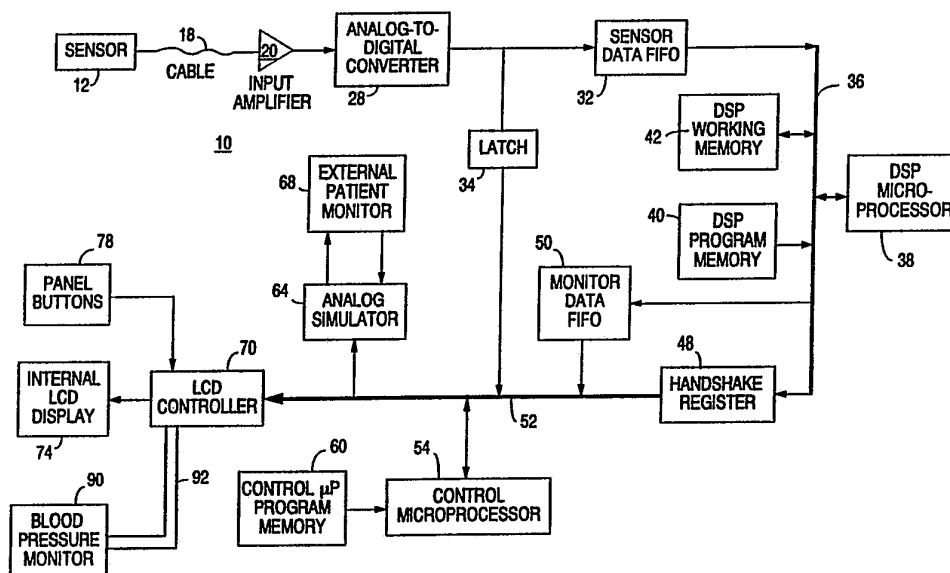




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(54) Title: METHOD AND APPARATUS FOR CONTINUOUSLY AND NON-INVASIVELY MEASURING THE BLOOD PRESSURE OF A PATIENT



## (57) Abstract

In the present invention, a method and an apparatus (10) for measuring the blood pressure of a patient continuously and non-invasively is disclosed. The blood pressure is determined by measuring the harmonic frequencies and displacements of an arterial wall of the patient using a sensor (12). The measurements are then calibrated against an absolute blood pressure determined using a blood pressure monitor (90) supplied by an operator at a particular time. The blood pressure is then the sum of the relative changes of the measurements and the absolute blood pressure.

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METHOD AND APPARATUS FOR CONTINUOUSLY AND NON-  
INVASIVELY MEASURING THE BLOOD PRESSURE OF A PATIENT

Technical Field

5 The present invention relates to a method and an  
apparatus for continuously and non-invasively measuring  
the blood pressure of a patient and, more particularly,  
the present invention relates to measuring the harmonic  
frequencies and displacement components of an arterial  
wall, and converting the measurements into a waveform  
10 signal emulating the signal from an invasive arterial  
sensor.

Background Of The Invention

Methods and apparatuses for the detection of the  
blood pressure of a patient are well-known in the art.  
15 One type of apparatus involves the use of an invasive  
arterial line. In an arterial line device, a sensor is  
inserted into the artery of a patient. The signal from  
the arterial sensor is supplied to a patient monitor  
which calculates, among other parameters, the blood  
20 pressure of a patient. The patient monitor is also  
capable of sending a signal to the sensor to determine  
if the sensor is electrically connected to the patient  
monitor. Although such apparatuses provide continuous  
and accurate blood pressure data, there are many  
25 disadvantages. First, there is a potential risk of  
infection. Further, the procedure is costly and  
consumes valuable health care takers' time, in that it  
is a surgical procedure involving the insertion of a  
catheter into a patient's artery. In addition, the  
30 patient experiences discomfort.

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In the area of non-invasive devices, one prior art teaches the automatic expansion of pressure cuffs and the measurement of the blood pressure based thereon. However, such devices can only be used intermittently. Prolonged and frequent use can lead to patient discomfort.

U. S. Patent 4,669,485 teaches using two cuffs to measure the blood pressure of a patient. The absolute blood pressure of the patient is initially measured. Thereafter, the second cuff is maintained at a low pressure, continuously, to monitor continuously the relative blood pressure of the patient. In this manner, the blood pressure of the patient can be continuously monitored. However, such a device still involves using blood pressure cuffs, which can be a source of discomfort to the patient and the device is subject to patient motion and as a consequence degraded results.

In column 4, lines 39 - column 5, line 3 of U.S. Patent 4,669,485, reference is made to yet another prior art non-invasive device which measured the arterial wall displacement and used those measurements to determine the relative blood pressure. However, the use of only the displacement measurement is subject to error.

Reference is also made to U.S. Patent No. 3,318,303. In that reference, the output of an external microphone sensor is used to determine the blood pressure of a patient. In particular, the so-called "Korotkoff" sound may be determined.

U.S. Patent No. 4,203,451 discloses that a data processor can be used to receive signals from the Carotid transducer.

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U.S. Patent No. 3,773,033 discloses the use of an arterial vibration sensor to monitor the performance of the heart and arteries during a succession of cardiac cycles. However, a pressure cuff also applies variable pressure to a specified artery being monitored by the arterial vibration sensor.

Heretofore, none of the prior art devices or references has suggested a non-invasive apparatus for continuously and accurately measuring the blood pressure of a patient using data obtained from the arterial wall, thereby providing for greater patient comfort. Further, none of the prior art devices has checked for the integrity of the sensor, apart from it being merely electrically connected to the monitor.

#### Summary Of The Invention

In the present invention, an instrument for continuously and non-invasively measuring the blood pressure of a patient is disclosed. The instrument has a non-invasive sensor for measuring continuously the frequencies and displacements of an arterial wall and for generating continuously a first signal in response thereto. The first signal is processed continuously to produce a continuous processed first signal. The instrument further receives a calibration signal which is indicative of the absolute blood pressure of the patient at a particular point in time. The processed first signal corresponding to the particular point in time is stored. Finally, the instrument compares continuously the continuous processed first signal to the stored processed first signal and generates continuously a signal indicative of the continuous blood pressure of the patient.

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### Brief Description Of The Drawings

Fig. 1 is a block level diagram of the apparatus of the present invention.

5 Fig. 2(a-d) are detailed schematic diagrams of various portions of the apparatus shown in Fig. 1.

Fig. 3(a-f) are schematic diagrams showing representative waveform signals processed and generated by the apparatus of the present invention.

### 10 Detailed Description Of The Drawings

Referring to Fig. 1, there is shown a block diagram of an apparatus 10 of the present invention. The apparatus 10 comprises a sensor assembly 12 which measures continuously, with time, the external  
15 displacement and the harmonic frequencies of the displacement caused by the lateral expansion of an arterial wall of a patient. The pattern of expansions and contractions is reflective of the changes in the pressure wave at this site in the arterial system. In  
20 one embodiment, the sensor assembly 12 has a piezo-electric sensing material, such as Kynar film 14. The Kynar film 14 generates a signal which is directly coupled to a sensing amplifier 16, located within the sensor assembly 12. The sensing amplifier 16  
25 transforms the impedance of the signal from a high value, on the order of several megohms of the sensing film 14, to a low value which can be transmitted over a cable 18 (typically on the order of 10 feet). The sensor assembly 12 is positioned at any site of a  
30 patient where the artery is close to the surface of the skin, such as over the radial artery.

