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(54) **CONTINUOUS ULTRASONIC MONITORING**

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(52) **U.S. Cl.**

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(58) **Field of Classification Search**

None

See application file for complete search history.

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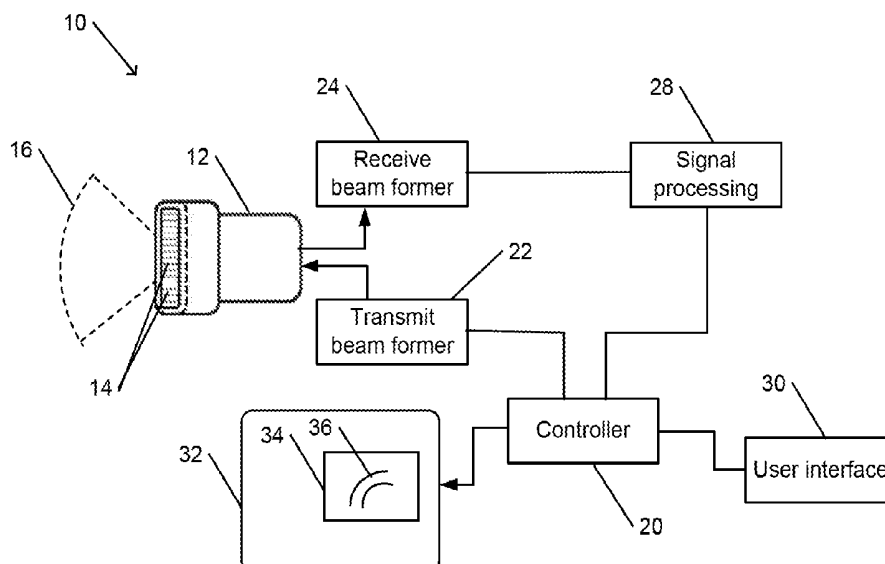
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ABSTRACT

A system for continuous ultrasonic monitoring includes an ultrasound transducer. The ultrasound transducer includes an array of ultrasonic transducer elements. The ultrasonic transducer elements are operable to steer and focus an ultrasound beam to a selectable location and to receive an ultrasound signal from the selectable location. A holder is provided for holding the ultrasound transducer, the holder being attachable to a skin surface of a patient. A controller is configured to repeatedly steer and focus the ultrasound beam to identify a target section of an object inside the body of the patient and to acquire an ultrasound image such that the target section is maintained within the acquired ultrasound image.

20 Claims, 6 Drawing Sheets



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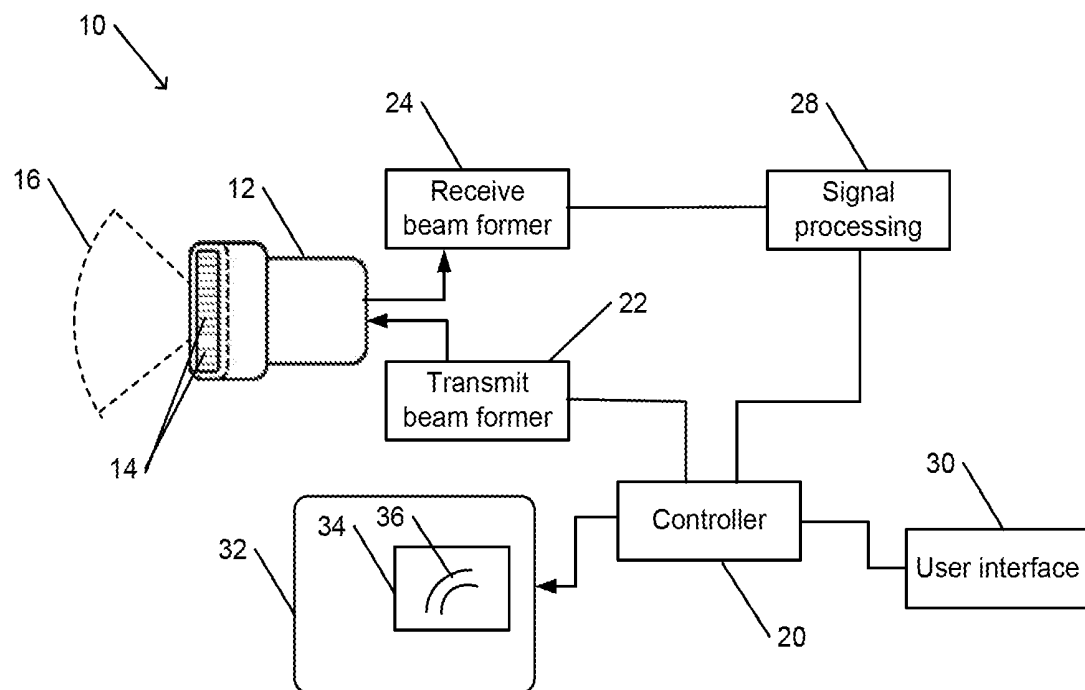


