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Hacker et al.

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[45] **Date of Patent:** **Aug. 8, 2000**

[54] **CASSETTE FOR MEASURING PARAMETERS OF BLOOD**

4,824,789 4/1989 Yafuso et al. .
4,830,013 5/1989 Maxwell .

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(List continued on next page.)

[73] Assignee: **Terumo Cardiovascular Systems Corporation**, Somerset, N.J.

FOREIGN PATENT DOCUMENTS

[21] Appl. No.: **09/031,415**

376 168 7/1990 European Pat. Off. .
380 664 8/1990 European Pat. Off. .
399227 11/1990 European Pat. Off. .
568 380 11/1993 European Pat. Off. .
575 712 12/1993 European Pat. Off. .
585212 3/1994 European Pat. Off. .
0 705 610 A1 10/1996 European Pat. Off. .
26 30 606 12/1978 Germany .
WO 87/04914 8/1987 WIPO .
WO 92/21281 12/1992 WIPO .
WO 93/06775 4/1993 WIPO .
WO 96/07268 8/1995 WIPO .
WO 96/10747 4/1996 WIPO .
WO 96/22730 8/1996 WIPO .

[22] Filed: **Feb. 26, 1998**

Related U.S. Application Data

[63] Continuation-in-part of application No. 08/810,954, Feb. 27, 1997, abandoned.

[51] **Int. Cl.**⁷ **A61B 5/00; G01N 33/48**

[52] **U.S. Cl.** **600/322; 356/39; 422/68.1; 436/68**

[58] **Field of Search** **600/310, 322; 356/39, 244, 246; 422/44, 68.1, 55, 102, 104; 436/63, 66, 68, 172**

OTHER PUBLICATIONS

P. 3-24 and 31-44, Operator's Manual, CDI Blood Gas Monitoring System 400, 3M Health Care (undated).

"Sensor Module" Research Disclosure, vol. 388, Aug. 1996, GB, pp. 520-525, XP000635453.

[56] **References Cited**

U.S. PATENT DOCUMENTS

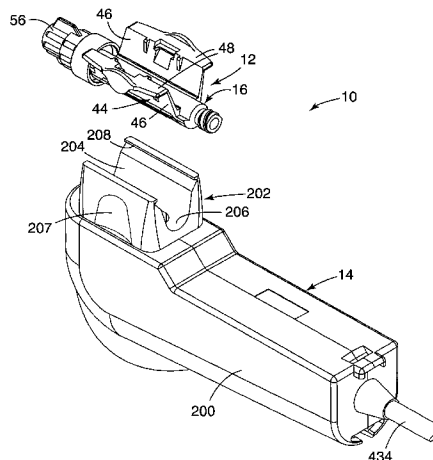
Re. 31,879 5/1985 Lubbers et al. .
D. 326,718 6/1992 Maxwell et al. .
2,927,582 3/1960 Berkman et al. .
3,639,829 2/1972 Harnoncourt .
3,639,830 2/1972 Harnoncourt .
3,718,568 2/1973 Neuwelt 204/195
3,799,672 3/1974 Vurek 356/41
3,826,730 7/1974 Watanabe et al. 204/195
4,041,932 8/1977 Fostick .
4,338,174 7/1982 Tamura 204/195
4,401,547 8/1983 Schinkmann et al. .
4,447,150 5/1984 Heinemann 356/41
4,469,659 9/1984 Carson et al. .
4,557,900 12/1985 Heitzmann .
4,640,820 2/1987 Cooper .
4,640,826 2/1987 Williams et al. .
4,745,279 5/1988 Karkar et al. 250/343
4,786,474 11/1988 Cooper .
4,798,738 1/1989 Yafuso et al. .

Primary Examiner—Eric F. Winakur
Attorney, Agent, or Firm—Burns, Doane, Swecker & Mathis, LLP

[57] **ABSTRACT**

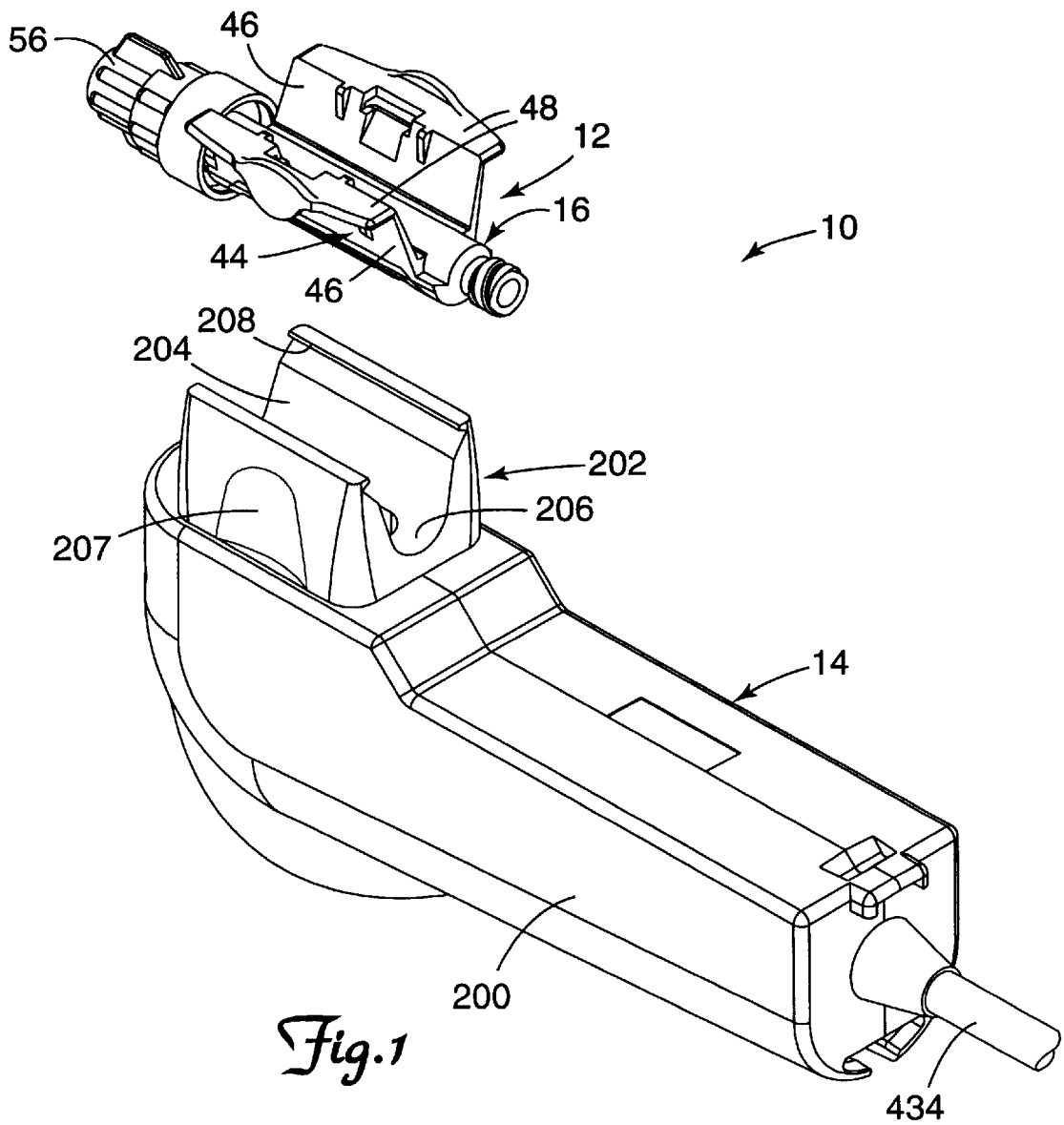
A cassette useful for measuring one or more parameters of blood in, for example, a cardiopulmonary bypass circuit has a convex coupling for releasable connection with a mating, concave coupling of a measuring device. The coupling of the cassette includes flexible leg portions that can be moved toward each other under the influence of finger pressure to detach the cassette from the measuring device. Optionally, the cassette includes a body carrying one or more sensors, and also includes a casing that can be detached from the body when desired. A shunt circuit arrangement enables the cassette to be connected to the bypass circuit either before or during a surgical procedure.

67 Claims, 28 Drawing Sheets



U.S. PATENT DOCUMENTS

4,849,172	7/1989	Yafuso et al. .	5,171,029	12/1992	Maxwell et al. .
4,867,919	9/1989	Yafuso et al. .	5,175,016	12/1992	Yafuso et al. .
4,886,338	12/1989	Yafuso et al. .	5,176,882	1/1993	Gray et al. .
4,919,891	4/1990	Yafuso et al. .	5,272,088	12/1993	Morlotti .
4,938,218	7/1990	Goodman et al. .	5,278,072	1/1994	Wall et al. .
4,951,669	8/1990	Maxwell et al. .	5,289,255	2/1994	Mullin et al. .
4,989,606	2/1991	Gehrich et al. .	5,296,381	3/1994	Yafuso et al. .
4,999,306	3/1991	Yafuso et al. .	5,304,492	4/1994	Klinkhammer .
5,006,314	4/1991	Gourley et al. .	5,308,320	5/1994	Safar et al. 604/4
5,043,285	8/1991	Surgi .	5,308,321	5/1994	Castro 604/74
5,048,525	9/1991	Maxwell .	5,348,706	9/1994	Abul-Haj et al. .
5,053,520	10/1991	Bieniarz et al. .	5,403,746	4/1995	Bentsen et al. .
5,057,278	10/1991	Maxwell et al. .	5,409,666	4/1995	Nagel et al. .
5,081,041	1/1992	Yafuso et al. .	5,453,248	9/1995	Olstein .
5,081,042	1/1992	Yafuso et al. .	5,462,879	10/1995	Bentsen .
5,094,820	3/1992	Maxwell et al. .	5,462,880	10/1995	Kane et al. .
5,104,623	4/1992	Miller .	5,474,743	12/1995	Trend et al. .
5,127,077	6/1992	Iyer et al. .	5,480,723	1/1996	Klainer et al. .
5,128,019	7/1992	Karpf et al. .	5,498,549	3/1996	Nagel et al. .
5,132,057	7/1992	Tomisaka et al. .	5,508,509	4/1996	Yafuso et al. .
5,136,033	8/1992	Masilamani et al. .	5,518,694	5/1996	Bentsen .
5,162,525	11/1992	Masilamani et al. .	5,591,400	1/1997	Dektar et al. .