The sensing film 14 can be excited by an excitation signal supplied from the control microprocessor 54 through the control microprocessor

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bus 52, through an amplifier 15 and through the cable 18 to a diode 13. The excitation signal causes the film 14 to mechanically transduce, i.e. flex. The excitation signal is applied momentarily (on the order of one microsecond), and is thereafter removed. Because the sensing film 14 will have been flexed by the excitation signal, upon removal of the excitation signal, the sensing film 14 will resonate and move in the opposite direction. This movement in the opposite direction causes the production of a response electrical signal which is supplied along the cable 18. The detection of this response signal indicates that the sensing film 14 is not only electrically connected and is adapted to transduce, but is also in fact electromechanically ready to transduce the arterial wall movements.

The sensor assembly 12 is then positioned to detect arterial wall motion. The signal detected by the sensor assembly 12, caused by the motion of the arterial wall, is then supplied to an input amplifier 20 which removes any DC component from the signal (Fig. 3a). The input amplifier 20 comprises an AC voltage amplifier and high pass filter 22 which receives the signal from the cable 18 and supplies an output thereof to a programmable gain DC voltage amplifier 24. The purpose of the programmable gain DC voltage amplifier 24 is to compensate for variations in the sensitivity of different sensors and physical variations among different patients. The output of the programmable gain DC voltage amplifier 24 is then supplied to a fixed gain DC amplifier and low pass filter 26. The DC amplifier and low pass filter 26 provides an output which is supplied to an analog-to-digital converter 28.

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The output of the input amplifier 20 is set to optimize the signal of the analog-to-digital converter 28.

5 The analog-to-digital converter 28 is a bipolar 10-bit A-to-D converter which receives a reference voltage 30 of 2.5 volts. The analog-to-digital converter 28 digitizes the input analog signal at the rate of approximately 250 hertz or one conversion each four milliseconds.

10 The output of the bipolar analog-to-digital converter 28 is then supplied to a first-in-first-out (FIFO) buffer 32. The data is stored in the FIFO buffer 32 until a predetermined number of samples are collected. In one embodiment, the apparatus 10 of the present invention waits until there are 16 samples in  
15 the FIFO buffer 32 before the data from the FIFO buffer 32 are read out and are supplied to the DSP microprocessor bus 36 and are then further processed.

Further, the eight most significant bits (MSB) of the ten bits from the analog-to-digital converter 28 of  
20 each sensor sample are stored in an 8-bit latch 34. The output of the 8-bit latch 34 is supplied to the control microprocessor bus 52 and can be supplied to an external or internal display screen 74. By outputting the relative signal strength of the sensor assembly 12  
25 and displaying it on the display 74, the apparatus 10 permits the operator to move the sensor assembly 12 to optimally place the sensor assembly 12 on the patient.

After the FIFO buffer 32 is filled with the predetermined number of digitized sensor samples, the  
30 digital signal processor (DSP) 38 is interrupted. The contents of the FIFO buffer 32 is then transferred to the DSP microprocessor bus 36 and into the DSP working memory 42. In one embodiment, the DSP working memory 42 comprises 16K by 16 bits of random access memory



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(RAM) 46 and two multiplexed memory address counters 44. One memory address counter addresses RAM 46 when writing data to the RAM 46. The other memory address counter addresses RAM 46 when reading data from RAM 46. Each counter can be set to automatically increment or decrement after a read or write operation is completed. This increases processing throughput, since the DSP microprocessor 38 does not have to address each operation. The memory address counters 44 supply 14 address lines to address the 16K by 16 RAM memory 46. The DSP microprocessor 38 is under the control of a program which is stored in a program memory 40. The DSP microprocessor 38 is a TMS-32010 made by Texas Instruments. The program memory 40 comprises 4K by 16 bits of PROM.

The program stored in the program memory 40 controls the DSP processor 38 to process the data from the FIFO buffer 32 to convert the sensor data into data values of a signal representative of the output blood pressure. A copy of the program is attached herewith as Exhibit A. The function of the program stored in the program memory 40 will be described hereinafter.

After the DSP microprocessor 38 has completed the processing of the sensor signal data from the FIFO buffer 32 and into a waveform signal data, the waveform signal data are passed to and stored in the monitor data FIFO buffer 50.

From the monitor data FIFO buffer 50, the computed points of the waveform signal may be directed to display on the display screen 74. The handshake register 48 links the DSP microprocessor 38 with the control microprocessor 54, by means of the DSP bus 36 and the control bus 52. Both data and control commands can be passed between the two processors by this

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register 48. Since the monitor data FIFO buffer 50 is loaded with and relieved of data at the same rate as the original sensor data FIFO buffer 32, the computed points which comprise the waveform are displayed at the rate of one data point for each four milliseconds.

The control microprocessor 54 also operates under a program which is stored in the program memory 60. A copy of that program is attached herewith as Exhibit B. The control microprocessor 54 is a 80C31 made by Intel Corporation.

The apparatus 10 of the present invention can also supply the waveform signal data from the DSP microprocessor 38 to an external patient monitor 68. When the apparatus 10 functions in this manner, it is acting as an arterial sensor emulator, i.e. to the patient monitor 68, the signal supplied thereto is no different than the signal generated by an arterial sensor. The data is supplied to an eight-bit multiplying digital-to-analog converter 64 which emulates a normal dome pressure transducer, which is conventionally used to convert invasive arterial line fluid pressure sensor to electrical signals for input to the patient monitor 68. The monitor 68 outputs its normal excitation voltage and, in turn, receives an input signal of the expected sensitivity expressed as microvolts output per volt of excitation per millimeter of mercury of pressure from the analog simulator 64.

The analog simulator 64 consists of an 8-bit multiplying digital-to-analog converter 66, which converts digital values received from the monitor data FIFO buffer 50 sent along the control bus 52, into an analog signal, required by the external patient monitor 68. The programmable amplifier/low pass filter 65 varies the output level of the analog signal in

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accordance with the input sensitivity of the particular patient monitor 68 being used. Thus, some of the commonly used patient monitors 68 require the following input sensitivities:

- 5 Hewlett-Packard: 5uv input/volt of excitation/ mm Hg.  
Marquette: 20uv input/volt of excitation/ mm Hg.

In the operation of the apparatus 10 of the present invention, the apparatus 10 is initially placed in a calibration mode. In this calibration mode, the  
10 operator identifies the patient monitor 68 with which the apparatus 10 is connected through the analog simulator 64. The selection of the particular patient monitor 68 sets the programmable amplifier 65 to the correct output voltages.

15 Thereafter, the apparatus 10 sends to the patient monitor 68 a signal that the patient monitor 68 expects during the venting of the arterial line dome that is normally connected to the arterial line port. When the operator confirms the completion of the above steps,  
20 the apparatus 10 then shifts into the next mode of operation, the acquire mode.

