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### (54) TISSUE IDENTIFYING METHOD IN ULTRASONOGRAPHY AND ULTRASONOGRAPH

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### Description

#### **Technical Field**

**[0001]** This invention relates to a method for identifying a living tissue in an ultrasonic diagnosis and an ultrasonic diagnosis system, and more particularly to a method for identifying a living tissue in an ultrasonic diagnosis, in which ultrasound is used to carry out a diagnosis of a lesion on a living tissue according to a noninvasive measurement, and an ultrasonic diagnosis system, thereby the kind of tissue such as a lipid-rich (or fatty) area, a thrombus area, an elastic fiber, a collagen fiber, and a calcified area is identified and is displayed in an effective manner for a diagnosis of a local lesion such as an atheroma in a blood vessel.

#### **Background Art**

[0002] Until now myocardial infarction, angina pectoris, cerebral infarction, and so on have been considered to be the development of a stenosis on a blood vessel which perfuses organs. However, it has become evident that these diseases are caused by the susceptibility of an atheroma to rupture. The atheroma is a blood wall disease. Actually according to a clinical examination of a cholesterol-lowering agent used worldwide, the remarkable effect of improving a survival rate and preventing myocardial infarction is obtained even through the stenosis of a blood vessel hardly changes. This is because the agent stabilizes an atheroma as a tissue. In consideration of this fact, a method for examining the susceptibility of an atheroma to rupture (susceptibility to rupture) has been demanded. A conventional method such as X-ray CT, MRI, and angiography cannot carry out such an examination.

[0003] For example, in view of an accurate measurement on a blood vessel disease, conventional echocardiography M-mode just has a resolution of 1 mm. Similarly when the vibration of an aorta is determined as a displacement velocity by conventional Doppler method, conditions for accuracy are theoretically satisfied but the pulsation of a blood vessel greatly affects in reality. Thus, it is difficult to extract a small vibration superimposed to a relatively large amplitude. Hence, researchers including the inventor have developed a phased tracking method whereby a small vibration on a beating heart and a large blood vessel is remotely measured by ultrasound and an elasticity modulus of a blood vessel wall can be calculated on a given spot. Thus, it is possible to accurately diagnose the susceptibility of an atheroma to rupture (Reference Documents 1 to 5).

[0004] The following are reference documents:

1. Kanai H, Hasegawa H, Chubachi N, Koiwa Y, Tanaka M. Noninvasive evaluation of local myocardial thickening and its color-coded Imaging. IEEE transaction UFFC. 1997; 44: 752-768; Hasegawa H, Kanai H, Hoshimiya N, Chubachi N, Koiwa Y. Accuracy evaluation in the measurement of a small change in the thickness of arterial walls and the measurement of elasticity of the human carotid artery. Jpn J Appl Phys 1998; 37: 3101-3105;
 Kanai H, Koiwa Y, J. Zhang Real-time measurements of local myocardium motion and arterial wall thickening. IEEE-transaction UFFC. 1999; 46: 1229-1241;

4. Japanese Patent Laid-Open No. 10-5226; and 5. Japanese Patent Laid-Open No. 12-229078.

[0005] The phased tracking method will be schematically described below. The phased tracking method is a 15 new bioinstrumentation for measuring a small vibration velocity on a cardiac wall and a blood vessel wall. This method makes it possible to accurately measure a vibration of 500 Hz or less and 0.01 mm and a change of 10 microns on a wall. With this method, for example, small 20 velocities on a plurality of measurement points between the layer (or layers) in the arterial wall or on the wall of an arterial vessel are determined by ultrasonic Doppler method, and the small velocities on the measurement points are subjected to time quadrature, so that a time 25 change in the positions of the measurement points can be calculated. Since a change in layer thickness can be determined by the time change in the positions of the measurement points, the elasticity modulus of the layer can be obtained, thereby estimating susceptibility to rup-30 ture.

**[0006]** Actually as shown in FIG. 17, an arterial intramural measurement point on an ultrasonic beam 91 is set at (i) and a measurement point with the subsequent depth is set at (i+1). Small vibration velocities  $v_i(t)$  and  $v_{i+1}(t)$  on the measurement points are determined and a difference between the small vibration velocities is subjected to time quadrature, so that a change  $\Delta h(t)$  in layer thickness between the measurement points (i) and (i+1) in the arterial wall is determined. Reference numeral 92 40 denotes a plaque.

$$\Delta h(t) = \int_{-\infty}^{t} \{ v_i(t) - v_{i+1}(t) \} dt$$

**[0007]** The simplest method for converting the change into an elasticity value for each layer in the arterial wall is performed as follows: a wall thickness is set at hd and a change in thickness is set at  $\Delta h$  at the lowest blood pressure where a wall thickness increases, a pulse pressure at a cuff pressure on brachial artery is set at  $\Delta P$ , and a wall elasticity modulus is measured for each layer in the following manner.

**[0008]** Based on a thickness change  $(\Delta h)$  of each layer from the intima to the adventitia of a blood vessel, an elasticity value (En) of each small part (n) in a blood ves-

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sel wall atheroma is determined by the formula below.

### $En = (\Delta P / (\Delta h / hd)n)$

**[0009]** With this phased tracking method, it is possible to measure an elasticity value for each layer along the depth direction in a blood vessel wall approximately every 0.75 mm to 0.075 mm on an ultrasonic beam, thereby displaying a tomogram based on the elasticity values.

**[0010]** In a clinical diagnosis using the phased tracking method, when an elasticity value is examined for each layer of a blood vessel wall, the elasticity value ranges from 100 to 2 MPa in a normal person. However, in the example of an atheroma, elasticity values are not evenly distributed. It is understood that fundamentally a physically soft portion is present in a thrombus and is covered with a hard portion. In addition, various patterns are present in the tomogram of an atheroma. For example, an extremely soft substance is exposed on the lumen of a blood vessel without continuity on a capsule of a hard surface, and a substance having a large elasticity value almost entirely covers a surface. Elasticity values in an atheroma are distributed from 0 to 4 MPa. Further, according to the examination results on the correspondence of a lipid-rich layer and a collage fiber in an extracted blood vessel and elasticity values in the phased tracking method, there is a probability that the tissue image of a collagen fiber covering an atheroma can be separately displayed from a tomogram of elasticity values, the tomogram being obtained in a noninvasive manner.

**[0011]** In the article "Kekkanheki soshiki seijo shindan chiryo system" by Hiroshi Kanai et al, published in "journal of Medical Ultrasonics", vol. 28, no. 3, 15 April 2001, there is disclosed a method for identifying a living tissue in an ultrasonic diagnosis. In Chapter 4 and FIG. 3 it is disclosed that the histogram of "healthy arterial wall" (upper part of FIG.3) differs from the histogram of "plark of arterial wall" (lower part of FIG. 3).

**[0012]** In the patent US-A-5 178 147 a method and an apparatus for elastographic measurement and imaging is disclosed. There, in column 19, lines 28-36 it is stated that the fat is much softer than the muscle.

**[0013]** The patent application WO 99 47046 Al also discloses a method for identifying a living tissue in an ultrasonic diagnosis. There FIG. 7 it is stated that the subcutaneous fat, parenchymal fat, fatty parecchyma, parecchyma, cancer, fibrocystic disease, and fibroadenoma are detected.

**[0014]** It is an object of the present invention to provide more specific information displayed in a tomogram and to identify the kind of living tissue such as a lipid rich area, a thrombus area, an elastic fiber, a collagen fiber, and a calcified area in an ultrasonic diagnosis of a living tissue such as a lesion on a blood vessel.

**[0015]** It is another object of the present invention to identify the kind of living tissue such as a lipid rich area,

a thrombus area, an elastic fiber, a collagen fiber, and a calcified area by using a shear elasticity modulus and a sheer viscosity in an ultrasonic diagnosis of a living tissue such as a lesion on a blood vessel.

5 [0016] The invention is defined by the method of claim1 and the system of claim 11. Preferred embodiments are as follows:

**[0017]** Further, the method for identifying a living tissue in an ultrasonic diagnosis preferably comprising the steps

10 of: exciting a vibration on a measured part of the living tissue to be diagnosed; measuring a shear elasticity and a shear viscosity of the excited living tissue to be diagnosed in the measurement using ultrasound; managing viscosity data for each kind of known living tissue such

as an adipose area, a thrombus area, an elastic fiber, a collagen fiber, and a calcified area in the management of the data library, the viscosity data being composed of a viscosity frequency histogram in each local small area for each kind of living tissue; and identifying the kind of
 living tissue to be diagnosed, regarding the elasticity data

and viscosity data serving as measurement results, by using the elasticity data and viscosity data for each kind of known living tissue in the identification of the kind of living tissue to be diagnosed.