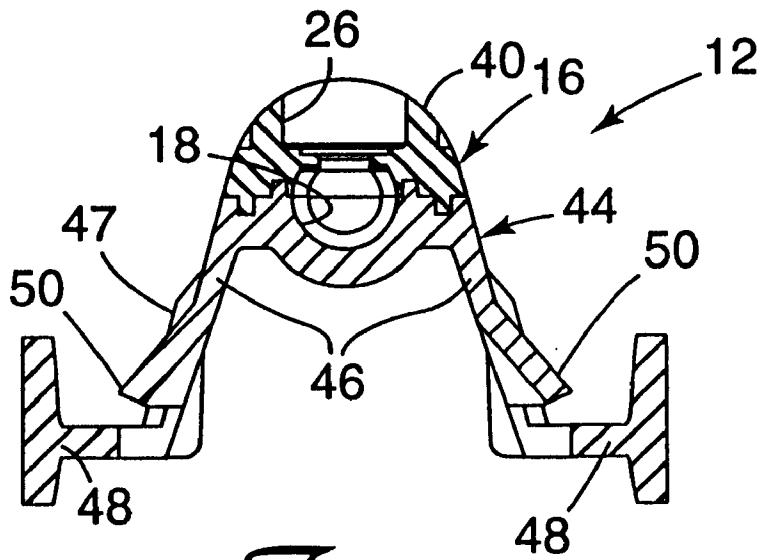


Fig. 2

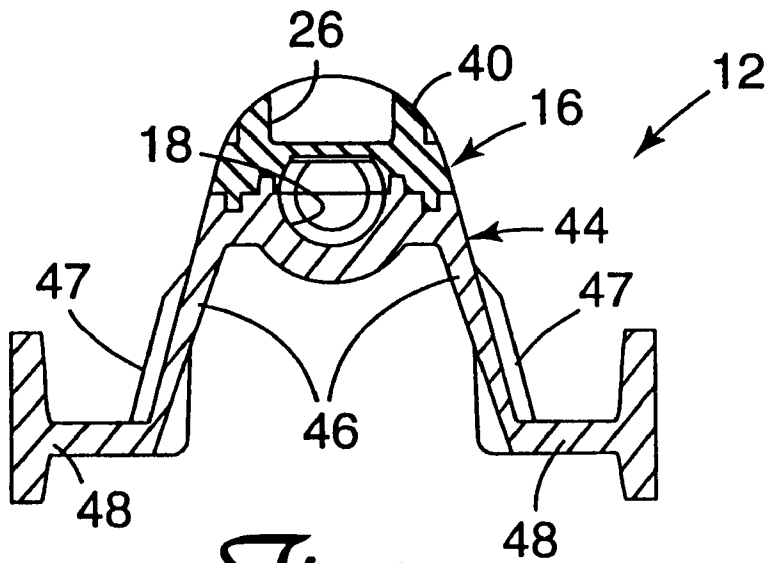


Fig. 3

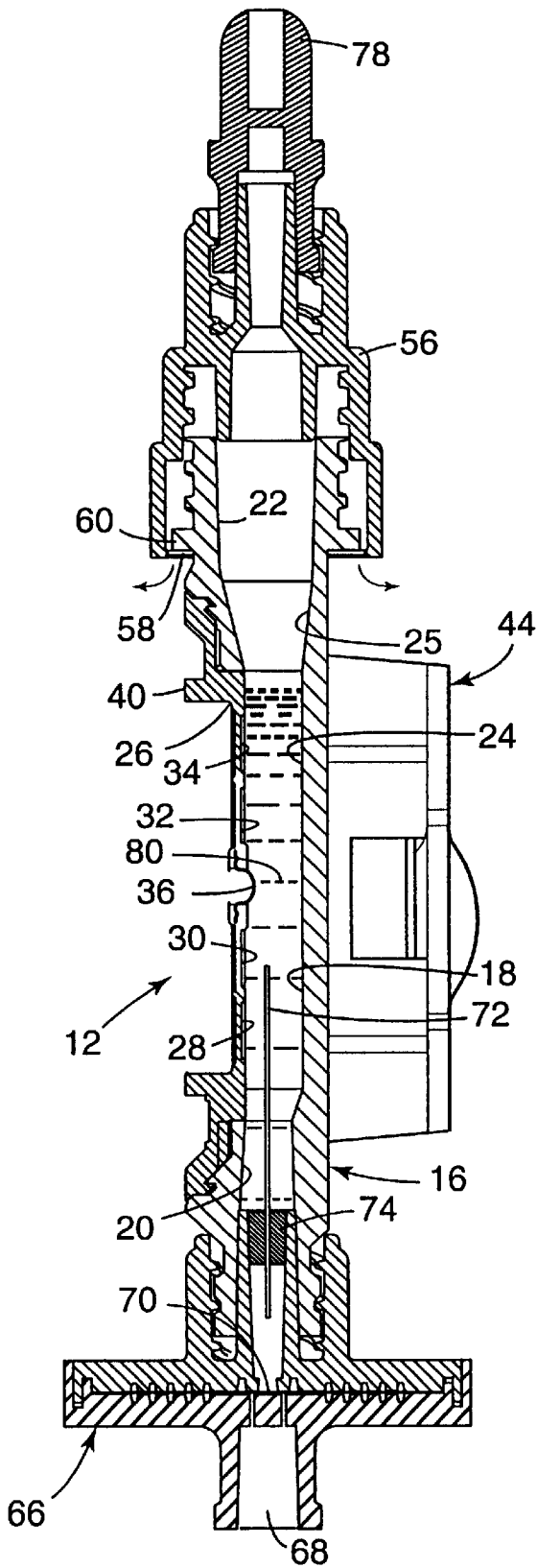


Fig. 6

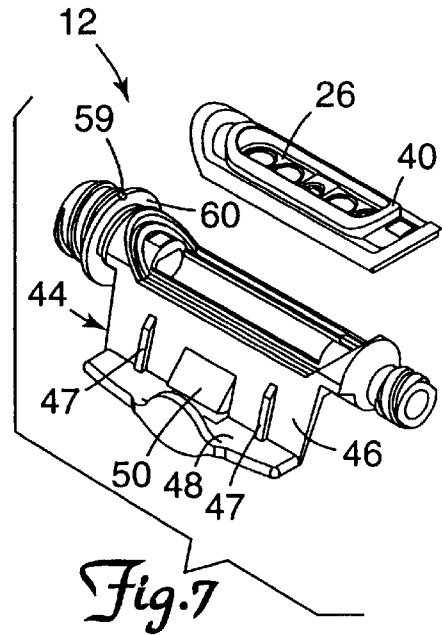


Fig. 7

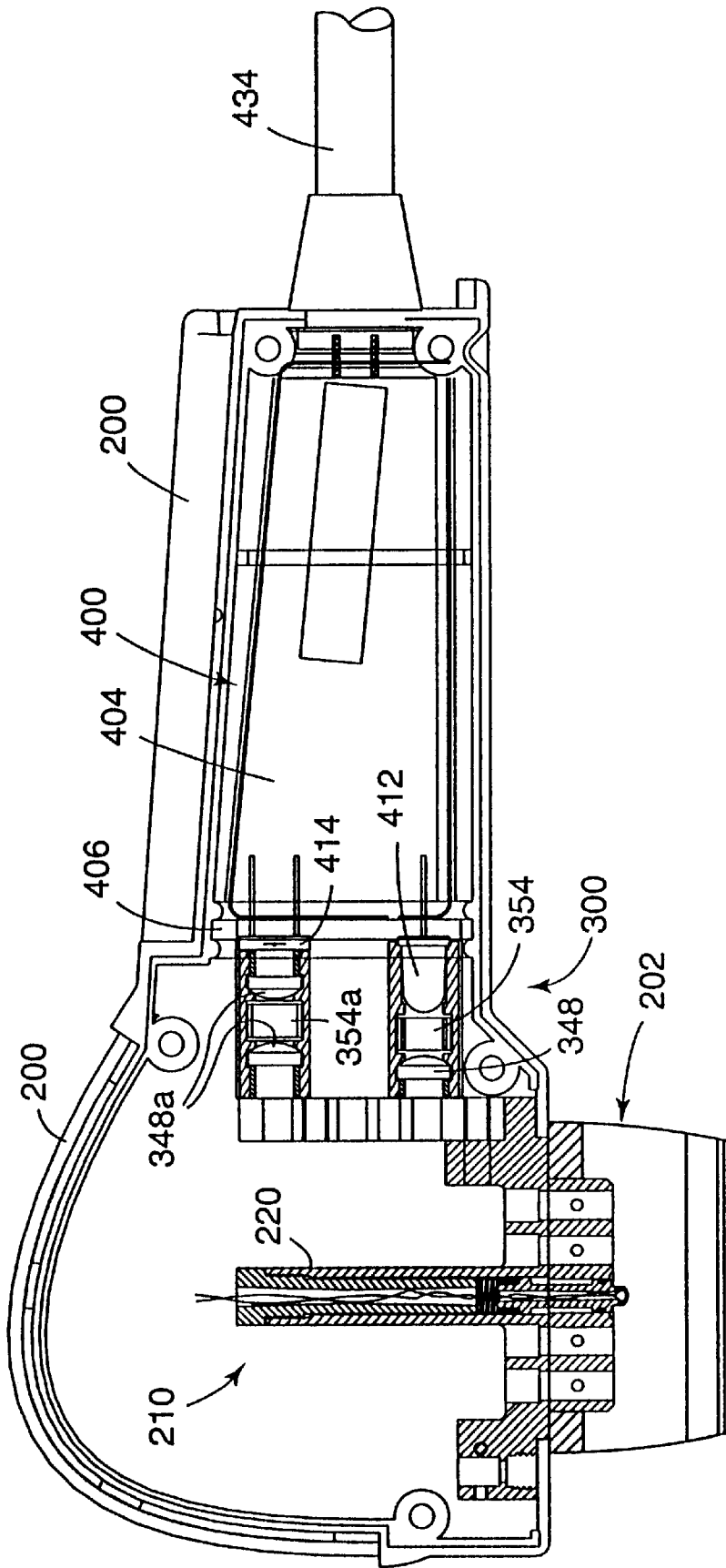


Fig. 8

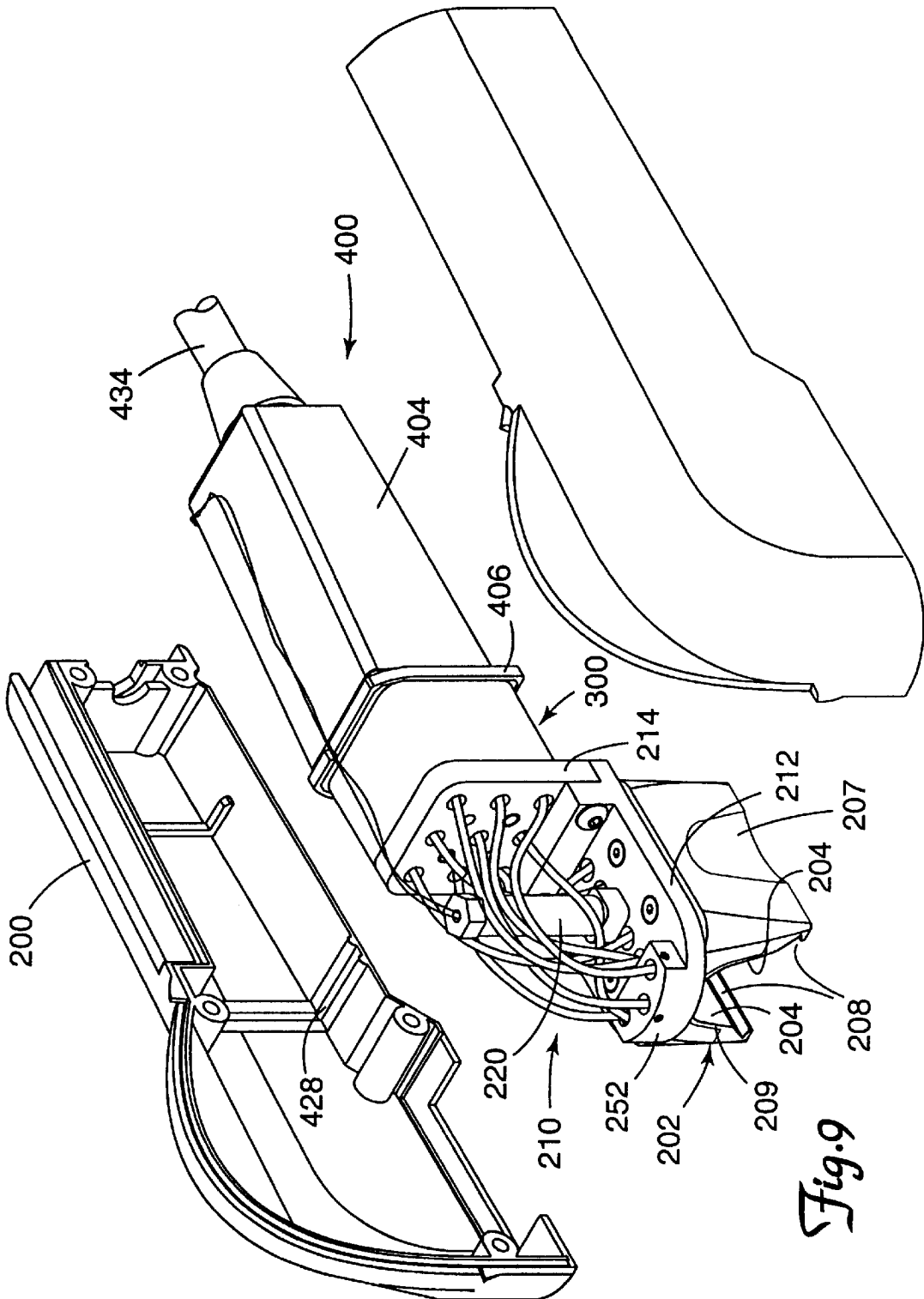
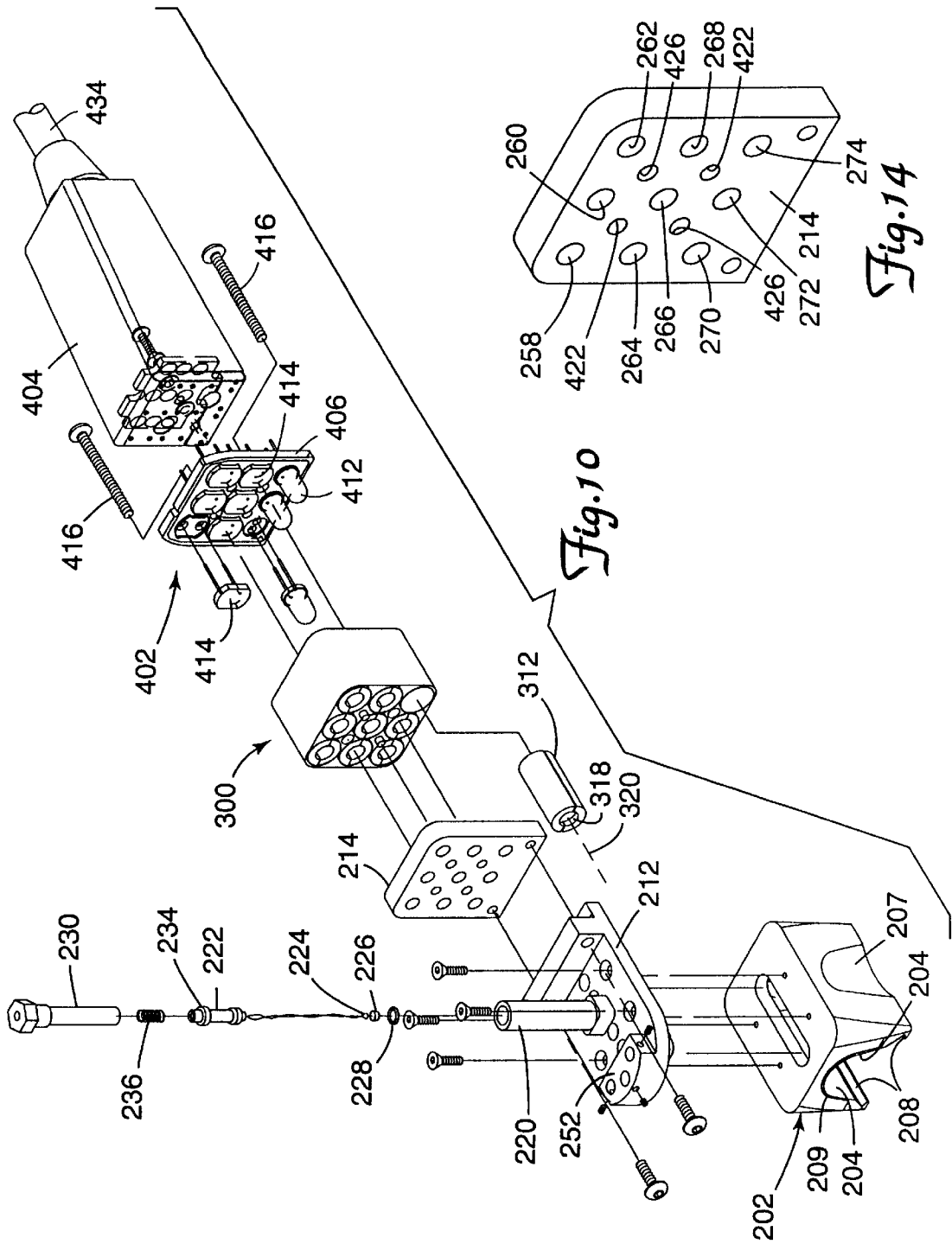


