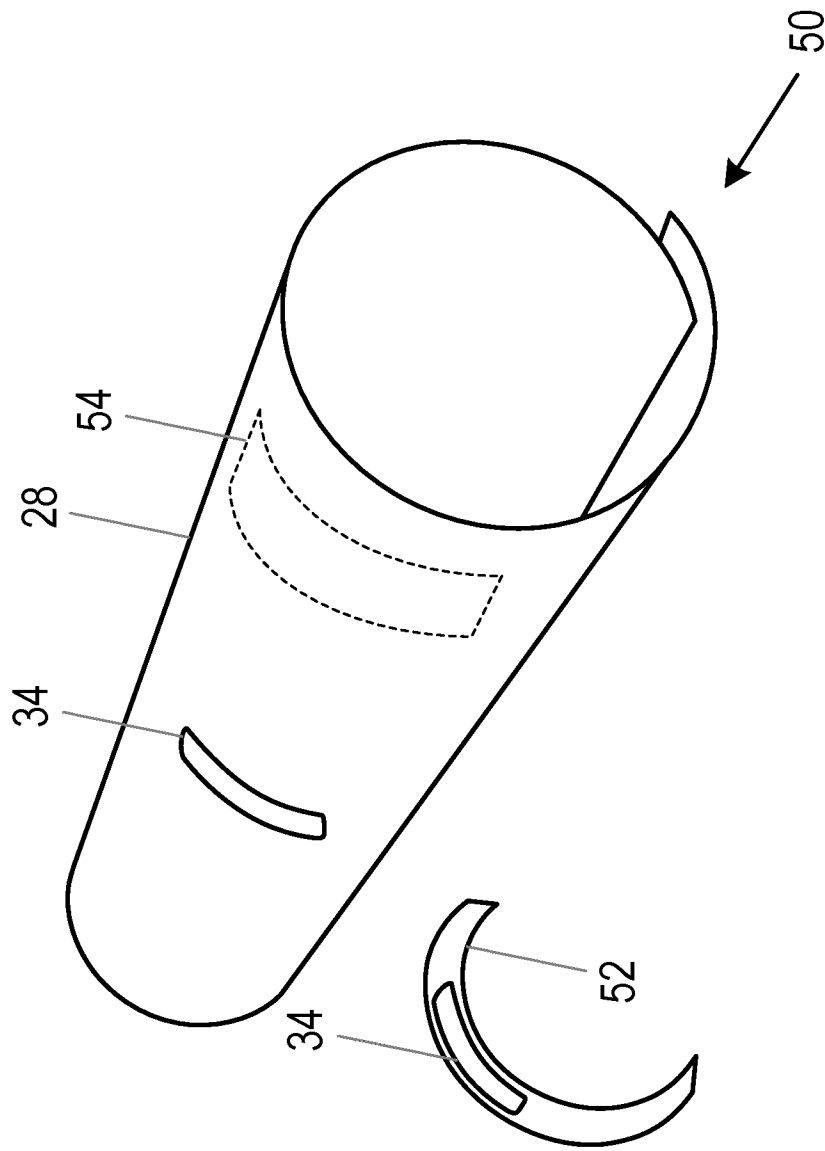


Fig. 3

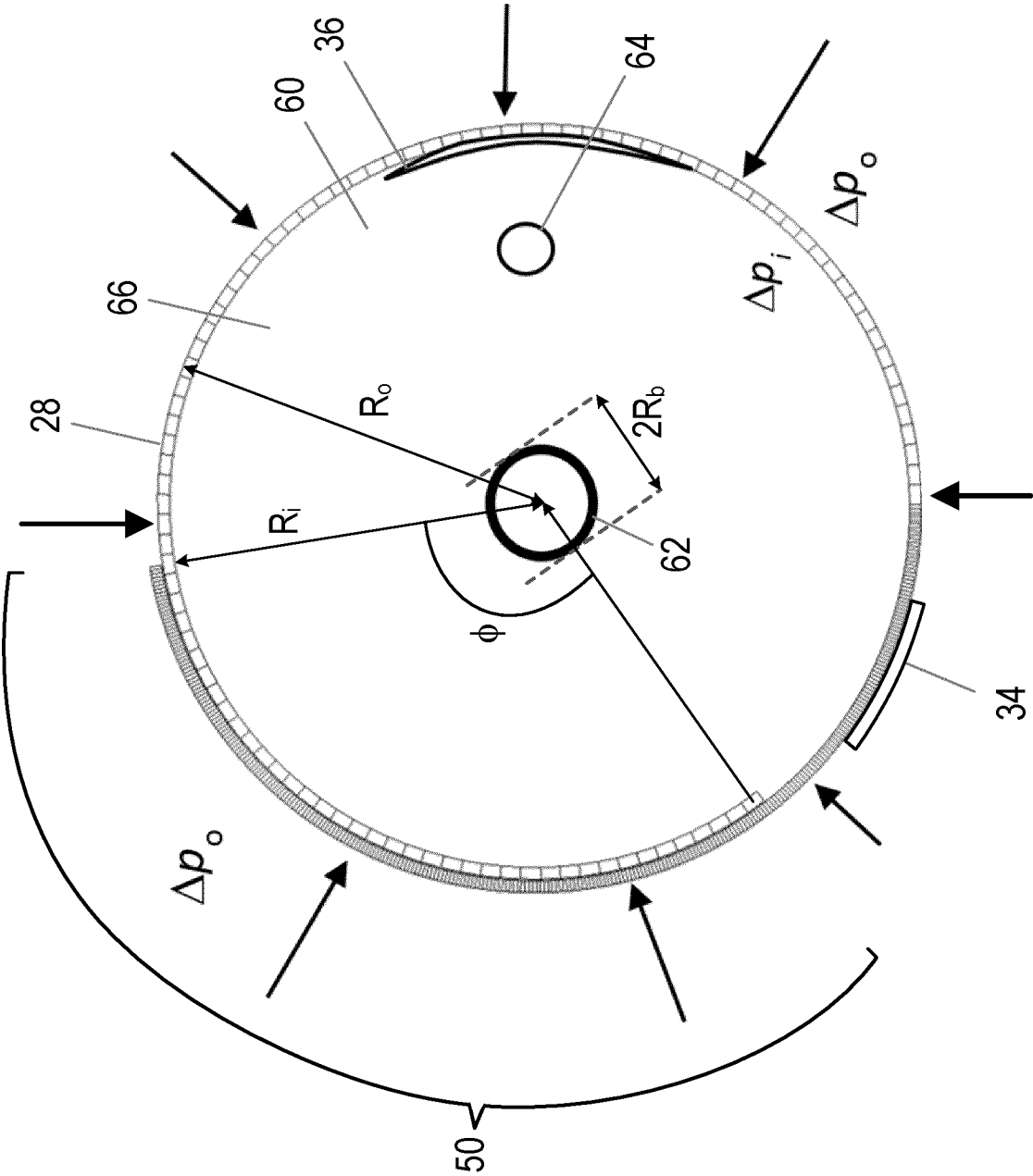


Fig. 4

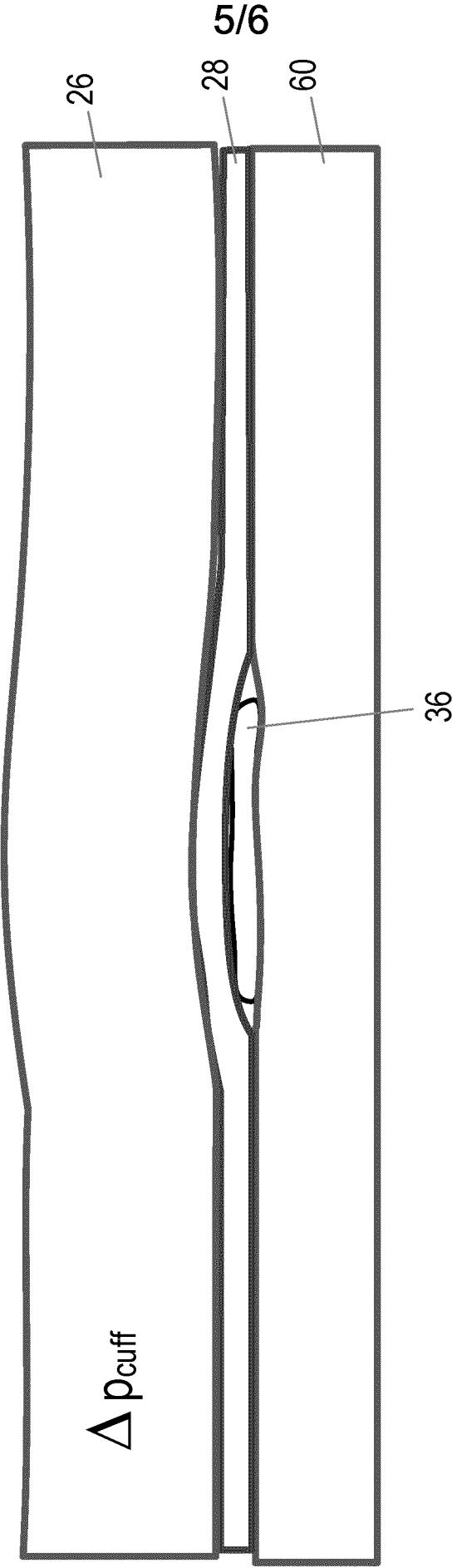


Fig. 5

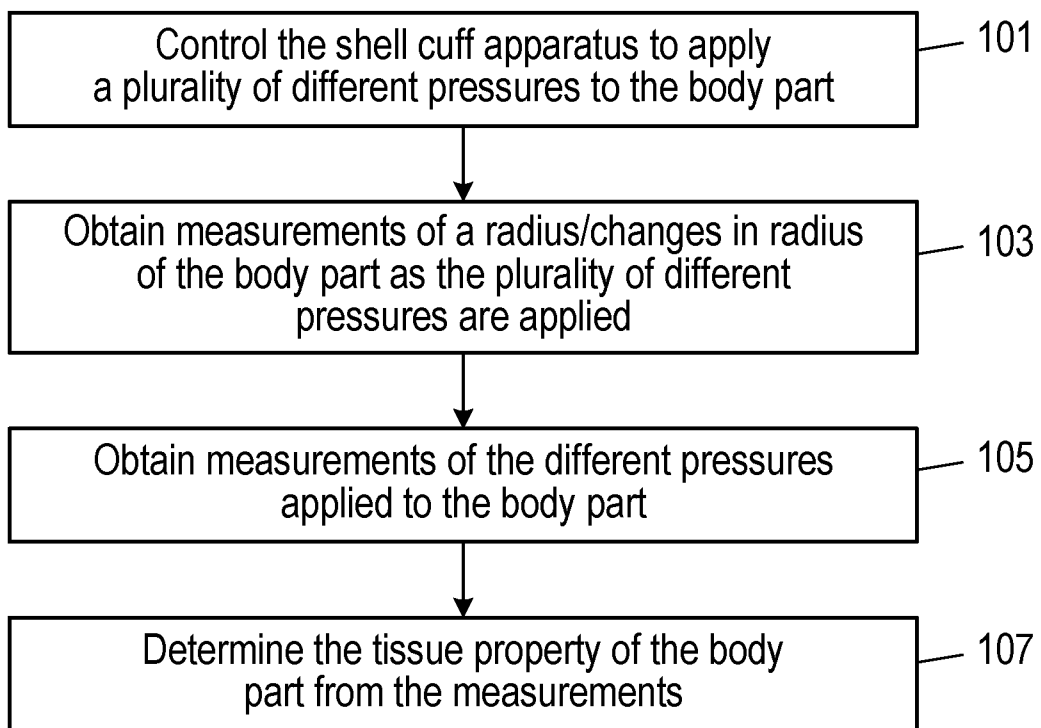


Fig. 6

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2022/075999

A. CLASSIFICATION OF SUBJECT MATTER INV. A61B5/00 A61B5/107 ADD.		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) A61B		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) EPO-Internal, BIOSIS, COMPENDEX, INSPEC, WPI Data		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 2014/121945 A1 (UP MED GMBH [DE]) 14 August 2014 (2014-08-14) cited in the application page 3, line 7 - line 22 page 22, line 28 - page 24, line 18 figures 2a, 2b, 3 -----	1-16
A	US 7 316 652 B2 (BANG & OLUFSEN MEDICOM AS [DK]) 8 January 2008 (2008-01-08) column 2, line 12 - line 46 column 5, line 37 - column 6, line 30 figures 1-3 -----	1-16
A	US 2013/158418 A1 (MIZUKAMI HIROMITSU [JP]) 20 June 2013 (2013-06-20) paragraph [0100] - paragraph [0104] paragraph [0115] figures 1-3, 5, 8 -----	1-16
-/--		
☒ Further documents are listed in the continuation of Box C. ☒ See patent family annex.		
* Special categories of cited documents : "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance;; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance;; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family		
Date of the actual completion of the international search		Date of mailing of the international search report
13 January 2023		23/01/2023
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer Völlinger, Martin

INTERNATIONAL SEARCH REPORT

International application No

PCT/EP2022/075999

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
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Information on patent family members

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