<sup>25</sup> [0018] Further, the method for identifying a living tissue in an ultrasonic diagnosis according to the present invention, wherein the vibration on the measured part of the living tissue to be diagnosed is preferably excited by a turbulent flow component or a vortex component, the
 <sup>30</sup> components being generated by a time change in a pulse

of a pulsatile flow in a blood vessel.

**[0019]** Moreover, the method for identifying a living tissue in an ultrasonic diagnosis according to the present invention, wherein the vibration on the measured part of the living tissue to be diagnosed is preferably excited by

a vibration percutaneously applied to a part away from the measured part from the outside of a body.

[0020] Additionally, the method for identifying a living tissue in an ultrasonic diagnosis according to the present
 invention, wherein in the management of the data library, each kind of known living tissue is preferably mapped on a two-dimensional plane having axes representing a

shear elasticity modulus and a shear viscosity. [0021] Besides, the method for identifying a living tis-

45 sue in an ultrasonic diagnosis according to the present invention, wherein the living tissue to be diagnosed is preferably a living tissue on a heart, a blood vessel, and so on.

**[0022]** Moreover, the method for identifying a living tissue in an ultrasonic diagnosis according to the present invention, wherein the local small area serving as a unit for measuring the elasticity of the living tissue to be diagnosed is preferably several tens to several hundreds square microns.

<sup>55</sup> [0023] Besides, the method for identifying a living tissue in an ultrasonic diagnosis according to the present invention, wherein the kind of living tissue to be diagnosed is identified using the elasticity data serving as

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measurement results, and the identification is preferably performed by referring to the elasticity data for each kind of known living tissue, the data being extracted from the data library, and determining the kind of known living tissue having the minimum distance.

[0024] Moreover, the method for identifying a living tissue in an ultrasonic diagnosis according to the present invention, wherein the kind of known living tissue having the minimum distance from the elasticity data serving as measurement results is preferably determined by using Bayes decision method for dispersion and an average value of the elasticity moduli determined from the histogram of elasticity data on each kind of known living tissue. [0025] Additionally, the method for identifying a living tissue in an ultrasonic diagnosis according to the present invention, wherein an electronic chromatic figure is preferably generated and displayed so as to identify in colors the kind of living tissue to be diagnosed, the kind of living

tissue being identified for each local small area.

**[0026]** Further, the ultrasonic diagnostic system preferably comprising exciting means for exciting a vibration on a measured part of a living tissue to be diagnosed, wherein the data library further includes viscosity data for each kind of known living tissue and manages the viscosity data for each kind of known living tissue such as an adipose area, a thrombus area, an elastic fiber, a collagen fiber, and a calcified area, the viscosity data being composed of a viscosity frequency histogram in each local small area for each kind of living tissue, and the tissue identifying unit identifies the kind of living tissue to be diagnosed, regarding the elasticity data and viscosity data serving as measurement results of the excited living tissue to be diagnosed, by using the elasticity data and viscosity data for each kind of known living tissue.

**[0027]** Moreover, the ultrasonic diagnostic system of the present invention, wherein each kind of known living tissue is preferably mapped on a two-dimensional plane having axes representing a shear elasticity modulus and a shear viscosity.

**[0028]** Besides, the ultrasonic diagnostic system of the present invention, wherein the living tissue to be diagnosed is preferably a living tissue on a heart, a blood vessel, and so on.

**[0029]** Further, the ultrasonic diagnostic system of the present invention, wherein the local small area serving as a unit for measuring the elasticity of the living tissue to be diagnosed is preferably several tens to several hundreds square microns.

**[0030]** Moreover, the ultrasonic diagnostic system of the present invention, wherein the tissue identifying unit preferably has identifying means which refers to the elasticity data for each kind of known living tissue, the data being extracted from the data library, and determines the kind of known living tissue having the minimum distance, for each elasticity data serving as measurement results in each local small area.

**[0031]** Besides, the ultrasonic diagnostic system of the present invention, wherein the identifying means prefer-

ably determines dispersion and an average value of the elasticity moduli from the histogram of elasticity data of each kind of known living tissue, the data being extracted from the data library, and the identifying means performs identification using Bayes decision method.

**[0032]** Additional, the ultrasonic diagnostic system of the present invention, wherein the tissue identifying unit has electronic chromatic figure generating means for generating an electronic chromatic figure so as to identify

<sup>10</sup> in colors the kind of living tissue to be diagnosed, the kind of living tissue being identified for each local small area.

### BRIEF DESCRIPTION OF THE DRAWINGS

### [0033]

FIG. 1 is a schematic structural diagram showing an ultrasonic diagnostic system according to the present invention.

FIG. 2 is an explanatory drawing showing the generation of an elasticity modulus tomogram, and especially FIG. 2A shows one scanning position of an ultrasonic beam, FIG. 2B shows the calculation of displacement waveforms, FIG. 2C shows a thickness change waveform, FIG. 2D shows an elasticity modulus tomogram, and FIG. 2E shows the overall flow of the generation of an elasticity modulus tomogram.

FIG. 3 is a schematic diagram showing the creation of an elasticity data library.

FIG. 4 is a flow showing tissue identification.

FIG. 5 is a diagram showing the measurement results of elasticity moduli of a fiber tissue.

FIG. 6 is a diagram showing an example of tissue identification (71-year-old man).

FIG. 7 is a diagram showing an example of tissue identification (52-year-old man).

FIG. 8 is a schematic structural diagram showing another ultrasonic diagnostic system according to the present invention.

FIG. 9 is a conceptual drawing showing a vibration excited on a blood vessel wall.

FIG. 10 is a conceptual drawing showing a vibration excited on a blood vessel wall.

FIG. 11 is a diagram showing an example of a waveform of a small vibration velocity v(t) on a blood vessel wall.

FIG. 12 is a diagram showing a power spectrum obtained by frequency analysis.

FIG. 13 is a diagram showing a tomogram, and especially FIG. 13A shows a tomogram of attenuation, and FIG. 13B shows a tomogram of a phase velocity FIG. 14 is a diagram showing a tomogram, and especially FIG. 14A shows a tomogram of a shear elasticity modulus, and FIG. 14B shows a tomogram of a shear viscosity.

FIG. 15 is a diagram showing an example of the data

library where tissues are mapped on a two-dimensional plane.

FIG. 16 is a conceptual drawing showing an example of a tomogram having undergone electronic dyeing and

FIG. 17 is a diagram showing an image of elasticity modulus distribution of an atheroma on a carotid wall.

Best Mode for Carrying Out the Invention

### [Embodiment 1]

**[0034]** In this example, the conventional problem is solved as follows: based on findings that tissues such as a lipid-rich area, a thrombus area, an elastic fiber, a collagen fiber, and a calcified area that are found on a blood vessel wall are different in hardness according to tissue characteristics, the frequency distribution (histogram) concerning an elasticity value for each kind of tissue is acquired beforehand and is managed in a data library, each local elasticity value obtained in an ultrasonic diagnosis is compared with the frequency distribution of the elasticity values of the tissues in the data library, a tissue is discriminated by determining the tissue having a frequency distribution of the closest elasticity value, and tissue type information is displayed, for example, in colors in a tomogram.

**[0035]** FIG. 1 is a schematic structural diagram showing an ultrasonic diagnostic system according to the present invention.

[0036] In FIG. 1, reference numeral 1 denotes a human body, reference numeral 2 denotes the surface of the body, and reference numeral 3 denotes a blood vessel including an artery to be measured. In the blood vessel, a blood vessel wall has a thickness of h(t), a lumen has a diameter of d(t), and a vibration is made at a movement velocity of v(t). Reference numeral 3a denotes the anterior wall of the blood vessel. Reference numeral 3b denotes the posterior wall of the blood vessel. Reference numeral 3c denotes a lesion such as an atheroma appearing in the blood vessel 3. Reference numeral 3d denotes a lipid rich tissue. Reference numeral 3e denotes the lumen of the blood vessel 3. Reference numeral 4 denotes an ultrasonic probe which can change the emitting direction of an ultrasonic beam to perform scanning over a certain range. Reference numeral 5 denotes an ultrasonic measuring unit which generates a transmission signal with an angular frequency of  $\omega_0$  at a fixed time interval  $\Delta T$  to drive the ultrasonic probe 4 and radiates an ultrasonic beam to the blood vessel 3. When the ultrasonic probe 4 receives a reflected wave from the blood vessel 3, the received signal of the probe is subjected to quadrature demodulation and AD conversion and a detection signal is outputted in a digital signal format.