Fig. 1

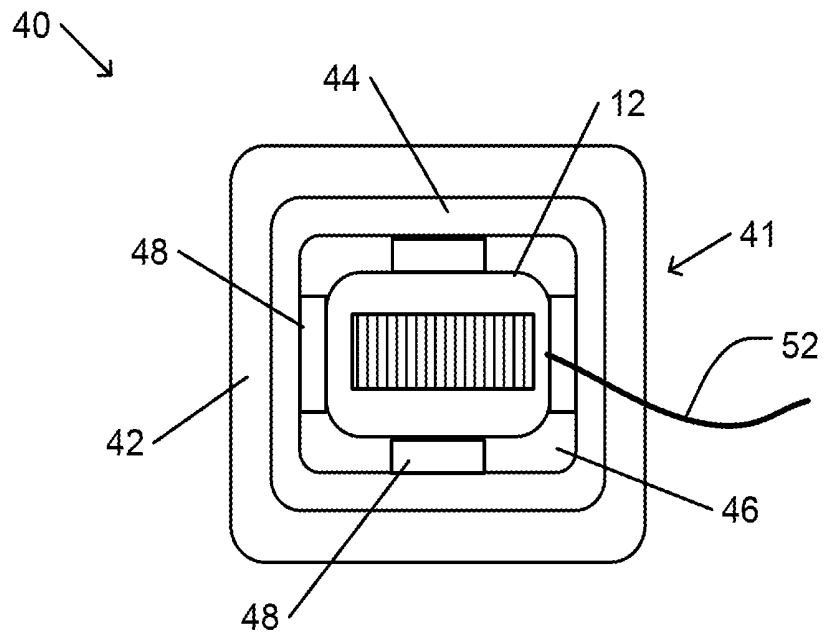


Fig. 2A

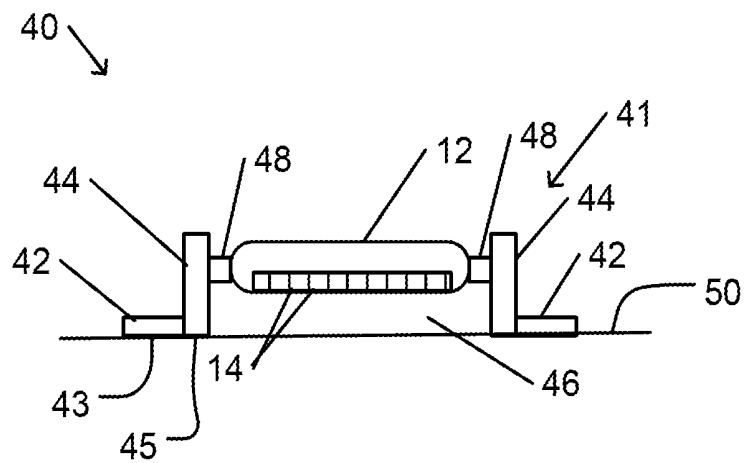


Fig. 2B

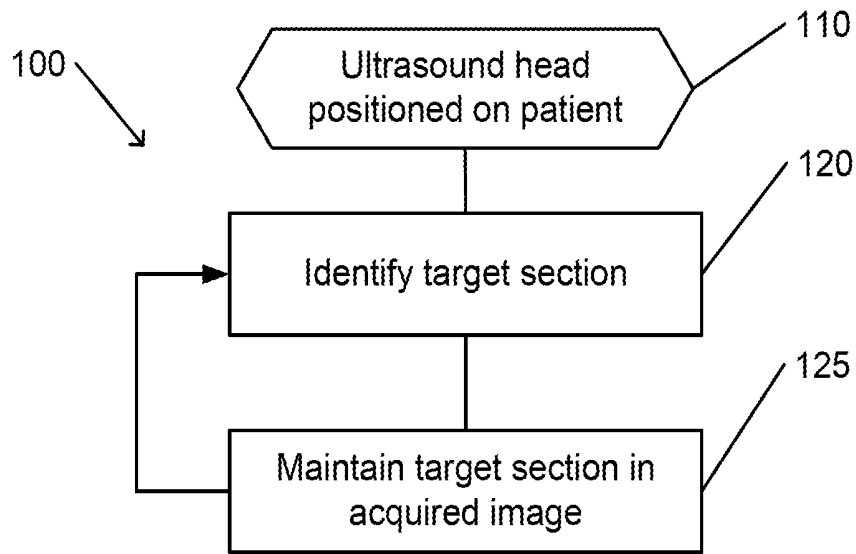


Fig. 3A

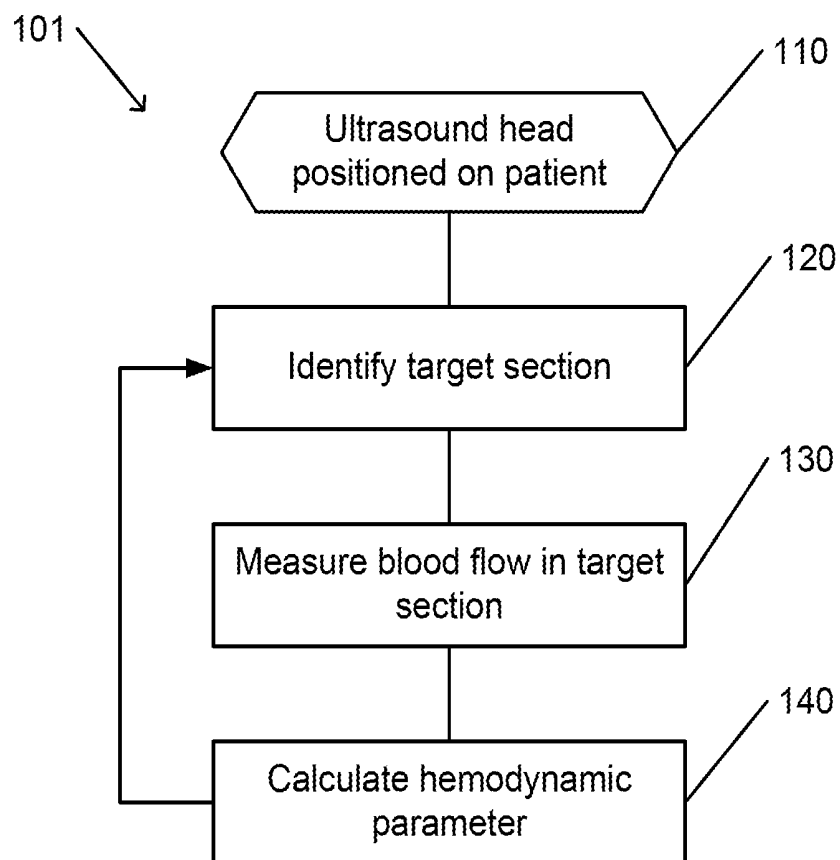


Fig. 3B

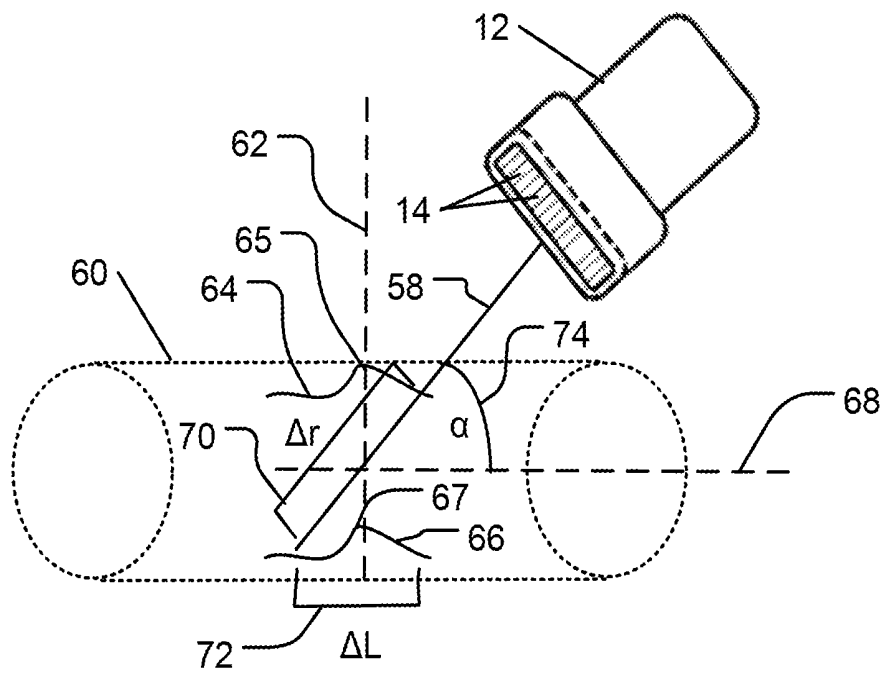


Fig. 4

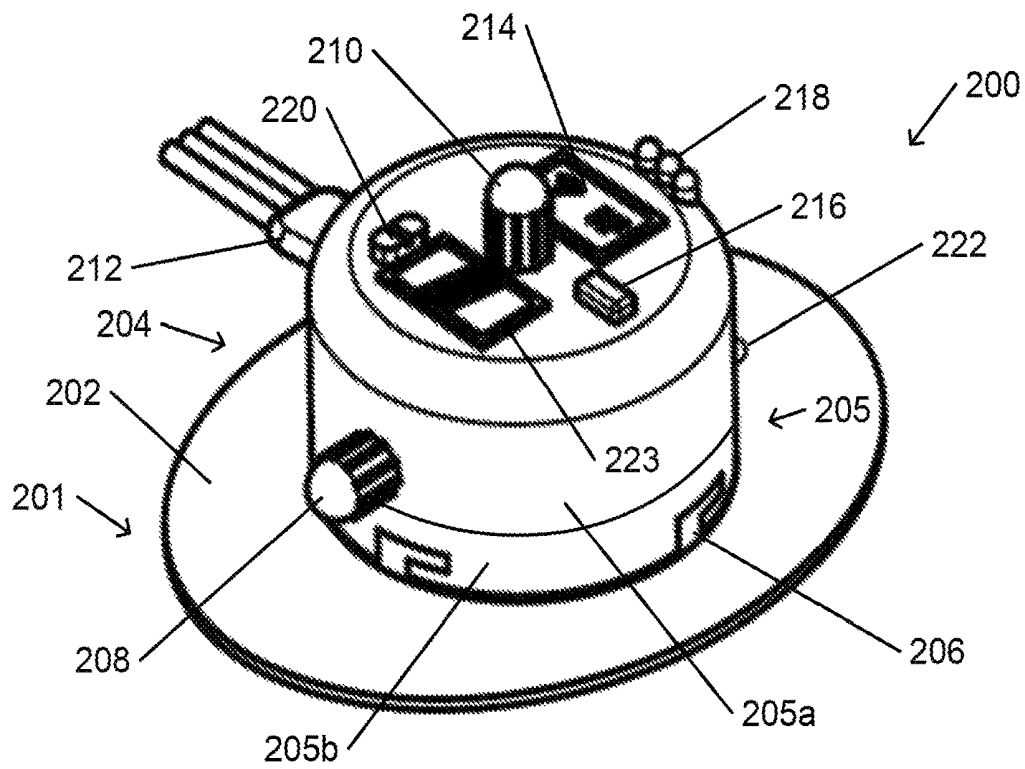


Fig. 5A

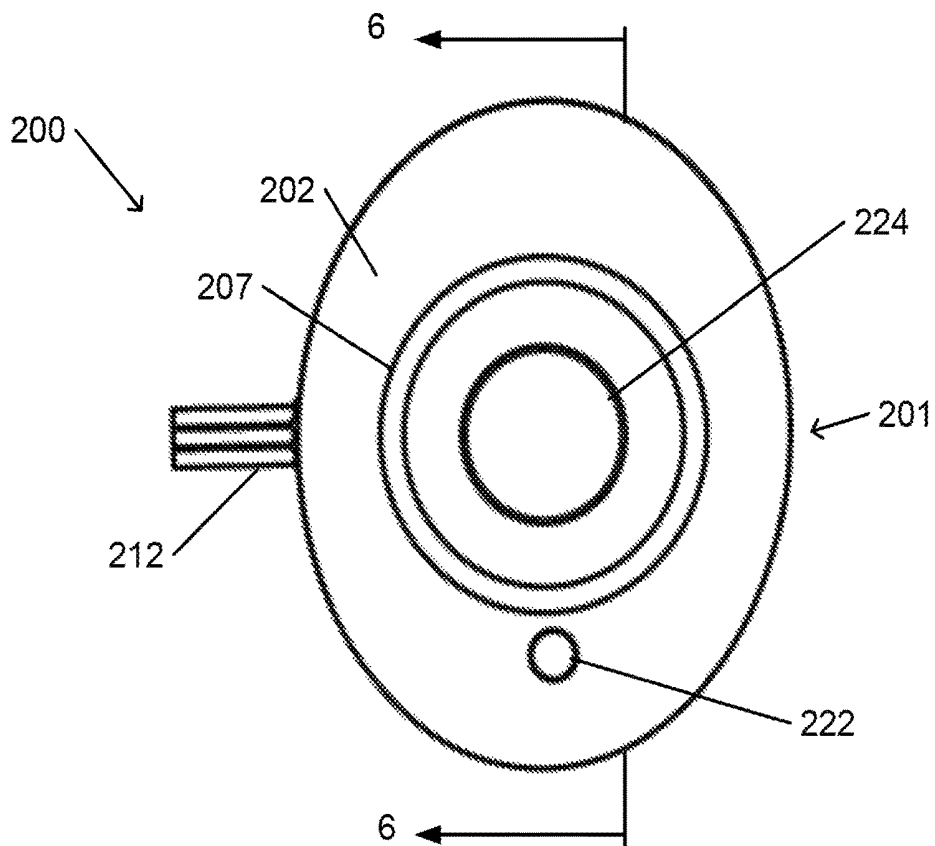


Fig. 5B

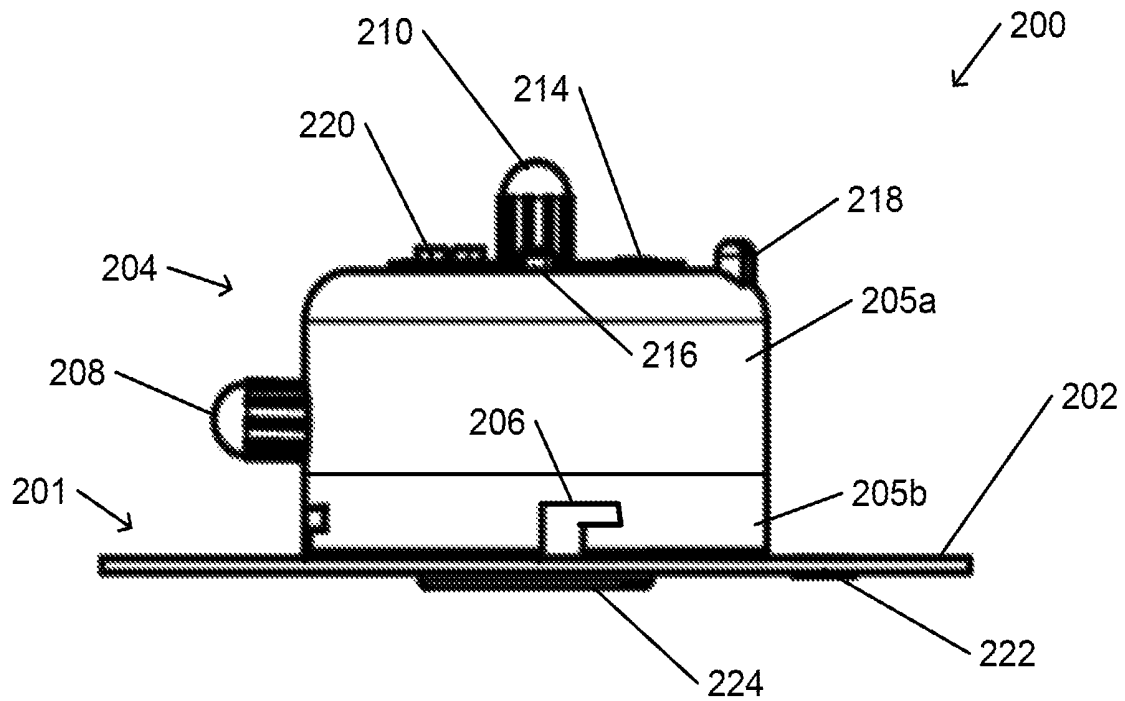


Fig. 5C

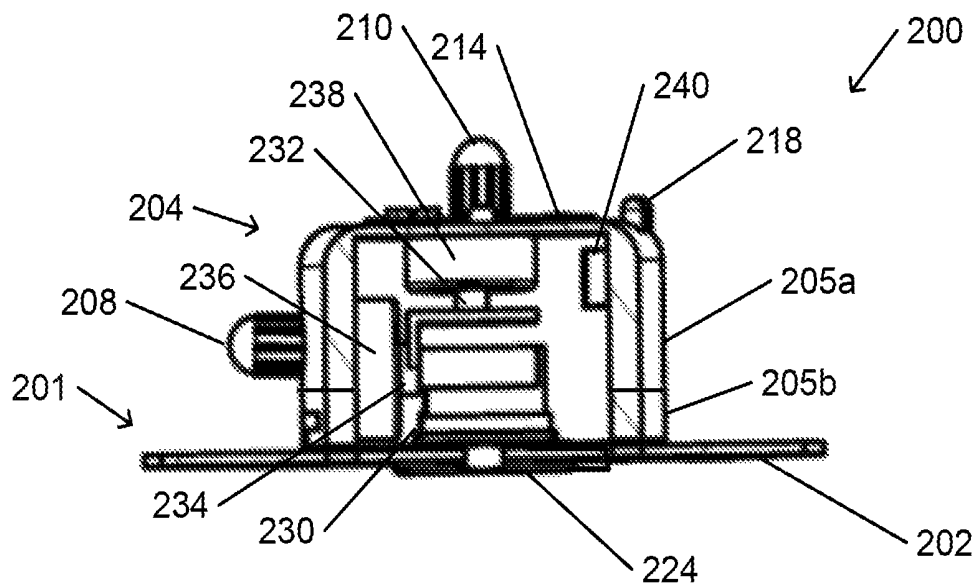


Fig. 6

CONTINUOUS ULTRASONIC MONITORING**CROSS REFERENCE TO RELATED APPLICATIONS**

The present invention is a continuation-in-part of International Patent Application No. PCT/IL2016/050664 with an international filing date of Jun. 22, 2016, which in turn claims the priority benefit of U.S. provisional patent application No. 62/183,278, filed on Jun. 23, 2015, both of which are incorporated in their entirety herein by reference.

FIELD OF THE INVENTION

The present invention relates to ultrasonic monitoring. More particularly, the present invention relates to continuous ultrasonic monitoring.

BACKGROUND OF THE INVENTION

Fluid management is typically required for all intensive care unit (ICU) patients. In the case of critically ill patients, fluid management is an important and difficult issue. Fluid resuscitation is an initial therapeutic intervention with patients after suffering a trauma, or when symptoms of hypotension, evidence of poor organ/tissue perfusion, or other significant hemodynamic challenges are present. Fluid resuscitation refers to the medical practice of replenishing body fluids lost through perspiration, bleeding, fluid shifts, or other pathological processes. The replenishing may include introduction of fluids orally, intravenously, or otherwise.

In some cases, suboptimal administration of fluid resuscitation may lead to an increase in morbidity and mortality in critically ill patients. Therefore, fluid management, in which fluid resuscitation is coordinated with hemodynamic monitoring, is an important component of the treatment of patients requiring fluid resuscitation. Various types of hemodynamic monitors may provide physicians with measurements of various hemodynamic parameters. Such parameters may include stroke volume (the volume of blood that is pumped from the left ventricle during each heartbeat), heart rate, cardiac output, respiration rate, or other relevant hemodynamic parameters. Knowledge of the values of such parameters may assist a physician in determining whether or not fluid resuscitation is necessary, and if it is, what quantity of fluid is to be administered.

The hemodynamic monitors that are most widely used are invasive devices. For example, an invasive device may include one or more catheters or sensors that are inserted into the patient's blood vessels.

Ultrasound imaging is widely used as a tool for medical diagnostics. Ultrasound imaging may provide high-resolution images of internal organs and biological structures deep inside the body, as well as functional information, e.g., with regard to cardiac function and blood flow. A typical ultrasound transducer includes an array of elements that emit and receive ultrasonic waves. When operated in brightness mode (B-mode), an ultrasound system may generate real time two-dimensional images based on the received waves. When operated in Doppler mode, the ultrasound system may provide measurements of the velocities of blood and tissue. Doppler ultrasound may be used to estimate blood velocity by transmitting streams of high-frequency sound waves and analyzing the signals reflected from circulating red blood cells. The local velocity profile is derived from measured changes in the phase of a received signal. Doppler scans may

be used in the diagnosis of such conditions as heart valve defects, congenital heart disease, artery occlusions, and aneurysms.

SUMMARY OF THE INVENTION

There is thus provided, in accordance with an embodiment of the present invention, a system for continuous ultrasonic monitoring, the system including: an ultrasound transducer, the ultrasound transducer including an array of ultrasonic transducer elements, the ultrasonic transducer elements being operable to steer and focus an ultrasound beam to a selectable location and to receive an ultrasound signal from the selectable location; a holder for holding the ultrasound transducer, the holder being attachable to a skin surface of a patient; and a controller that is configured to repeatedly steer and focus the ultrasound beam to identify a target section of an object inside the body of the patient and to acquire an ultrasound image such that the target section is maintained within the acquired ultrasound image.

Furthermore, in accordance with an embodiment of the present invention, the controller is configured to operate the ultrasound transducer in Doppler mode.

Furthermore, in accordance with an embodiment of the present invention, the target section includes a section of a blood vessel, the controller being configured to operate the ultrasound transducer in Doppler mode to acquire a measurement of a flow rate of blood in the target section and to calculate a value of a hemodynamic parameter using the measured flow rate.

Furthermore, in accordance with an embodiment of the present invention, the target section includes a section of the ascending aorta, the aortic arch, or the descending aorta.

Furthermore, in accordance with an embodiment of the present invention, the hemodynamic parameter is selected from a group of hemodynamic parameters consisting of stroke volume, cardiac output, stroke volume variation and systemic vascular resistance.

Furthermore, in accordance with an embodiment of the present invention, the holder includes a patch that includes one or a plurality of adhesive surfaces for attaching the holder to the skin surface.

Furthermore, in accordance with an embodiment of the present invention, the one or a plurality of adhesive surfaces includes an adhesive rim at a periphery of the patch.

Furthermore, in accordance with an embodiment of the present invention, the holder includes a retaining structure for retaining an ultrasound coupling material between the plurality of ultrasonic transducers and the skin surface.

Furthermore, in accordance with an embodiment of the present invention, the ultrasound coupling material includes a gel or a grease.