Fig. 9



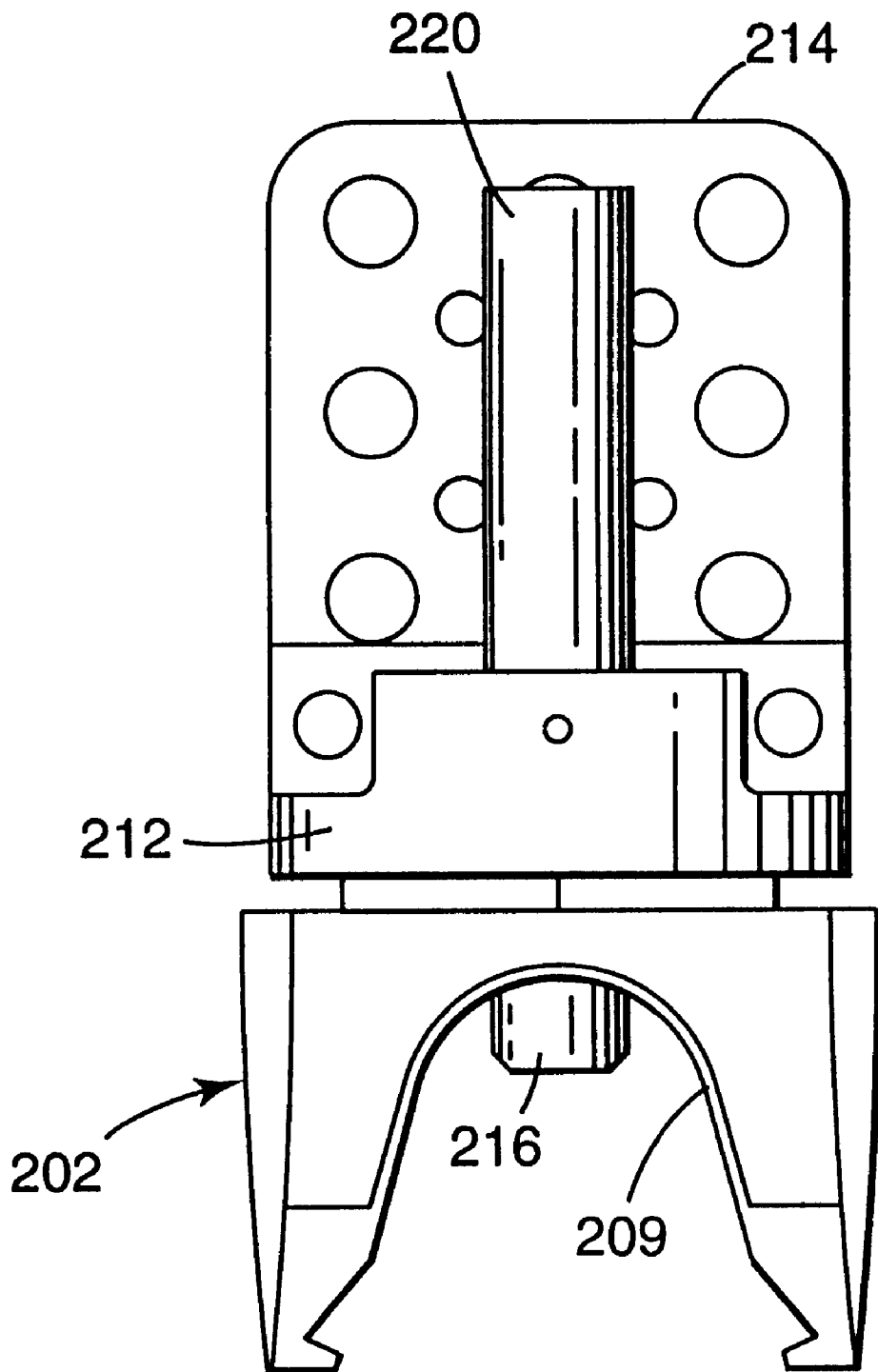


Fig. 11

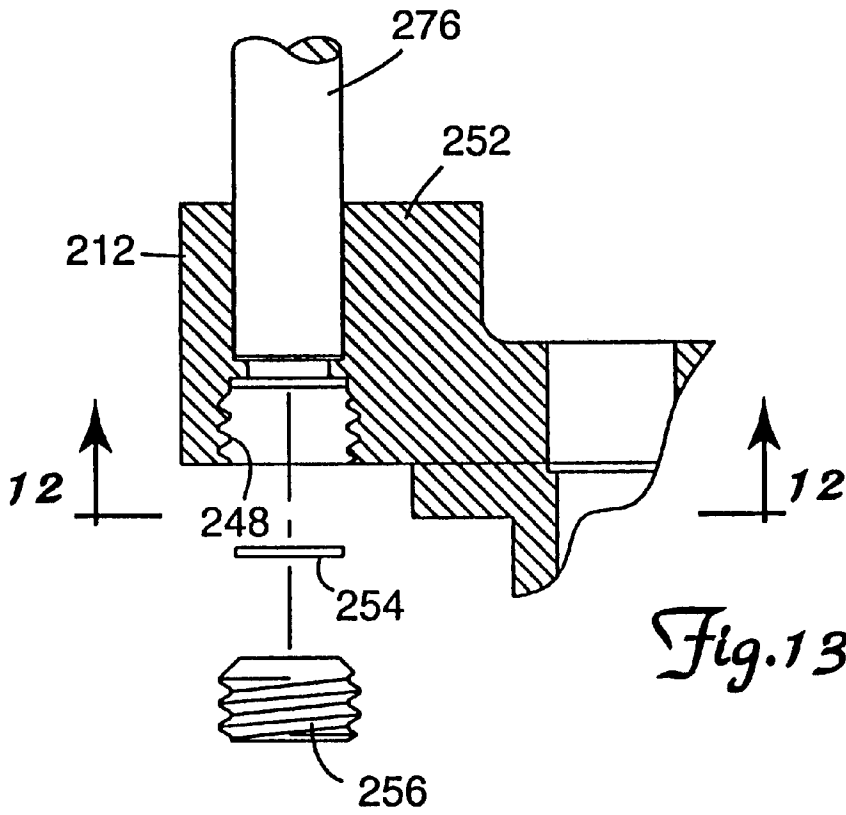


Fig. 13

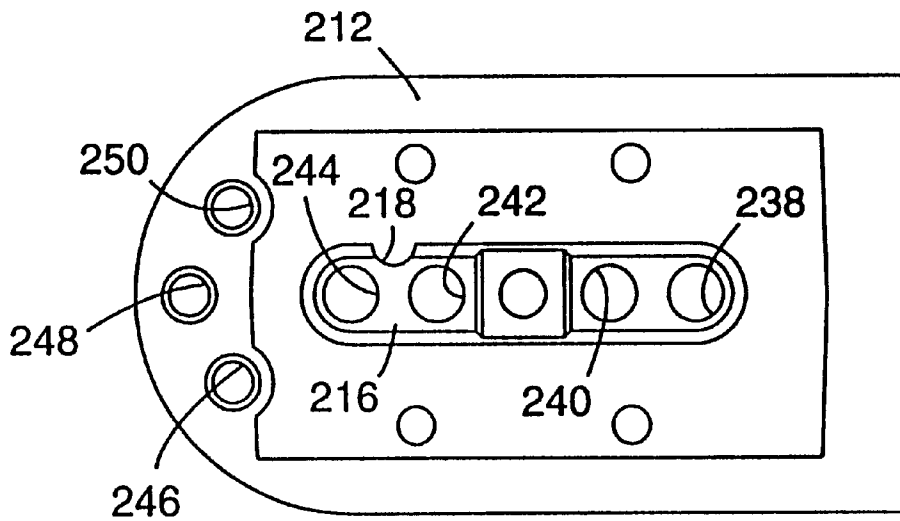


Fig. 12

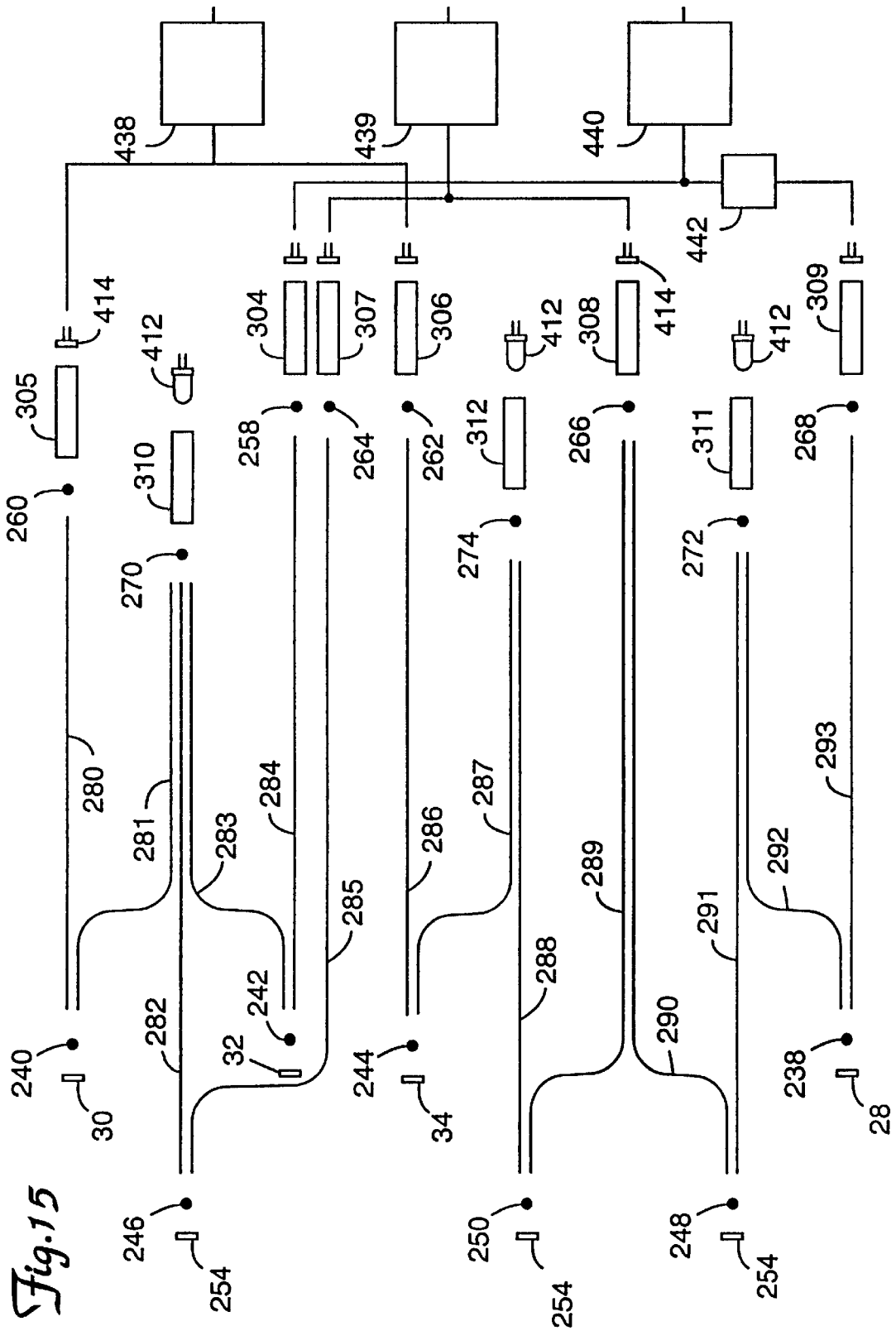


Fig. 15

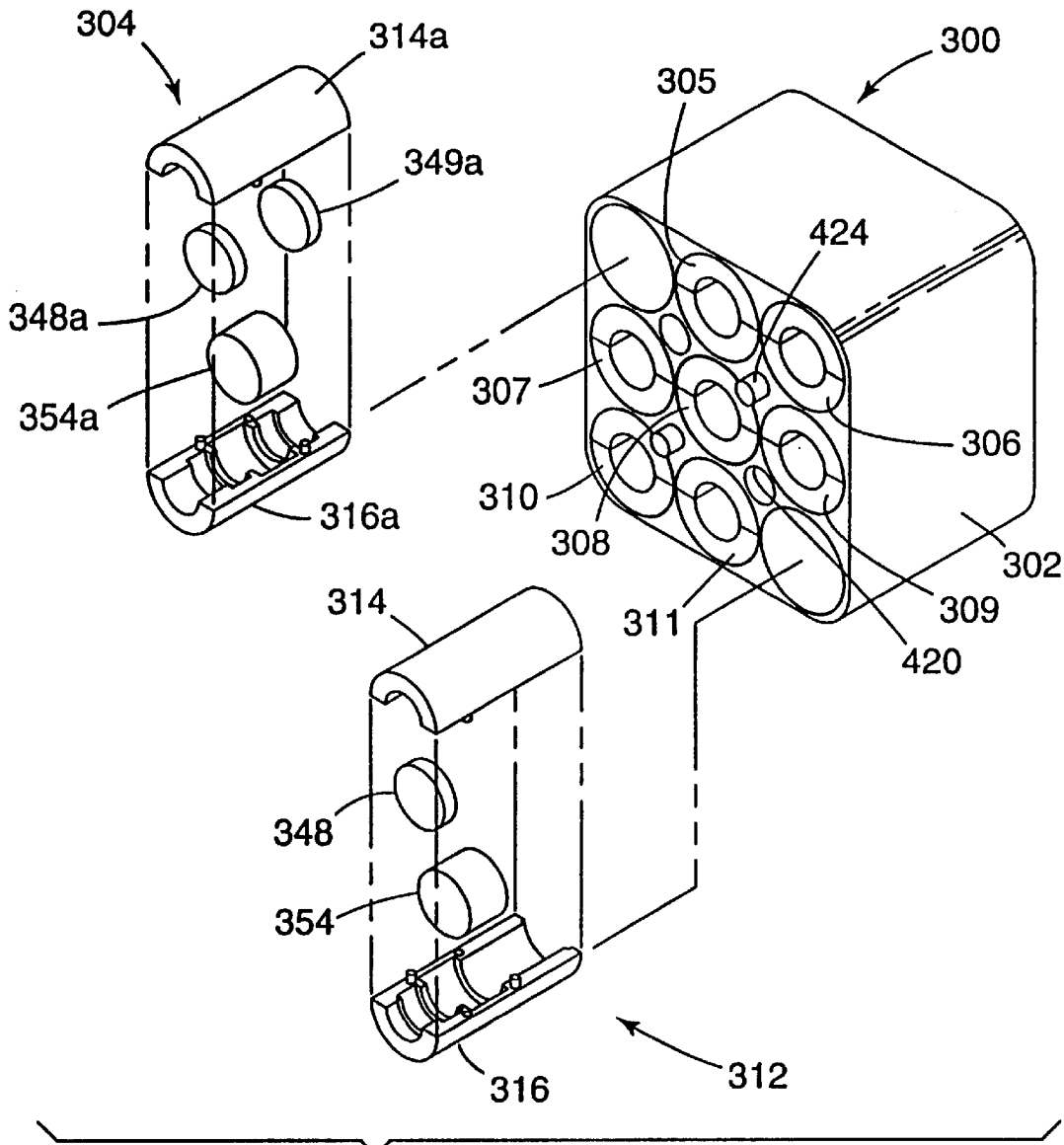


Fig. 16

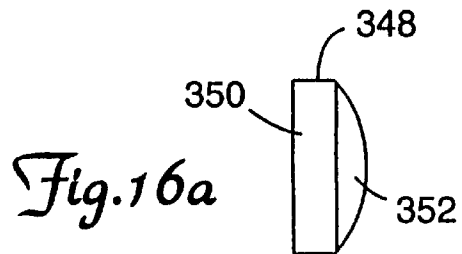
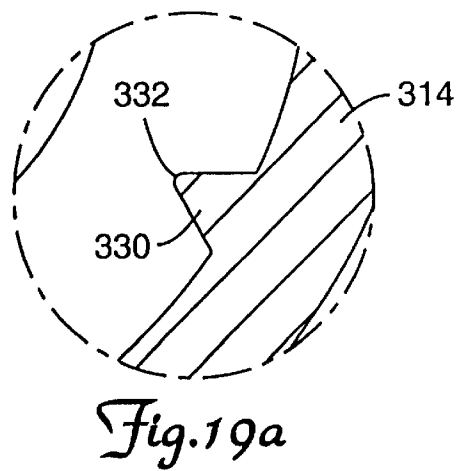
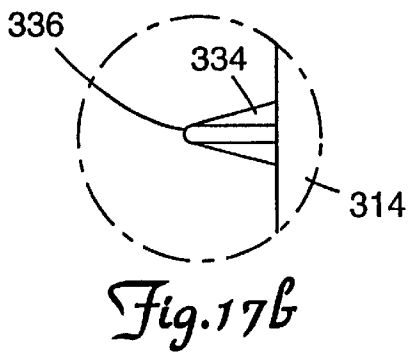
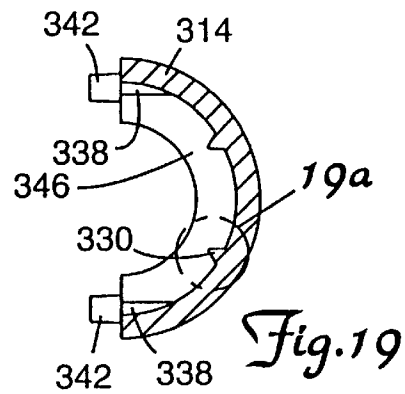
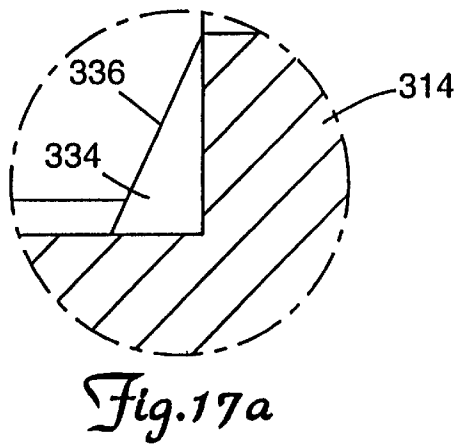
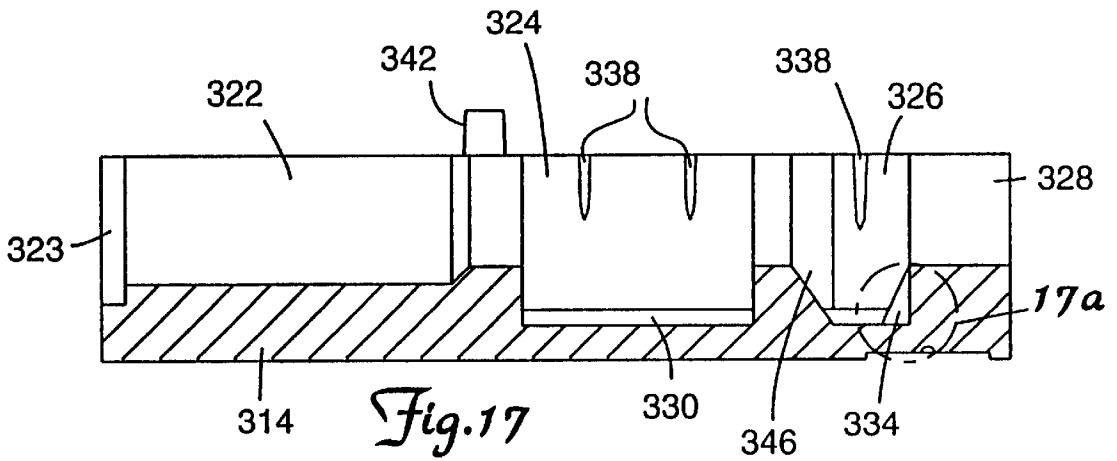


Fig. 16a



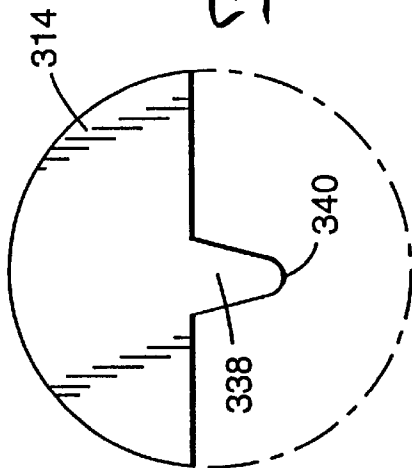


Fig. 17c

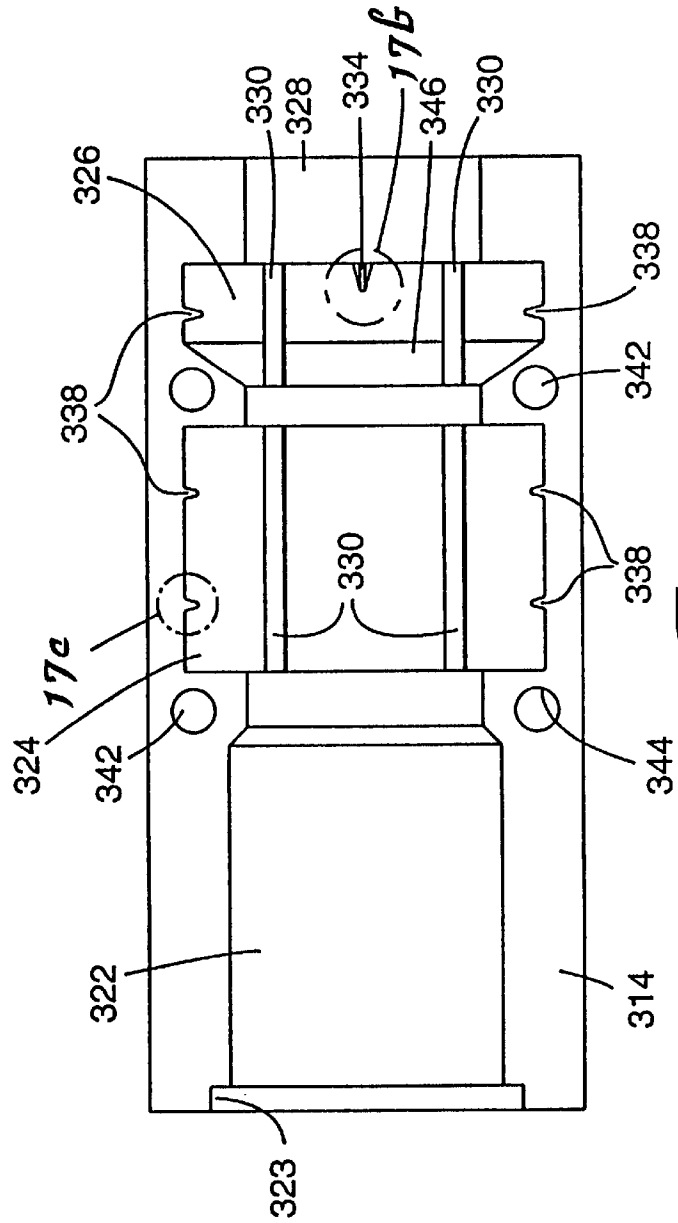
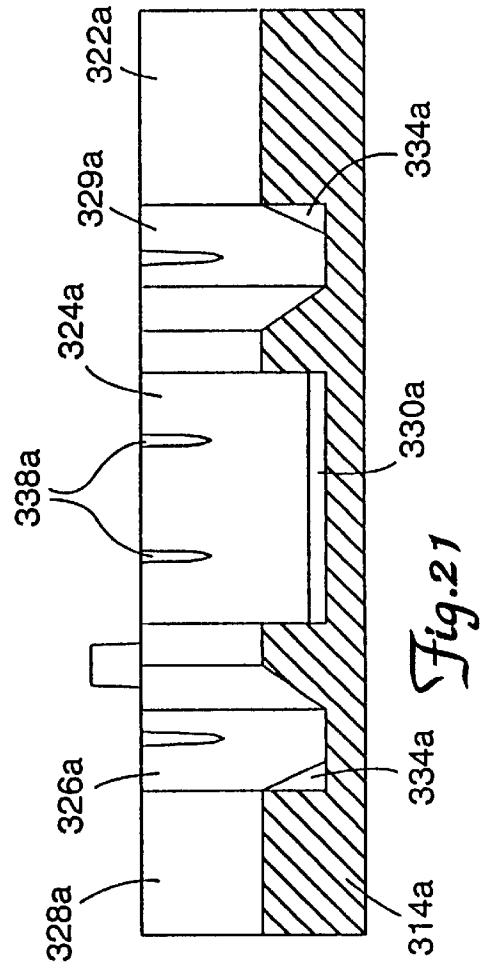
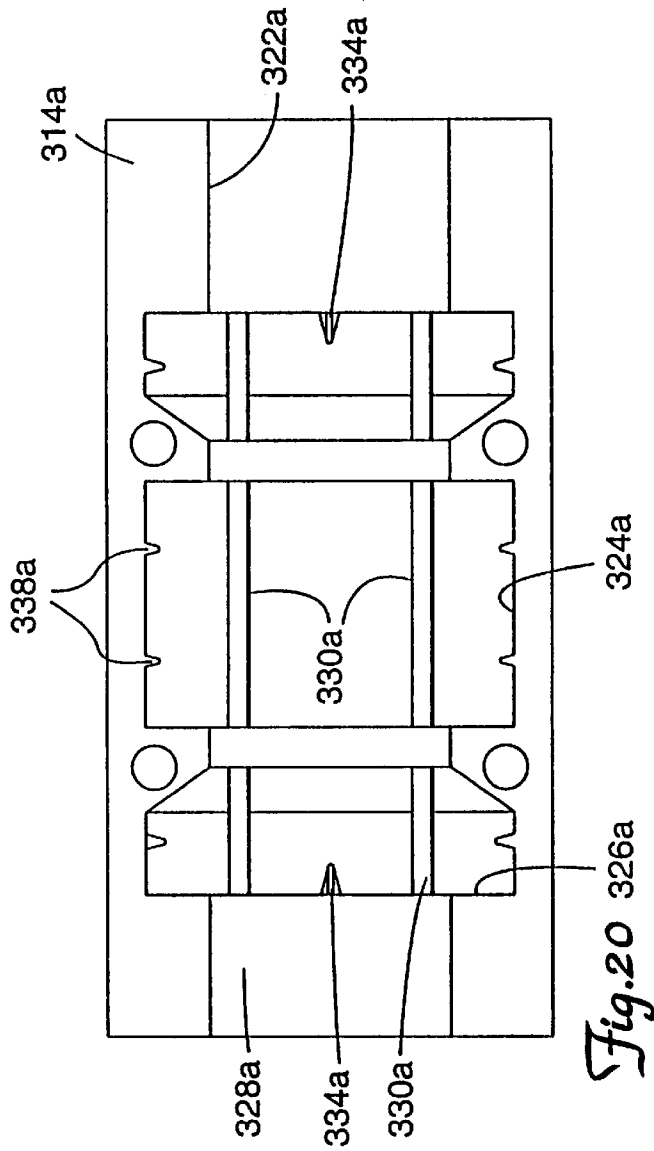