**[0037]** Reference numeral 6 denotes a data analyzing unit. The data analyzing unit 6 analyzes the detection signal, which has been outputted in a digital signal format

from the ultrasonic measuring unit 5, according to phased tracking method, performs tracking on the track of the large amplitude displacement of the blood vessel 3, and determines a small vibration velocity of each reflection point on the surface of the blood vessel wall and between

layers on the track. According to the result, a time change is calculated for the thickness of the blood vessel wall or each of the layers. An elasticity modulus of a tissue on the blood vessel wall or each of the layers is accurately

10 calculated based on the time change in each small area determined by a resolving power, and elasticity modulus tomogram data is generated. In this tracking, the displacing motion of a small vibration in the layers of the blood vessel is analyzed under the constraint that a cumulative

<sup>15</sup> displacement in a pulse is set at 0 so as to make a return to the original position at each pulse of the heart, thereby stabilizing an image.

[0038] Reference numeral 7 denotes an elasticity data library used as the reference (a kind of dictionary) of tissue identification. The elasticity data library 7 stores elasticity data concerning kinds of living tissues. The data has been inputted beforehand and serves as the reference. To be specific, the elasticity data library 7 stores

frequency distribution (histogram) data concerning an
 elasticity value of each kind of known living tissue. Further, an average value and dispersion may be calculated
 from the frequency distribution data and may be replaced with the frequency distribution.

**[0039]** Reference numeral 8 denotes a tissue identifying unit, in which regarding an elasticity value on each point of the elasticity modulus tomogram data generated by the data analyzing unit 6, the elasticity data concerning various kinds of living tissues is extracted from the elasticity data library 7, the elasticity data serving as the ref-

<sup>35</sup> erence, a comparison is made, the closest kind of living tissue is discriminated, and electronic chromatic figure data is generated. In the electronic chromatic figure data, the area of the elasticity modulus tomogram is displayed in colors corresponding to the kinds of tissues.

40 [0040] Reference numeral 9 denotes a display device which displays an electronic chromatic figure based on the electronic chromatic figure data generated by the tissue identifying unit 9.

[0041] Reference numeral 10 denotes an elasticity data library creating unit, in which regarding data effectively acting as the reference in the elasticity modulus tomogram data generated by the data analyzing unit 6, a comparison is made with a pathologic chromatic figure generated in advance, so that the kind of living tissue is iden-

<sup>50</sup> tified in each local area of the tomogram, and attribute data indicating the kind of living tissue is inputted and set in the elasticity modulus tomogram data. Then, attribute data in each local area is examined, and the frequency distribution of an elasticity modulus is determined for <sup>55</sup> each kind of living tissue and is registered in the elasticity data library 7.

**[0042]** The embodiment of the ultrasonic diagnostic system shown in FIG. 1 will be discussed in detail below.

**[0043]** FIG. 2E shows the overall flow of the generation of an elasticity modulus tomogram that corresponds to conventional processing performed by the ultrasonic measuring unit 5 and the data analyzing unit 6 of FIG. 1. As shown in FIG. 2A, a blood vessel area to be diagnosed on a living tissue is scanned by the ultrasonic probe along an arrow S, an ultrasonic pulse is transmitted at each scanning position (beam position P) at a regular interval (step S11), and reflected waves are collected from the anterior wall and posterior wall of a blood vessel and boundaries between the layers of the blood vessel (step S12). Step S11 and step S12 are repeated by the ultrasonic measuring unit 5 at each scanning position of an ultrasonic beam position.

[0044] Subsequently, data of the reflected waves collected by the data analyzing unit 6 is analyzed. First, as shown in FIG. 2B, regarding points (indicated by circles) on the surface of the blood vessel wall and boundaries between the layers of the blood vessel on one beam, displacement waveforms are calculated and tracking is performed to track the large amplitude motion generated on the blood vessel by heart stroke (step S13). Step S13 is repeated on all the points of one beam. Then, between adjacent points of FIG. 2A, a thickness change waveform is calculated as shown in FIG. 2C (step S14). In this example, at the largest diameter of the lumen in FIG. 2B, a thickness change in FIG. 2C has the maximum value Max, which indicates the smallest thickness. The maximum value of the thickness change in a pulse is calculated (step S15). By using the value and a difference between the maximum blood pressure and the minimum blood pressure of a cuff pressure measured on an upper arm, a local elasticity modulus is calculated (step S16). Steps S14 to S16 are repeated on all the points. When local elasticity moduli are calculated for layers between all the adjacent points on one beam scanning position (step S17), a shift is made to the subsequent beam scanning position, steps S13 to S17 are repeated, and local elasticity moduli are similarly calculated for layers between all the adjacent points. In this way, local elasticity moduli are calculated on all the beam scanning positions and an elasticity modulus tomogram is calculated, in which the local elasticity modulus of a diagnostic area is indicated by a pixel value or a color attribute as shown in FIG. 2D (step S18).

**[0045]** FIG. 3 is a schematic diagram showing the generation of the elasticity data library that corresponds to the processing of the elasticity data library creating unit 10 shown in FIG. 1. In the generation of the elasticity data library, regarding a suitable living tissue to be used as the reference of tissue identification, an area of an elasticity modulus tomogram is classified for each kind of tissue, and the frequency distribution (histogram) of an elasticity modulus is calculated for each kind of tissue and is registered in the elasticity data library. Hence, an ultrasonic measurement part, that is the elasticity modulus tomogram calculated by the flow of FIG. 2 is displayed on a screen (step S21) and another sample tissue

is compared with an actual chromatic figure, in which a specific kind of tissue is dyed by pathologic dyeing, so that each kind of tissue area is identified in the elasticity modulus tomogram and area specification is performed

- <sup>5</sup> on each kind of tissue (step S22). The example of FIG. 3 has five kinds of tissues including a lipid-rich area, a thrombus area, an elastic fiber, a collagen fiber, and a calcified area. In this way, after the setting of an area for each kind of tissue required for the elasticity modulus
- <sup>10</sup> tomogram, the frequency distribution of an elasticity modulus is calculated for each resolution area such as a pixel in each area for each kind of tissue (step S22) and the results are registered in the elasticity data library (step S23).

<sup>15</sup> [0046] FIG. 4 shows the flow of tissue identification that corresponds to the processing of the tissue identifying unit 8 shown in FIG. 1. In tissue identification, the following processing is repeated: on each point of the elasticity modulus tomogram of a measured living body,

20 the elasticity modulus is classified to the closest kind of tissue with reference to elasticity modulus frequency distribution data of tissues registered in the elasticity data library. In this example, Bayes decision method (Bayes classification method) is used as a mathematical method

- for tissue identification. Bayes decision method is one of statistical estimation methods. In this method, regarding the frequency distribution of a kind of tissue that is most likely to have an elasticity modulus on one point of the elasticity modulus tomogram, an estimation is made so that a defined loss function has the minimum expected
- value. This estimation corresponds to the selection of a similar pattern of the minimum distance in pattern recognition.

[0047] First, the elasticity modulus tomogram of a measured living body is determined by the processing of FIG. 2 (step S31). On the other hand, from the elasticity data library generated based on the calculation of the elasticity modulus tomogram (step S36) in the processing of FIG. 3, elasticity modulus frequency distribution data

- 40 is read regarding five kinds including a lipid-rich area, which serves as a candidate of tissue classification, and an average value and dispersion are calculated for each elasticity modulus distribution (Step S37). Then, elasticity moduli on the points of the elasticity modulus tomo-
- <sup>45</sup> gram are sequentially extracted (step S32), Bayes decision is performed by using values determined in step S37 (step S33), and the elasticity modulus is classified as a minimum-distance tissue in processing results (step S34). Steps S32 to S34 are repeated on all the points of
- 50 the elasticity modulus tomogram. In this way, an elasticity modulus on each point of the elasticity modulus tomogram is classified as a proper tissue and an electronic chromatic figure is generated (step S35).

**[0048]** Subsequently, FIGS. 5, 6, and 7 show the ex-<sup>55</sup> amples of actual measurements.

**[0049]** In FIG. 5, reference numerals 51 to 53 denote the examples of elasticity modulus measurement results of a fibrous tissue. An image 51 on the left shows an

elasticity modulus tomogram and an image 52 shows the tomogram of a fibrous tissue. A graph 53 on the right shows the histogram of a fibrous tissue. In FIG. 5, reference numerals 54 to 56 denote the examples of elasticity modulus measurement results of a lipid-rich tissue. An image 54 on the left shows an elasticity modulus tomogram and an image 55 shows the tomogram of a lipid-rich tissue. A graph 56 on the right shows the histogram of a lipid-rich tissue.