Furthermore, in accordance with an embodiment of the present invention, the holder includes a holding structure for holding the ultrasound transducer, the holding structure being configured to engage corresponding structure of the ultrasound transducer.

Furthermore, in accordance with an embodiment of the present invention, the controller is configured to repeatedly steer and focus the ultrasound beam to identify the target section by operating the ultrasound transducer in B-mode to acquire an image of a region of the patient that includes the target section.

Furthermore, in accordance with an embodiment of the present invention, the controller is configured to continuously alternate operation of the ultrasound transducer between B-mode and Doppler mode.

Furthermore, in accordance with an embodiment of the present invention, the holder is configured to be attached to the skin surface of the suprasternal notch or the upper abdomen of the patient.

Furthermore, in accordance with an embodiment of the present invention, the controller is configured to calculate an angle between an axis of the target section and a direction of the ultrasound beam.

Furthermore, in accordance with an embodiment of the present invention, the controller is configured to measure a diameter of the target section.

There is further provided, in accordance with an embodiment of the present invention, a method for automatic operation of a continuous ultrasonic monitoring system, the method including: with an ultrasound transducer of the system held to a skin surface of a patient and positioned to steer and focus an ultrasound beam to a location within a body of the patient, the location including a target section, and to receive an ultrasound signal from the location, automatically operating the ultrasound transducer by a controller of the system to repeatedly steer and focus the ultrasound beam to identify the target section and to acquire an ultrasound image such that the target section is maintained within the acquired ultrasound image.

Furthermore, in accordance with an embodiment of the present invention, automatically operating the ultrasound transducer includes operating the ultrasound transducer in Doppler mode.

Furthermore, in accordance with an embodiment of the present invention, automatically operating the ultrasound transducer includes operating the ultrasound transducer alternately in B-mode and Doppler mode.

Furthermore, in accordance with an embodiment of the present invention, automatically operating the ultrasound transducer includes automatically operating the ultrasound transducer to steer and focus a Doppler measurement beam on the target section.

Furthermore, in accordance with an embodiment of the present invention, automatically operating the ultrasound transducer includes modifying a field of view of the ultrasound transducer to maintain the target section within the acquired ultrasound image.

BRIEF DESCRIPTION OF THE DRAWINGS

In order for the present invention, to be better understood and for its practical applications to be appreciated, the following Figures are provided and referenced hereafter. It should be noted that the Figures are given as examples only and in no way limit the scope of the invention. Like components are denoted by like reference numerals.

FIG. 1 shows a block diagram of a system for continuous ultrasonic monitoring, in accordance with an embodiment of the present invention.

FIG. 2A schematically illustrates an ultrasound measurement unit of the system shown in FIG. 1.

FIG. 2B is a schematic sectional view of the ultrasound measurement unit shown in FIG. 2A.

FIG. 3A is a flowchart depicting a method for continuous ultrasonic monitoring, in accordance with an embodiment of the present invention.

FIG. 3B is a flowchart depicting a method for ultrasonic continuous hemodynamic monitoring, in accordance with an embodiment of the present invention.

FIG. 4 illustrates determination of a flow direction from a measurement made with the ultrasound measurement unit shown in FIG. 2A.

FIG. 5A schematically illustrates an ultrasound measurement assembly with a mechanically orientable transducer, in accordance with an embodiment of the present invention.

FIG. 5B schematically illustrates a bottom surface of the ultrasound measurement assembly shown in FIG. 5A.

FIG. 5C schematically illustrates a side view of the ultrasound measurement assembly shown in FIG. 5A.

FIG. 6 schematically illustrates a sectional view of the of the ultrasound measurement assembly shown in FIG. 5C.

DETAILED DESCRIPTION OF THE INVENTION

In the following detailed description, numerous specific details are set forth in order to provide a thorough understanding of the invention. However, it will be understood by those of ordinary skill in the art that the invention may be practiced without these specific details. In other instances, well-known methods, procedures, components, modules, units and/or circuits have not been described in detail so as not to obscure the invention.

Although embodiments of the invention are not limited in this regard, discussions utilizing terms such as, for example, “processing,” “computing,” “calculating,” “determining,” “establishing,” “analyzing,” “checking,” or the like, may refer to operation(s) and/or process(es) of a computer, a computing platform, a computing system, or other electronic computing device, that manipulates and/or transforms data represented as physical (e.g., electronic) quantities within the computer’s registers and/or memories into other data similarly represented as physical quantities within the computer’s registers and/or memories or other information non-transitory storage medium (e.g., a memory) that may store instructions to perform operations and/or processes. Although embodiments of the invention are not limited in this regard, the terms “plurality” and “a plurality” as used herein may include, for example, “multiple” or “two or more”. The terms “plurality” or “a plurality” may be used throughout the specification to describe two or more components, devices, elements, units, parameters, or the like. Unless explicitly stated, the method embodiments described herein are not constrained to a particular order or sequence.