In the acquire mode, the apparatus 10 provides a check that the sensor assembly 12 is connected to the apparatus 10. As previously discussed, this includes  
25 supplying the excitation signal and the detection of the response signal. The detection of the response signal from the sensing film 14 indicates that the sensor assembly 12 is electrically connected and the sensing film 14 is electromechanically operational.  
30 The apparatus 10 acquires and displays the raw pulse vibration signal that the sensor assembly 12 detects for the purpose of positioning the sensor assembly 12 optimally. Upon command by the operator, the apparatus 10 acquires and stores the previous four-second segment

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of the pulse vibration signal for correlation with the starting blood pressure values.

The DSP microprocessor 38 then processes the signal corresponding to the four-second segment from the FIFO buffer 32. The DSP microprocessor 38 determines the characteristic components of the input signal in accordance with the following steps:

1. The input signal comprising approximately four (4) seconds of data at a sample frequency of 4 milliseconds is stored in the working memory 42 (Fig. 3a).
2. The input signal is fast fourier transformed at approximately .25 Hz. interval from approximately 0-250 Hz., thereby generating 1024 frequency values, with each frequency value having an amplitude value (Fig. 3d).
3. The frequency that corresponds to the maximum amplitude value is determined.
4. All the amplitudes of the spectrum data are normalized to the frequency found in step 3, with that amplitude as 100.
5. The normalized fast fourier transformed signal is stored in the working memory 42.

After the DSP microprocessor 38 has completed the processing of the sensor signal from the FIFO buffer 32 the apparatus 10 shifts to the BP input mode.

In the BP input mode, the apparatus 10 requests the operator to input the patient's absolute blood pressure values as measured by an independent blood pressure apparatus. The absolute blood pressure values may be manually determined by a health care worker and entered into the apparatus 10 via the panel buttons 78. Alternatively, the absolute blood pressure values may be supplied from a conventional, well known, automatic

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blood pressure monitor 90, such as a pressure cuff measuring instrument 90. The data from the blood pressure monitor 90 is supplied to the apparatus 10 along an RS-232 input line 92, to the LCD controller 5 70, just like the inputs from the panel buttons 78. Since the apparatus 10 of the present invention 10 requires correlation with absolute blood pressure value only during the initial stage, the automatic blood pressure monitor 90, even if it were of pressure cuff 10 design, would not pose the disadvantages of continual use.

The absolute blood pressure values correspond to the correlated, stored four-second segment sensed by the sensor assembly 12. When the operator confirms the 15 conclusion of the input blood pressure value, the DSP microprocessor 38 performs the following steps:

- a. The four-seconds of data stored in memory 42 (from step 1, above) are integrated (Fig. 3b).
- b. A first scale factor is calculated. The first 20 scale factor is chosen such that the maximum absolute amplitude of the waveform signal found in step a times the first scale factor would equal the pulse pressure (pulse pressure = systolic - diastolic) (Fig. 3c).
- c. An offset (equal to the diastolic) is added to the 25 waveform of step b (Fig. 3c).
- d. The processed waveform signal from step c is stored in the memory 42.

In sum, during the input mode, the DSP microprocessor 38 calculates the first scale factor and the offset. 30 The waveform generated by the apparatus 10 (from step c), after passing through the analog simulator 64, would be as if the waveform signal were generated by an invasive arterial sensor. Thus, the apparatus 10

emulates an invasive arterial sensor. The apparatus 10 then enters into the monitor mode.

In the monitoring mode, the apparatus 10 continuously receives the signal from the output of the sensor assembly 12 and continuously processes that signal via the input amplifier 20, digital-to-analog converter 28 and the sensor data FIFO buffer 32. The DSP microprocessor 38 continuously processes the digitized signal in the following manner.

- 6. The digitized signal from the FIFO buffer 32 is processed in accordance with steps 1-5 above.
- 7. The normalized frequency spectrum of the current signal from step 6 is subtracted from the normalized frequency spectrum of the initial signal, as stored in working memory 42 (Fig. 3e or 3f).
- 8. A comparison is made between the maximum frequency of the normalized frequency of the current signal and of the initial signal.
- 9. The change in the amount of the maximum frequency is noted and is supplied to the following look up table, and the corresponding second scale factor is determined:

|    | <u>Change of Maximum Frequency</u> | <u>Second Scale Factor</u> |
|----|------------------------------------|----------------------------|
| 25 | 50%                                | 85%                        |
|    | 67%                                | 90%                        |
|    | 83%                                | 95%                        |
|    | 100%                               | 100%                       |
|    | 133%                               | 105%                       |
| 30 | 167%                               | 110%                       |
|    | 200%                               | 115%                       |

- 10. A comparison is made between the frequency of the maximum amplitude of the current signal to the initial signal. For each shift in frequency (+ or -) of 1 Hz., the offset (determined from step c) is adjusted correspondingly to yield a change of 5

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mm Hg. Thus, if the frequency of the maximum amplitude of the current signal is increased 1 Hz. from the frequency of the maximum amplitude of the initial signal, the offset for the initial signal is increased by an amount equal to 5 mm Hg.

5

11. The data points from the FIFO buffer 32 are integrated.

12. The data points from step 11 are multiplied by the first scale factor and by the second scale factor, and the new offset (from step 10) is added to the resultant waveform signal.

10

The points that represent the waveform signal (step 12) are supplied to the monitor data FIFO buffer 50 and then to the analog simulator 64 or LCD controller 70 for display on the external patient monitor 68 or internal LCD display 74. The steps of 6-12 are repeated for each new four-second segment of data.

15

In another embodiment of the present invention, the health care worker can supply operating limits to the apparatus 10 by way of the input panel buttons 78. When the current signal, processed by the DSP microprocessor 38, is outside of the operating limits, an alarm can be sounded to alert the health care worker. Alternatively, the LCD controller 70 can initiate a control signal to the automatic blood pressure monitor 90 to re-initiate the apparatus 10.

20

25

From the foregoing, it can be seen that a highly accurate and efficient blood pressure measuring apparatus and method has been disclosed.

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WHAT IS CLAIMED IS:

1. An instrument for continuously and non-invasively measuring the blood pressure of a patient, said instrument comprising:

5 non-invasive sensor means for continuously measuring the frequencies and displacements of an arterial wall of said patient, and for generating continuously a first signal in response thereto;

10 means for processing continuously said first signal to produce a continuous processed first signal;

means for receiving a calibration signal indicative of the absolute blood pressure of said patient at a particular point in time;

15 means for storing said continuous processed first signal corresponding to said particular point in time; and

20 means for comparing continuously said continuous processed first signal to said stored processed first signal and for generating continuously a signal indicative of the continuous blood pressure of said patient.

25

2. The instrument of Claim 1, wherein said non-invasive sensor means comprises a piezo-electric sensor.

30 3. The instrument of Claim 1, wherein said processing means further comprises:

means for filtering continuously said first signal to produce a continuous filtered first signal;



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programmable gain means for amplifying continuously said continuous filtered first signal to produce a continuous amplified first signal;

5 digitizing means for digitizing continuously said continuous amplified first signal to produce a continuous digital first signal; and

10 fourier transform means for transforming said continuous digital first signal into said continuous processed first signal.

4. The instrument of Claim 3, wherein said comparing means further comprising:

15 means for subtracting said continuous processed first signal to said stored continuous processed first signal and for generating a scale factor and an offset in response thereto;

20 means for integrating said first signal to produce an integrated signal; and

means for adjusting said integrated signal by said scale factor and said offset to generate said signal indicative of the continuous blood pressure of said patient.