Fig. 18



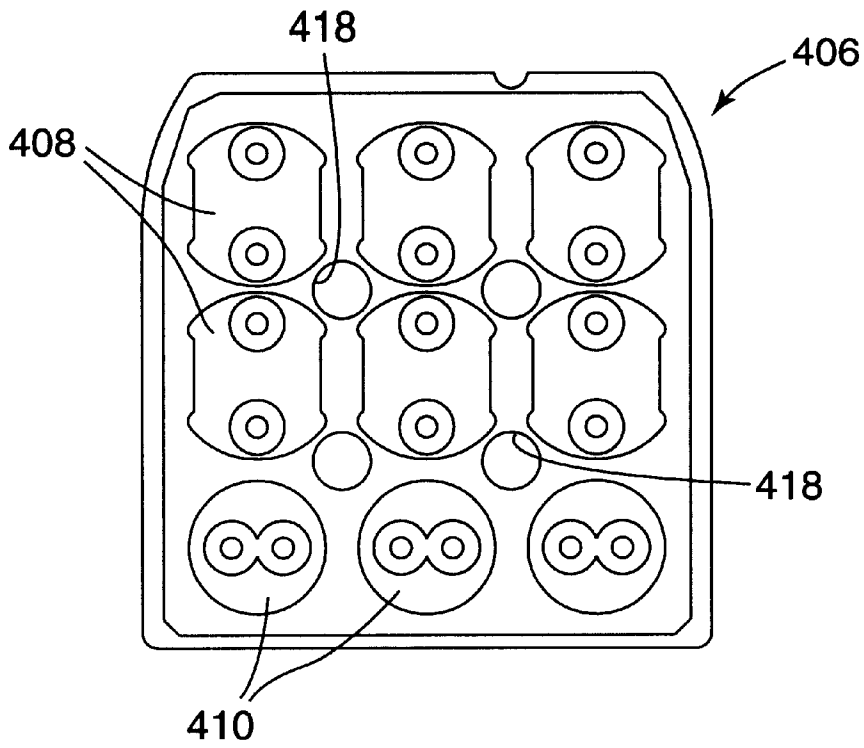


Fig. 22

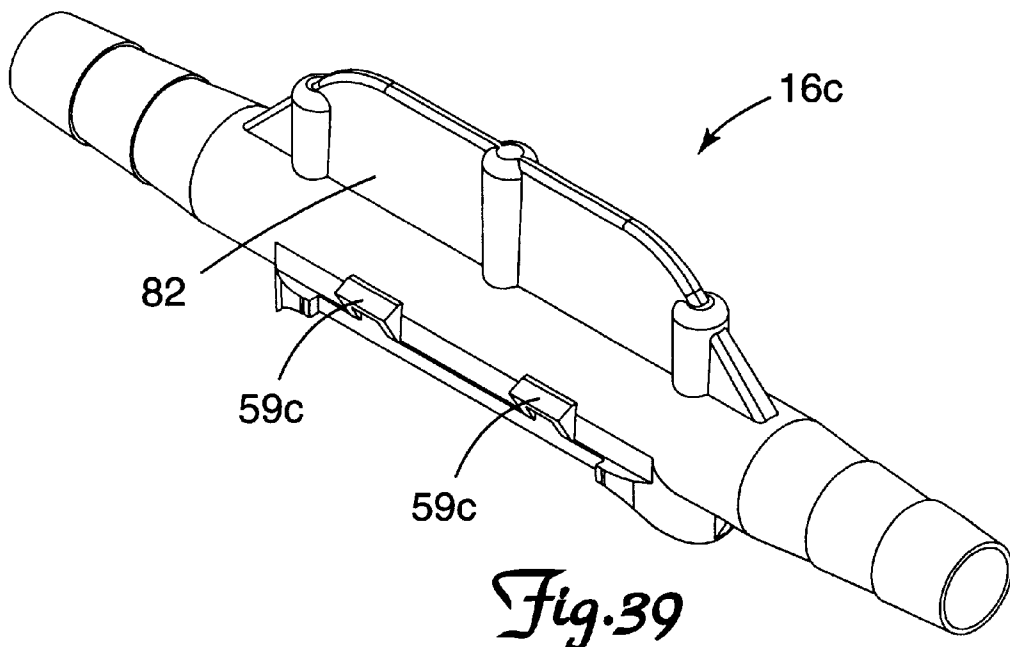


Fig. 39

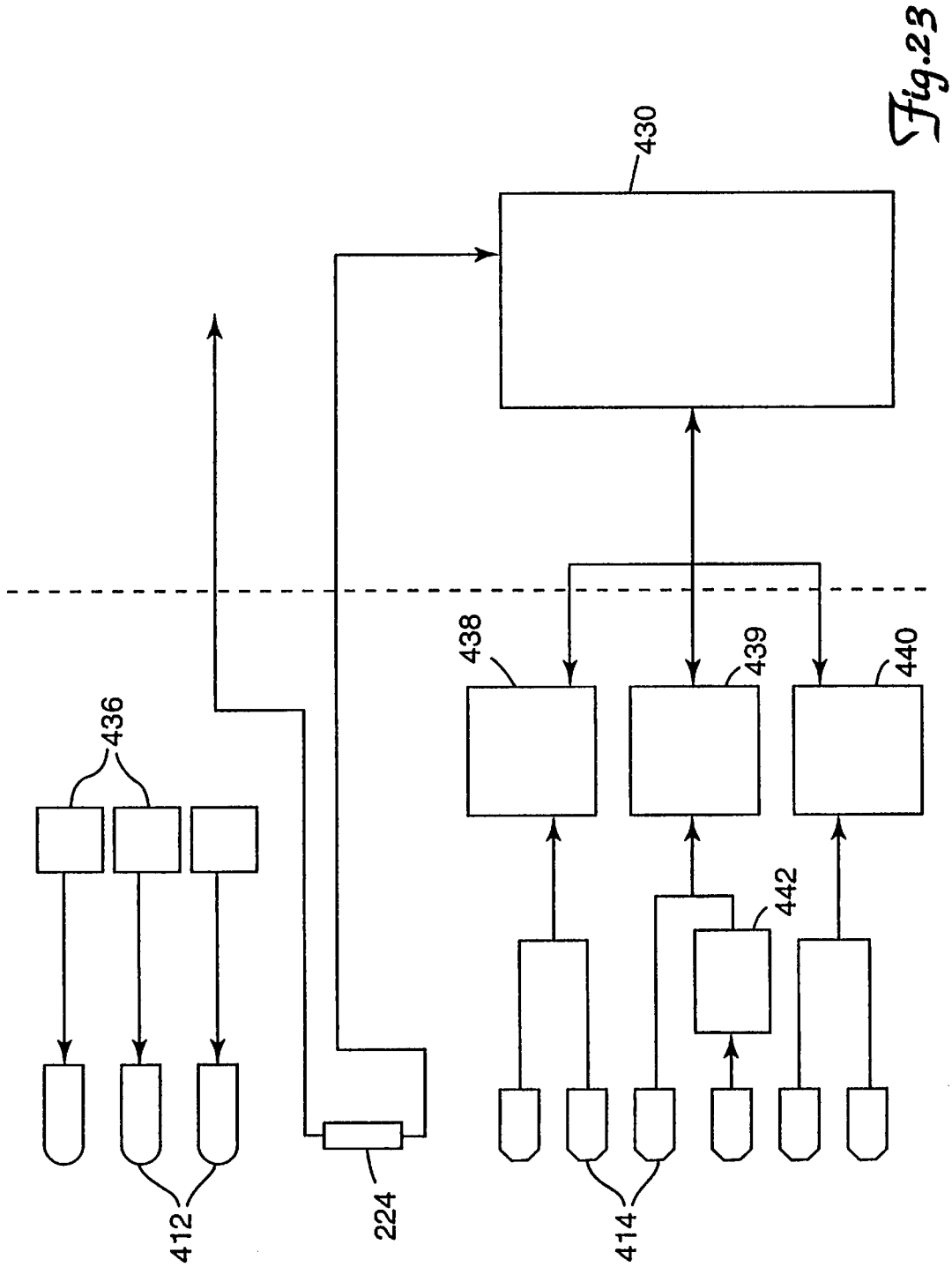


Fig. 23

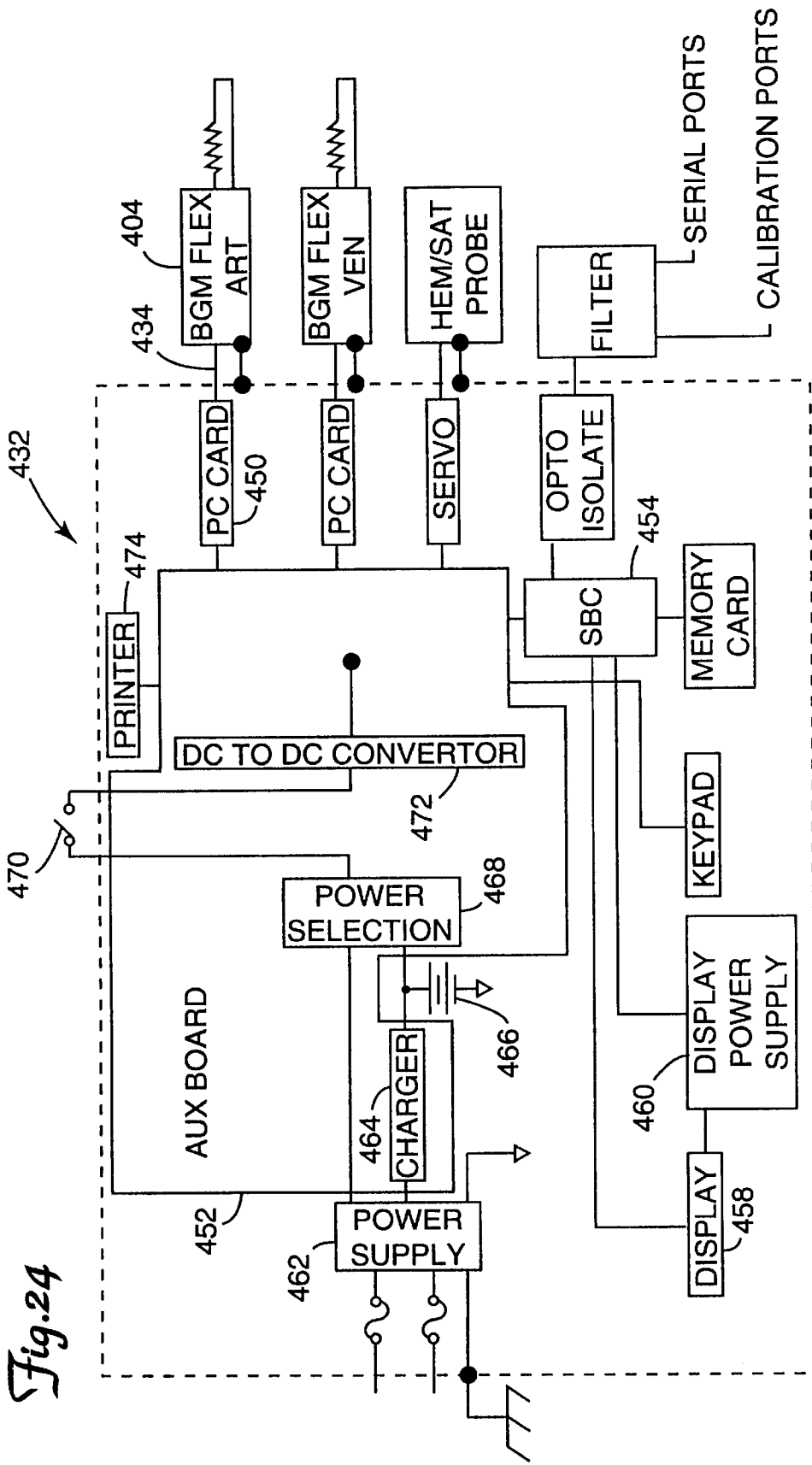


Fig. 24

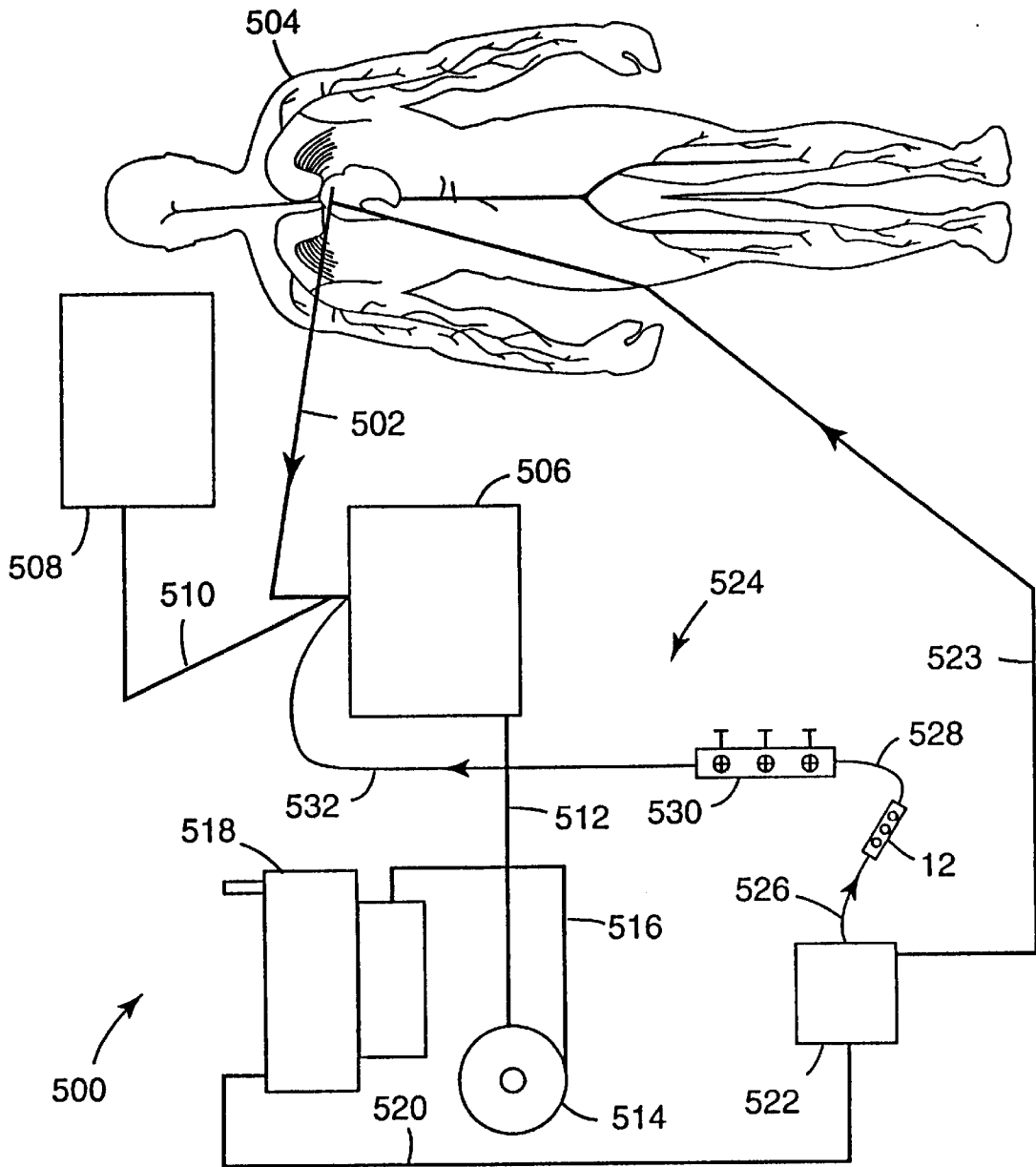


Fig.25

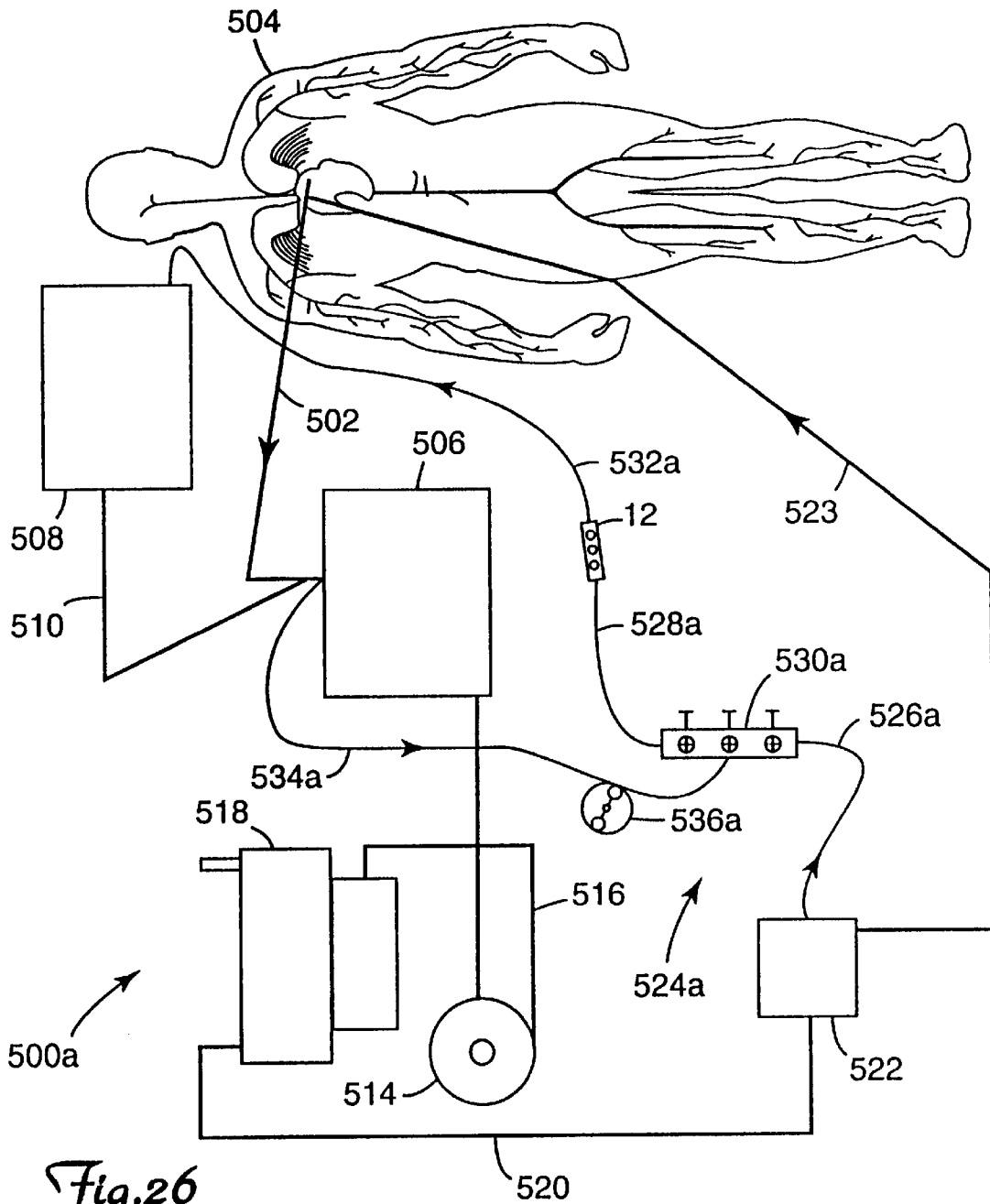


Fig.26

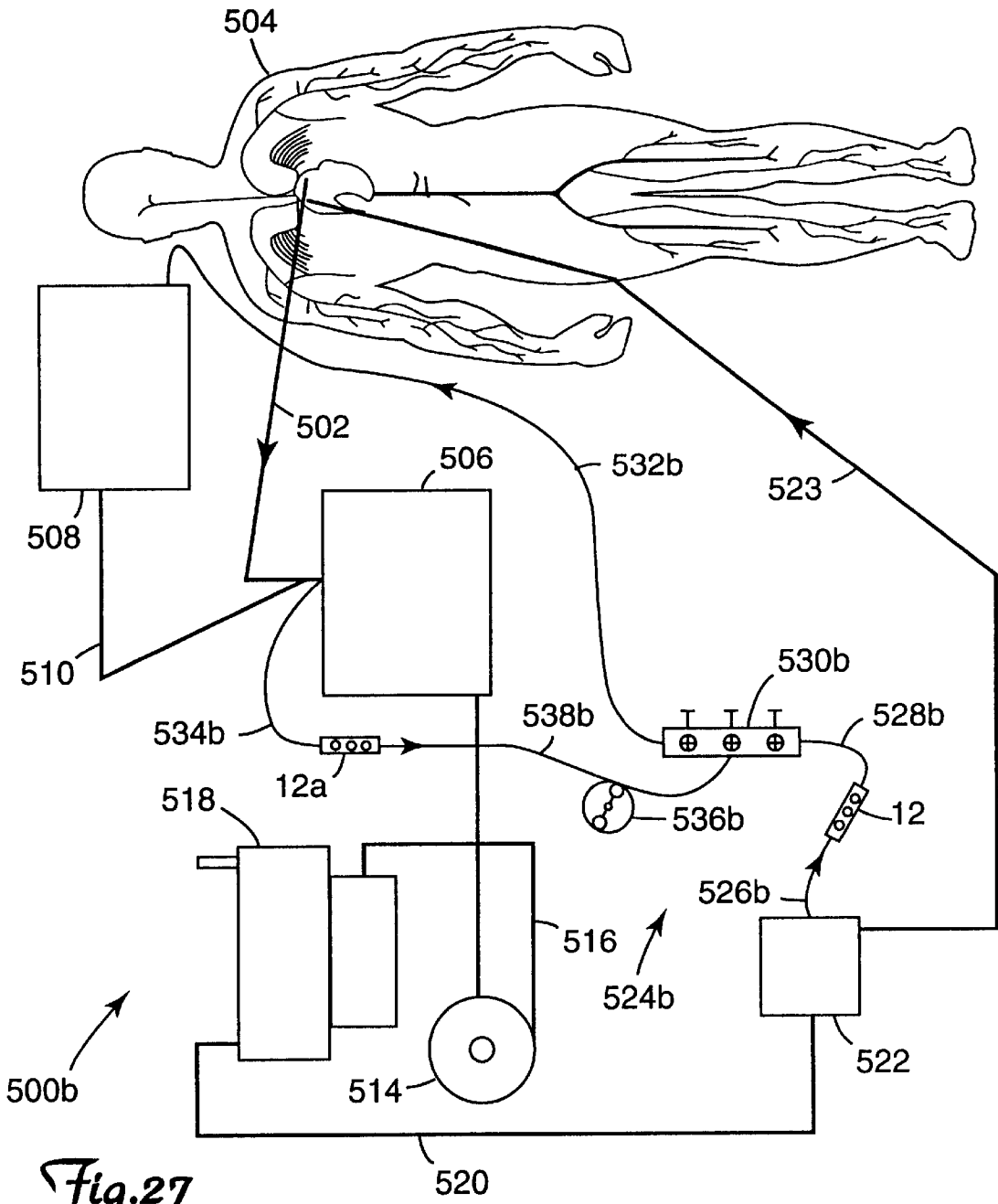


Fig.27

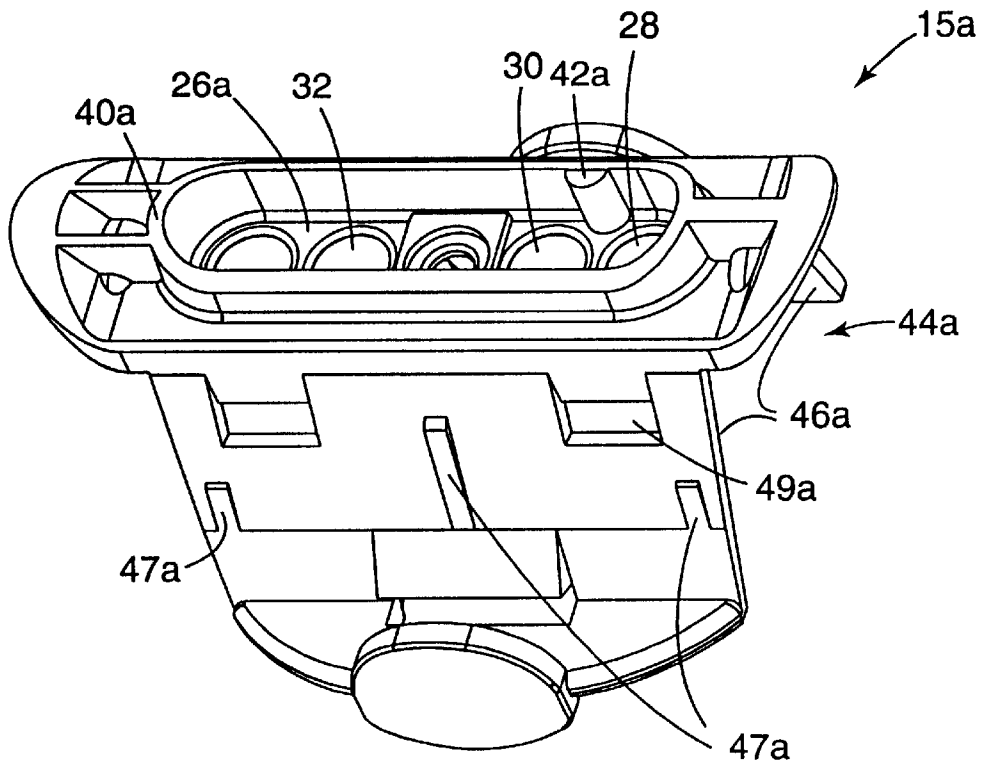


Fig. 28

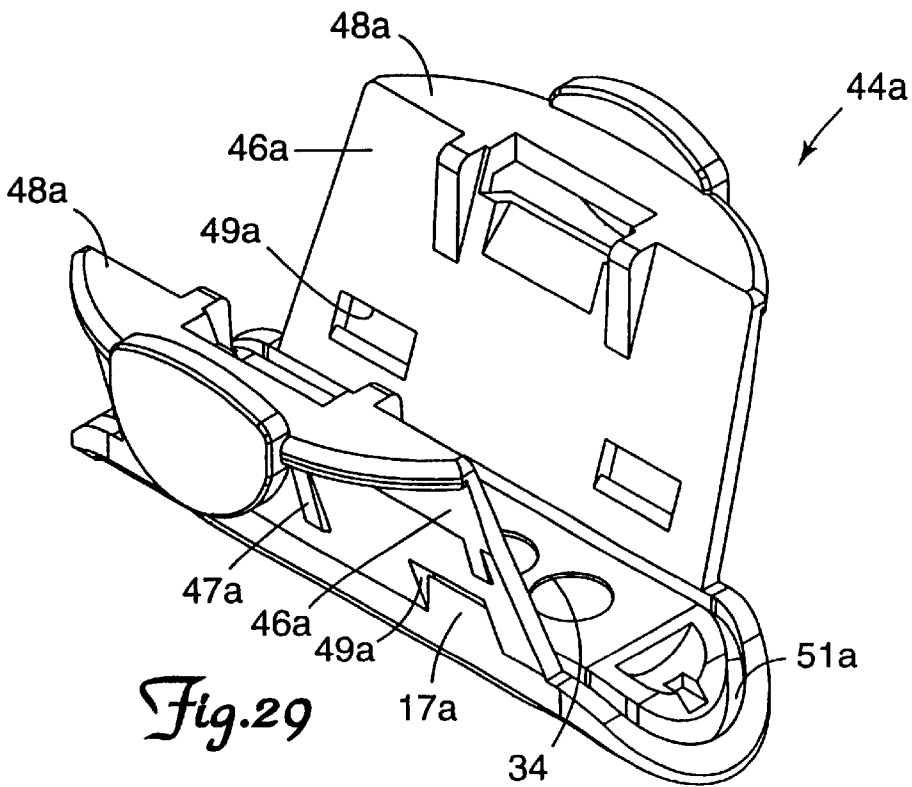


Fig. 29

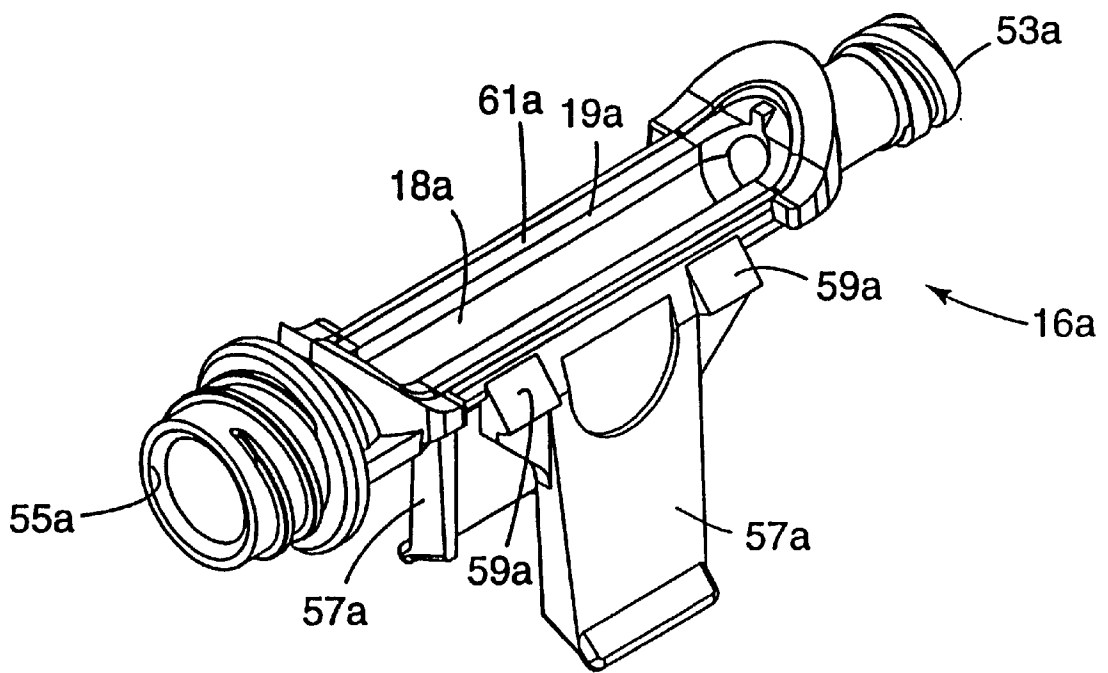
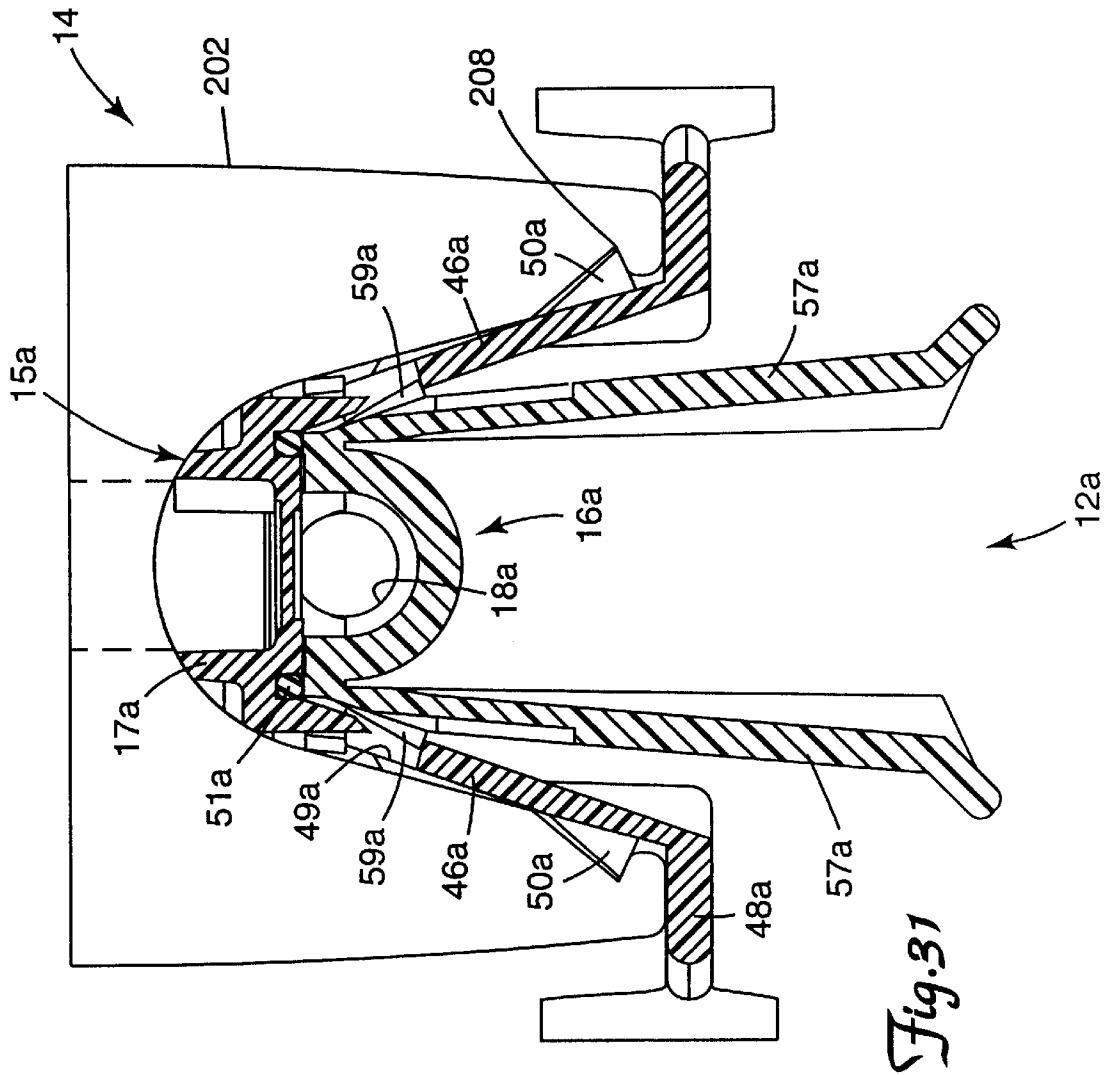