**[0050]** FIGS. 6 and 7 show tissue discrimination examples of a 71-year-old man and a 52-year-old man. In FIGS. 6 and 7, reference numerals 61 and 71 denote B-mode images, reference numerals 62 and 72 denote elasticity modulus tomograms, reference numerals 63 and 73 denote the tomograms of a lipid-rich tissue, reference numerals 64 and 74 denote the tomograms of a fibrous tissue, and reference numerals 65 and 75 denote the electronic chromatic figures of a tissue.

### [Embodiment 2]

[0051] In this example, the conventional problem is solved as follows: based on findings that tissues such as a lipid-rich area, a thrombus area, an elastic fiber, a collagen fiber, and a calcified area that are found on a blood vessel wall are different in viscosity in addition to hardness according to each tissue characteristic, the frequency distribution (histogram) of viscosity in addition to an elasticity value for each kind of tissue is acquired beforehand and is managed in a data library, each local elasticity value and viscosity value that are obtained in an ultrasonic diagnosis are compared with the frequency distribution of the elasticity values and viscosity values of the tissues in the data library, a tissue is discriminated by determining a tissue having a frequency distribution of the closest elasticity value and viscosity, and tissue type information is displayed, for example, in colors in a tomogram.

**[0052]** A motion such as the pulsation of a living tissue and a change in intravascular pressure have a low frequency band of 30 Hz or less. Thus, when a motion such as the pulsation of a living tissue and a change in intravascular pressure serve as external forces applied to a tissue, a measurement is considered to be static, though an amplitude is large. In the case of such a static measurement, it is necessary to measure a deformation of a tissue before and after a change in pressure and calculate an elasticity modulus by the calculation of (pressure change)/(distortion), (so called static technique). However, in reality, it is difficult to measure a pressure applied inside and a change in pressure in vivo.

**[0053]** Hence, the following method is proposed: a lowfrequency sine wave oscillation (generally a single frequency component) is applied from the outside by using an oscillator and so on, the propagation velocity of a vibration (shear elasticity) is measured in vivo, and a parameter for a hardness of a tissue is calculated (external oscillation + dynamic method, Jpn J. Med Ultrasonics, Vol. 16, No. 3, pp. 221-229, 1989). In this method, it is not necessary to measure a pressure and a change in pressure. When a blood vessel wall is oscillated, although a bending oscillation can be readily excited so as

<sup>5</sup> to entirely bend a blood vessel wall, it is difficult to propagate a vibration so as to penetrate the wall from the outside to the inside of the blood vessel wall.

**[0054]** Hence, the present invention focuses attention on a fact that components such as a "turbulent flow" and

<sup>10</sup> a "vortex" that are generated by a time change in a pulse of a pulsatile flow in a blood vessel "excite vibrations of components over a wide frequency band of several tens Hz to several hundreds Hz on a blood vessel wall with a small amplitude of several mm/s or less." The wall vibra-

<sup>15</sup> tion is accurately measured by using ultrasound and a waveform is analyzed, so that a propagation velocity and a propagation loss of a vibration (shear elastic wave) in vivo can be calculated at each frequency f.

[0055] Besides, (1) the vibrations excited by the "turbulent flow" and the "vortex" are small in amplitude (several mm/s or less) and (2) the position of the wall is moved by about several mm due to the arrival of pulsation. Thus, measurement could not be performed in vivo.

**[0056]** For example, in a conventional ultrasonic Doppler measurement technique, an average blood flow velocity of an area (several mm width) set in the lumen of a blood vessel is measured by frequency analysis on ultrasound reflected in a blood flow or correlation method. However, a target velocity in a blood flow measurement

<sup>30</sup> is several tens cm/s, which is larger than the vibration velocity of a blood vessel wall by two digits or more. Although the momentary magnitude of a blood flow velocity is considered important in a diagnosis, the waveform of the obtained blood flow velocity was not subjected to
 <sup>35</sup> waveform analysis (frequency analysis) and so on.

**[0057]** In this example, an ultrasonic measurement method invented by the inventor (Japanese Patent Laid-Open No. 8-163418, U.S. Patent No. 5,840,028) is adopted. With this method, a waveform with a small vibration

40 velocity generated by pulsation on an arterial wall can be measured by using ultrasound. In the present invention, a waveform of wall vibration is measured by the ultrasonic measurement method simultaneously on a plurality of points set at intervals of several hundreds microns from

<sup>45</sup> the intima to the adventitia of a blood vessel wall on one ultrasonic beam.

**[0058]** FIG. 8 shows a schematic structural diagram showing another ultrasonic diagnostic system according to the present invention.

50 [0059] FIG. 8 shows that the ultrasonic diagnostic system is further provided with a data library 7A instead of the elasticity data library 7, a data library creating unit 10A instead of the elasticity data library creating unit 10, and exciting means 11, in the structure of FIG. 1.

<sup>55</sup> [0060] In addition to the elasticity data, the data library 7A stores viscosity data of living tissues that serves as previously inputted reference data, that is frequency distribution (histogram) data of a viscosity value for each

kind of known living tissue.

**[0061]** In addition to the elasticity data, the data library creating unit 10A registers the frequency distribution of a viscosity for each kind of living tissue. That is, regarding data effectively acting as the reference data in viscosity tomogram data generated by a data analyzing unit 6, a comparison is made with a previously generated pathologic chromatic figure, so that the kind of living tissue is identified in each local area of the tomogram, and attribute data indicating the kind of living tissue is inputted and set in the viscosity tomogram data. Then, attribute data of each local area is examined, the distribution frequency of viscosities is obtained for each kind of living tissue, and the frequency distribution is registered in the data library 7A.

**[0062]** The embodiment of the ultrasonic diagnostic system of FIG. 8 will be described in detail. The following explanation will mainly discuss differences from Embodiment 1.

**[0063]** In accordance with the processing flow of FIG. 2E, for example, a waveform with a small vibration velocity v(t) that is generated on an arterial wall due to pulsation is measured (steps S11 to S13) by using the ultrasonic measurement method above-described (U.S. Patent No. 5,840,028). Thus, a waveform of wall vibration is measured simultaneously on a plurality of points set at intervals of several hundreds microns from the intima to the adventitia of a blood vessel wall on one ultrasonic beam. The amplitude is several mm/s or less and the frequency is a band of 0 Hz to several hundreds Hz.

**[0064]** At the measurement, as shown in FIG. 9, a pulse wave 11a caused by pulsation propagates through a lumen 3e of a blood vessel 3 and a lesion 3c causes a turbulent flow 11b or a vortex 11c. The turbulent flow 11b and the vortex 11c, each serving as a vibration source, vibrate the wall of the blood vessel 3 in the above-described manner. Thus, a vibration v(t) is measured on each point (white circle) by an ultrasonic beam B. Reference numeral 3f denotes the adventitia of the blood vessel wall. Reference numeral 3g denotes the intima of the blood vessel wall. The measured vibration v(t) is shown in FIG. 11. FIG. 11 shows measurement results in a period of four pulses. It is found that each point vibrates according to pulsation.

**[0065]** A pulse wave 11a may be generated by pulsation. That is, any structure is applicable as long as an inner product is changed by excited bending oscillation (of blood vessel 3) and thus an internal pressure change  $\Delta P(f)$  is excited as shown in FIG. 10B. Reference character "f" denotes a frequency at which a vibration source vibrates. The inner pressure change  $\Delta P(f)$  propagates as the pulse wave 11a. Therefore, for example, a vibration on a measured part 3h (FIG. 10) is excited by the component of the turbulent flow 11b or the vortex 11c which are generated by a time change in a pulse of a pulsatile flow in the blood vessel 3.

**[0066]** Further, as shown in FIG. 10A, the pulse wave 11a may be generated by the exciting means 11. The

vertical vibration of the exciting means 11 at the frequency "f" excites the internal pressure change  $\Delta P(f)$ . Therefore, for example, the vibration on the measured part 3h is excited by a vibration percutaneously applied to a part away from the measured part 3h from the outside of the body.