25

5. The instrument of Claim 4 further comprising:

means for storing said integrated signal corresponding to said particular point in time.

30

6. The instrument of Claim 5 wherein said subtracting means further comprising:

means for determining the change of the maximum frequency from said continuous processed

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first signal to said stored continuous processed first signal;

5 look up table means for converting said change of the maximum frequency into said scale factor;

means for determining the change of the frequency of the maximum amplitude of said integrated signal to said stored integrated signal; and

10 means for calculating said offset in response to said determining means.

7. A method for continuously and non-invasively measuring the blood pressure of a patient, said method comprising:

15 non-invasively measuring continuously the frequencies and displacements of an arterial wall of said patient and for generating a first signal in response thereto;

20 processing continuously said first signal to produce a processed first signal; calibrating said processed first signal at a particular point in time by:

25 receiving a calibration signal indicative of the absolute blood pressure of said patient at said particular point in time;

30 storing said processed first signal corresponding to said particular point in time;

comparing continuously said processed first signal to said stored processed first signal; and

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generating continuously a signal indicative of the continuous blood pressure of said patient based upon said comparison.

- 5           8. The method of Claim 7, wherein said processing step further comprising:  
            filtering continuously said first signal to produce a filtered first signal;  
            amplifying continuously said filtered first signal to produce an amplified first  
10           signal;  
            digitizing continuously said amplified first signal to produce a digital first signal; and  
            fourier transforming said digital first  
15           signal into said processed first signal.
9. The method of Claim 8, wherein said comparing step further comprising:  
20           subtracting said processed first signal to said stored processed first signal and generating a scale factor and an offset in response thereto; and  
            integrating said first signal to produce an  
25           integrated signal.
10. The method of Claim 9, wherein said generating step further comprising:  
            adjusting said integrated signal by said  
30           scale factor and said offset to generate said signal indicative of the continuous blood pressure of said patient.
11. The method of Claim 10 further comprising:

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storing said integrated signal corresponding to said particular point in time.

5           12. The method of Claim 11 wherein said  
subtracting step further comprising:  
            determining the change of the maximum  
            frequency from said processed first signal to said  
            stored processed first signal;  
            converting said change of the maximum  
10           frequency into said scale factor;  
            determining the change of the frequency of  
            the maximum amplitude of said integrated signal to  
            said stored integrated signal; and  
            calculating said offset in response to said  
15           determining step.

            13. A non-invasive arterial sensor emulator for  
            supplying a signal to a patient monitor indicative of  
            the continuous blood pressure of a patient, said  
20           emulator comprising:  
            non-invasive sensor means for  
            continuously measuring the frequencies and  
            displacements of an arterial wall of said  
            patient, and for generating continuously a  
25           first signal in response thereto;  
            means for processing continuously said  
            first signal to produce a processed first  
            signal;  
            means for receiving a calibration signal  
30           indicative of the absolute blood pressure of  
            said patient at a particular point in time;  
            means for storing said processed first  
            signal corresponding to said particular point  
            in time; and

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means for comparing continuously said processed first signal to said stored processed first signal and for generating said signal to said patient monitor.

5

14. The emulator of Claim 13, wherein said non-invasive sensor means comprises a piezo-electric sensor.

10

15. The emulator of Claim 13, wherein said processing means further comprises:

means for filtering continuously said first signal to produce a filtered first signal;

15

programmable gain means for amplifying continuously said filtered first signal to produce an amplified first signal;

20

digitizing means for digitizing continuously said amplified first signal to produce a digital first signal; and

fourier transform means for transforming said digital first signal into said processed first signal.

25

16. The emulator of Claim 15, wherein said comparing means further comprising:

means for subtracting said processed first signal to said stored processed first signal and for generating a scale factor and an offset in response thereto;

30

means for integrating said first signal to produce an integrated signal; and

means for adjusting said integrated signal by

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said scale factor and said offset to generate said signal for said patient monitor.

5 17. The emulator of Claim 16 further comprising:  
means for storing said integrated signal  
corresponding to said particular point in time.

10 18. The instrument of Claim 17 wherein said  
subtracting means further comprising:  
means for determining the change of the  
maximum frequency from said processed first signal  
to said stored processed first signal;  
look up table means for converting said  
change of the maximum frequency into said scale  
15 factor;  
means for determining the change of the  
frequency of the maximum amplitude of said  
integrated signal to said stored integrated  
signal; and  
20 means for calculating said offset in response  
to said determining means.

25 19. The emulator of Claim 1 further comprising:  
means for generating said calibration signal  
indicative of the absolute blood pressure of said  
patient; and  
means for controlling said automatic means to  
receive said calibration signal automatically.

30 20. A medical instrument comprising:  
an electromechanical sensing means for  
generating a signal in response to mechanical  
motion detected by said sensing means, and for

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moving in response to an electrical signal  
supplied thereto;

means for supplying an excitation signal to  
said sensing means to cause motion in said sensing  
means; and

5

means for detecting a response signal  
generated by said sensing means moving in response  
to said excitation signal supplied thereto,

10

wherein said detection of said response  
signal is indicative of the operational readiness  
of said sensing means.

## AMENDED CLAIMS

[received by the International Bureau  
on 30 April 1990 (30.04.90);  
original claims 1-20 replaced by amended claims 1-14 (8 pages)]

1. An instrument for non-invasively  
measuring the blood pressure of a patient, said  
instrument comprising:

non-invasive sensor means for measuring the  
5 frequencies and displacements of an arterial  
wall of said patient, and for generating a first  
signal in response thereto;

means for filtering said first signal to  
produce a filtered first signal;

10 programmable gain means for amplifying said  
filtered first signal to produce a amplified  
first signal;

digitizing means for digitizing said  
15 amplified first signal to produce a digital  
first signal;

fourier transform means for transforming  
said digital first signal into said processed  
first signal;

20 means for receiving a calibration signal  
indicative of the absolute blood pressure of  
said patient at a particular point in time;

means for storing said processed first  
signal corresponding to said particular point in  
time;

25 means for subtracting said processed first  
signal from said stored processed first signal



and for generating a scale factor and an offset  
in response thereto;

means for integrating said first signal  
from said sensor means to produce an integrated  
signal; and

means for adjusting said integrated signal  
by said scale factor and said offset to generate  
said signal indicative of the blood pressure of  
said patient.

2. The instrument of Claim 1, wherein said  
non-invasive sensor means comprises a piezo-electric  
sensor.

3. The instrument of Claim 1 further  
comprising:

means for storing said integrated signal  
corresponding to said particular point in time.

4. The instrument of Claim 3 wherein said  
subtracting means further comprising:

means for determining the change of the  
maximum frequency from said continuous processed  
first signal to said stored continuous processed  
first signal;

look up table means for converting said change of the maximum frequency into said scale factor;

5 means for determining the change of the frequency of the maximum amplitude of said integrated signal to said stored integrated signal; and

means for calculating said offset in response to said determining means.