Additionally, some of the described method embodiments or elements thereof can occur or be performed simultaneously, at the same point in time, or concurrently. Unless otherwise indicated, the conjunction “or” as used herein is to be understood as inclusive (any or all of the stated options).

Some embodiments of the invention may include an article such as a computer or processor readable medium, or a computer or processor non-transitory storage medium, such as for example a memory, a disk drive, or a USB flash memory, encoding, including or storing instructions, e.g., computer-executable instructions, which when executed by a processor or controller, carry out methods disclosed herein.

In accordance with an embodiment of the present invention, a continuous ultrasonic monitoring system is provided to continuously monitor an anatomical structure by maintaining an object within the body of a patient within an ultrasound image. For example, the object may include an anatomical structure such as an organ, blood vessel, or tumor, or a foreign object (e.g., an implanted or ingested object, or an object that is introduced into the body as a result of injury). The system includes an ultrasound measurement unit. The ultrasound measurement unit is configured to attach to the body of a patient, to transmit ultrasound

pulses into the patient's body, and to receive ultrasound signals that are reflected from internal structures in the patient's body.

In particular, the ultrasound monitoring system may be utilized to continuously and non-invasively monitor stroke volume, cardiac output (blood volume per unit time, e.g., calculated by multiplying stroke volume by heart rate), stroke volume variations, systemic vascular resistance, or another hemodynamic parameter.

The ultrasound measurement unit includes an ultrasound transducer that includes an array of ultrasonic transducer elements. The system includes a beam former that may introduce phase delays between the signals that are transmitted or received by each of the ultrasonic transducer elements. The beam former may control the phases to do one or more of focus the transmitted ultrasonic beam in the plane of the array to a selectable distance from the ultrasound transducer, receive ultrasound signals from point at a selectable distance from the ultrasound transducer, to aim the ultrasound beam in a selectable direction in the plane of the array, to receive ultrasound from a selectable direction in the plane of the array, or to produce a plane wave. The ultrasonic transducer elements of the ultrasound transducer may be operated in an imaging mode (B-mode) to produce an image, or in Doppler mode to measure a velocity of a measured object or flow.

The ultrasound transducer may be attached to a part of the body of a patient so as to acquire measurements within a region of the patient's body. For example, the region may include the thoracic region or upper abdomen of the patient, or another region of the patient (e.g., the leg, neck, or another part of the body). For example, the ultrasound transducer may be attached to the suprasternal notch, upper abdomen, or elsewhere on the patient. The ultrasound transducer may be attached to the patient's body such that the ultrasonic transducer is aimed toward one or more major blood vessels, or another anatomical structure. For example, an ultrasound transducer that is attached to the suprasternal notch may be aimed toward the ascending aorta, the descending aorta or the aortic arch of the patient. An ultrasound transducer that is attached to the upper abdomen may be aimed toward the descending aorta of the patient. In some cases, the ultrasound transducer may have dimensions that are less than 3 centimeters. For example, a length or width of the ultrasound transducer may be in the range of 1 cm to 1.5 cm.

An adhesive patch may be provided to hold the ultrasound transducer to the patient's body. The adhesive patch may include a probe holder that includes a frame, receptacle, socket, or other structure for holding the ultrasound transducer. For example, one or both of the ultrasound transducer and the probe holder may include clips, latches, magnets, pins, or other structure that is configured to engage corresponding structure (e.g., tabs, loops, ferromagnetic plates, bores, sockets, or other corresponding structure) of the other.

The adhesive patch includes one or more adhesive surfaces. The adhesive surfaces are configured to adhere to the patient's skin when placed or pressed against the skin surface. For example, the adhesive surfaces may form an outer boundary of the adhesive patch, e.g., in the form of a circular, rectangular, or otherwise shaped ring. The size and shape of the adhesive surfaces may be configured to effectively hold the adhesive patch to a particular region of the patient's skin. For example, the size and shape may be configured to hold the adhesive patch to the patient's suprasternal notch or to the patient's upper abdomen.

Alternatively or in addition to an adhesive patch, the probe holder may be attached to the patient's body using suction, straps, clips, or otherwise.

The adhesive patch includes a structure for containing an acoustic coupling material between the ultrasound transducer and the patient's skin. For example, the acoustic coupling material may include an acoustic coupling gel, grease, or other material that may enable efficient propagation of ultrasound signals between the ultrasonic transducers and the skin surface. The structure for holding the acoustic coupling material may include walls to form a bath or chamber that may be fully or partially filled with the acoustic coupling material.

The system may be configured to operate the ultrasonic transducer elements to operate concurrently or alternately in B-mode and in Doppler mode. For example, Doppler and B-mode and measurements may be interleaved with instantaneous switching between Doppler and B-mode pulse sequences. During operation in B-mode, an ultrasound B-mode image of the interior of the patient may be acquired. The image may be expected to include an image of one or more sections of an internal object such as the patient's aorta. For example, when the ultrasound transducer is attached to the suprasternal notch, the image may include part of the ascending aorta or the aortic arch. When the ultrasound transducer is attached to the upper abdomen, the image may include part of the descending aorta.

One or more image processing techniques may be applied to the acquired image to automatically detect and identify one or more target sections. For example, automatic detection of the target section may be based on one or more of an expected size of the target section, a characteristic shape of the target section, a position of the target section relative to other identifiable anatomic features, or other detected or measured features. Various ultrasound imaging and display modes may be utilized, such as B-mode, motion mode (M-mode), color Doppler, spectral Doppler, or other modes.

For example, detection may be based on performing a template matching procedure. An acquired image may be compared to a bank of typical ultrasound images of the region that includes the target section. Correspondence of the acquired image with each image in the bank may be assessed by comparing the image correlation scores of the acquired image with different images in the image bank. Other techniques may be applied.

Upon identification in the image of the target section, one or more parameters of the target section may be derived from the image. For example, a diameter of the cross section of a target section of the aorta may be automatically measured from identified boundaries of the aorta. In some cases, at least an approximate orientation of the identified target section relative to the ultrasound transducer may be automatically determined, e.g., by comparison with images neighboring or other identified anatomical features, or by other characteristics (e.g., from analysis of blood flow within a target section of the aorta).

Continued operation in B-mode may be controlled such that the target section remains in subsequent acquired images. For example, an orientation of an imaged sector may be adjusted such that the target section remains at an approximately constant position within the acquired images.

When the ultrasonic transducer is operated in Doppler mode, the Doppler measurement ultrasound beam may be steered toward, focused on, or both the identified target section. For example, the ultrasound transducer may be operated to obtain a measurement of a velocity of the blood in an identified target section of a blood vessel such as the

aorta. By alternating between operation of the ultrasound transducer in B-mode and operation in Doppler mode, the ultrasonic beam during operation in Doppler mode may be kept aimed at and focused on the target section. For example, the period of alternation may be sufficiently short such that typical anticipated body movements (e.g., breathing, voluntary movement of a limb, head, or other body part, or other relatively slow movements) do not significantly change the position of the ultrasound transducer relative to the target section. Alternatively or in addition, a sensor measurement may be used for gating the ultrasonic measurements in accordance with detected movements (e.g., heartbeat, breathing, or other cyclic motions).

For example, when the ultrasound transducer is operated in Doppler mode, ultrasound beam may be scanned over the identified target section. Thus, the velocity of the blood flow at different points on a cross section of the target section of the aorta may be measured. One or more of the measured blood flow velocity profile, measured dimensions (e.g., diameter or other dimension) of the target section, a measured heartbeat rate (e.g., measured using the ultrasound transducer or measured otherwise), or other measured quantities may be utilized to calculate a stroke volume, or one or more other hemodynamic parameters. In some cases, e.g., when the ultrasound transducer is placed at the suprasternal notch and the target section is the ascending aorta, the direction of the blood flow may be assumed to be approximately parallel to the direction of the ultrasound beam. For example, measurements may be limited to a plane (or spherical cap) that is approximately perpendicular to the ultrasound beam.

In some cases, e.g., when the ultrasound transducer is placed on the upper abdomen and the target section is the descending aorta, the measured blood flow profile may be utilized to calculate an angle between the direction of blood flow in the target section and the direction of the ultrasound beam. The calculated angle may be used in calculating the stroke volume.

A continuous ultrasonic monitoring system in accordance with an embodiment of the present invention may be advantageous over other techniques or devices for hemodynamic monitoring. A continuous ultrasonic monitoring system in accordance with an embodiment of the present invention enables continuous noninvasive hemodynamic monitoring. The continuous ultrasonic monitoring system may be prepared for operation and operated quickly and easily by a technician who lacks extensive medical training. Once the continuous ultrasonic monitoring system is prepared for use, the system may be left alone to continue monitoring, with no further attention or handling on the part of any personnel (or entail, at most, minimal adjustment).

For example, invasive hemodynamic monitors require insertion of a catheter into a blood vessel. Using an invasive monitor, requiring skill and time to insert the catheters and obtain the measurements, could entail delays that could lead to increased complications in treatment. Furthermore, the high cost per use of invasive hemodynamic monitors may limit the number of patients that may be monitored at the same time. Administering fluid treatment without hemodynamic monitoring, and thus without determining the responsiveness of a patient to treatment, could result in increased in morbidity, mortality, or a longer stay in a hospital.

Typical ultrasound systems may be utilized to estimate blood flow velocities by transmitting streams of high-frequency sound waves and analyzing the signals reflected from circulating red blood cells. Currently, most ultrasound Doppler techniques measure the component of blood veloc-

ity that is parallel to the direction of the beam. These one-dimensional measurements depend on the spatial angle between the transmitted beam and the direction of the flow. Development of vector Doppler methods that apply apodization may create a transverse modulation of the ultrasound beam, thus enabling the measurement of the flow velocity within an entire two-dimensional imaging plane. Extension of vector Doppler imaging to three dimensions entails using expensive two dimensional matrix transducers and high-end scanners. Current ultrasound imaging methodology requires manual handling, e.g., positioning and aiming, of the ultrasound probe. Thus, the use of ultrasound in fluid resuscitation management may be limited in a situation (e.g., an operating room or emergency room) where access to the patient is required from several directions, or where trained personnel are required for the performance of more urgent tasks.

FIG. 1 shows a block diagram of a system for continuous ultrasonic monitoring, in accordance an embodiment of the present invention.