**[0067]** Moreover, instead of steps S14 to S16 of FIG. 2E, waveforms are cut out at the same timing in a pulse and frequency analysis is performed (while a window is

<sup>10</sup> applied in a diastole), so that a transfer function H(f) from a vibration waveform vm(t) at m point on the intima to a vibration waveform vn(t) at n-th point of the adventitia is calculated. That is, the power spectrum of FIG. 12 is obtained by the frequency analysis. When a distance is d

<sup>15</sup> between the two points in the power spectrum, an attenuation constant  $\alpha(f) = |H(f)|/d$  associated with propagation between the two points can be calculated from an amplitude |H(f)|. Moreover, a phase velocity v(f) =  $2\pi f/(\angle H(f)/d)$  of propagation between the two points can be calculated from a phase <H(f) of the transfer function.

**[0068]** A tomogram on the attenuation of FIG. 13A can be obtained from the attenuation constant  $\alpha$ (f). Further, a tomogram of the phase velocity of FIG. 13B can be obtained from the phase velocity v(f). Comparison between the tomograms proves that the single lesion 3c includes both of a lesion shown in the tomogram of attenuation and a lesion show in the tomogram of phase velocity.

**[0069]** Based on the two parameters, a complex propagation constant  $\gamma(f) = \alpha(f) + j\beta(f)$  is determined where "j" represents an imaginary unit and  $\beta(f) = \omega/v(f)$  and  $\omega = 2\pi f$  are established.  $\gamma(f)$  indicates that a propagation per unit length has an attenuation of  $\alpha(f)$  and a phase rotation of  $\beta(f)$ .

<sup>35</sup> **[0070]** According to a document (J. Biomechanics, vol. 28, No. 10, pp. 1145 to 1154, 1995), the complex propagation constant  $\gamma$ (f) has the relationship of

$$\mu + j\omega \eta = -\rho \omega^2 / \gamma^2(f)$$

between a shear (or shearing) elastic constant (a modulus of rigidity)  $\mu$  and a shear viscosity constant  $\eta$ . When a measured complex transfer constant  $\gamma$ (f) is substituted into the right side of the formula and a real item and an imaginary item are arranged, the shear elastic constant  $\mu$  and the shear viscosity constant  $\eta$  are expressed by the formulas below.

$$\mu = -\rho \omega^2 (\alpha^2(f) - \beta^2(f)) / \Delta$$

$$\eta = 2\rho \omega \alpha(f) \beta(f) / \Delta$$

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art.

### Where,

### $\Delta = (\alpha^2(\mathbf{f}) \cdot \beta^2(\mathbf{f}))^2 + (2\alpha(\mathbf{f}) \cdot \beta(\mathbf{f}))^2$

is established. Thus, a shear elastic constant  $\mu$  and a shear viscosity constant  $\eta$  are calculated for a small area between the point m and the point n on one ultrasonic beam (step S 16).

**[0071]** The above-described measurement is performed on each point of the lesion in and on the arterial wall on one ultrasonic beam (step S17). The point is set with the order of a waveform of a used ultrasonic wave, for example, at intervals of 375 microns. Further, an ultrasonic beam is scanned in real time along the axial direction of the blood vessel wall at intervals of 600 microns (FIG. 9). Thus, a tomogram with the shear elastic constant  $\mu$  and the shear viscosity constant  $\eta$  is extracted in each local area of 375 microns × 600 microns (step S18).

**[0072]** A tomogram of a shear elasticity modulus shown in FIG. 14A can be obtained from the shear elastic constant (i.e., shear elasticity modulus)  $\mu$ . Moreover, a tomogram of a shear viscosity shown in FIG. 14B can be obtained from the shear viscosity constant (i.e., shear viscosity)  $\eta$ . The tomogram of the shear elasticity modulus shown in FIG. 14A corresponds to the tomogram of a phase velocity shown in FIG. 13B, and the tomogram of the shear viscosity shown in FIG. 14B corresponds to the tomogram of the tomogram of a the tomogram of attenuation shown in FIG. 13A.

**[0073]** As with the processing of FIG. 3, regarding a representative lesion, the shear elastic constant  $\mu$  and the shear viscosity constant  $\eta$  that are obtained on each point are compared with a pathologic chromatic figure and data of elasticity and viscosity of tissues such as lipid-rich, thrombus, elastic fiber, collagen fiber and calcification is registered in the data library 7A. At this point of time, as shown in FIG. 15, the tissues are mapped on a two-dimensional plane where the vertical axis represents a shear viscosity (Pa·s) and the horizontal axis represents a shear elasticity modulus (Pa), so that registration is performed. In this way, classification can be performed on the two-dimensional plane by using an elastic characteristic and a viscous characteristic, thereby increasing the accuracy of the data library 7A.

**[0074]** Then, as with the processing of FIG. 4, on the tomogram of the shear elastic constant  $\mu$  and shear viscosity constant  $\eta$  to be actually measured, a tissue characteristic is identified on each local area with reference to the data library 7A and electronic dyeing is performed. Thus, as shown in FIG. 16, tissues such as a lipid-rich area 3h, a thrombus area 3i, an elastic fiber 3j, a collagen fiber 3k, and a calcified area 31 can be clearly displayed on a screen by electronic dyeing.

### Industrial Applicability

**[0075]** According to the present invention, in an ultrasonic diagnosis of a living tissue such as a lesion in a blood vessel, it is possible to readily identify the kind of living tissue such as a lipid rich area, a thrombus area, an elastic fiber, a collagen fiber, and a calcified area on an ultrasonic diagnostic image. Thus, a noninvasive diagnosis can be performed in a clinical diagnosis more properly and quickly as compared with the conventional

**[0076]** For example, the present invention makes it possible to predict the rupture of an atheroma (arterio-sclerotic plaque) that is a main cause of an acute cardi-

<sup>15</sup> ovascular event such as unstable angina, acute myocardial infarction, and sudden death. Such a prediction is impossible in a noninvasive manner in the conventional art.

**[0077]** It is considered that a serious cardiovascular disease such as myocardial infarction and cerebral infarction is caused by the clogging of a thrombus in the lumen of a blood vessel, the thrombus being formed by the rupture of an arteriosclerotic plaque (atheroma), which is formed on an arterial wall and is rich in lipid-rich.

<sup>25</sup> The present invention can evaluate the stability of an atherosclerotic lesion in a noninvasive manner.[0078] Further, the present invention can evaluate the curative effect of a cholesterol lowering treatment.

### 30 Claims

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1. A method for identifying a living tissue in an ultrasonic diagnosis, the method comprising the steps of:

measuring at least an elasticity modulus frequency distribution of a living tissue to be diagnosed in each local small area by using ultrasound (S11 - S18);

managing, in a data library (7; 7A), at least elasticity data for an elastic fiber, a collage fiber, a lipid-rich area, a thrombus area, and a calcified fiber which are kinds of known living tissues, the elasticity data being composed of an elasticity modulus frequency histogram in each local small area (3h - 31) for each kind of living tissue (S22 - S23); and identifying the kind of living tissue to be diagposed using the frequency distribution of the

nosed, using the frequency distribution of the measured elasticity data and using the frequency distribution of the elasticity data for each kind of known living tissue (S31-S37).

 The method for identifying a living tissue in an ultrasonic diagnosis according to claim 1, the method further comprising the steps of:

exciting a vibration on a measured part of the

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living tissue to be diagnosed (S11 - S13); measuring a shear elasticity and a shear viscosity of the excited living tissue to be diagnosed in the measurement using ultrasound (S16 - S17); managing viscosity data for an elastic fiber, a collagen fiber, a lipid-rich area, a thrombus area, and a calcified fiber which are kinds of known living tissues, in the management of the data library (7; 7A), the viscosity data being composed of a viscosity frequency histogram in each local small area (3h - 31) for each kind of living tissue (S18); and

identifying the kind of living tissue to be diagnosed, regarding the elasticity data and viscosity data serving as measurement results, by using the elasticity data and viscosity data for each kind of known living tissue in identification of the kind of living tissue to be diagnosed (s31 - S37).