10 5. A method for and non-invasively measuring the blood pressure of a patient, said method comprising:

15 non-invasively measuring the frequencies and displacements of an arterial wall of said patient and for generating a first signal in response thereto;

filtering said first signal from said measuring step to produce a filtered first signal;

20 amplifying said filtered first signal to produce an amplified first signal;

digitizing said amplified first signal to produce a digital first signal;

25 fourier transforming said digital first signal into said processed first signal;

calibrating said processed first signal at  
a particular point in time by:

receiving a calibration signal

indicative of the absolute blood pressure  
of said patient at said particular point in  
time;

storing said processed first signal  
corresponding to said particular point in  
time;

subtracting said processed first signal to  
said stored processed first signal and  
generating a scale factor and an offset in  
response thereto;

integrating said first signal from the  
measuring step to produce an integrated signal;  
and

generating a signal indicative of the blood  
pressure of said patient based upon said  
comparison.

6. The method of Claim 5, wherein said  
generating step further comprising:

adjusting said integrated signal by said  
scale factor and said offset to generate said  
signal indicative of the blood pressure of said  
patient.

7. The method of Claim 6 further comprising:  
storing said integrated signal  
corresponding to said particular point in time.

8. The method of Claim 7 wherein said  
5 subtracting step further comprising:

determining the change of the maximum  
frequency from said processed first signal to  
said stored processed first signal;

10 converting said change of the maximum  
frequency into said scale factor;

determining the change of the frequency of  
the maximum amplitude of said integrated signals  
to said stored integrated signal; and

15 calculating said offset in response to said  
step of determining the change of the frequency  
of the maximum amplitude.

9. A non-invasive arterial sensor emulator for  
supplying a patient monitor signal to a patient  
monitor indicative of the blood pressure of a  
20 patient, said emulator comprising:

non-invasive sensor means for measuring the  
frequencies and displacements of an arterial  
wall of said patient, and for generating a first  
signal in response thereto;

25 means for filtering said first signal to  
produce a filtered first signal;

programmable gain means for amplifying said filtered first signal to produce an amplified first signal;

5 digitizing means for digitizing said amplified first signal to produce a digital first signal;

fourier transform means for transforming said digital first signal into said processed first signal;

10 means for receiving a calibration signal indicative of the absolute blood pressure of said patient at a particular point in time;

means for storing said processed first signal corresponding to said particular point in time;

15 means for subtracting said processed first signal from said stored processed first signal and for generating a scale factor and an offset in response thereto;

20 means for integrating said first signal from said sensor means to produce an integrated signal; and

25 means for adjusting said integrated signal by said scale factor and said offset to generate said signal for said patient monitor.

10. The emulator of Claim 9, wherein said non-invasive sensor means comprises a piezo-electric sensor.

5 11. The emulator of Claim 9 further comprising:  
means for storing said integrated signal  
corresponding to said particular point in time.

12. The instrument of Claim 11 wherein said  
subtracting means further comprising:

10 means for determining the change of the  
maximum frequency from said processed first  
signal to said stored processed first signal;

look up table means for converting said  
change of the maximum frequency into said scale  
factor;

15 means for determining the change of the  
frequency of the maximum amplitude of said  
integrated signal to said stored integrated  
signal; and

20 means for calculating said offset in  
response to said determining means of the change  
of the frequency of the maximum amplitude.

13. The emulator of Claim 9 further comprising:  
means for generating said calibration  
signal indicative of the absolute blood pressure  
of said patient; and

25

means for controlling said automatic means  
to receive said calibration signal  
automatically.

14. A medical instrument comprising:

5 an electromechanical sensing means for  
generating a signal in response to mechanical  
motion detected by said sensing means, and for  
moving in response to an electrical signal  
supplied thereto;

10 means for supplying an excitation signal to  
said sensing means to cause motion in said  
sensing means; and

15 means for detecting a response signal  
generated by said sensing means moving in  
response to said excitation signal supplied  
thereto,

wherein said detection of said response  
signal is indicative of the operational  
readiness of said sensing means.

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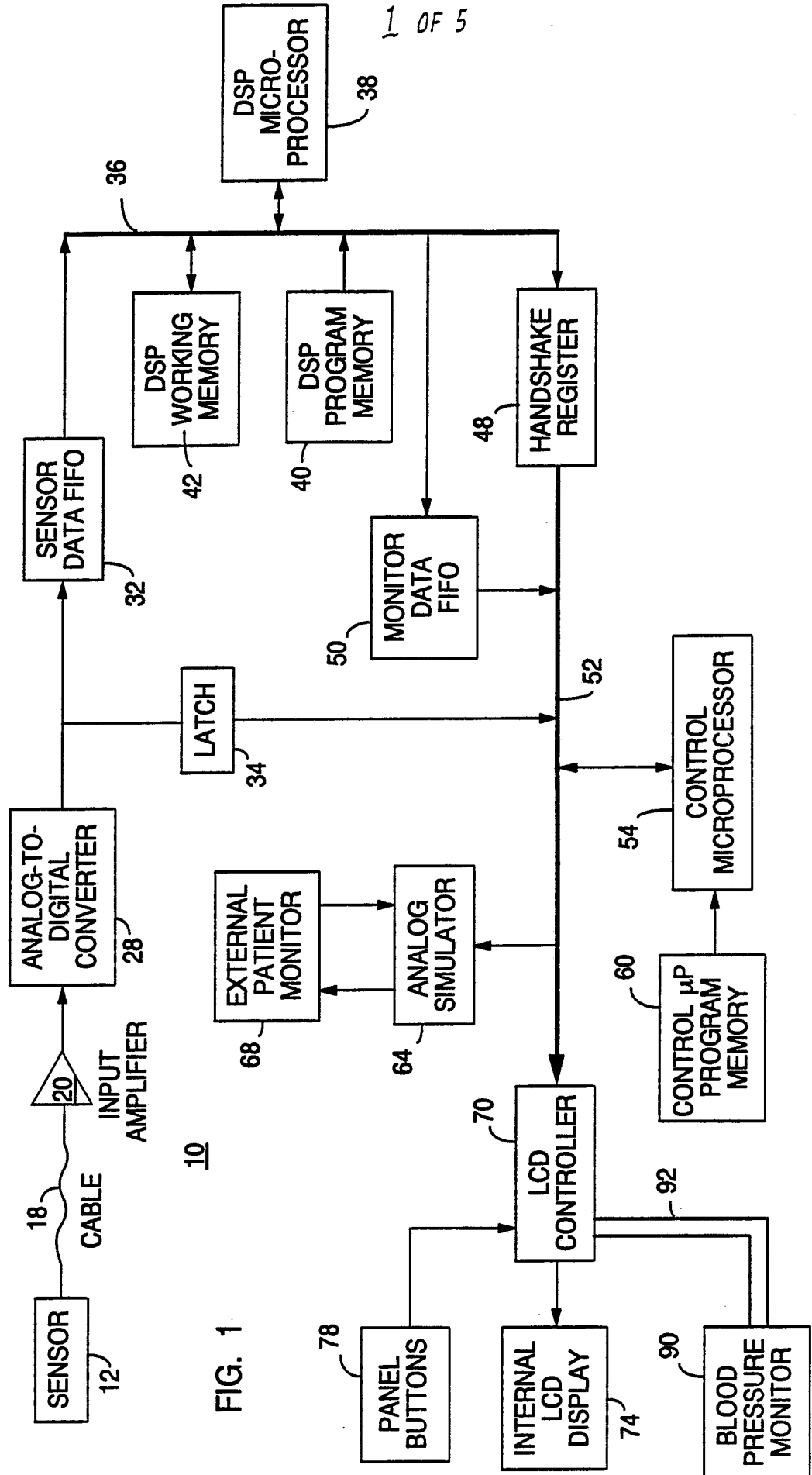