Fig. 30



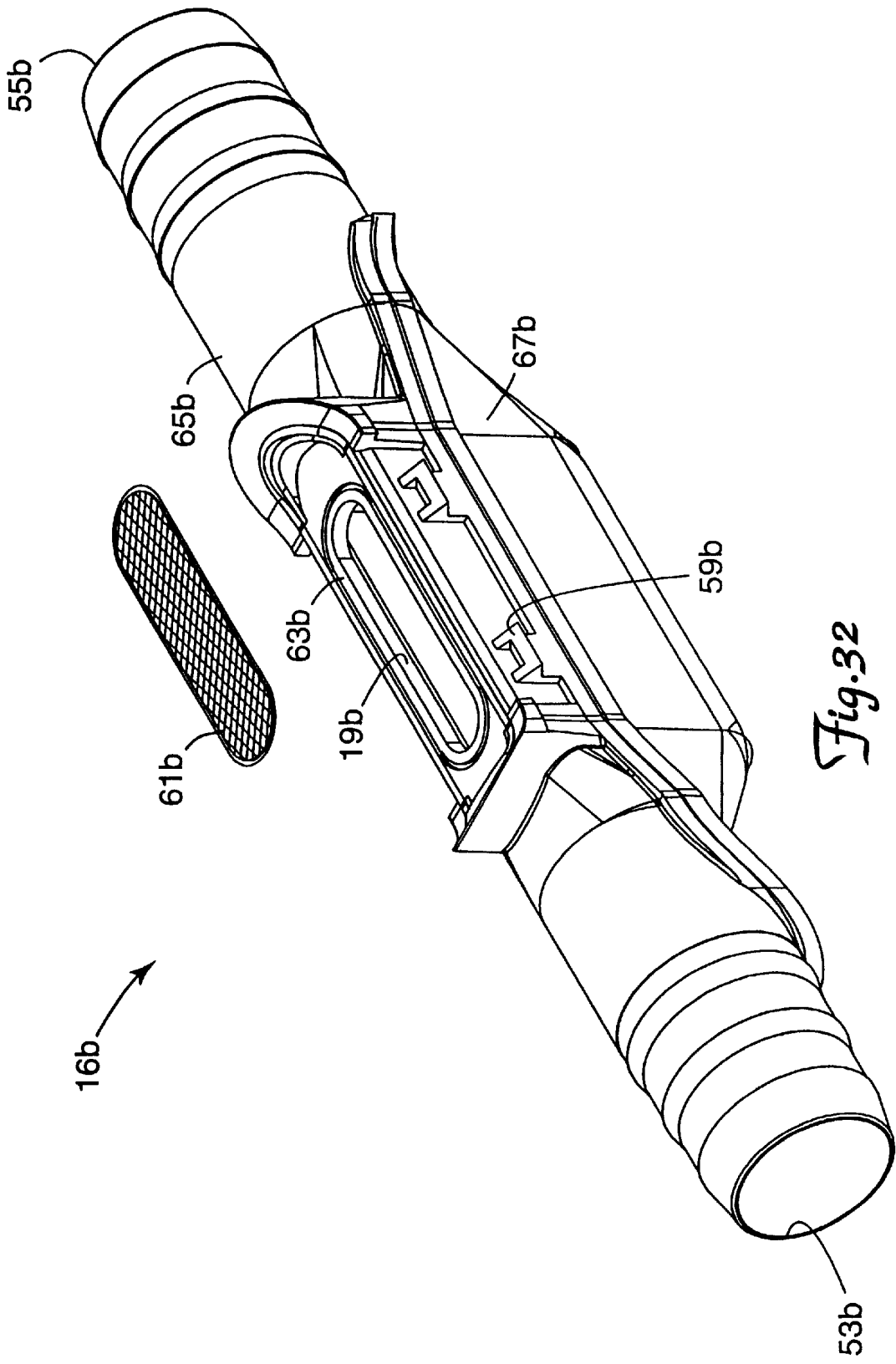


Fig. 32

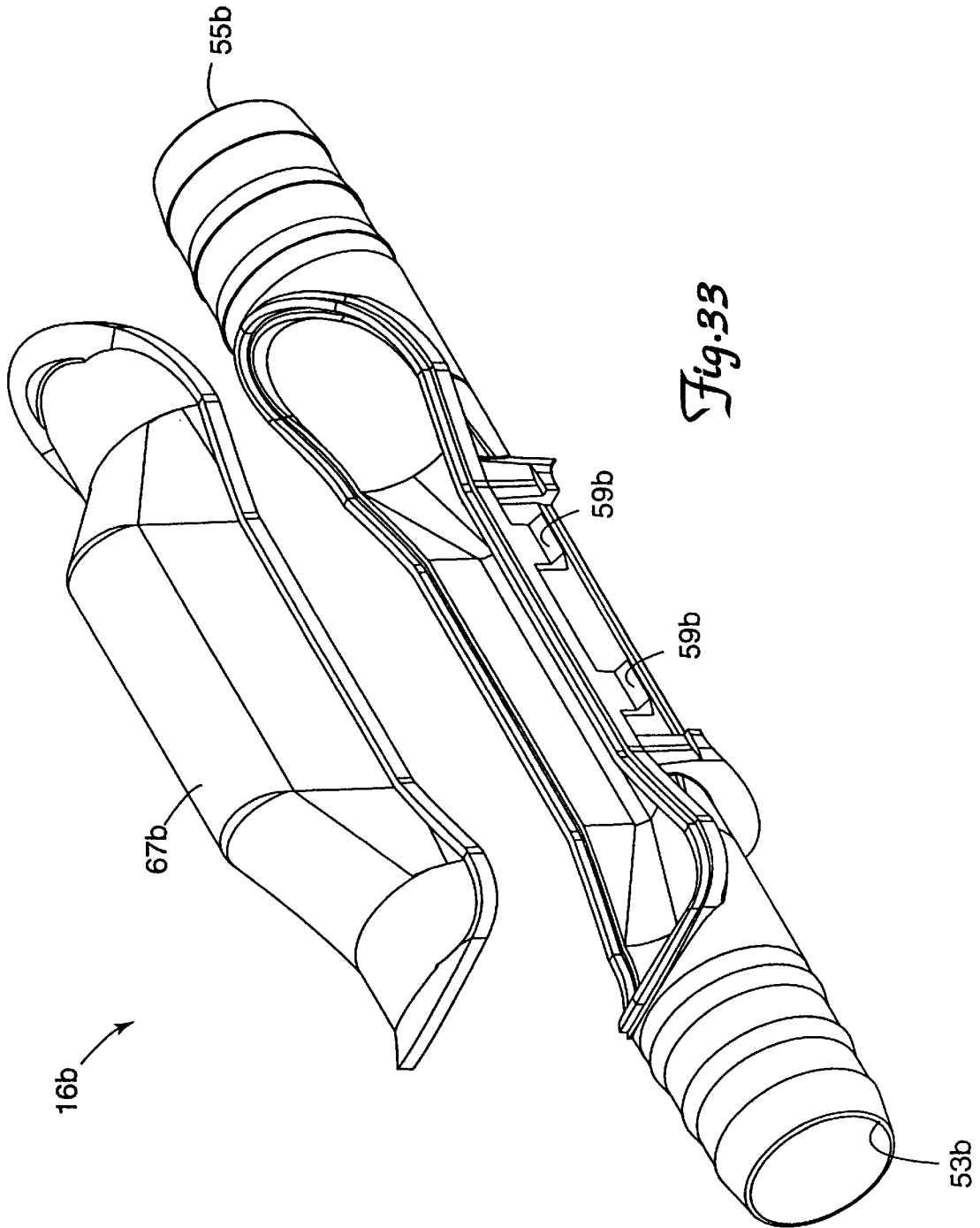


Fig. 33

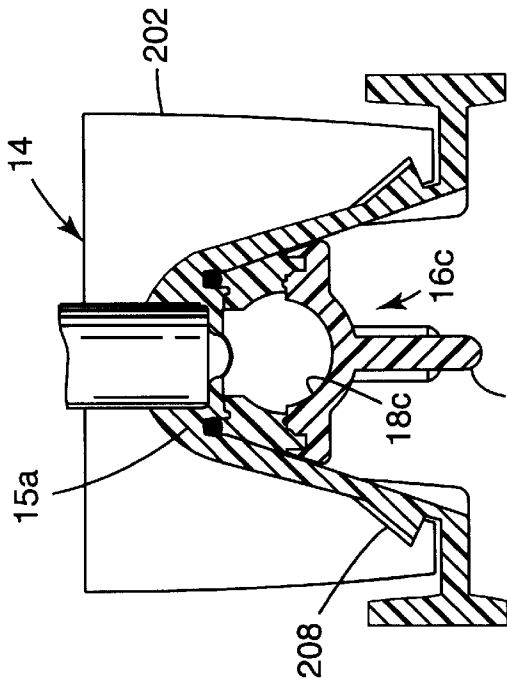


Fig. 37

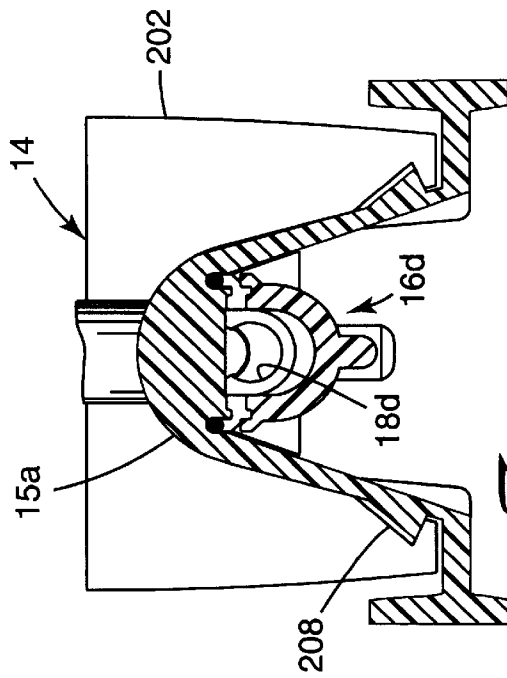


Fig. 38

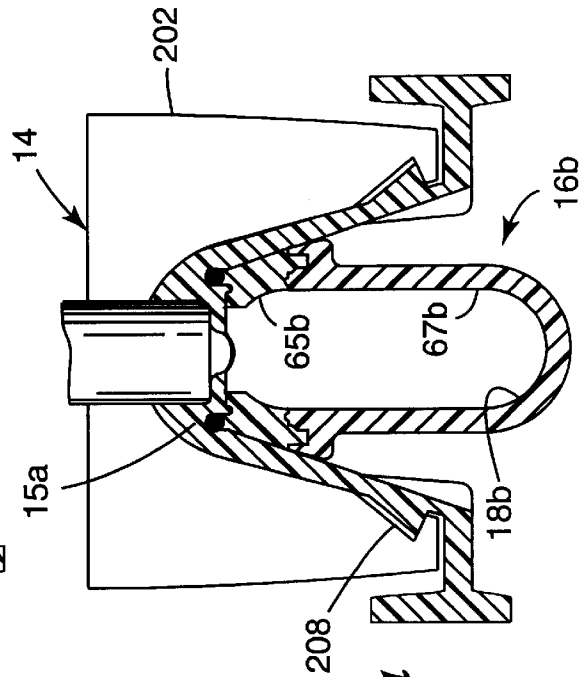


Fig. 34

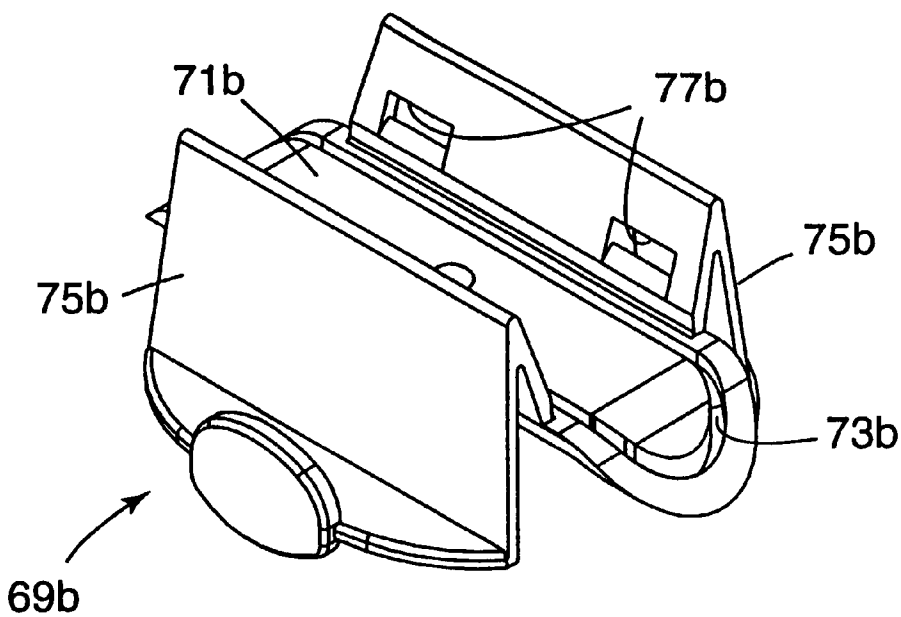


Fig. 35

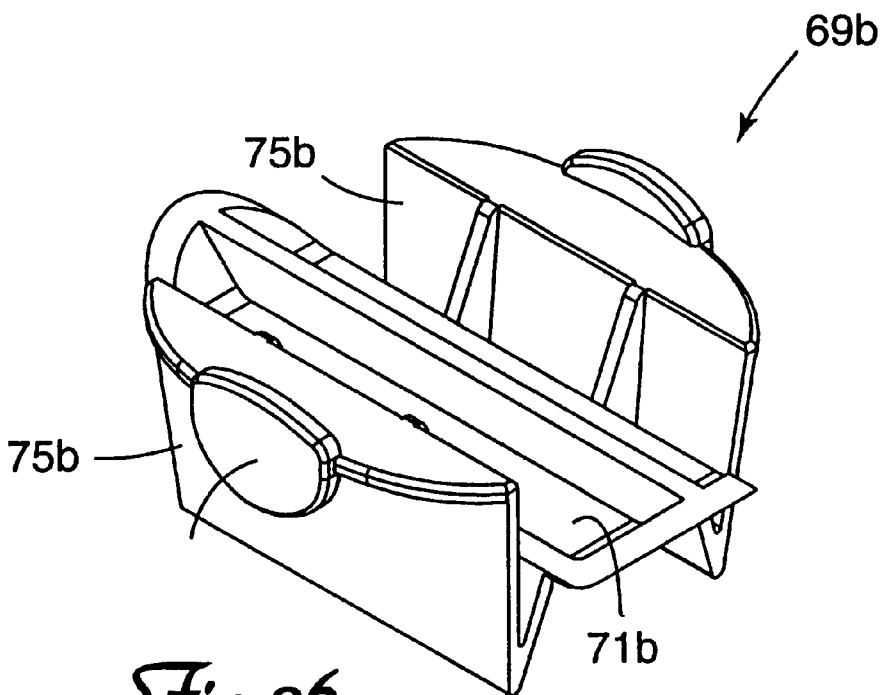


Fig. 36

CASSETTE FOR MEASURING PARAMETERS OF BLOOD

This is a continuation-in-part of application Ser. No. 08/810,954 filed Feb. 27, 1997 now abandoned.