- **3.** The method for identifying a living tissue in an ultrasonic diagnosis according to claim 2, wherein the vibration on the measured part of the living tissue to be diagnosed is excited by a turbulent flow component or a vortex component, the components being generated by a time change in a pulse of a pulsatile flow in a blood vessel.
- 4. The method for identifying a living tissue in an ultrasonic diagnosis according to claim 2, wherein the vibration on the measured part of the living tissue to be diagnosed is excited by a vibration percutaneously applied to a part away from the measured part from an outside of a body.
- The method for identifying a living tissue in an ultrasonic diagnosis according to claim 2, wherein in management of the data library (7; 7A), each kind of known living tissue is mapped on a two-dimensional plane having axes representing a shear elasticity modulus and a shear viscosity.
- **6.** The method for identifying a living tissue in an ultrasonic diagnosis according to claim 1 or 2, wherein the living tissue to be diagnosed is a living tissue on a heart, a blood vessel, and so on.
- The method for identifying a living tissue in an ultrasonic diagnosis according to claim 1 or 2, wherein the local small area (3h 31) serving as a unit to measure an elasticity and/or a viscosity of the living tissue to be diagnosed is several tens to several hundreds square microns.
- The method for identifying a living tissue in an ultrasonic diagnosis according to claim 1 or 2, wherein 55 the kind of living tissue to be diagnosed is identified using the elasticity data and/or the viscosity data serving as measurement results, and the identifica-

tion is performed by referring to the elasticity data and/or the viscosity data for each kind of known living tissue, the data being extracted from the data library (7; 7A), and determining the kind of known living tissue having a minimum distance.

- **9.** The method for identifying a living tissue in an ultrasonic diagnosis according to claim 8, wherein the kind of known living tissue having a minimum distance from the elasticity data serving as a measurement result is determined by using Bayes decision method for dispersion and an average value of elasticity moduli determined from the histogram of the elasticity data on each kind of known living tissue.
- 10. The method for identifying a living tissue in an ultrasonic diagnosis according to claim 1 or 2, wherein an electronic chromatic figure is generated and displayed so as to identify in colors the kind of living tissue to be diagnosed, the kind of living tissue being identified for each local small area (3h 31).
- **11.** An ultrasonic diagnostic system configured to measure at least a frequency distribution of the elasticity modulus of a living tissue to be diagnosed in each local small area by using ultrasound and to display a tomogram, the system comprising:

a data library (7; 7A) comprising at least elasticity data for an elastic fiber, a collagen fiber, a lipid-rich area, a thrombus area, and a calcified fiber which are kinds of known living tissues, the elasticity data being composed of an elasticity modulus frequency histogram in each local small area (3h - 31) for each kind of living tissue; and

a tissue identifying unit (8) configured to identify the kind of living tissue to be diagnosed using the frequency distribution of the measured elasticity data and using the frequency distribution of the elasticity data for each kind of know living tissue.

12. The ultrasonic diagnostic system according to claim11, further comprising:

exciting means(11) for exciting a vibration on a measured part of a living tissue to be diagnosed, wherein the data library (7; 7A) further includes viscosity data for an elastic fiber, a collagen fiber, a lipid-rich area, a thrombus area, and a calcified fiber which are kinds of known living tissues, and manages the viscosity data for said kinds of known living tissues, the viscosity data being composed of a viscosity frequency histogram in each local small area (3h - 31) for each kind of living tissue, and

the tissue identifying unit (8) identifies the kind

of living tissue to be diagnosed, regarding the elasticity data and viscosity data serving as measurement results of the excited living tissue to be diagnosed, by using the elasticity data and viscosity data for each kind of known living tissue.

- 13. The ultrasonic diagnostic system according to claim
  12, wherein each kind of known living tissue is
  mapped on a two-dimensional plane having axes
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  representing a shear elasticity modulus and a shear
  viscosity.
- 14. The ultrasonic diagnostic system according to claim
   11 or 12, wherein the living tissue to be diagnosed
   15 is a living tissue on a heart, a blood vessel, and so on.
- **15.** The ultrasonic diagnostic system according to claim 11 or 12, wherein the local small area (3h-31) serving as a unit to measure an elasticity of the living tissue to be diagnosed is several tens to several hundreds square microns.
- 16. The ultrasonic diagnostic system according to claim 11 or 12, wherein the tissue identifying unit (8) has identifying means which refers to the elasticity data for each kind of known living tissue, the data being extracted from the data library, and determines the kind of known living tissue having a minimum distance, for each elasticity data serving as a measurement result in each local small area.
- 17. The ultrasonic diagnostic system according to claim
  16, wherein the identifying means determines dispersion and an average value of elasticity moduli
  35 from the histogram of elasticity data of each kind of known living tissue, the data being extracted from the data library, and the identifying means performs identification using Bayes decision method.
- The ultrasonic diagnostic system according to claim 11 or 12, wherein the tissue identifying unit (8) has electronic chromatic figure generating means for generating an electronic chromatic figure so as to identify in colors the kind of living tissue to be diagnosed, the kind of living tissue being identified for each local small area (3h - 31).

### Patentansprüche

1. Verfahren zur Gewebs- Identifikation in einer Ultraschall-Diagnose, umfassend die Schritte:

> Messen mindestens einer Elastizitäts- Modul-Häufigkeits- Verteilung eines zu diagnostizierenden Gewebes in jedem lokal kleinen Bereich durch Verwendung von Ultraschall (S11-S18);

Verwalten in einer Datenbibliothek (7, 7A) von zumindest Elastizitäts- Daten für eine elastische Faser, eine Kollagen- Faser, einen Lipid- reichen Bereich, einen Thrombus- Bereich und eine verkalkte Faser, welche Arten von bekanntem lebendem Gewebe darstellen, wobei die Elastizitäts- Daten von einem Elastizität- Modul-Häufigkeits- Histogramm in jedem lokal kleinen Bereich (3h-31) für jede Art von lebendem Gewebe (S22-S23) zusammengesetzt werden; und

Identifizieren der Art von lebendem zu diagnostizierendem Gewebe durch Verwenden der Häufigkeitverteilung der gemessenen Elastizitäts- Daten und durch Verwenden der Häufigkeit- Verteilung der Elastizitäts-Daten für jede Art von bekanntem lebendem Gewebe (S31-S37).

 20 2. Verfahren zur Gewebs- Identifikation in einer Ultraschall-Diagnose nach Anspruch 1, wobei das Verfahren weiter die Schritte umfasst:

> Anregen einer Vibration auf einem gemessenen Abschnitt des lebenden zu diagnostizierenden Gewebes (S11-S13);

Messen einer Scherungs- Elastizität und Scherungs-Viskosität des angeregten lebenden zu diagnostizierenden Gewebes in der Messung mittels Ultraschall (S16-S17);

Verwalten von Viskositäts- Daten für eine elastische Faser, eine Kollagen- Faser, einen Lipidreichen Bereich, einen Thrombus- Bereich und eine verkalkte Faser, welche Arten von bekanntem lebendem Gewebe sind, in der Verwaltung der Datenbibliothek (7; 7A), wobei die Viskositäts- Daten von einem Viskositäts- Häufigkeits-Histogramm in jedem lokal kleinen Bereich (3h-31) für jede Art von lebendem Gewebe (S18) zusammengesetzt werden; und

Identifizieren der Art von lebendem zu diagnostizierendem Gewebe, wobei die Elastizitäts-Daten und Viskositäts- Daten als Messergebnisse dienen, durch Verwenden der Elastizitäts-Daten und Viskositäts- Daten für jede Art von bekanntem lebendem Gewebe bei Identifikation der Art von lebendem zu diagnostizieren Gewebe (S31-S37).

50 3. Verfahren zur Gewebsidentifikation in einer Ultraschall-Diagnose nach Anspruch 2, wobei die Vibration auf dem gemessenen Abschnitt des lebenden zu diagnostizieren Gewebes durch Anregen einer turbulenten Strömungs-Komponente oder einer Wirbel- Komponente angeregt wird, wobei die Komponenten durch eine zeitliche Änderung eines Impulses einer pulsierenden Strömung in einem Blutgefäß erzeugt werden.