FIG. 1



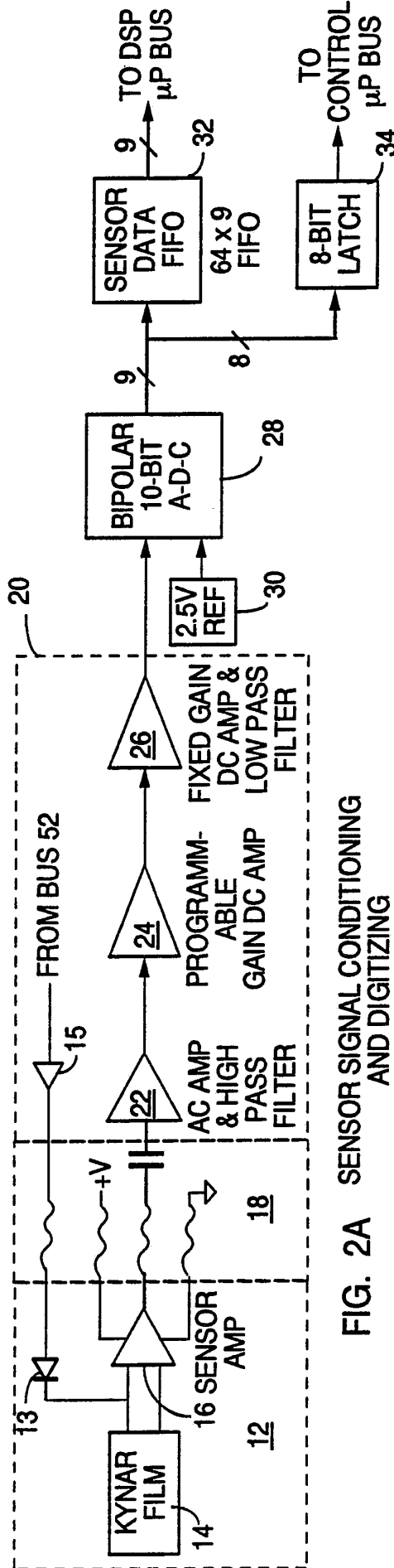


FIG. 2A SENSOR SIGNAL CONDITIONING AND DIGITIZING

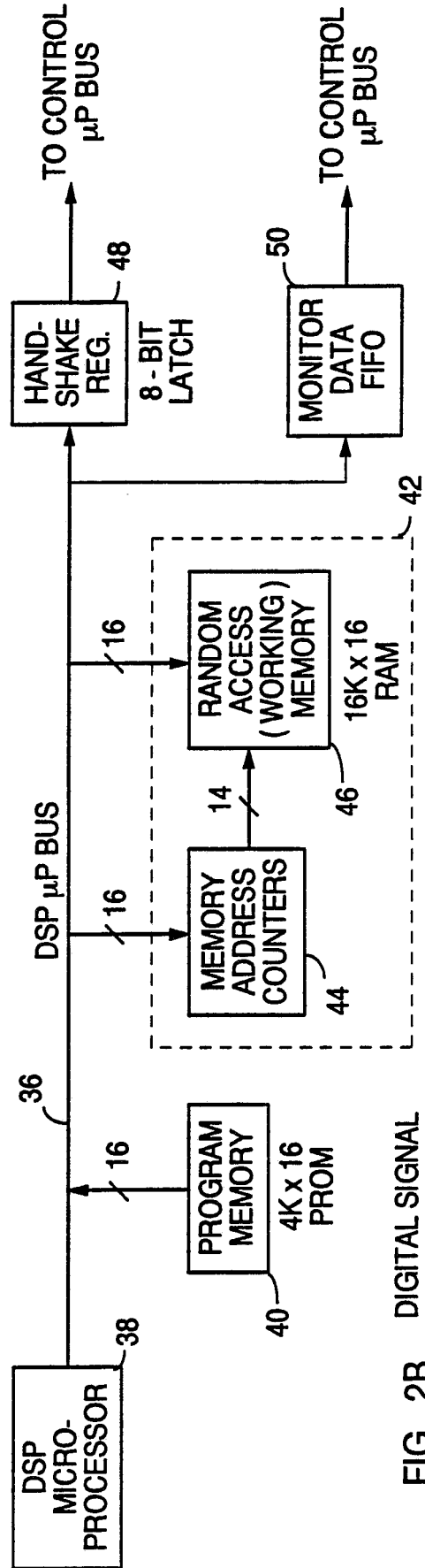


FIG. 2B DIGITAL SIGNAL MICROPROCESSOR

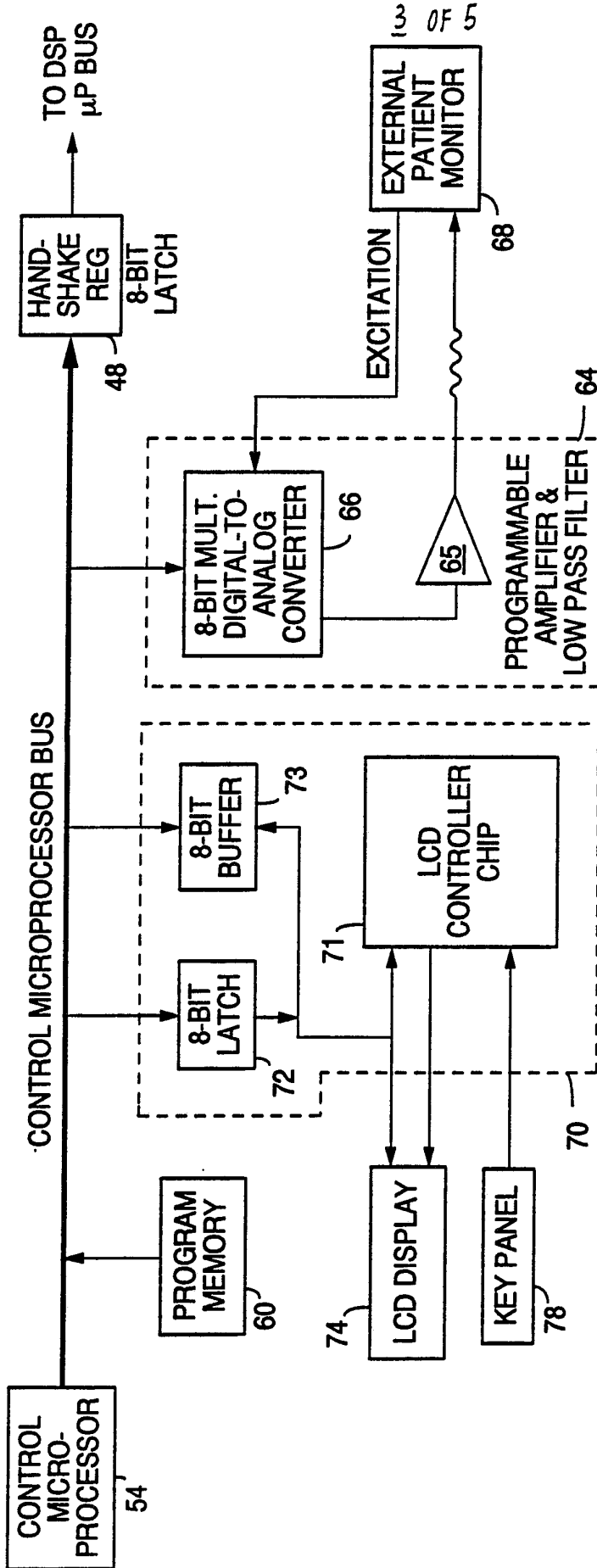


FIG. 2C CONTROL MICROPROCESSOR

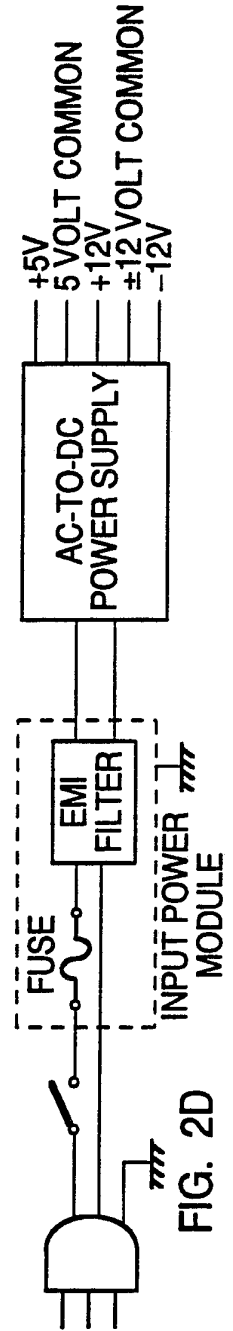