Continuous ultrasonic monitoring system 10 is configured to monitor one or more hemodynamic parameters of a patient. Ultrasound transducer 12 of continuous ultrasonic monitoring system 10 is configured to be attached to a surface of the skin of the patient.

Ultrasound transducer 12 includes an array of ultrasonic transducer elements 14. The array may be linear, curved, or two dimensional. Each ultrasonic transducer element 14 may include a piezoelectric transducer, a capacitive micro-machined ultrasonic transducer (CMUT), or another suitable ultrasonic transducer element.

For example, the array of ultrasonic transducer elements 14 may be controlled to function as a phased array to produce or receive a steerable ultrasound signal. For example, the signal may be in the form of a pulsed ultrasound signal (e.g., equivalent to a series of continuous waves having a particular spectrum of wavelengths and phases). Ultrasound transducer 12 may be configured to generate ultrasound signals that may be transmitted to, and whose reflections may be received from, points within ultrasound field of view 16 of ultrasound transducer 12. Ultrasound field of view 16 represents a substantially planar range, whose thickness is much smaller than its other dimensions.

Controller 20 is configured to control operation of ultrasound transducer 12 and of associated electronics to as to obtain a desired measured result. For example, the measured result may include a B-mode image of ultrasound that is reflected from various structures within a patient's body, or a Doppler mode measurement of a velocity of an object or collection of objects (e.g., blood cells) within the patient's body.

Controller 20 may include a standalone unit that includes processing capability and is dedicated to operation of continuous ultrasonic monitoring system 10. Alternatively or in addition, controller 20 may include a software module, circuit board, or other software or hardware components that are incorporated within, connected to, or otherwise hosted by a general purpose computer.

A processor of controller 20 may be configured to operate in accordance with programmed instructions. For example, the programmed instructions may be stored in a memory or data storage device with which the processor is configured to communicate.

One or more of transmit beam former 22, receive beam former 24, and a digital processing unit may include a separate or distinguishable hardware component or circuitry within controller 20, a software module of programmed

instructions for operation of a processor or processing unit of controller 20, or may represent a functionality of controller 20.

Transmit beam former 22 is configured to individually control transmission by each ultrasonic transducer element 14 so as to produce an ultrasonic beam with particular characteristics. The relative phases, amplitudes, and pulse shapes of ultrasound signals that are produced by each ultrasonic transducer element 14 may be controlled to form an ultrasonic beam having particular characteristics. For example, the beam may be aimed in a selectable direction within ultrasound field of view 16, or to modify ultrasound field of view 16 (e.g., rotate ultrasound field of view 16 or change the distance to the near and far boundaries of ultrasound field of view 16). Alternatively or in addition, the relative phases may be controlled to form a beam that is focused at a selectable distance from ultrasonic transducer elements 14 within ultrasound field of view 16. Ultrasonic transducer elements 14 may be controlled to form plane waves.

The transmitted signal may be reflected back toward ultrasonic transducer elements 14. For example, the signal may be reflected from blood cells in the aorta or another blood vessel, from walls of the aorta or another blood vessel, or from another structure, surface, or interface within the patient's body. Ultrasonic transducer elements 14 may detect the reflected signal.

Receive beam former 24 may be operated to receive signals from a selectable direction within ultrasound field of view 16. For example, receive beam former 24 may control synchronization of the signals that are received by each ultrasonic transducer element 14 so as to detect (e.g., reflected) ultrasound signals that originate from a selectable direction relative to ultrasound transducer 12. Similarly, receive beam former 24 may control synchronization of the signals that are received by each ultrasonic transducer element 14 so as to detect ultrasound signals that originate from a selectable distance from ultrasound transducer 12.

The received signals may be processed by signal processing unit 28. Signal processing unit 28 may perform one or more functions for extracting a desired measurement from the received signal.

For example, signal processing unit 28 may be configured to distinguish a received reflected ultrasound pulse from a carrier wave (e.g., demodulation) or background noise (e.g. filtering).

Signal processing unit 28 may be configured to determine an intensity or amplitude ("brightness") of an ultrasound pulse that is received at each point within ultrasound field of view 16. Thus, signal processing unit 28 may produce a B-mode image of structure within ultrasound field of view 16.

Signal processing unit 28 may be configured to operate in Doppler mode to calculate the phase shifts of consecutive ultrasound pulses that are reflected from a point or region within ultrasound field of view 16. These phase shifts may be analyzed to yield a component of velocity of the structure that reflected the pulses toward or away from ultrasound transducer 12. For example, if the structure includes blood cells, the measurement may yield a velocity of a component of the blood flow toward or away from ultrasound transducer 12.

A processing capability of controller 20 may analyze results from measurements using ultrasound transducer 12 to yield one or more hemodynamic parameters. For example,

results of B-mode operation and Doppler mode operation may be utilized to calculate a stroke volume, or another hemodynamic parameter.

A user of continuous ultrasonic monitoring system 10 may operate controller 20 via user interface 30. For example, user interface 30 may include one or more screen controls or input devices (e.g., switches, pushbuttons, levers, dials, knobs, touch screens, pointing devices, keyboards, keypads, or other input devices) that enable a user to input one or more instructions to controller 20. For example, instructions may include an instruction to begin or stop operation, operating parameters, patient information, or other instructions.

Results of operation of continuous ultrasonic monitoring system 10 may be displayed on monitor 32. For example, alphanumeric characters may be displayed on monitor 32 to indicate the value of one or more hemodynamic parameters, such as stroke volume, heartbeat rate, or other hemodynamic parameters. For Monitor 32 may be operated to display warnings or notifications to a user, results of analysis (e.g., a stroke volume or other hemodynamic parameter), patient information, or other information.

Monitor 32 may display an ultrasound image 34. Ultrasound image 34 may include an image of one or more anatomical structures 36 (e.g., the aortic arch, descending aorta, heart, or other structure of interest). For example, ultrasound image 34 may include a B-mode image of anatomical features within ultrasound field of view 16. In some cases, a B-mode image may be modified to display additional information. For example, parts of ultrasound image may be color coded or otherwise modified to display velocity results of Doppler mode measurements, e.g., within images of the aorta, or where Doppler mode measurements were made.

In some cases, continuous ultrasonic monitoring system 10 may include, or may be configured to communicate with, other types of sensors. Such additional sensor may include electrocardiogram electrodes, a pulse oximeter, a respiration monitor, a blood pressure monitor, or another type of sensor. In some cases, one or more sensors, or one or more electrodes of a sensor, may be incorporated into ultrasound measurement assembly 40 (e.g., in adhesive outer rim 42 or elsewhere). Results of such other sensors may be utilized in calculating values of additional hemodynamic parameters. Alternatively or in addition, results of such sensor measurements may be utilized to correct or adjust the ultrasound measurements (e.g., by enabling the filtering of noise, or otherwise).

FIG. 2A schematically illustrates an ultrasound measurement unit of the system shown in FIG. 1. For example, FIG. 2A may schematically represent the ultrasound measurement unit as viewed from the bottom (viewing the side that is configured for attachment to the skin surface). FIG. 2B is a schematic sectional view of the ultrasound measurement unit shown in FIG. 2A.

Ultrasound measurement assembly 40 includes ultrasound transducer 12 mounted on ultrasound transducer holder 41. Ultrasound transducer holder 41 is configured to hold ultrasound transducer 12 such that ultrasonic waves may pass efficiently between ultrasonic transducer elements 14 and skin surface 50 (and tissue below skin surface 50). Part or all of ultrasound transducer holder 41 may be designed to be disposable after a single use. For example, ultrasound transducer holder 41 may be in the form of a patch that may be caused to adhere to the patient's skin. In some cases, one or more components of ultrasound trans-

ducer holder **41** may be designed to be cleanable, sterilizable, or otherwise configured to enable single or multiple reuse of those components.

Ultrasound transducer holder **41** includes one or more adhesive surfaces that are coated with a material that is designed to adhere to skin surface **50**. The adhesive material may include a standard medical or surgical adhesive, or another material that includes one or more properties that make it suitable for use on skin, such as adherence to skin, being hypoallergenic, being removable from skin without discomfort or without leaving excessive residue, or other properties.

Alternatively or in addition to adhesive surfaces, ultrasound transducer holder **41** may include other structure for holding ultrasound transducer holder **41** to skin surface **50**. For example, ultrasound transducer holder **41** may include one or more suction cups, vacuum ports, straps, clamps, or other components for holding ultrasound transducer holder **41** to skin surface **50**.

The adhesive surfaces may be arranged so as to avoid interfering with the transmission of ultrasound waves from or to ultrasonic transducer elements **14**. For example, the adhesive strips may be arranged in the form of adhesive outer rim **42** that partially or fully surrounds other components at the periphery of ultrasound transducer holder **41**. Adhesive outer rim **42** may have an approximately square or rectangular form (or a rounded square or rectangular form), as shown, or may have another form (e.g., circular, oval, polygonal, or another form). Adhesive outer rim **42** may have a closed form as shown, or may include one or more gaps. In some cases, additional strips or pieces of adhesive may be located interior to adhesive outer rim **42**.

For example, prior to use, adhesive surface **43** of adhesive outer rim **42** may be covered with a removable layer of paper, plastic, or another type of material. After removal of the cover, adhesive surface **43** may be placed at a user-selected location on skin surface **50**. Application of pressure to the side of adhesive outer rim **42** opposite adhesive surface **43** may cause adhesive surface **43** to adhere to skin surface **50**.

Ultrasound transducer holder **41** includes coupling material retaining structure **44**. Coupling material retaining structure **44** may include walls or similar structure that is configured to retain an ultrasound coupling material (e.g., typically in the form of a gel or grease) within coupling material retaining structure **44**. For example, walls of coupling material retaining structure **44** may be made of plastic, or another material, that is impermeable to the ultrasound coupling material. Walls of coupling material retaining structure **44** may be held against skin surface **50**, e.g., when adhesive surface **43** of adhesive outer rim **42** adheres to skin surface **50**. When held against skin surface **50**, the walls of coupling material retaining structure **44** and skin surface **50** form a container or tub with interior space **46**. Interior space **46** of coupling material retaining structure **44** may be filled with the ultrasound coupling material. Thus the coupling material may be retained in interior space **46** to fill the volume between ultrasonic transducer elements **14** of ultrasound transducer **12** and skin surface **50**.

In some cases, end surface **45** of the walls of coupling material retaining structure **44** may be designed to form a seal with skin surface **50**. For example, end surface **45** may include a flexible material (e.g., rubber, silicone, or another flexible plastic or material) that functions as a gasket when end surface **45** is held against skin surface **50**. Thus, when end surface **45** of coupling material retaining structure **44** is held against skin surface **45**, the flexible material may

prevent seepage or the ultrasound coupling material may be configured to retain the ultrasound coupling material retaining structure **44** and skin surface **50**. (Prevention of such seepage may be desirable, in addition to prevention of loss of the ultrasound coupling material, in order to prevent interference of the ultrasound coupling material with adhesion of adhesive surface **43** to skin surface **50**.)