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- 4. Verfahren zur Gewebsidentifikation in einer Ultraschall-Diagnose nach Anspruch 2, wobei die Vibration auf dem gemessenen Abschnitt des lebenden zu diagnostizieren Gewebes durch eine vibration angeregt wird, die perkutan auf einen Abschnitt abseits von dem gemessenen Abschnitts von Außerhalb des Körpers angewendet wird.
- 5. Verfahren zur Gewebsidentifikation in einer Ultraschall-Diagnose nach Anspruch 2, wobei beim Verwalten der Datenbibliothek (7, 7A) jede Art von bekanntem lebenden Gewebe auf einer zwei- dimensionalen Ebene mit Achsen abgebildet wird, die ein Scherungs- Elastizitäts- Modul und eine Scherungs-Viskosität darstellen.
- Verfahren zur Gewebsidentifikation in einer Ultraschall-Diagnose nach Anspruch 1 oder 2, wobei das lebende zu diagnostizierende Gewebe lebendes Gewebe des Herzens, eines Blutgefäßes usw. ist. 20
- Verfahren zur Gewebsidentifikation in einer Ultraschall-Diagnose nach Anspruch 1 oder 2, wobei der lokal kleine Bereich (3h - 31), der als eine Einheit zum Messen einer Elastizität und/oder einer Viskosität des lebenden zu diagnostizierenden Gewebes dient, mehrere Zehn bis mehrere Hundert Quadrat-Mikrometer groß ist.
- Verfahren zur Gewebsidentifikation in einer Ultraschall-Diagnose nach Anspruch 1 oder 2, wobei die Art des lebenden zu diagnostizierenden Gewebes durch Verwenden der Elastizitäts- Daten und/oder Viskositäts- Daten identifiziert wird, die als Messergebnisse dienen, und wobei die Identifikation durchgeführt wird durch Bezugnahme auf die Elastizitäts-Daten und/oder Viskositäts- Daten für jede Art von bekanntem lebendem Gewebe, wobei die Daten aus der Datenbibliothek (7; 7A) bezogen werden, und wobei die Art von bekanntem lebendem Gewebe bestimmt wird, die einen minimale Abstand aufweist.
- Verfahren zur Gewebsidentifikation in einer Ultraschall-Diagnose nach Anspruch 8, wobei die Art des bekannten lebenden Gewebes, die einen minimalen 45 Abstand zu den Elastizitäts- Daten aufweist und als Messergebnis dient, bestimmt wird durch Verwenden der Bayes- Entscheidungs-Methode für Dispersion und einen Durchschnittswert von Elastizitäts-Modulen, die von dem Histogramm der Elastizitäts-50 Daten von jeder Art bekanntem lebendem Gewebes bestimmt werden.
- 10. Verfahren zur Gewebsidentifikation in einer Ultraschall-Diagnose nach Anspruch 1 oder 2, wobei eine elektronische chromatische Figur erzeugt und angezeigt wird, um somit in Farben das lebende zu diagnostizierende Gewebe zu identifizieren, wobei die

Art von lebendem Gewebe für jeden lokal kleinen Bereich (3h - 31) identifiziert wird.

11. Ultraschall- Diagnose- System, das beschaffen ist, zumindest eine Häufigkeits- Verteilung des ElastizitätsModuls eines lebenden zu diagnostizierenden Gewebes in jedem lokal kleinen Bereich durch Verwenden von Ultraschall zu messen und ein Tomogramm anzuzeigen, wobei das System umfasst:

eine Datenbibliothek (7; 7A), die mindestens Elastizitäts- Daten für eine elastische Faser, eine Kollagen- Faser, einen Lipid- reichen Bereich, einen Thrombus- Bereich und eine verkalkte Faser aufweist, welche Arten von bekanntem lebendem Gewebe darstellen, wobei die Elastizitäts- Daten von einem Elastizitäts-Modul-Häufigkeits- Histogramm in jedem lokal kleinen Bereich (3h-31) für jede Art von lebendem Gewebe zusammengesetzt werden; und eine Gewebs- Identifikations- Einheit (8), die beschaffen ist, die Art von lebendem zu diagnostizierendem Gewebe zu identifizieren durch Verwenden der Häufigkeit-Verteilung der gemessenen Elastizitäts- Daten und durch Verwenden der Häufigkeits- Verteilung der Elastizitäts-Daten für jede Art von bekanntem lebendem Gewebe.

30 12. Ultraschall- Diagnose- System nach Anspruch 11, das weiterhin umfasst:

Anregungs- Mittel (11) zum Anregen einer Vibration auf einem gemessenen Abschnitt des lebenden zu diagnostizierenden Gewebes; wobei die Datenbibliothek (7; 7A) weiterhin Viskositäts- Daten für eine elastische Faser, eine Kollagen- Faser, einen Lipid- reichen Bereich, einen Thrombus- Bereich und eine verkalkte Faser enthält, welche Arten von bekanntem lebendem Gewebe sind, und die Viskositäts- Daten für die Arten von bekanntem lebendem Gewebe sind, und die Viskositäts- Daten für die Arten von bekanntem lebendem Gewebe verwaltet, wobei die Viskositäts- Daten von einem Viskositäts- Häufigkeits- Histogramm in jedem lokal kleinen Bereich (3h- 31) für jede Art von lebendem Gewebe zusammengesetzt sind, und

die Gewebs- Identifikations- Einheit (8) die Art von lebendem zu diagnostizierendem Gewebe identifiziert, wobei auf die Elastizitäts- Daten und Viskositäts- Daten, die als Messergebnisse des angeregten lebenden zu diagnostizierenden Gewebes dienen, Bezug genommen wird durch Verwenden der Elastizitäts- Daten und ViskositätsDaten für jede Art von bekanntem lebendem Gewebe.

13. Ultraschall- Diagnose- System nach Anspruch 12,

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- Ultraschall- Diagnose- System nach Anspruch 11 oder 12, wobei das lebende zu diagnostizierende Gewebe lebendes Gewebe des Herzens, eines Blutgefäße usw. ist.
- 15. Ultraschall- Diagnose- System nach Anspruch 11 oder 12, wobei der lokal kleine Bereich (3h-31), der als eine Einheit zum Messen einer Elastizität und/ oder einer Viskosität des lebenden zu diagnostizierenden Gewebes dient, mehrere Zehn bis mehrere Hundert Quadrat- Mikrometer groß ist.
- 16. Ultraschall- Diagnose- System nach Anspruch 11 oder 12, wobei Gewebs- Identifikations- Einheit (8) IdentifikationsMittel aufweist, die auf die Elastizität- 20 Daten und/oder Viskositäts- Daten für jede Art von bekanntem lebendem Gewebe Bezug nehmen, wobei die Daten aus der Datenbibliothek (7; 7A) bezogen werden, und die die Art von bekanntem lebendem Gewebe bestimmt, die einen minimale Abstand 25 aufweist für jede Elastizitäts- Daten, die als Messergebnis in jedem lokal kleinem Bereich dienen.
- 17. Ultraschall- Diagnose- System nach Anspruch 16, wobei die Identifikations- Mittel Dispersion und einen <sup>30</sup> Durchschnittswert für Elastizitäts- Module von dem Histogramm der Elastizitäts- Daten von jeder Art bekanntem lebendem Gewebes bestimmt, wobei die Daten aus der Datenbibliothek bezogen sind und die IdentifikationsMittel eine Identifikation durch Ver-<sup>35</sup> wenden der Bayes-Entscheidungs- Methode durchführt.
- 18. Ultraschall- Diagnose- System nach Anspruch 11 oder 12, wobei die Identifikations- Einheit (8) ein Mit- 40 tel zum Erzeugen einer elektronischen chromatischen Figur aufweist, um eine elektronische chromatische Figur zu erzeugen, um somit in Farben das lebende zu diagnostizierende Gewebe zu identifizieren, wobei die Art von lebendem Gewebe für jeden 45 lokal kleinen Bereich (3h-31) identifiziert wird.

### Revendications

1. Procédé d'identification d'un tissu vivant dans un diagnostic par ultrasons, le procédé comprenant les étapes consistant à :

> - mesurer au moins une distribution de fréquence de module d'élasticité d'un tissu vivant devant être diagnostiqué dans chaque petite zone locale à l'aide d'ultrasons (S11-S18);

- gérer, dans une bibliothèque de données (7 ; 7A), au moins des données d'élasticité pour une fibre élastique, une fibre de collagène, une zone riche en lipides, une zone de thrombus et une fibre calcifiée qui sont des sortes de tissus vivants connus, les données d'élasticité étant composées d'un histogramme de fréquence de module d'élasticité dans chaque petite zone locale (3h-31) pour chaque sorte de tissu vivant (S22-S23) ; et

- identifier la sorte de tissu vivant devant être diagnostiqué, à l'aide de la distribution de fréquence des données d'élasticité mesurées et à l'aide de la distribution de fréquence des données d'élasticité pour chaque sorte de tissu vivant connu (S31-S37).

2. Procédé d'identification d'un tissu vivant dans un diagnostic par ultrasons selon la revendication 1, le procédé comprenant en outre les étapes consistant à :

> exciter une vibration sur une partie mesurée du tissu vivant devant être diagnostiqué (S11-S13);

> - mesurer une élasticité de cisaillement et un coefficient de viscosité de cisaillement du tissu vivant excité devant être diagnostiqué dans la mesure à l'aide d'ultrasons (S16-S17);

- gérer des données de viscosité pour une fibre élastique, une fibre de collagène, une zone riche en lipides, une zone de thrombus et une fibre calcifiée qui sont des sortes de tissus vivants connus, dans la gestion de la bibliothèque de données (7;7A), les données de viscosité étant composées d'un histogramme de fréquence de viscosité dans chaque petite zone locale (3h-31) pour chaque sorte de tissu vivant (S18) ; et - identifier la sorte de tissu vivant devant être diagnostiqué, au vu des données d'élasticité et des données de viscosité qui servent de résultats de mesure, à l'aide des données d'élasticité et des données de viscosité pour chaque sorte de tissu vivant connu dans l'identification de la sorte de tissu vivant devant être diagnostiqué (S31-S37).