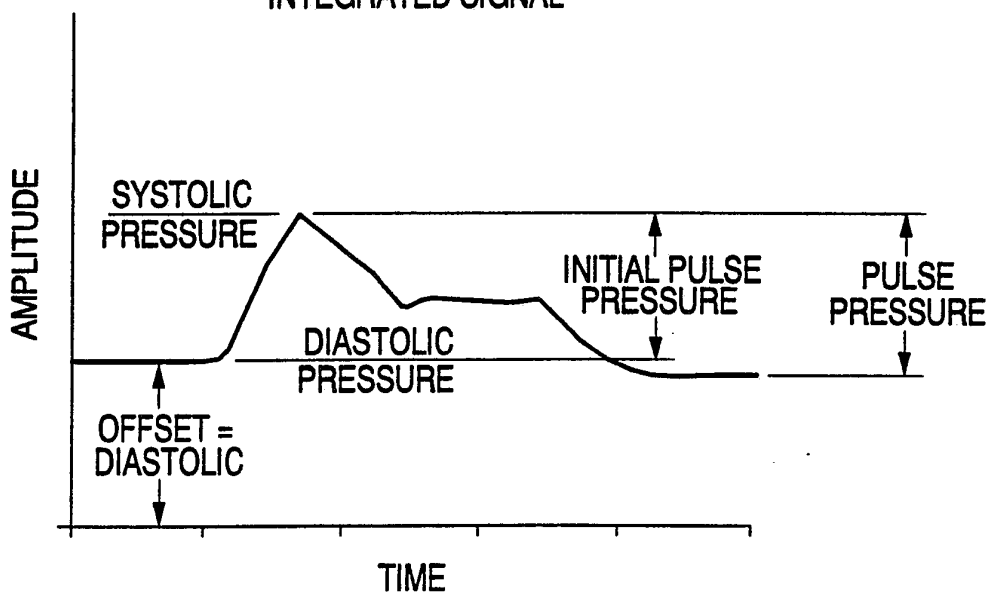
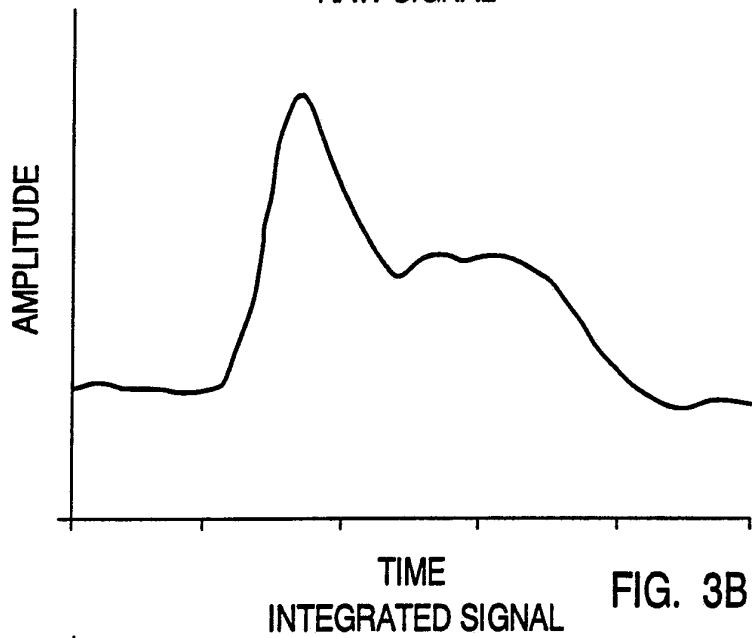
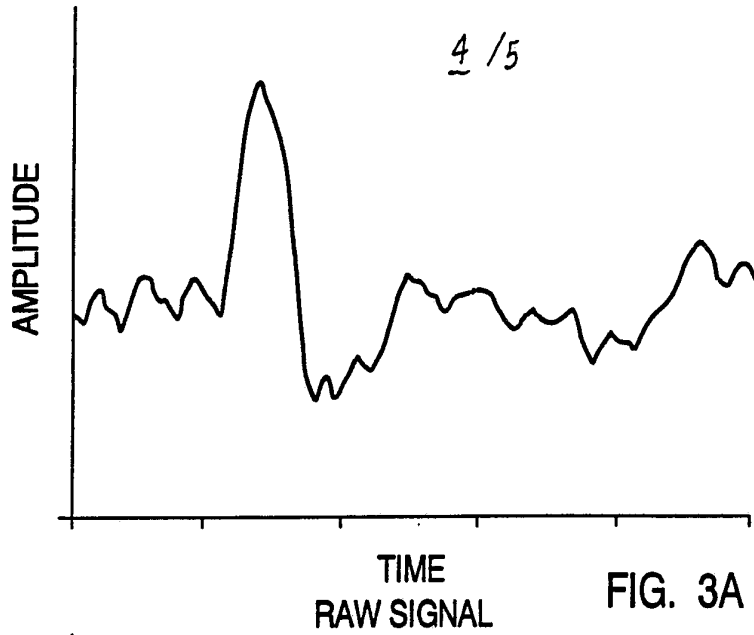


FIG. 2D

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INTEGRATED SIGNAL ADJUSTED FOR INITIAL SECOND SCALE FACTOR AND INITIAL OFFSET FIG. 3C