BACKGROUND OF THE INVENTION

1. Field of the Invention

This invention broadly relates to a cassette for measuring one or more parameters of blood, and is especially suitable for use during surgical procedures.

2. Description of the Related Art

Various characteristics or parameters of blood are often monitored in real time during certain surgical procedures. For example, during open heart surgery the surgeon and other members of the surgical team often monitor the pH of the patient's blood as well as the concentration of certain blood gases such as carbon dioxide and oxygen. In many instances, the parameters of the patient's blood are monitored not only as the surgery is progressing, but also in the time period before and after the surgical procedure.

Measurement of blood parameters is often accomplished using an extracorporeal blood circuit having lengths of flexible tubing with passageways that are in fluid communication with the vascular system of the patient. In many extracorporeal blood circuits, one or more sensors that are useful for determining blood parameters are placed adjacent the passageway and are connected to a processing unit. The processing unit is typically connected to a display device such as a monitor so that the surgical team can review the parameters of interest when desired. Optionally, the processing unit is connected to a recording device such as a printer to provide a log of the parameters over a period of time.

Optical sensors are often used for sensing blood parameters in real time. For example, U.S. Pat. No. Re. 31,879 to Lubbers et al. and U.S. Pat. No. 5,403,746 to Bentsen et al. describe fluorescent sensors that respond to light in accordance to the partial pressure of oxygen, the partial pressure of carbon dioxide and the pH of blood. Sensors that function on the principles of light absorbance are described, for example, in U.S. Pat. No. 4,041,932 to Fostick.

Extracorporeal blood circuits having sensors for determining blood parameters may be arranged in various manners, and the manner selected for use in a particular instance often depends upon the preferences of the surgical team. In some cases, the sensors are mounted in a housing located along a length of relatively small-diameter tubing that is connected at only one end to the patient's blood supply, and a device such as a syringe is used to draw a sample of blood past the sensors. Examples of such circuits are described in U.S. Pat. No. 4,989,606 to Gehrich et al. Such circuits are an advantage in that the length of tubing connected to the sensors can be coupled to the blood supply at any convenient time before or during the surgical procedure, but are a disadvantage in that the blood does not continuously circulate in the length of tubing connected to the sensors and as a result the accuracy of the sensed parameter may be adversely affected.

Another type of extracorporeal blood circuit has sensors located along tubing that is part of an arterial or a venous passageway connected to an oxygenator. The sensors in this type of circuit are often connected to an element known as a flow-through cell that has fitting on opposite sides for coupling to the circuit tubing. Flow-through cells are described, for example, in U.S. Pat. No. 4,640,820 to

Cooper. Flow-through cells are often favored because a fresh supply of blood is continuously circulated past the sensors. Unfortunately, flow-through cells are in series relation to the circuit tubing and hence cannot be installed after blood has begun to flow through the tubing.

Optical sensors for measuring blood parameters are often optically coupled to a remote measuring device that includes a source of light for directing light to the sensors as well as apparatus for analyzing the light returned from the sensors. In many systems, a bundle of optical fibers extends from the remote device to a transmission block or retainer, and a releasable coupling is provided to detachably connect the retainer or block to the cell or housing that supports the sensors. For example, U.S. Pat. No. 4,989,606 to Gehrich et al. describes a transmission block that is detachably secured to a housing by a movable securement assembly having a swingable member and a threaded screw for clamping the housing between an outer end of the member and the block.

While many of the components known in the art for measuring blood parameters are satisfactory in certain instances, there is a continuing need to improve the components in order to facilitate ease of use without adversely affecting the accuracy of the measurements or unduly increasing the cost of the components. Moreover, it would be desirable to provide components that can be used in a variety of blood circuit configurations.

SUMMARY OF THE INVENTION

The present invention concerns in one aspect a system for measuring one or more parameters of blood. The system includes a casing having walls defining a chamber for receiving a quantity of blood, and at least one sensor connected to the casing adjacent the chamber for sensing at least one parameter of blood in the chamber. The system also includes a measuring device having a housing, at least one light source connected to the housing and at least one light detector connected to the housing. A connector releasably couples the casing to the housing. The connector includes a first coupling connected to the casing and a second coupling connected to the housing. The first coupling has a convex configuration and the second coupling has a concave configuration.

Another aspect of the present invention is directed toward a cassette for measuring one or more parameters of blood. The cassette includes a casing having walls defining a chamber for receiving a quantity of blood, and at least one sensor connected to the casing adjacent the chamber for sensing at least one parameter of blood in the chamber. The cassette also includes a coupling for connecting the casing to a blood parameter measurement device. The coupling has a convex configuration with a central section. The at least one sensor is located in the central section, and the coupling includes at least one flexible leg portion with a tab for releasable connection to the blood parameter measurement device. The at least one leg portion extends in a direction away from the central section, and the tab extends outwardly in a lateral direction from the at least one leg portion.

Another aspect of the present invention is directed to a cassette for measuring one or more parameters of blood. The cassette includes a casing having walls defining a chamber for receiving a quantity of blood, and at least one sensor connected to the casing adjacent the chamber for sensing at least one parameter of blood in the chamber. The cassette also includes a coupling for connecting the casing to a blood parameter measurement device. The coupling has a convex configuration and has two outer surfaces that extend in

respective reference planes oriented at an angle in the range of about 28 degrees to about 32 degrees relative to each other.

In another embodiment, the present invention is directed toward a cassette for measuring one or more parameters of blood. The cassette includes a casing having walls defining a chamber for receiving a quantity of blood and at least one sensor connected to the casing adjacent the chamber for sensing at least one parameter of the blood in the chamber. The cassette also includes a coupling having a convex configuration for connecting the casing to a blood parameter measurement device. The chamber has a longitudinal axis and includes an inlet portion, an outlet portion and a middle portion between the inlet portion and the outlet portion. The inlet portion and the outlet portion each have generally circular configurations in reference planes perpendicular to the longitudinal axis. The middle portion has a non-circular configuration in reference planes perpendicular to the longitudinal axis. The circular configurations of the inlet portion and the outlet portion have respective areas that are each approximately equal to the area of the non-circular configuration of the middle portion.

The present invention is also directed toward a cassette for measuring one or more parameters of blood. The cassette includes a body and at least one sensor connected to the body for sensing at least one parameter of the blood. The cassette further includes a casing having wall sections at least partially defining a chamber, an inlet for admitting blood to the chamber and an outlet for discharging blood from the chamber. A connector detachably couples the body to the casing in such a manner that the at least one sensor is located next to the chamber for sampling one or more parameters of blood flowing through the chamber.

Another embodiment of the present invention is directed toward a cassette for measuring one or more parameters of blood. The cassette includes a body and at least one sensor connected to the body for sensing at least one parameter of the blood. The cassette also includes a casing detachably coupled to the body and having walls defining an elongated chamber for receiving a quantity of blood. The casing has an external rib that extends along its length at a location remote from the body.

In another aspect, the invention is directed toward a cassette for measuring one or more parameters of blood. The cassette includes a casing having walls defining a chamber for receiving a quantity of blood and at least one sensor connected to the casing adjacent the chamber for sensing at least one parameter of blood in the chamber. The cassette also includes a coupling for connecting the casing to a blood measurement device, and the coupling includes a generally oval-shaped recess. At least one sensor is located in the recess.

The present invention in another embodiment concerns a cardiopulmonary by-pass circuit. The circuit includes a venous passageway having an inlet and an outlet, and the inlet is adapted for connection to a patient's venous blood vessel. The circuit includes an arterial passageway having an inlet and an outlet, and the outlet is adapted for connection to a patient's arterial blood vessel. The circuit also includes a device for providing oxygen to blood, and the device interconnects the outlet of the venous passageway and the inlet of the arterial passageway. A pump is provided for moving blood through the venous passageway and the arterial passageway. The circuit also includes a shunt passageway having an inlet and an outlet, and the inlet of the shunt passageway communicates with one of the venous

passageway and the arterial passageway and the outlet of the shunt passageway communicates with one of the venous passageway and the arterial passageway. At least one fluorescing sensor is located adjacent the shunt passageway for sensing the one or more parameters of blood flowing through the shunt passageway.

The connector and the coupling as set out in some of the embodiments above provides a quick yet reliable connection that can be readily uncoupled when desired. The concave configuration of the device coupling partially surrounds the sensors of the cassette and thereby provides protection to the sensors from the influence of ambient light or localized changes in temperature. The cardiopulmonary by-pass circuit of the invention is a particular advantage, in that the shunt passageway can be conveniently connected or disconnected from the venous or arterial passageway without disrupting the flow of blood between the patient and the device for providing oxygen to the blood.

These and other aspects of the invention are described in further detail below in the text that follows as well as in the accompanying drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a perspective view of a calibration and fluid parameter measurement cassette of the invention along with a fluid parameter measuring device, showing one example of how the cassette and the measuring device are oriented with respect to each other before being coupled together;

FIG. 2 is an enlarged, longitudinally transverse cross-sectional view through the cassette alone that is shown in FIG. 1;

FIG. 3 is a view somewhat similar to FIG. 2 except that FIG. 3 is taken along a different reference plane;

FIG. 4 is an enlarged elevational view in partial section of the cassette shown in FIGS. 2-3, looking toward a side of the cassette that faces the measuring device when the cassette and the measuring device of FIG. 1 are coupled together;

FIG. 5 is an enlarged side cross-sectional view of the cassette and part of the measuring device of FIG. 1 but shown as coupled together, and additionally showing connectors and tubing for coupling the cassette to a fluid circuit such as a cardiopulmonary by-pass circuit;

FIG. 6 is a view somewhat similar to FIG. 5 except that the measuring device is not shown and additional components have been connected to the cassette for calibration of sensors of the cassette;

FIG. 7 is an exploded, perspective view of the cassette alone that is shown in FIG. 1 but from a different view, illustrating a two-piece construction of the cassette for exemplary purposes;

FIG. 8 is an enlarged side cross-sectional view of the measuring device alone that is illustrated in FIG. 1, except that optical fibers within the device have not been shown;

FIG. 9 is an exploded perspective view of the measuring device shown in FIG. 1;

FIG. 10 is an enlarged, exploded perspective view of the measuring device of FIG. 1 except that a housing of the measuring device has been removed;

FIG. 11 is an enlarged end view of part of the measuring device of FIG. 1, looking in a direction parallel to a longitudinal axis of the device housing and showing among other items a fiber terminal block assembly;

FIG. 12 is an enlarged bottom view of the fiber terminal block assembly shown in FIG. 11;

FIG. 13 is an enlarged side cross-sectional view of a portion of the fiber terminal block assembly that is depicted in FIG. 11;

FIG. 14 is an enlarged perspective view of an insert plate that is part of the fiber terminal block assembly shown in FIG. 11;

FIG. 15 is a schematic illustration showing among other things the various paths of optical fiber bundles of the measuring device illustrated in FIG. 1;

FIG. 16 is an enlarged perspective view in partially exploded form of part of an optics assembly of the measuring device of FIG. 1;

FIG. 16a is an enlarged side elevational view of a lens of the optics assembly illustrated in FIG. 16;

FIG. 17 is an enlarged cross-sectional view taken along a longitudinal axis of one optical retainer of the optics assembly depicted in FIG. 16;

FIGS. 17a, 17b and 17c are enlarged views of various portions of the optical retainer shown in FIG. 17;

FIG. 18 is a plan view of the optical retainer shown in FIG. 17;

FIG. 19 is a cross-sectional view of the retainer illustrated in FIGS. 16–17, taken along a reference plane perpendicular to the longitudinal axis of the retainer;

FIG. 19a is an enlarged view of a portion of the retainer shown in FIG. 19;

FIG. 20 is an enlarged plan view of another optical retainer of the optics assembly depicted in FIG. 16;

FIG. 21 is a cross-sectional view taken along a longitudinal axis of the optical retainer shown in FIG. 20;

FIG. 22 is an enlarged elevational view of an electro-optics coupling plate of the measuring device shown in FIG. 1;

FIG. 23 is a schematic block diagram of an electrical assembly of the device of FIG. 1 as well as part of a monitor;

FIG. 24 is a schematic block diagram of the monitor, also showing connection to the device of FIG. 1 along with connection to other devices;

FIG. 25 is a schematic diagram of a cardiopulmonary bypass circuit using the cassette and device of FIG. 1;

FIG. 26 is a view somewhat similar to FIG. 1 but in accordance with another embodiment of the invention;

FIG. 27 is a view somewhat similar to FIG. 25 but in accordance with another embodiment of the invention;

FIG. 28 is an enlarged perspective view of a body of a blood parameter measurement cassette constructed in accordance with another embodiment of the invention;

FIG. 29 is a view somewhat similar to FIG. 28 except looking in another direction toward the body;

FIG. 30 is an enlarged perspective view of an exemplary cassette casing for detachable connection with the cassette body shown in FIGS. 28–29;

FIG. 31 is an enlarged end cross-sectional view of the cassette casing of FIG. 30 shown as attached to the cassette body of FIGS. 28–29, and additionally illustrating a coupling of the blood parameter measuring device of FIG. 1 coupled to the cassette;

FIG. 32 is an enlarged perspective view in partially exploded form of another cassette casing that can be attached to the cassette body shown in FIGS. 28–29 if desired to present a somewhat larger cross-sectional area for the passage of blood;

FIG. 33 is a view somewhat similar to FIG. 32 except taken in another direction toward the cassette casing and depicting another part of the casing in exploded form;

FIG. 34 is a view somewhat similar to FIG. 31 except showing the cassette casing of FIGS. 32–33 in place of the cassette casing of FIG. 30;

FIG. 35 is an enlarged perspective view of a shipping cap for use with the cassette casing of FIGS. 32–33;

FIG. 36 is a view somewhat similar to FIG. 35 except looking in another direction toward the cap;

FIG. 37 is a view somewhat similar to FIG. 34 except showing another cassette casing that is connected to the cassette body in accordance with another embodiment of the invention;

FIG. 38 is a view somewhat similar to FIG. 34 except showing another cassette casing that is coupled to the cassette body in accordance with yet another embodiment of the invention; and

FIG. 39 is a perspective view of the cassette casing alone that is illustrated in FIG. 37.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

A system 10 for measuring one or more characteristics or parameters of fluid such as blood is illustrated in FIG. 1. The system 10 broadly includes a cassette 12 that receives the fluid along with a measuring device 14 for measuring parameters of fluid in the cassette 12.

The cassette 12 is shown in more detail in FIGS. 2–6, and includes an elongated casing 16 having wall sections defining an elongated, internal, flow-through fluid chamber 18 that extends along the longitudinal axis of the casing 16. As illustrated in FIGS. 5 and 6, the fluid chamber 18 includes a first portion 20 having a first or “inlet” port for admitting fluid into the chamber 18, a second portion 22 having a second or “outlet” port for allowing fluid to exit the fluid chamber 18 and a central portion 24 located between the portions 20, 22. (Although the description that follows refers to fluid flowing into the chamber 18 through the first portion 20 and discharged from the chamber 18 through the second portion 22, it should be understood that the fluid may also flow if desired in an opposite direction through the chamber 18 such that the fluid enters the chamber 18 through the second port and exits through the first port).

The fluid chamber 18 also includes a frustoconical expansion region or portion 25 that is located between the central portion 24 and the second portion 22. The expansion portion 25 has free area that is larger than the free area of the central portion 24 in reference planes perpendicular to the longitudinal axis of the casing 16 and that increases in size as the second portion 22 is approached. The portions 20, 22, 24 and 25 communicate with each other and have circular cross-sections in reference planes perpendicular to the longitudinal axis of the cassette 16. Preferably, at least the wall sections defining the central portion 24 include a hydrophilic surface, and more preferably the wall sections defining all of the portions 20, 22, 24 and 25 include a hydrophilic surface. Optionally, the hydrophilic surface comprises a coating of heparin.

An external side of the casing 16 includes a central section with a generally oval-shaped recess 26. At least one sensor for determining one or more parameters of fluid in the chamber 18 is carried by the casing 16. In the embodiment shown, a series of four sensors are located between the recess 26 and the central portion 24 of the fluid chamber 18, and the sensors are placed in four cavities that are arranged in aligned, spaced-apart relationship along the longitudinal axis of the casing 16. As depicted in FIG. 4, the sensors

include an ion (potassium) sensor **28**, a pH sensor **30**, and carbon dioxide sensor **32** and an oxygen sensor **34** that are received in cavities **27**, **29**, **31**, **33** respectively.

If desired additional sensors may be employed as described below. Sensors useful in the apparatus of the invention preferably comprise a multi-layer assembly that can be adhesively attached to the cassette casing **16**.

The ion sensor **28** preferably comprises the following layers: (i) a backing membrane, (ii) a pressure-sensitive adhesive (PSA) coated on the backing membrane, (iii) a sensing element comprising an ion sensing compound bound to a substrate, the substrate being attached to the membrane (e.g., by a non-interfering adhesive), and (iv) an outermost opacifying overcoat layer on the exposed surface of the substrate.

Useful pressure-sensitive adhesives include silicone adhesives and polyurethane adhesives and others that are capable of bonding a membrane (described below) to the cassette. Preferably, the adhesives are essentially transparent to wavelengths of light used in sensing cassettes of the invention, and are chemically non-interfering with useful ion sensors. Useful silicone adhesives include PSA-518™ (General Electric Co., Schenectady, N.Y.), described in Example 2 of U.S. Pat. No. 5,508,509, which is incorporated herein by reference. Useful polyurethane adhesives include FLEXOBOND 431™ (Bacon Co., Irvine, Calif.), described in Example 3 of U.S. Pat. No. 5,591,400, which is incorporated herein by reference.