- 3. Procédé d'identification d'un tissu vivant dans un diagnostic par ultrasons selon la revendication 2, dans lequel la vibration sur la partie mesurée du tissu vivant devant être diagnostiqué est excitée par une composante d'écoulement turbulent ou une composante tourbillonnaire, les composantes étant générées par un changement de temps dans une pulsation d'un écoulement pulsatoire dans un vaisseau sanguin.
- 4. Procédé d'identification d'un tissu vivant dans un diagnostic par ultrasons selon la revendication 2, dans

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lequel la vibration sur la partie mesurée du tissu vivant devant être diagnostiqué est excitée par une vibration appliquée par voie percutanée sur une partie à distance de la partie mesurée, depuis l'extérieur d'un corps.

- Procédé d'identification d'un tissu vivant dans un diagnostic par ultrasons selon la revendication 2, dans lequel, dans la gestion de la bibliothèque de données (7 ; 7A), chaque sorte de tissu vivant connu est tracée sur un plan en deux dimensions ayant des axes représentant un module d'élasticité de cisaillement et un coefficient de viscosité de cisaillement.
- Procédé d'identification d'un tissu vivant dans un diagnostic par ultrasons selon l'une des revendications 1 ou 2, dans lequel le tissu vivant devant être diagnostiqué est un tissu vivant sur un coeur, un vaisseau sanguin, etc.
- 7. Procédé d'identification d'un tissu vivant dans un diagnostic par ultrasons selon l'une des revendications 1 ou 2, dans lequel la petite zone locale (3h-31) servant d'unité pour mesurer une élasticité et/ou une viscosité du tissu vivant devant être diagnostiqué est de plusieurs dizaines à plusieurs centaines de micromètres carrés.
- 8. Procédé d'identification d'un tissu vivant dans un diagnostic par ultrasons selon l'une des revendications 1 ou 2, dans lequel la sorte de tissu vivant devant être diagnostiqué est identifiée à l'aide des données d'élasticité et/ou des données de viscosité servant de résultats de mesure, et l'identification est réalisée par référence aux données d'élasticité et/ou aux 35 données de viscosité pour chaque sorte de tissu vivant connu, les données étant extraites de la bibliothèque de données (7 ; 7A), et par détermination de la sorte de tissu vivant connu ayant une distance minimale.
- 9. Procédé d'identification d'un tissu vivant dans un diagnostic par ultrasons selon la revendication 8, dans lequel la sorte de tissu vivant connu ayant une distance minimale à partir des données d'élasticité servant de résultat de mesure est déterminée à l'aide de la méthode de décision de Bayes pour la dispersion et une valeur moyenne de modules d'élasticité déterminée à partir de l'histogramme des données d'élasticité sur chaque sorte de tissu vivant connu.
- 10. Procédé d'identification d'un tissu vivant dans un diagnostic par ultrasons selon l'une des revendications 1 ou 2, dans lequel une figure chromatique électronique est générée et affichée de façon à identifier en couleurs la sorte de tissu vivant devant être diagnostiqué, la sorte de tissu vivant étant identifiée pour chaque petite zone locale (3h-31).

11. Système de diagnostic par ultrasons configuré pour mesurer au moins une distribution de fréquence du module d'élasticité d'un tissu vivant devant être diagnostiqué dans chaque petite zone locale à l'aide d'ultrasons et pour afficher un tomogramme, le système comprenant :

- une bibliothèque de données (7 ; 7A) comprenant au moins des données d'élasticité pour une fibre élastique, une fibre de collagène, une zone riche en lipides, une zone de thrombus et une fibre calcifiée qui sont des sortes de tissus vivants connus, les données d'élasticité étant composées d'un histogramme de fréquence de module d'élasticité dans chaque petite locale (3h-31) pour chaque sorte de tissu vivant ; et - une unité d'identification de tissu (8) configurée pour identifier la sorte de tissu vivant devant être diagnostiqué à l'aide de la distribution de fréquence des données d'élasticité mesurées et à l'aide de la distribution de fréquence des données d'élasticité pour chaque sorte de tissu vivant connu.

25 12. Système de diagnostic par ultrasons selon la revendication 11, comprenant en outre :

> - des moyens d'excitation (11) pour exciter une vibration sur une partie mesurée d'un tissu vivant devant être diagnostiqué,

 la bibliothèque de données (7 ; 7A) comprenant en outre des données de viscosité pour une fibre élastique, une fibre de collagène, une zone riche en lipides, une zone de thrombus et une fibre calcifiée qui sont des sortes de tissus vivants connus, et gérant les données de viscosité pour lesdites sortes de tissus vivants connus, les données de viscosité étant composées d'un histogramme de fréquence de viscosité dans chaque petite zone locale (3h-31) pour chaque sorte de tissu vivant, et

 l'unité d'identification de tissu (8) identifiant la sorte de tissu vivant devant être diagnostiqué, au vu des données d'élasticité et des données de viscosité qui servent de résultats de mesure du tissu vivant excité devant diagnostiqué, à l'aide des données d'élasticité et des données de viscosité pour chaque sorte de tissu vivant connu.

- 13. Système de diagnostic par ultrasons selon la revendication 12, dans lequel chaque sorte de tissu vivant connu est tracé sur un plan en deux dimensions ayant des axes représentant un module d'élasticité de cisaillement et un coefficient de viscosité de cisaillement.
- 14. Système de diagnostic par ultrasons selon l'une des

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revendication 11 ou 12, dans lequel le tissu vivant devant être diagnostiqué est un tissu vivant sur un coeur, un vaisseau sanguin, etc.

- 15. Système de diagnostic par ultrasons selon l'une des revendications 11 ou 12, dans lequel la petite zone locale (3h-31) servant d'unité pour mesurer une élasticité du tissu vivant devant être diagnostiqué est de plusieurs dizaines à plusieurs centaines de micromètres carrés.
- 16. Système de diagnostic par ultrasons selon l'une des revendications 11 ou 12, dans lequel l'unité d'identification qui se réfèrent aux données d'élasticité pour chaque <sup>15</sup> sorte de tissu vivant connu, les données étant extraites de la bibliothèque de données, et déterminant la sorte de tissu vivant connu ayant une distance minimale, pour chaque donnée d'élasticité servant de résultat de mesure dans chaque petite zone lo-<sup>20</sup> cale.
- 17. Système de diagnostic par ultrasons selon la revendication 16, dans lequel les moyens d'identification déterminent la dispersion et une valeur moyenne des d'élasticité à partir de l'histogramme de données d'élasticité de chaque sorte de tissu vivant connu, les données étant extraites de la bibliothèque de données, et les moyens d'identification réalisant une identification à l'aide de la méthode de décision 30 de Bayes.
- 18. Système de diagnostic par ultrasons selon l'une des revendications 11 ou 12, dans lequel l'unité d'identification de tissu (8) a des moyens de génération de figure chromatique électronique pour générer une figure chromatique électronique de façon à identifier en couleurs la sorte de tissu vivant devant être diagnostiqué, la sorte de tissu vivant étant identifiée pour chaque petite zone locale (3h-31).

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FIG. 2D



















FIG. 6













FIG. 11



FIG. 12





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# FIG. 13B







FIG. 14B







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FIG. 17

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### **REFERENCES CITED IN THE DESCRIPTION**

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# patsnap

专利名称(译)	超声和超声检查中的组织识别方法			
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优先权	2001249398 2001-08-20 JP			
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外部链接	<u>Espacenet</u>			

摘要(译)

在用于在超声诊断中识别活组织的方法中,通过使用超声测量每个局部 小区域的待诊断的活组织的弹性。每种已知活组织的弹性数据由每种活 组织的每个局部小区域中的弹性模量频率直方图组成,并在数据库中管 理。关于用作测量结果的弹性数据,通过使用每种已知活组织的弹性数 据来识别待诊断的活组织的种类。

 $\Delta h(t) = \int_{0}^{t} \{ v_{i}(t) - v_{i+1}(t) \} dt$