The walls of coupling material retaining structure **44**, or other structure of ultrasound transducer holder **41**, may include holding structure **48** for holding ultrasound transducer **12**. For example, holding structure **48** may include one or more clips, tabs, slots, pins, or other structure that is configured to engage corresponding structure of ultrasound transducer **12**. For example, a clip or pin of holding structure **48** may be elastically bendable, spring loaded, or otherwise configured such that when ultrasound transducer **12** is pushed into a socket or other part of ultrasound transducer holder **41**, the clip or pin is extended to engage a corresponding slot, socket, or hole on ultrasound transducer **12**.

Holding structure **48** may be configured such that when ultrasound transducer **12** is held by holding structure **48**, and when coupling material retaining structure **44** is filled with ultrasound coupling material, ultrasonic transducer elements **14** of ultrasound transducer **12** is submerged in the ultrasound coupling material. Furthermore, holding structure **48** may be configured so as to aim ultrasonic transducer elements **14** toward structure of interest (e.g., a section of the aorta or another structure) below skin surface **50** (e.g., when ultrasound transducer holder **41** is placed at one or more predetermined positions on skin surface **50**, e.g., at the suprasternal notch or on the upper abdomen). In some cases, part of ultrasound transducer holder **41** (e.g., adhesive outer rim **42**, or another part) may be provided with markings to facilitate placement of ultrasound transducer holder **41** at one or more standard positions on skin surface **50**. Thus, ultrasonic waves may be transmitted efficiently and effectively between ultrasonic transducer elements **14** and structure of interest (e.g., the aortic arch or the descending aorta) beneath skin surface **50** of a patient.

Ultrasound transducer **12** may be connected to transmit beam former **22**, receive beam former **24**, controller **20**, or other components of continuous ultrasonic monitoring system **10** via cable **52**. Alternatively of in addition, ultrasound transducer **12** may be connected to other components of continuous ultrasonic monitoring system **10** via a wireless connection.

Continuous ultrasonic monitoring system **10** may be configured to execute a method for continuous ultrasonic monitoring, in accordance an embodiment of the present invention.

FIG. 3A is a flowchart depicting a method for continuous ultrasonic monitoring, in accordance with an embodiment of the present invention.

It should be understood with respect to any flowchart referenced herein that the division of the illustrated method into discrete operations represented by blocks of the flowchart has been selected for convenience and clarity only. Alternative division of the illustrated method into discrete operations is possible with equivalent results. Such alternative division of the illustrated method into discrete operations should be understood as representing other embodiments of the illustrated method.

Similarly, it should be understood that, unless indicated otherwise, the illustrated order of execution of the operations represented by blocks of any flowchart referenced herein has been selected for convenience and clarity only.

13

Operations of the illustrated method may be executed in an alternative order, or concurrently, with equivalent results. Such reordering of operations of the illustrated method should be understood as representing other embodiments of the illustrated method.

Continuous ultrasonic monitoring method **100** may be executed by a processor of controller **20** of continuous ultrasonic monitoring system **10**. Execution of continuous ultrasonic monitoring method **100** may be initiated by a user or operator of continuous ultrasonic monitoring system **10**.

Controller **20** may be instructed to initiate execution of continuous ultrasonic monitoring method **100** after ultrasound measurement assembly **40**, with ultrasound transducer holder **41** and ultrasound transducer **12**, has been positioned on, and attached to, skin surface **50** of a patient (block **110**). Ultrasonic transducer elements **14** of ultrasound transducer **12** may be aimed in the general direction of a target section of a suitable anatomical structure (e.g., an anatomical structure with features that are identifiable in an ultrasound image).

For example, continuous ultrasonic monitoring system **10** may be operated in B-mode to obtain one or more images of a sector of the interior of the patient. The acquired images may be analyzed to identify the location of the target section relative to ultrasound transducer **12** (block **120**).

Continuous ultrasonic monitoring system **10** may be operated to repeatedly steer and focus the ultrasound beam to maintain the target section within an acquired ultrasound image (block **125**). For example, ultrasound field of view **16** may be moved such that an image of the target section remains within a subsequently acquired image, or at an approximately constant position within the acquired image.

The operations of identifying the target section (e.g., in an acquired image) and maintaining the image of the target section within acquired images may be repeated at frequent intervals (repeating continuously the operations of blocks **120** and **125**). Thus, the target section may be continuously monitored using ultrasound measurements.

In particular, continuous ultrasonic monitoring system **10** may be configured to execute a method for ultrasonic continuous hemodynamic monitoring, in accordance with an embodiment of the present invention.

FIG. **3B** is a flowchart depicting a method for ultrasonic continuous hemodynamic monitoring, in accordance with an embodiment of the present invention.

Continuous hemodynamic monitoring method **101** may be executed by a processor of controller **20** of continuous ultrasonic monitoring system **10**. Execution of continuous hemodynamic monitoring method **101** may be initiated by a user or operator of continuous hemodynamic ultrasonic monitoring system **10**.

Controller **20** may be instructed to initiate execution of continuous hemodynamic monitoring method **101** after ultrasound measurement assembly **40**, with ultrasound transducer holder **41** and ultrasound transducer **12**, has been positioned on, and attached to, skin surface **50** of a patient (block **110**). For example, ultrasound transducer holder **41** may be attached to the suprasternal notch, upper abdomen, or another suitable location on the patient. Ultrasonic transducer elements **14** of ultrasound transducer **12** may be aimed in the general direction of a target section of the aorta, or at another suitable anatomical structure.

Continuous ultrasonic monitoring system **10** may be operated in B-mode to obtain one or more images of the interior of the patient. The acquired images may be analyzed to identify the location of the target section relative to ultrasound transducer **12** (block **120**).

14

Continuous ultrasonic monitoring system **10** may be operated in Doppler mode to measure the velocity of the blood flow at various parts of the target section (block **130**). For example, the direction and focal distance of the transmitted and received ultrasound beam may be directed to various points of the target section, as determined from analysis of the B-mode images.

One or more hemodynamic parameters, such as stroke volume, may be calculated on the basis of the measured blood flow and geometry of the target section (block **140**). For example, flow rates may be summed over a cross section of the target section of the aorta and integrated over time to yield at least an initial estimate of the stroke volume. Adjustments and corrections to the calculation may be required.

In some cases, measurements in an approximately straight section of the target section (e.g., the descending aorta) may be analyzed to yield an angle of the target section relative to the ultrasound beam. Knowledge of the angle may enable derivation of an actual (total or absolute) blood velocity from an ultrasound measurement of a component of the velocity that is parallel to the beam direction.

FIG. **4** illustrates determination of a flow direction from a measurement made with the ultrasound measurement unit shown in FIG. **2A**.

In this case, value α of angle **74** may be calculated. Angle **74** is the angle between beam direction **58** and vessel axis **68** of target section **60** (e.g., of a blood vessel such as the descending aorta). Measurements of velocity profiles **64** and **66** may be made at two points that are equidistant from vessel axis **68** (and assumed to have the same actually flow velocity, assuming laminar flow of the blood in target section **60**, which implies an axially symmetric velocity profile). Velocity profile **64** includes peak **65**, and velocity profile **66** includes peak **67** which is assumed to correspond to peak **65** with an additional time delay due to the increased distance from ultrasonic transducer elements **14**.

Measured velocity $x(t)$ as a function of time t may be written for velocity profile **64** as $x(t)=s(t)+n_1(t)$, and measured velocity $y(t)$ for velocity profile **66** at a more distant point may be written as $y(t)=s(t+d)+n_2(t)$, where $s(t)$ is the ultrasound signal, $n(t)$ is noise that is added to each signal, and d is a time delay due to the distance between the measurement points.

The cross correlation between the two signals $R_{xy}(\tau)=R_{ss}(\tau+d)$, the correlation between the signal and the signal after time delay d . Since the cross correlation $R_{xy}(\tau)$ reaches its maximum value at $\tau=-d$, the time delay d may be derived from the cross correlation.

During time delay d , the blood moving at (actual unknown) velocity V travels a distance **72** with value $\Delta L=d \cdot V$.

The component of velocity V that is measured along beam direction **58** may be written as $v=V \cos(\alpha)$. Distance component **70**, measured to be Δr , between the points of measurement of velocity profiles **64** and **66** is $\Delta r=\Delta L/\cos(\alpha)$.

Therefore, $\Delta r=\Delta L/\cos(\alpha)=d \cdot V/\cos(\alpha)=d \cdot v/\cos^2(\alpha)$.

Therefore, value α of angle **74** may be derived from measured values (for the assumed symmetry of the velocity profile):

$$\alpha=\arccos(\sqrt{d \cdot v/\Delta r})$$

In other cases, e.g., where the target section includes the aortic arch, it may be possible to perform measurements on a part of the target section whose cross section is close to perpendicular to the beam direction.

The operations of alternating B-mode image acquisition and Doppler mode velocity measurements may be repeated at frequent intervals. During each interval, the hemodynamic parameters may be calculated (repeating continuously the operations of blocks **120** to **140**). Thus, the hemodynamic parameters may be continuously monitored using ultrasound measurements.

For example, hemodynamic parameters may be monitored both prior to and after application of fluid resuscitation or of fluid infusion to a patient. In this way, a physician or other healthcare administrator may be able to treat a patient with reduced risk of improper treatment.

In accordance with an embodiment of the present invention, an ultrasound measurement assembly may include a built-in layer of ultrasound coupling material. For example, the ultrasound measurement assembly may include a capsule that is filled with an ultrasound gel, a pad that is impregnated with an ultrasound gel or grease, or another ultrasound coupling structure.

A holding structure of the ultrasound measurement assembly for holding an ultrasound transducer may enable manual or actuated movement of the ultrasound transducer. The movement may include rotation, translation, or both. For example, the ultrasound transducer may be enclosed in an enclosure that includes a mechanism for altering an orientation of the ultrasound transducer about one or more axes, or for translating the ultrasound transducer along one or more axes. For example, linear translation may include pressing the ultrasound transducer toward the skin surface so as to improve the imaging of structure inside the body.

An ultrasound transducer, and, in some cases, a motorized or actuated orientation adjusting mechanism, translation mechanism, or both, may be connected to an external controller by a wireless connection. For example, an enclosure of the ultrasound transducer may include one or more components that enable wireless communication with an external device.

FIG. 5A schematically illustrates an ultrasound measurement assembly with a mechanically movable transducer, in accordance with an embodiment of the present invention. FIG. 5B schematically illustrates a bottom surface of the ultrasound measurement assembly shown in FIG. 5A. FIG. 5C schematically illustrates a side view of the ultrasound measurement assembly shown in FIG. 5A.