A release liner may be useful when manufacturing the ion sensors to protect the exposed surface of the adhesive layer. These liners can be any that are in common use in industry for the purpose, and are selected according to the adhesive from which they are to release. Examples of useful release liners include poly(ethyleneterephthalate) (PET) that may optionally be coated with, e.g., silicone or a fluoropolymer, for increased ease of release from the adhesive. One useful liner is Scotch Pack 1022™ (3M Company, St. Paul, Minn.), a PET film coated with perfluoro polyether, described in U.S. Pat. No. 5,508,509, Example 2.

The backing membrane provides support (e.g., stiffness and handling capability) for the multilayer assemblies. Preferably, the backing membrane is transparent and essentially impermeable to, or much less permeable than the sensing substrate to, the solution in which a target ion is present, such as blood or a calibrating solution. The membrane preferably allows the signal or signals, preferably the optical signals, from the sensing element and substrate, to pass therethrough. Particularly useful materials of construction for this backing membrane include polymeric materials, such as polyesters, polycarbonates, polysulfones including but not limited to polyethersulfones and polyphenylsulfones, polyvinylidene fluoride, polymethylpentenes, and the like. In a presently preferred embodiment for ion sensor **28**, the backing membrane is polycarbonate.

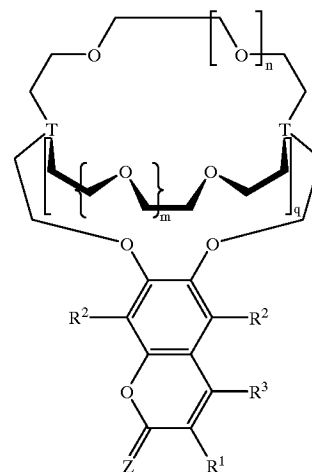
Suitable ion sensors which may be used as the potassium sensor **28** are described in U.S. Pat. Nos. 5,474,743 (Trend et al.), 5,176,882 (Gray et al.), 5,136,033, and 5,162,525 (Masilamani et al.); U.S. patent application Ser. No. 08/521,869; and U.S. patent application Docket No. 52630USA7A (filed on even date herewith and assigned to the assignee of the present invention), which are herein incorporated by reference.

Preferred sensors **28** comprise a fluorescent ionophoric compound ("the ionophore") that contains a complexing moiety for binding an ion and a fluorescing moiety. The compound has a wavelength of maximum absorbance of at

least about 350 nm. Suitable fluorescing moieties preferably contain close-lying $\pi\pi^*$ and $\pi\pi^*$ excited states. Suitable fluorescing moieties, when coupled to an appropriate complexing moiety, preferably are capable of ion dependent out-of-plane puckering. Also, the $\pi\pi^*$ state of suitable fluorescing moieties preferably is sufficiently high in energy that ion dependent mixing dominates non-radiative coupling to the ground state. Particularly preferred fluorescing moieties include coumarin moieties, although other aromatic carbonyls or nitroaromatics or N-heterocyclic moieties may be employed. Suitable ion complexing moieties include cyclic "cage" moieties capable of binding an ion. Preferably the cage is capable of selective binding of an ion. Preferred ion complexing moieties include cryptand and crown ether moieties, with cryptand moieties being particularly preferred.

Ions which may be sensed using the ionophore include, for example, Ag^+ , Ba^{+2} , Ca^{+2} , Ce^+ , Cd^{2+} , Fr^+ , Hg^{2+} , K^+ , Li^+ , Mg^{+2} , Mn^{2+} , Na^{+} , Pb^{+2} , Ru^+ , Sr^{+2} , Ti^+ , and Zn^{2+} . If desired the ionophore may be used in conjunction with an ion selective membrane. Preferred sensors comprise ionophores that sense for K^+ , Na^+ , and Ca^{+2} .

Suitable fluorescent ionophoric compounds include compounds having the following general formula (Formula "A"):



wherein

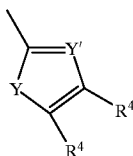
T is O or N, with the proviso that when T is O q is 0 and n is 0 to 2, and when T is N q is 1 and m and n are independently 0 or 1;

each R^2 independently is a sterically non-interfering group, including moieties such as hydrogen, halogen, a hydrocarbyl-containing group, a hetero-acyclic group, or a group having the formula $(\text{CH}_2\text{X})_a\text{E}$ in which X is **0**, NH, or a single bond, E is a functional group that includes active hydrogen, and a is a whole number from 1 to 100;

R^3 preferably is a non-electron withdrawing group, including non-electron withdrawing moieties such as hydrogen, a hydrocarbyl-containing group, a hetero-acyclic group, a heterocyclic group, or a group having the formula $(\text{CH}_2\text{X})_b\text{E}$ in which X and E are defined as above and b is a whole number from 0 to 100;

R^1 is an electron withdrawing or polarizable group, including moieties such as carboxyl, carboxamide, sulfonylaryl, ester, keto-alkyl ester, heterocyclic moieties and aromatic groups (preferably substituted at one

or more positions), most preferred R^1 groups include substituted heterocyclic moieties having the general formula (Formula "C"):



wherein

Y and Y' independently are O, S, NH_x , or CH_y , where x is 0 or 1 and y is 1 or 2, with the proviso that at least one of Y and Y' must be O, S, or NH_x , each R^4 group is independently hydrogen, halogen, a hydrocarbyl-containing group, a hetero-acyclic group, a heterocyclic group, or a group having the formula $(CH_2X)_cE$ in which X and E are defined as above and c is a whole number from 0 to 100, or both R^4 groups together with the carbon atoms to which they are attached form a 5- or 6-membered ring which optionally can have one or more further R^4 groups attached; and

Z is O or NR^5 , where R^5 is hydrogen or a hydrocarbyl-containing group, more preferably R^5 is H or a C1 to C4 alkyl group, and most preferably R^5 is H.

In general, compounds of Formula A have a wavelength of excitation of at least about 350 nm and a wavelength of emission preferably of no more than about 500 nm. Preferred compounds, (e.g., wherein R^1 is a heterocyclic moiety having the general formula of Formula "C") have a wavelength of excitation of at least about 380 nm and a wavelength of emission of no more than about 500 nm. In a particularly preferred embodiment, substituent groups and their position on the coumarin ring have been chosen so as to ensure that the excitation (i.e., absorption) maximum of the ionophore of the present invention is centered at a wavelength greater than 380 nm. This allows the ionophore of the present invention to be used with solid state light sources such as, for example, blue LEDs and lasers. The wavelengths of excitation and emission of these compounds are preferably at least about 10 nm apart, which allow these compounds to be useful in fluorescence-based cation concentration measurement techniques. Substituent groups and their positions are also preferably chosen to keep the emission wavelength below 500 nm, thereby preserving ionophore response for this class of indicators. Finally, substituent groups and their positions are preferably chosen to provide the option for covalent attachment to substrates. Preferably, the substrate to which the indicator is attached is chosen to support uniform and reproducible ionophore response and to minimize the effect of physiological pH changes on ionophore response. Suitable coupling agents for covalent attachment are described in U.S. Pat. No. 5,053,520, which is herein incorporated by reference. Homobifunctional and/or heterobifunctional coupling agents are described in World Pat. No. WO 96/07268 and WO 96/10747, which are herein incorporated by reference.

Preferably, the ionophore is covalently bonded to a suitable substrate that can be attached to the backing membrane, as described below. The substrate preferably is a polymeric material that is water-swellaible and permeable to the ionic species of interest, and is preferably insoluble in the medium to be monitored. Particularly useful substrate polymers include, for example, ion-permeable cellulosic materials, high molecular weight or crosslinked polyvinyl alcohol

(PVA), dextran, crosslinked dextran, polyurethanes, quaternized polystyrenes, sulfonated polystyrenes, polyacrylamides, polyhydroxyalkyl acrylates, polyvinyl pyrrolidones, hydrophilic polyamides, polyesters and mixtures thereof. In a particularly useful embodiment, the substrate is cellulosic, especially ion-permeable crosslinked cellulose. In a presently preferred embodiment, the substrate comprises a regenerated cellulose membrane (CUPROPHAN™, Enka A G, Ohderstrasse, Germany) that is crosslinked with an epoxide, such as butanediol diglycidyl ether, further reacted with a diamine to provide amine functionality pendant from the cellulosic polymer, as described in U.S. Pat. No. 5,591,400, Example 4, incorporated herein by reference.

The above-described ionophore is preferably covalently bonded to the amine-functional cellulose substrate by any useful reactive technique, which may depend upon the chemical functionality of the ionophore.

The ionophore-functionalized cellulose substrate optionally can be adhesively bonded to the above-described backing membrane by any non-interfering adhesive. Preferably, the adhesive is essentially transparent to light used in excitation of the ionophore and to light emitted therefrom. One such useful adhesive is FLEXOBOND 431™ urethane adhesive (Bacon Co., Irvine, Calif.).

Alternatively, the functionalized substrate can be thermally fused to the membrane, providing the conditions necessary for thermal bonding are not detrimental to functioning of the ionophore, sensor and backing membrane.

The outermost layer of the multilayer sensing assembly, that is, the layer in immediate contact with the fluid to be monitored, preferably comprises an opacifying layer that optically isolates the ionophore in the sensing assembly. The opacifying agent can be applied before the ionophore-substrate component is applied to the backing membrane, as described in U.S. Pat. Nos. 5,081,041 and 5,081,042, incorporated herein by reference, or after the sensing component is attached to the substrate. It can be directly attached to the sensing element or it can be separate from the sensing element. In preferred embodiments, it is applied after the sensing component is attached to the backing membrane.

The overcoat preferably is a material that is permeable to the analyte of interest, such as a polymeric material as described above, containing an opaque agent such as carbon black, or carbon-based opaque agents, ferric oxide, metallic phthalocyanines, and the like. Such opaque agents are preferably substantially uniformly dispersed in the polymer in an amount effective to provide the desired degree of opacity to provide the desired optical isolation. A particularly useful opaque agent is carbon black. The overcoat can also be an ink coating on the sensing element applied using a variety of techniques, such as an inkjet technique or an ink-screening technique. The overcoat can also be a black membrane stapled or heat-staked to the cassette holding the sensing element. For example, it can be a black DURAPORE™ membrane (available from Millipore as a white membrane which is then treated with black ink) and heat sealed to the cassette, as described in U.S. Pat. Nos. 5,508,509 and 5,591,400, incorporated herein by reference. A presently preferred embodiment comprises carbon black dispersed in a matrix of epoxy-crosslinked dextran, as described in U.S. Pat. No. 4,919,891, incorporated herein by reference.

A presently preferred embodiment of sensor 28 comprises a sensing layer that includes 6,7-[2.2.2]-cryptando-3-[2"-(5"-carboxy)furyl]coumarin covalently bonded to a crosslinked amine functional cellulose membrane

(CUPROPHAN™, Enka A G, Ohderstrasse, Germany), the sensing layer being adhered to a polycarbonate backing membrane by FLEXOBOND 430™ urethane adhesive and the backing membrane having coated thereon CW14™ pressure-sensitive adhesive on a release liner.

Suitable pH sensors **30** are described in U.S. Reissue Pat. No. Re 31,879 (Lubbers), U.S. Pat. Nos. 4,798,738 (Yafuso), 4,824,789 (Yafuso), 4,886,338 (Yafuso), 4,999,306 (Yafuso), 5,081,041 (Yafuso), 5,081,042 (Yafuso), 5,127,077 (Iyer), 5,132,057 (Tomisaka), 5,403,746 (Bentsen), 5,508,509 (Yafuso), and 5,591,400 (Dektar et al.), the teachings of which are incorporated herein by reference.

The pH sensor **30** preferably comprises the following layers: (i) a backing membrane, (ii) a pressure-sensitive adhesive (PSA) coated on the backing membrane, (iii) a sensing element comprising a pH sensing component bound to a substrate, the substrate being attached to the membrane (e.g., by a non-interfering adhesive), and (iv) an outermost opacifying overcoat layer on the exposed surface of the substrate. With the exception of the pH sensor, these layers and the multilayer construction are essentially as described above for the potassium-ion sensor **28**.

Suitable pH sensing components include many well known pH indicators and/or functionalized derivatives of such indicators. Among the useful pH sensing components are hydroxypyrenetrisulfonic acid ("HPTS") and derivatives, e.g., salts, thereof, phenolphthalein, fluorescein, phenol red, cresol red, pararosaniline, magenta red, xylenol blue, bromocresol purple, bromphenol blue, bromothymol blue, metacresol purple, thymol blue, bromphenol blue, bromothymol blue, tetrabromophenol blue, bromchlorophenol blue, bromocresol green, chloropheno red, o-cresolphthalein, thymolphthalein, metanil yellow diphenylamine, N,N-dimethylaniline, indigo blue, alizarin, alizarin yellow GG, alizarin yellow R, congo red, methyl red, methyl violet 6B, 2,5-dinitrophenol, and/or the various functionalized derivatives of the above species. Sensing components for other ionic species can be made from organic species which include fluorescein, diiodofluorescein, dichlorofluorescein, phenosafranin, rose bengal, cosin I bluish, cosin yellowish, magneson, tartrazine, eriochrome black T, coumarin, alizarin, and others. The preferred pH sensing component is hydroxypyrenetrisulfonic acid (HPTS), derivatives of hydroxypyrenetrisulfonic acid, and mixtures thereof.

Additional suitable indicator components for use in the present invention include: 9-amino-6-chloro-2-methoxyacridine; 2',7'-bis-(2-carboxyethyl)-5-(and -6)-carboxyfluorescein; 2',7'-bis-(2-carboxyethyl)-5-(and -6)-carboxyfluorescein, acetoxymethyl ester; 2',7'-bis-(2-carboxyethyl)-5-(and -6)-carboxyfluorescein, acetoxymethyl ester; 5-(and -6)-carboxy-2',7'-dichlorofluorescein; 5-(and -6)-carboxy-2',7'-dichlorofluorescein diacetate; 5-(and -6)-carboxy-4',5'-dimethylfluorescein; 5-(and -6)-carboxy-4',5'-dimethylfluorescein diacetate; 5-carboxyfluorescein; 6-carboxyfluorescein; 5-(and -6)-carboxyfluorescein; 5-carboxyfluorescein diacetate; 6-carboxyfluorescein diacetate; 5-carboxyfluorescein diacetate, acetoxymethyl ester; 5-(and -6)-carboxyfluorescein diacetate; 5-(and -6)-carboxynaphthofluorescein; 5-(and -6)-carboxynaphthofluorescein diacetate; 5-(and -6)-carboxySNAFL®-1, succinimidyl ester {5'(and 6')-succinimidyl ester-3,10-dihydroxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; 5-(and -6)-carboxySNAFL®-2, succinimidyl ester {5'(and 6')-succinimidyl ester-9-chloro-3,10-dihydroxy-spiro[7H-

benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; carboxySNAFL®-1 {5'(and 6')-carboxy-3,10-dihydroxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; carboxySNAFL®-1 diacetate {5'(and 6')-carboxy-3,10-diacetoxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; carboxySNAFL®-2 {5'(and 6')-carboxy-9-chloro-3,10-dihydroxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; carboxySNAFL®-2 diacetate {5'(and 6')-carboxy-9-chloro-3,10-diacetoxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; carboxySNARF®-1 {5'(and 6')-carboxy-10-dimethylamino-3-hydroxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; carboxySNARF®-1, AM acetate {3-acetoxy-5'-acetoxymethoxycarbonyl-10-dimethylamino-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; carboxySNARF®-2 {5'(and 6')-carboxy-10-diethylamino-3-hydroxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; carboxySNARF®-2, AM acetate {3-acetoxy-5'-acetoxymethoxycarbonyl-10-diethylamine-3-hydroxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; carboxySNARF®-6 {5'(and 6')-carboxy-10-diethylamino-3-hydroxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; carboxySNARF®-X {5'(and 6')-carboxy-3-hydroxy-tetrahydroquinolizino[1,9-hi]spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; 5-chloromethylfluorescein diacetate; 4-chloromethyl-7-hydroxycoumarin; C1-NERF {4-[2-chloro-6-(ethylamino)-7-methyl-3-oxo-3H-xanthen-9-yl]-1,3-benzene-dicarboxylic acid}; dextran, BCECF, 10,000 MW, anionic {dextran, 2,7'-bis(2-carboxyethyl)-5-(and 6)-carboxy-fluorescein, anionic}; dextran, BCECF, 40,000 MW, anionic; dextran, BCECF, 70,000 MW, anionic; dextran, C1-NERF, 10,000 MW, anionic; dextran, C1-NERF, 70,000 MW, anionic; dextran, C1-NERF, 10,000 MW, anionic, lysine fixable; dextran, DM-NERF, 10,000 MW, anionic {dextran, 4-[2,7-dimethyl-6-(ethylamino)-3-oxo-3H-xanthen-9-yl]-1,3-benzene dicarboxylic acid, anionic}; dextran, DM-NERF, 70,000 MW, anionic; dextran, DM-NERF, 10,000 MW, anionic, lysine fixable; dextran, 7-hydroxycoumarin, 10,000 MW, neutral; dextran, 7-hydroxycoumarin, 70,000 MW, neutral; dextran, b-methylumbelliferone, 10,000 MW, neutral; dextran, b-methylumbelliferone, 70,000 MW, neutral; dextran, SNAFL®-2, 10,000 MW, anionic {dextran, 9-chloro-3,10-dihydroxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]3'-one, anionic}; dextran, SNAFL®-2, 70,000 MW, anionic {dextran, 10-dimethylamino-3-hydroxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one, anionic}; dextran, SNARF®-1, 10,000 MW, anionic; dextran, SNARF®-1, 70,000 MW, anionic; 1,4-dihydroxyphthalonitrile; DM-NERF {4-[2,7-dimethyl-6-ethylamino)-3-oxo-3H-xanthen-9-yl]1,3-benzene dicarboxylic acid}; fluorescein diacetate; 8-hydroxypyrene-1,3,6-trisulfonic acid, trisodium salt; naphthofluorescein; naphthofluorescein diacetate; SNAFL®-1 {3,10-dihydroxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}; and SNAFL®-1, diacetate {3,10-diacetoxy-spiro[7H-benzo[c]xanthene-7,1'(3'H)-isobenzofuran]-3'-one}.

In a presently preferred embodiment, HPTS indicator is covalently bonded to an amine-functional CUPROPHAN™ substrate that is bonded by FLEXOBOND 431™ polyurethane adhesive to a polycarbonate backing membrane. The sensing substrate is overcoated with an epoxy-crosslinked dextran matrix having carbon black dispersed therein.

Suitable carbon dioxide sensors **32** are described in U.S. Reissue Pat. No. Re 31,879 (Lubbers), U.S. Pat. Nos.

4,557,900 (Heitzmann), 4,824,789 (Yafuso), 4,849,172 (Yafuso), 4,867,919 (Yafuso), 4,919,891 (Yafuso), 5,127,077 (Iyer), 5,175,016 (Yafuso), 5,272,088 (Morlotti), 5,403,746 (Bentsen), 5,453,248 (Olstein), and 5,508,509 (Yafuso), incorporated herein by reference.

The carbon dioxide sensor **32** may also be in the form of a multilayer assembly. In one presently preferred embodiment, the sensing substrate layer of sensor **32** comprises a hydrophobic matrix in which is dispersed a plurality of hydrophilic particles or beads carrying a carbon dioxide sensing indicator. The indicator can be attached to or in the beads in any effective manner.

Because the beads are hydrophilic, they are adapted to receive and contain an aqueous solution of the indicator. "Hydrophilic" means a material, such as a polymeric substance, that retains a large fraction (e.g. greater than 20% of its weight) of water within its structure but does not dissolve in water. Hydrophilic materials useful as beads in carbon dioxide sensors include glass beads or hydrogels, polyacrylamide, cross-linked dextran, agarose, poly (hydroxyalkyl methacrylate), sulfonated polystyrene, and the like. A presently preferred hydrophilic bead material is SEPHADEX 75G™ cross-linked dextran (Pharmacia Biotech, Inc., Piscataway, N.J.).

In sensing carbon dioxide concentrations, examples of absorbance indicators that can be used include chlorophenyl red, bromo cresol purple, nitrophenol, bromo thymol blue, penachlorome, phenol red and the like. Useful fluorescence indicators for carbon dioxide include the sensors listed above that are useful for pH sensing, beta-methylumbelliferone, fluorescein and the like. A particularly useful carbon dioxide sensor is hydroxypyrene 3,6,8-trisulfonic acid, herein referred to as HPTS or hydroxypyrene trisulfonic acid and derivatives, e.g., salts of HPTS. The more preferred sensing component, particularly for sensing the concentration of carbon dioxide in blood, is selected from HPTS, derivatives of HPTS and mixtures thereof. The alkali and alkaline earth metal salts of HPTS are useful HPTS derivatives.