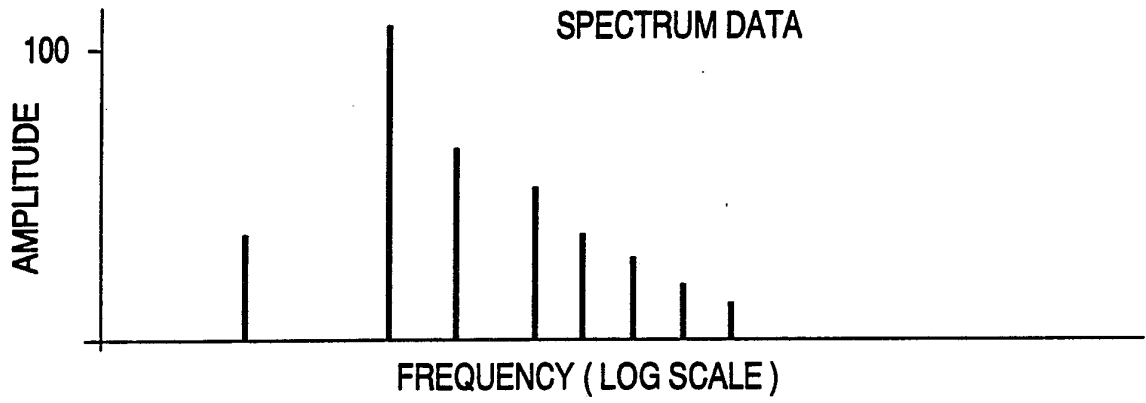


FIG. 3D

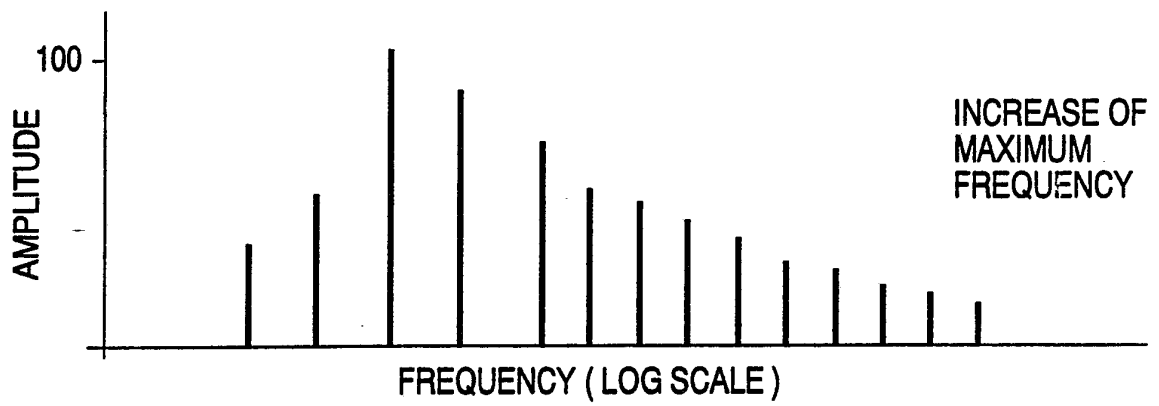


FIG. 3E

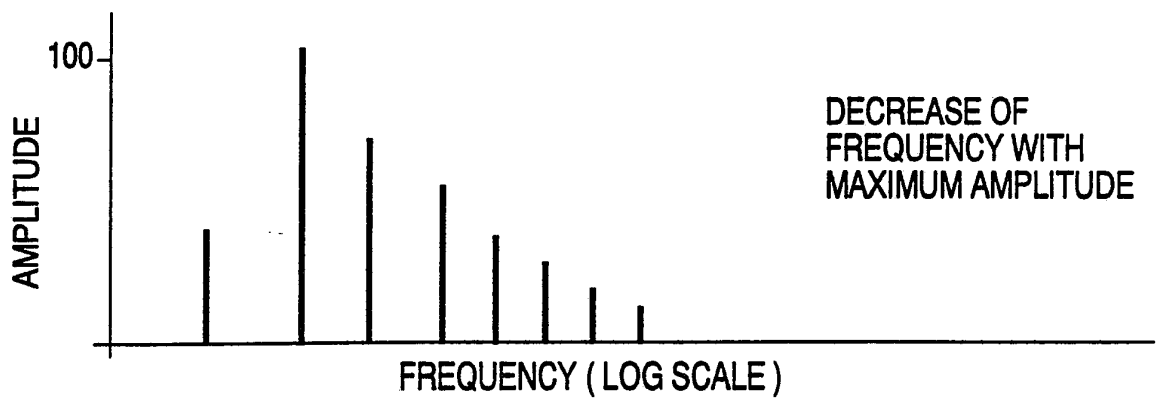
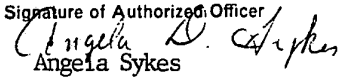


FIG. 3F

# INTERNATIONAL SEARCH REPORT

International Application No. PCT/US89/05216

|  |  |                                     |
|--|--|-------------------------------------|
| <b>I. CLASSIFICATION OF SUBJECT MATTER</b> (if several classification symbols apply, indicate all) <sup>6</sup>  |  |                                     |
| According to International Patent Classification (IPC) or to both National Classification and IPC  |  |                                     |
| IPC (5): A61B 5/02   |  |                                     |
| U.S. CL: 128/677   |  |                                     |
| <b>II. FIELDS SEARCHED</b>   |  |                                     |
| Minimum Documentation Searched <sup>7</sup>  |  |                                     |
| Classification System  | Classification Symbols   |                                     |
| U.S.   | 128/672, 677, 678, 679, 680, 681, 682, 683, 684, 685, 686, 687, 688, 689, 690                                  |                                     |
| Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched <sup>8</sup>   |  |                                     |
| <b>III. DOCUMENTS CONSIDERED TO BE RELEVANT</b> <sup>9</sup>   |  |                                     |
| Category *   | Citation of Document, <sup>11</sup> with indication, where appropriate, of the relevant passages <sup>12</sup> | Relevant to Claim No. <sup>13</sup> |
| Y  | US,A, 4,669,485 (RUSSELL), 02 JUNE 1987.<br>See the entire document.   | 1-3,7-8<br>13-15, 19-20             |
| Y  | US,A, 4,423,738 (NEWGARD), 03 JANUARY 1984.<br>See the entire document.  | 2, 14                               |
| A,P  | US,A, 4,802,488 (ECKERLE), 07 FEBRUARY 1989.<br>See the entire document.                                       | 1-19                                |
| A  | US,A, 4,562,843 (DJORDJEVICH), 07 JANUARY 1986.<br>See the entire document.                                    | 1-19                                |
| <p>* Special categories of cited documents: <sup>10</sup></p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"&amp;" document member of the same patent family</p> |  |                                     |
| <b>IV. CERTIFICATION</b>   |  |                                     |
| Date of the Actual Completion of the International Search  | Date of Mailing of this International Search Report  |                                     |
| 14 January 1990  | 01 MAR 1990  |                                     |
| International Searching Authority  | Signature of Authorized Officer  |                                     |
| ISA/US   | <br>Angela Sykes           |                                     |