Ultrasound measurement device **204** of ultrasound measurement assembly **200** includes an ultrasound transducer **230** (shown in FIG. 6) and components to enable movement or aiming of, control of, and communication with ultrasound transducer **230**. Ultrasound measurement device **204** may be enclosed in enclosure **205**. For example, enclosure **205** may be constructed of a suitable plastic or metal material.

Fixed enclosure section **205b** of enclosure **205** may be attached to enclosure seat **207** of adhesive patch **201** by attachment structure **206** of adhesive patch **201**. For example, enclosure seat **207** may include a rigid section, socket, or rack structure that is shaped (e.g., molded, machined, or otherwise shaped) to receive fixed enclosure section **205b**. Enclosure seat **207** may be molded from plastic or otherwise formed. Attachment structure **206** may include clips, pins, latches, or another attachment structure. Enclosure seat **207** and attachment structure **206** may be configured to hold enclosure **205** and ultrasound measurement device **204** at a fixed position and orientation relative to adhesive patch **201**.

Adhesive patch **201** may be configured to hold ultrasound measurement assembly **200** to a skin surface. For example, an adhesive section **202** of adhesive patch **201** may include

an adhesive surface that may be exposed by removing (e.g., peeling off) a protective cover layer (e.g., made of a flexible plastic sheet). The exposed adhesive may be placed on or pressed onto a skin surface in order to hold ultrasound measurement assembly **200** to the skin surface.

Adhesive patch **201** may be disposable, including only relatively inexpensively made standard components. All reusable (e.g., electronic and relatively expensive) components of ultrasound measurement assembly **200** may be included in ultrasound measurement device **204**. Direct contact of ultrasound measurement device **204** with the skin surface may thus be prevented by adhesive patch **201**. Since ultrasound measurement device **204** does not directly contact the skin of a patient, ultrasound measurement device **204** may be reusable with different patients after light cleaning, without requiring sterilization. In some cases, ultrasound measurement device **204** may be constructed of inexpensive hardware components. For example, ultrasound transducer **230** may include a capacitive micro-machined ultrasonic transducer. In such a case, ultrasound measurement device **204** may also be disposable, such that all of ultrasound measurement assembly **200** may be disposable.

Adhesive patch **201** may include ultrasound coupler **224**. For example, ultrasound coupler **224** may include a capsule that is filled with, or made of, an ultrasound gel or grease. Thus, ultrasound measurement assembly **200** may be used to make subcutaneous ultrasound measurements without application of an ultrasound gel or grease. Therefore, preparation of ultrasound measurement assembly **200** for use, and reuse of ultrasound measurement device **204** and enclosure **205**, may be expedited by eliminating any requirement for extensive cleaning to remove an externally applied ultrasound coupling medium.

Adhesive patch **201** may include an electrode **222**. For example, electrode **222** may enable connection of adhesive patch **201** to an electrocardiography device or similar device configured to measure electric or electromagnetic signals. For example, connection to an electrocardiography device may enable ultrasound measurements that are synchronized with a patient's heartbeat or cardiac cycle. The electrocardiography signal may be processed by a controller that is associated with ultrasound measurement assembly **200** (e.g., device controller **240**, controller **20** of continuous ultrasonic monitoring system **10**, or another controller).

Ultrasound measurement device **204** is configured to enable manual rotation or aiming of an ultrasound transducer. For example, axial knob **210** may be rotated by a user of ultrasound measurement assembly **200** to axially rotate rotatable enclosure section **205a**, including an enclosed ultrasound transducer, relative to fixed enclosure section **205b**. Tilt knob **208** may be rotated by the user to tilt the enclosed ultrasound transducer relative to fixed enclosure section **205b** and adhesive patch **201**, and thus relative to the skin surface. In this manner, two-axis aiming of ultrasound transducer **230** may be enabled.

In some cases, axial knob **210** may be configured to be pushed inward. The inward pushing on axial knob **210** may press ultrasound transducer **230** toward adhesive patch **201** and the skin surface. For example, pressing ultrasound transducer **230** toward the skin surface may enable improved image by ultrasound transducer **230** of internal structures of the patient's body.

Ultrasound measurement device **204** may be configured to be connected by wire connector **212** to an external source of electric power, to a controller, or to other devices or components, e.g., of a continuous hemodynamic ultrasonic monitoring system **10** or other system. Wire connector **212**

may enable connecting ultrasound measurement device **204** to a fixed or portable external computer, to an external display or control unit, or to another type of device.

Ultrasound measurement device **204** may be configured to communicate wirelessly with one or more external devices (e.g., another ultrasound measurement device **204**), via wireless communication device **214**. For example, wireless communication device **214** may include a radiofrequency antenna, as shown, or another type of wireless communication device. The wireless connection may include a radiofrequency identification (RFID) patch, a barcode, or other structure to enable remote or automatic identification of a particular ultrasound measurement device **204**. For example, such identification may be used to ensure that an ultrasound measurement device **204** is compatible with a particular ultrasound transducer **230** or adhesive patch **201**, or to provide specific identification for each ultrasound measurement assembly **200** that is placed on a patient.

Ultrasound measurement device **204** may include one or more sensors **216** to sense one or more conditions. For example, a sensor **216** may include one or more motion and/or position sensors, such as accelerometers or proximity sensors (e.g., capacitive, potentiometric, or magnetic sensors), in order to ascertain the position or movement of the patch either relative to the patient body, with respect to other objects or devices (e.g., another ultrasound measurement assembly **200**, a surgical or other tool, or another object or device) or in absolute coordinates. For example, the sensors may be utilized to predict and correct an ultrasound measurement based on motion or position, or to provide additional clinical information regarding patient motion, such as due to breathing or other motion.

Ultrasound measurement device **204** may be configured to enable direct operation of monitoring by a user. For example, ultrasound measurement device **204** may include one or more user-operable controls **220** (e.g., pushbuttons or other controls, e.g., to select between modes of operation or operation parameters). Ultrasound measurement device **204** may include indicator lights **218** (e.g., light emitting diodes or other types of indicators), a display **223** (e.g., a liquid crystal display), or other components or structure to enable monitoring by a user (e.g., to indicate a mode of operation, status, warning, or other indication).

FIG. 6 schematically illustrates a sectional view of the of the ultrasound measurement assembly shown in FIG. 5C.

Ultrasound transducer **230** is mounted to tilt axis **234**. Tilt axis **234** together with ultrasound transducer **230** may be tilted by manual rotation of tilt knob **208**. Alternatively or in addition, ultrasound transducer **230** may be tilted about tilt axis **234** by operation of tilt actuator **236**. For example, tilt actuator **236** may be operated by a user via wired or wireless communication with device controller **240**. In some cases, an ultrasound measurement device may include only one of tilt knob **208** (for manual operation only) or tilt actuator **236** (for motorized or actuated operation only). In some cases, an ultrasound measurement device may include additional motors or actuators, or manual operation controls.

Tilt axis **234**, tilt knob **208**, and ultrasound transducer **230**, together with rotatable enclosure section **205a** and tilt actuator **236**, are rotatable about axial rotation axis **232**. For example, axial rotation axis **232** may be rotated by manual rotation of axial knob **210**. Alternatively or in addition, axial rotation axis **232** and ultrasound transducer **230** may be axially rotated by operation of axial actuator **238**. For example, axial actuator **238** may be operated by a user via wired or wireless communication with device controller **240**. In some cases, an ultrasound measurement device may

include only one of axial knob **210** (for manual operation only) or axial actuator **238** (for motorized operation only).

Device controller **240** may control one or more of operation of ultrasound transducer **230**, axial actuator **238**, tilt actuator **236**, indicator lights **218**, display **223**, or another component of ultrasound measurement device **204**. Device controller **240** may operate one or more sensors **216** and control operation of one or more components in accordance with a quantity that is sensed by a sensor **216**. Device controller **240** may include circuitry, or one or more processors. Device controller may include a battery or other internal power supply (e.g., an energy producing mechanism or wireless charging mechanism).

Thus, an orientation of ultrasound transducer **230** may be mechanically adjustable along one or two orthogonal axes by manual operation of one or both of axial knob **210** and tilt knob **208**. Alternatively or in addition to operation of knobs, an orientation of ultrasound transducer **230** may be mechanically adjusted by operation of one or more levers, dials, and knobs.

An orientation of ultrasound transducer **230** may be remotely, automatically, or otherwise electrically controlled by operation of one or both of axial actuator **238** and tilt actuator **236**. Axial actuator **238** or tilt actuator **236** may include a direct current motor or alternating current motor that includes a geared motor, stepper motor, servomotor, actuator, another type of motor, or any combination of the above. Alternatively or in addition, an orientation adjusting mechanism may include one or more magnets, whose orientation may be controlled by a magnetic or electromagnetic field.

In some cases, axial actuator **238** (or another axially oriented linear actuator) may be configured to linearly translate ultrasound transducer **230** along, or parallel to, axial rotation axis **232**. For example, linear translation of ultrasound transducer **230** by axial actuator **238** may press ultrasound transducer **230** toward adhesive patch **201** and the skin surface. For example, pressing ultrasound transducer **230** toward the skin surface may enable improved image by ultrasound transducer **230** of internal structures of the patient's body.

A motorized, hydraulic, or other remotely or electrically controllable orientation adjusting mechanism, e.g., that includes tilt actuator **236**, axial actuator **238**, both, or additional or alternative controllable mechanisms, may be operated to mechanically scan ultrasound transducer **230** over a range of orientations, or to mechanically translate ultrasound transducer **230** in one or more directions. For example, a scanning mechanism may enable scanning with one or more additional degrees of freedom (e.g., up to six degrees of freedom, e.g., by incorporating a Stewart platform mechanism, or otherwise). These degrees of freedom may include, or may be in addition to, the degrees of freedom that are attainable by electronically steering and focusing the ultrasound beam using phased array capabilities of ultrasound transducer **230**. The scanning mechanisms may be automatically controlled by a scanner (e.g., incorporated into or communicating with device controller **240**, controller **20** of continuous ultrasonic monitoring system **10**, or another controller) to continuously modify the orientation of ultrasound transducer **230** to track a target object inside the body, such as an organ, blood vessel, surgical instrument, robot, or another object that is inside the body or that was inserted into the body.

For example, automatic tracking may include acquiring a sequence of ultrasound images, and processing the images to detecting a target object in the images. The scanner may

measure the motion (e.g., speed, acceleration, curvature, or other parameter of the motion) of the object in each acquired frame of the ultrasound scan, and control the orientation of ultrasound transducer **230** so as to accommodate the measured motion.

In some cases, ultrasound measurement assembly **200** or ultrasound measurement device **204** may include a plurality of ultrasound transducers. An ultrasound transducer **230** may include a one- or two-dimensional array of ultrasonic transducer elements **14**.

In some cases, an adhesive patch **201** or ultrasound measurement device **204** may be configured to hold and orient a standard commercially available ultrasound probe and handle. In this case, adhesive patch **201** may enable affixing the probe to a body to enable provide automatically adjustable, hands-free and stable operation.

Ultrasound measurement assembly **200** may be utilized to automatically monitor blood flow in a selected blood vessel inside a patient's body. For example, the blood vessel may be automatically detected in an ultrasound image by applying image processing techniques. Device controller **240** (or an external controller, such as controller **20**, which is in communication with device controller **240**) may track motion of the blood vessel and automatically adjust the orientation of ultrasound transducer **230** to maintain its focus on the target vessel and to aim at the blood vessel. This tracking may enable continuous monitoring of clinical blood flow parameters such as flow volume, vascular resistance, pressure and rate.

Ultrasound measurement assembly **200** may be used to monitor minimally invasive surgery, e.g., where surgical tools are operated without a direct line of sight by the surgeon. In this case, ultrasound measurement assembly **200** may be placed on the patient's body near the region of the operation, and may provide a continuous ultrasound image of the region of the operation inside the body. The positions of the surgical tools may be tracked, and the orientation and focus of ultrasound transducer **230** may be adjusted accordingly. For example, the image of a surgical tool may be automatically identified and tracked. Similarly, ultrasound measurement assembly **200** may be used to track other foreign objects inserted into the body, such as miniature robots, pill cameras, or endoscopes.

Ultrasound measurement assembly **200** may be used to monitor post-operative and postpartum internal bleeding and/or fluid leakage. For example, ultrasound measurement assembly **200** may be placed on the body surface adjacent to the region of the operation following the operation. Ultrasound measurement assembly **200** may be set up to automatically detect and track irregular blood or fluid concentrations or accumulations, and to monitor their sizes and locations.

Ultrasound measurement assembly **200** may be used to monitor a previously detected tumor inside a patient's body. Following tumor detection or initial treatment, an ultrasound measurement assembly **200** may be periodically and repeatedly placed on the patient's body over the general location of the tumor. Ultrasound measurement assembly **200** may then automatically detect the tumor, and measure its size and accurate location. For example, changes in the tumor's size, shape, and location may be monitored following different treatments in order to evaluate their effectiveness.

Ultrasound measurement assembly **200** may be utilized in combination with a high intensity focused ultrasound (HIFU) system to provide automatic detection, tracking, and ablation of malignant tumors inside the body. In this configuration, both components, the imaging and HIFU trans-

ducers, may be co-aligned so as to be focused at the same region. Ultrasound measurement assembly **200** may scan, detect, and track the location and position of a tumor, and the HIFU transducer may focus onto the detected tumor and use ablation to treat or remove it. Ultrasound measurement assembly **200** may then validate the ablation results by tracking the tumor's size and shape after treatment.

Ultrasound measurement assembly **200** may be used to monitor a specific organ failure or insufficiency, e.g., as caused by a reduction in blood flow to the organ. For example, ultrasound measurement assembly **200** may be placed on the body surface near a failing kidney or other failing organ to monitor blood flow to the organ before and after treatment.

Imaging data generated using ultrasound measurement assembly **200** may be registered with previously acquired images of a patient using other imaging modalities (e.g., computed tomography, magnetic resonance imaging, or another modality). Thus, information that is acquired from the different modalities may be overlaid in a single image. For example, an image of a tumor that was identified using a different modality may be overlaid on acquire ultrasound images to assist in identifying a target section.

Data from two or more ultrasound measurement assemblies **200** may be synchronized and combined. For example, data from multiple ultrasound measurement assemblies **200** that are measuring blood flow in different blood vessels, e.g., central and peripheral blood vessels, may be analyzed to determine blood distribution in the circulatory system, calculate a blood pulse transit time (PTT), blood flow into and out of a particular organ, or another result. Ultrasound images that are acquired by two or more ultrasound measurement assemblies **200** at different angles may be processed to construct a three-dimensional representation of a volume within the body.

An ultrasound measurement assembly **200** containing a transducer array and mechanical scanning mechanism may be utilized to generate a three-dimensional representation of a volume inside the body, by mechanically sweeping the ultrasound transducer along an axis that is out of (e.g., perpendicular to) the imaging plane of the ultrasound transducer.

An orientation of an ultrasound measurement assembly **200**, or other parameters, may be stored in a data storage unit of, or accessible by, a controller. The stored information may be utilized at a later time to enable another ultrasound measurement assembly **200** to repeat previous measurements (e.g., at the same location or from the same viewing angle as in the previous measurement).

Different embodiments are disclosed herein. Features of certain embodiments may be combined with features of other embodiments; thus certain embodiments may be combinations of features of multiple embodiments. The foregoing description of the embodiments of the invention has been presented for the purposes of illustration and description. It is not intended to be exhaustive or to limit the invention to the precise form disclosed. It should be appreciated by persons skilled in the art that many modifications, variations, substitutions, changes, and equivalents are possible in light of the above teaching. It is, therefore, to be understood that the appended claims are intended to cover all such modifications and changes as fall within the true spirit of the invention.

While certain features of the invention have been illustrated and described herein, many modifications, substitutions, changes, and equivalents will now occur to those of ordinary skill in the art. It is, therefore, to be understood that

the appended claims are intended to cover all such modifications and changes as fall within the true spirit of the invention.

The invention claimed is:

1. A system for continuous ultrasonic monitoring, the system comprising:
 - (A) an ultrasound transducer, the ultrasound transducer comprising an array of ultrasonic transducer elements;
 - (B) an orientation adjusting mechanism including:
 - (i) an axial actuator configured for linear translation of the ultrasound transducer along an axial rotation axis for pressing the ultrasound transducer towards a skin surface,
 - (ii) a tilt actuator configured for tilting of the ultrasound transducer along a tilt axis, and
 - (iii) at least one motor physically connected to the axial actuator and the tilt actuator;
 - (C) an enclosure enclosing the orientation adjusting mechanism and the ultrasound transducer;
 - (D) a holder including an enclosure seat that is shaped to receive a fixed enclosure section of the enclosure for holding the ultrasound transducer, the holder being attachable to a skin surface of a patient; and
 - (E) a controller that is configured to execute programmed instructions for:
 - generating instructions for operating the at least one motor connected to the axial actuator and the tilt actuator of the orientation adjusting mechanism to automatically track a target section of an object inside a body of the patient by repeatedly operating the orientation adjusting mechanism for steering and focusing the ultrasound transducer to scan an ultrasound beam over a range of orientations to acquire a sequence of images of a sector, process the sequence of images of the sector to identify the target section, track motion of the target section in each image of the sequence of images, and operate the at least one motor connected to the axial actuator and the tilt actuator of the orientation adjusting mechanism to accommodate the tracked motion for acquiring an ultrasound image such that the target section is maintained within the acquired ultrasound image.
2. The system of claim 1, wherein the controller is configured to operate the ultrasound transducer in Doppler mode.
3. The system of claim 1, wherein the controller is configured to operate the ultrasound transducer in Doppler mode at a plurality of times and spatial locations during the automatic tracking to acquire a measurement of a flow rate of blood in the target section and to calculate a value of a hemodynamic parameter using the measured flow rate when the target section comprises a section of a blood vessel.
4. The system of claim 3, wherein the target section comprises a section of an ascending aorta, an aortic arch, or a descending aorta.
5. The system of claim 3, wherein the hemodynamic parameter is selected from a group of hemodynamic parameters consisting of stroke volume, cardiac output, stroke volume variation and systemic vascular resistance.
6. The system of claim 1, wherein the holder comprises a mechanism for altering an orientation and enabling move-

ment of the enclosure, and comprises a patch that includes one or a plurality of adhesive surfaces for attaching the holder to the skin surface.

7. The system of claim 1, wherein the holder is configured to enable rotation or translation of the ultrasound transducer.

8. The system of claim 1, wherein the holder includes a retaining structure for retaining an ultrasound coupling material between the ultrasonic transducer and the skin surface.

9. The system of claim 1, wherein the holder includes a capsule that is filled with an ultrasound coupling material.

10. The system of claim 1, wherein the controller is configured to repeatedly steer and focus the ultrasound beam to identify the target section by operating the ultrasound transducer in B-mode to acquire an image of a region of the patient that includes the target section.

11. The system of claim 10, wherein the controller is configured to continuously alternate operation of the ultrasound transducer between B-mode and Doppler mode.

12. The system of claim 1, wherein the holder is configured to be attached to the skin surface of the suprasternal notch and wherein the target section comprises an aorta.

13. The system of claim 1, wherein the controller is configured to calculate an angle between an axis of the target section and a direction of the ultrasound beam for calculation of a total blood velocity using a single array of ultrasonic transducer elements.

14. The system of claim 1, wherein the controller is configured to measure a diameter of the target section for integration of an entire blood flow through said target section over time to calculate flow volume parameters.

15. The system of claim 1, wherein the controller is configured to alternate between operation of the ultrasound transducer in B-mode and operation in Doppler mode, the ultrasonic beam during operation in Doppler mode is kept aimed at and focused on the target section computed from images obtained during B-mode, wherein during a period of alternation typical anticipated body movements do not significantly change the position of the ultrasound transducer relative to the target section.

16. The system of claim 6, wherein the controller is further configured to operate the mechanism for altering the orientation and enabling movement of the enclosure enclosing the orientation adjusting mechanism.

17. The system of claim 6, wherein the mechanism for altering the orientation and enabling movement of the enclosure enclosing the orientation adjusting mechanism is configured for at least one of: rotation, translation, altering an orientation of the enclosure about one or more axes, translating the enclosure along one or more axes, and pressing the enclosure towards the skin surface.

18. The system of claim 1, wherein the tilt axis is set for rotation about the axial rotational axis.

19. The system of claim 1, wherein the at least one motor is selected from the group consisting of direct current motor, alternating current motor, geared motor, stepper motor, servomotor, and combinations of the aforementioned.

20. The system of claim 1, wherein the tilt axis and the axial rotation axis are orthogonal.

* * * * *

专利名称(译)	连续超声波监测		
公开(公告)号	US10610193	公开(公告)日	2020-04-07
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[标]申请(专利权)人(译)	hemonitor医疗有限公司		
申请(专利权)人(译)	hemonitor医疗有限公司		
[标]发明人	BAR ZION AVINOAM MAYBLUM TOM TOUME SAMER		
发明人	BAR-ZION, AVINOAM MAYBLUM, TOM TOUME, SAMER		
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摘要(译)

用于连续超声监测的系统包括超声换能器。超声换能器包括超声换能器元件的阵列。超声换能器元件可操作以将超声束转向和聚焦到可选位置，并从可选位置接收超声信号。提供了用于保持超声换能器的保持器，该保持器可附接到患者的皮肤表面。控制器被配置为重复操纵和聚焦超声束以识别患者体内的对象的目标部分并获取超声图像，使得目标部分保持在所获取的超声图像内。