The hydrophobic matrix material in which the beads bearing a suitable indicator are dispersed is preferably transparent to excitation and emission wavelengths of light useful in sensing cassettes of the invention, and is otherwise inert to carbon dioxide, the absorbance or fluorescence indicator, and the beads. The hydrophobic matrix serves to isolate the indicator, while allowing carbon dioxide to diffuse therethrough. Suitable hydrophobic matrix materials include numerous silicones, such as silicone elastomer, room temperature vulcanizable (RTV) silicone rubber, heat vulcanizable silicone rubber, polydimethylsiloxane, poly (vinyl siloxane), silicone-polycarbonate copolymer, and the like, as well as perfluorinated (polyether) urethanes. Particularly preferred silicone matrix materials include PS 443™ vinyl-terminated dimethylsiloxane and PE1055™ polydimethylsiloxane, both commercially available from Petrarch Systems, Inc.

In a presently preferred embodiment, the carbon dioxide sensor **32** comprises a sensing layer including HPTS sensing dye on SEPHADEX 75G™ crosslinked dextran beads in a silicone matrix adhered to a polycarbonate backing membrane and overcoated with a opacifying layer comprising iron oxide pigment dispersed in a silicone matrix.

In an alternative embodiment, a solution of a suitable indicator dye can be formed in an aqueous buffer, and the solution can be emulsified with a liquid precursor of the hydrophobic polymeric matrix. Upon polymerization of the precursor, the emulsified indicator is essentially uniformly

dispersed throughout the polymer matrix. Indicator dyes and silicone polymers described above can be useful in the embodiment.

Suitable oxygen sensors **34** are described in U.S. Pat. Nos. 4,557,900 (Heitzmann), 4,849,172 (Yafuso), 4,867,919 (Yafuso), 4,919,891 (Yafuso), 5,043,285 (Surgi), 5,127,077 (Iyer), 5,296,381 (Yafuso), 5,409,666 (Nagel et al.), 5,453,248 (Olstein), 5,462,879 (Bentsen), 5,462,880 (Kane), 5,480,723 (Klainer), 5,498,549 (Nagel et al.), and 5,508,509 (Yafuso), incorporated herein by reference, and European Patent Application EP 585,212.

The oxygen sensor **34** may be in the form of a multilayer sensing assembly. In particular, the construction of the oxygen sensor **34** may closely resemble that of the carbon dioxide sensor **32**, in that the sensing layer comprises a sensing dye or indicator in a gas-permeable (e.g. silicone) matrix, and the overcoat layer may comprise a pigment in a silicone matrix. The gas-permeable matrix materials useful in the oxygen sensor **34** preferably can be the same as those previously described.

Useful oxygen sensing indicators comprise fluorescence indicators including one or more polynuclear aromatic compounds, derivatives of polynuclear aromatic compounds and the like. Examples of such polynuclear aromatic compounds include decacyclene, benzo-ghi-perylene and coronene. Oxygen indicators may include a mixture of tertiary butyl derivatives of such polynuclear aromatic compounds. Such indicators are more fully described in Yafuso, et al U.S. Pat. No. 4,849,172 which is incorporated in its entirety herein by reference.

Additional useful oxygen indicators include complexes of ruthenium(II), osmium(II), iridium(III), rhodium, rhenium, and chromium(III) with 2,2'-bipyridine, 1,10-phenanthroline, 4,7-diphenyl-1,10-phenanthroline, 4,7-dimethyl-1,10-phenanthroline, 4,7-disulfonated-diphenyl-1,10-phenanthroline, 2,2'-bi-2-thiazoline, 2,2'-bithiazole, 5-bromo-1,10-phenanthroline, and 5-chloro-1,10-phenanthroline, and complexes of Co(II), Cu(II), Pt(II), Pd(II) and Zn(II) with porphyrin, etioporphyrin, tetraphenylporphyrin, tetrafluorophenylporphyrin, tetrabenzporphyrin, tetrafluorobenzporphyrin, tetrachlorobenzporphyrin, mesoporphyrin IX diester, protoporphyrin IX dimethyl ester, and octaethylporphyrin. Ruthenium complexes are preferred, among the metal complexes.

The oxygen indicators may be covalently bonded to the polymeric materials or matrix materials included in the sensing composition. Such covalent bonding is preferably accomplished by providing an oxygen indicator component including a reactable group which reacts with a reactable group, preferably a different reactable group, present in one of the components of the precursor of the polymeric matrix material. Thus, during the formation of the polymeric matrix material, the above-noted reactable groups also react to covalently bond the oxygen indicator to the matrix material. Particularly useful oxygen indicator components include the above-noted polynuclear aromatic compounds derivatized to include a reactable group, such as a reactable group with functional carbon-carbon unsaturation. Vinyl derivatives of such compounds are particularly preferred.

Alternatively, the oxygen sensor can comprise a sensing element, an excitation means, and a detecting means, wherein the sensing means includes one or more, preferably one or two, monomeric indicator components, preferably located in, more preferably covalently bonded to, a matrix material, preferably a solid matrix material. Each of these monomeric indicator components is capable of providing a first emitted signal of a given wavelength in response to

exposure to a first excitation signal. Further, this sensing element is capable of providing a second emitted signal (due to emission by the excited state complex), preferably having a longer wavelength than the first emitted signal or signals, in response to a second excitation signal.

In a particularly useful embodiment, the indicator component is sensitive to the concentration of oxygen in a fluid and comprises one or more polynuclear aromatic compounds and/or one or more derivatives thereof. The polynuclear aromatic compound is preferably any fluorescent or absorbent, more preferably fluorescent, optical indicator of the polynuclear aromatic class. The polynuclear aromatic compound from which the indicator component is derived is still more preferably selected from the group consisting of perylene, decacyclene, benzoperylene (e.g., benzo[ghi]perylene), coronene, pyrene, porphycine, porphyrin, chlorin, phthalocyanine and derivatives and mixtures thereof. Since perylene and derivatives of perylene have a relatively reduced sensitivity to oxygen, other polynuclear aromatic compounds, such as those noted herein, are preferably employed when the analyte is oxygen. When an excimer component is to be utilized, the monomeric indicator component is preferably selected from one polynuclear aromatic compound, derivatives of the same one polynuclear aromatic compound and mixtures thereof. Excellent results are achieved if the polynuclear aromatic compound is benzo[ghi]perylene.

If desired, the basic polynuclear aromatic compound may be derivatized with one or more other groups, e.g., non-functional substituent groups such as alkyl groups, provided such derivatization does not substantially interfere with excited state complex provided emitted signal generation. Such derivatives are discussed in Nagel et al. U.S. Pat. No. 5,409,666 which is incorporated by reference. For example, the monomeric indicator component of a sensing element derived by covalently bonding vinyl benzo[ghi]perylene in an addition cure silicone polymer is said to be a derivative of benzo[ghi]perylene.

Monomeric components useful in sensor **34** may include, for example, two or more similar monomeric indicator components, two or more different monomeric indicator components, or one or more, preferably one, monomeric indicator components and one or more, preferably one, monomeric non-indicator components. Preferably, such monomeric components have no substantial detrimental effect on the sensing element, on the sensor system, on the analyte or on the medium to which the sensing element is exposed.

Examples of monomeric components which produce more preferred oxygen-sensing components include: (1) polynuclear aromatic monomeric components; (2) aliphatic or aromatic amine-containing or aromatic ether-containing monomeric components; and (3) aromatic nitrile monomeric components. More preferred exciplex components comprise at least one monomeric component selected from group (1) and at least one monomeric component selected from group (2). Alternatively, another more preferred component comprises at least one monomeric component selected from group (3) and at least one monomeric component selected from either group (1) or (2).

Examples of useful aromatic monomeric components (group 1) include biphenyl, naphthalene, phenanthrene, p-terphenyl, chrysene, benzpyrene, pyrene, dibenzanthrene, benzanthrene, anthracene, perylene, benzperylene, fluoranthene, coronene, quinoline, phenylquinoline, benzquinoline, quinoxaline, dibenzquinoxaline, benzquinoxaline, phthalimide, pyridine, phenazine,

dibenzphenzine, acridine, benzacridine and derivatives of these compounds. Examples of useful aliphatic or amine-containing or aromatic ether-containing monomeric components (group 2) include tetramethyl-p-phenylenediamine, dimethoxydimethylaniline, methoxydimethylaniline, diethylaniline, diphenylmethylamine, triethylamine, indole, dimethyltoluidine, tri-p-anisylamine, ditolylmethylamine, tritolylamine, triphenylamine, ethylcarbazole, trimethoxybenzene, tetramethoxybenzene and derivatives of these compounds. Examples of aromatic nitrile acceptor monomeric components (group 3) include benzonitrile, cyanonaphthalene, dicyanobenzene and derivatives of these compounds.

Any of these monomeric component pairs can be tethered and/or covalently bonded to a matrix material, e.g., silicone.

In a presently preferred embodiment, the oxygen sensor **34** comprises a sensing layer including vinyl benzo[ghi]perylene covalently bonded to a silicone matrix comprising crosslinked polyalkyl(aryl)hydrosiloxane, adhered to a polycarbonate backing membrane and overcoated with a silicone matrix of dispersed carbon black.

In a particularly presently preferred embodiment, the sensor **28** is provided as a multilayered laminate attached to the cassette **12** in the cavity **27**. The ion sensor **28** and the pH sensor **30** are preferably placed near the first "inlet" portion **20** of the fluid chamber **18** so that they will be positioned at the bottom half of the cassette during calibration. This assures that the sensors **28** and **30** will be exposed to liquid during calibration. The sensors **32** and **34** are less sensitive to the need for immersion in liquid during calibration.

Alternatively, the cassette **12** can include sensors for potassium, sodium, calcium and glucose, wherein these sensors use essentially the same chemistry as described above. For example, potassium, sodium, and calcium ion detection can use suitable ionophoric coumarocryptands according to Formula A, wherein the size of the cryptand cage is specific for each ion. A suitable glucose sensor can comprise any one of the oxygen sensors described above, modified by the presence of the enzyme glucose oxidase. Glucose detection can be based upon depletion of oxygen during enzymatic oxidation of glucose, as described, e.g., in U.S. Pat. No. 5,518,694, incorporated herein by reference. With minor modifications to the optical train described below, the measuring device **14** can be adapted to accommodate these alternate sensors. In some applications, e.g., cardiovascular blood parameter monitoring, it may be advantageous to use both types of measuring devices.

Other sensors that can be useful in the cassette **12** may include, e.g., a fluorescence-based temperature sensor, such as can be prepared by immobilizing a ruthenium based indicator such as ruthenium(II)(diphenylphenanthroline)₃(dimethylsilylpropanesulfonate)₂ in an oxygen-impermeable matrix, e.g., poly(methylmethacrylate).

A hole in the casing **16** is located between the pH sensor **30** and the carbon dioxide sensor **32**. A thermistor-receiving well **36** is fixed to the casing **16** and extends over the hole. The well **36** has a hat-shaped configuration with a brim that is bonded by an adhesive to wall sections of the casing **16** that face the central portion **24** of the fluid chamber **18**. A suitable adhesive is an acrylic urethane adhesive such as "UV Cure" brand adhesive from Loctite Corporation. The well **36** is preferably made of a corrosion-resistant material having a thermal conductivity similar to metal, such as 0.004 inch (0.1 mm) thick titanium. As illustrated for example in FIGS. **5** and **6**, the well **36** protrudes into the central portion **24** of the fluid chamber **18** to provide intimate thermal contact with fluid therein.

The casing 16 also includes a generally oval-shaped rim 40 that circumscribes the recess 26 and extends outwardly in a direction away from the longitudinal axis of the casing 16. As can be appreciated by reference to FIG. 4, the major axes of the oval-shaped recess 26 and the surrounding rim 40 coincide and extend across the center of the sensors 28, 30, 32, 34 and the well 36 and are also parallel with the longitudinal axes of the casing 16 and the fluid chamber 18.

A semi-cylindrical alignment key 42 is integrally connected to an inner wall of the rim 40. Preferably, the alignment key 42 is oriented such that a reference plane that is perpendicular to the longitudinal axis of the casing 16 and extends equidistant between the sensor 32 and the sensor 34 also bisects the key 42 along its central diametrical plane.

The cassette 12 further includes a first, male coupling 44 for detachably connecting the casing 16 to the measuring device 14. The coupling 44 has a convex, generally U-shaped configuration in directions perpendicular to the longitudinal axis of the casing 16 as depicted in FIGS. 2 and 3. The coupling 44 includes the aforementioned central section of the casing 16 and opposed leg portions 46 that extend outwardly from the casing 16 in a direction away from the direction of outward extension of the rim 40. Each leg portion 46 includes a pair of support sections having flat, coplanar outer surfaces 47 (see, e.g., FIGS. 2, 3 and 7; omitted in FIG. 4) that are parallel to the outer side of the respective leg portion 46. Preferably, the outer surfaces 47 of the opposed leg portions 46 converge as the casing 16 is approached and extend along respective reference planes that are oriented at an angle in the range of about 28 degrees to about 32 degrees relative to each other. More preferably, the outer surfaces 47 extend along respective reference planes that are oriented at an angle of about 30 degrees relative to each other.

A flange 48 is integrally connected to the outer end of each leg portion 46. The flanges 48 lie in a common reference plane that is parallel to the longitudinal axis of the casing 16. The leg portions 46 are somewhat flexible and can be moved slightly toward each other under the influence of finger pressure, but also have sufficient memory to quickly and repeatedly return to their original, normal orientation as shown in the drawings once finger pressure is released.

An outer, central end region of each leg portion 46 is integrally connected to a wedge-shaped tab 50 that lies between the support sections. The tabs 50 extend away from each other and outwardly from the respective leg portions 46 along respective reference planes that are oriented at an angle of about 80 degrees relative to each other. Additionally, a distal edge of each tab 50 extends in a reference plane that is oriented at an angle of 25 degrees relative to the direction of extension of the flanges 48. Outermost edges of the tabs 50 are spaced outwardly relative to adjacent regions of respective leg portions 46 and lie in a common reference plane that is between the longitudinal axis of the casing 16 and the aforementioned reference plane containing the flanges 48.

Preferably, the casing 16 is made of a relatively clear plastic material such as medical grade polycarbonate, and is constructed of two or more initially separate pieces that are injection-molded and then joined together. An example of a suitable two-piece construction is shown in FIG. 7. In FIG. 7, one piece of the casing 16 includes the recess 26 and the rim 40, and carries the four sensors 28, 30, 32, 34 and the second piece includes the leg portions 47, the inlet and outlet ports and other elements as shown. The pieces may be connected together by ultrasonic welding, solvent welding or adhesive bonding. Of course, other constructions (such as

an integral, one-piece construction or a three-piece construction) are also possible.

As illustrated in FIGS. 1 and 4-6, the casing 16 has a first external threaded section that surrounds the inlet port of the first portion 20. The first threaded section is preferably constructed to matingly connect to an internally threaded Luer-type connector such as the male Luer connector 52 shown in FIG. 5 when the cassette 12 is in use for measuring parameters of fluid flowing through the chamber 18. The connector 52 has a ribbed portion for providing an interference-fit coupling to a section of flexible tubing 54 that directs fluid toward the chamber 18.

A second external threaded section surrounds the outlet port of the second fluid chamber portion 22. As shown in FIG. 5, a fitting 56 has an internal threaded section that matingly receives the second threaded section. The fitting 56 optionally includes a rearwardly extending collar having a radially inwardly extending rib 58. The casing 16 has a circumscribing, radially outwardly extending rib 60 adjacent the second threaded section that functions as a stop and provides a physical interference to the rib 58 in order to prevent detachment of the fitting 56 under normal circumstances whenever the fitting 56 is partially unthreaded from the casing 16.

The fitting 56 also includes another internal threaded section that is constructed to matingly receive a female Luer connector 62 (FIG. 5) when the cassette 12 is used with the measuring device 14 for measuring parameters of fluid flowing through the chamber 18. A section of flexible tubing 64 is connected in interference fit relation to a ribbed section of the connector 62 for directing the flow of fluid that exits the chamber 18.

FIG. 6 is a depiction for exemplary purposes of the cassette 12 during calibration of the sensors 28, 30, 32 and 34. During calibration, a gas filter assembly 66 replaces the connector 52 that is illustrated in FIG. 5 and has an internal threaded section that matingly receives the threaded section surrounding the inlet port of the first fluid chamber portion 20. An opposite end of the gas filter assembly 66 has a gas inlet opening 68 that is provided with a connector constructed with a partial screw thread. This connector is adapted to couple with a tubing connector (not shown) and the tubing, in turn, is connected to a source of calibration gas.

The gas filter assembly 66 has an enlarged, cylindrical central housing section that contains a disk-shaped section of filtering membrane 70. Preferably, the membrane 70 is made of a hydrophobic material (such as polytetrafluoroethylene) that is sterilized by autoclaving or a material (such as a modified acrylic) that is sterilized by radiation. A suitable modified acrylic material is VERSA-PORE "H" brand membrane from Gelman Sciences. A network of nested, concentric circular channels and intersecting radial channels is provided throughout walls facing both sides of the membrane 70 to facilitate the passage of calibration gas through substantially all of the various regions of the membrane 70.

The gas filter assembly 66 also includes an outlet that receives a section of sparger tube 72. An example of a suitable tube 72 is a tube made of polyetheresterketone, having a 0.003 inch (0.075 mm) inner diameter and 0.012 inch (0.3 mm) outer diameter, from Zeus Products. A plug 74 surrounds the sparger tube 72 and secures the sparger tube 72 to the outlet of the filter assembly 66 in sealed relation. Examples of suitable material for the plug 74 include polycarbonate if the cassette 12 is to be sterilized by radiation and acrylic resin if the cassette 12 is to be sterilized by autoclaving.

When packaged for shipment to the end user, the cassette **12** preferably is provided with the filter assembly **66** as well as a cap **78** (FIG. **6**) and a quantity of calibration fluid **80** received in the fluid chamber **18**. For shipping purposes, the cap **78** is tightly threaded into the outlet section of the fitting **56** in sealed relation and the fitting **56** is tightened against the casing **16** (as shown in its orientation depicted in FIG. **5**) to provide a fluid seal between the fitting **56** and the casing **16**. Although not shown in the drawings, a shipping cap is securely coupled to the outer end of the filter assembly **66** for substantially preventing contaminants from entering the inlet opening **68** during transit and initial handling.

During calibration, the cassette **12** is oriented in an upright, preferably vertical position as shown in FIG. **6** and the fitting **56** is partially unthreaded by rotating the fitting **56** relative to the casing **16** in an arc about the longitudinal axis of the latter. During such motion, the fitting **56** is moved from a first position closing a gas outlet port and to a second position for opening or venting the gas outlet port. The ribs **58**, **60** prevent inadvertent separation of the fitting **56** from the casing **16**. Loosening of the fitting **56** relative to the casing **16** enables gas to flow from the second fluid chamber portion **22** to the atmosphere through the gas outlet port as depicted by the arrows in FIG. **6**. Optionally, the outer peripheral wall of the rib **60** is provided with one or more channels **59** (see FIGS. **1** and **7**) that extend in a direction parallel to the longitudinal axis of the casing **16** in order to facilitate the discharge of gas from the upper outlet port of the fluid chamber **18**.

The quantity of calibration fluid **80** is preferably selected so that the level of fluid **80** in the chamber **18** during calibration extends across a lower section of the expansion zone portion **25** as illustrated in FIG. **6**. Such a level decreases the likelihood that a portion of the calibration fluid **80** may escape through the upper outlet port, and yet assures that the fluid **80** fully covers the sensors **28**, **30**, **32**, **34**. Advantageously, the frustoconical configuration of the expansion zone portion **25** facilitates the rupture of calibration gas bubbles that pass through the fluid **80** in order to further decrease the likelihood of escape of the fluid **80** from the chamber **18**. In addition, the hydrophilic surface on the wall sections of the chamber central portion **24** facilitates the smooth passage of calibration gas bubbles past the sensors **28**, **30**, **32**, **34**. Optionally, an anti-foaming agent may be used in addition to or in place of the hydrophilic surface.

The calibration steps are described in further detail below. Once calibration is complete, the filter assembly **66** is removed from the casing **16** and replaced with the connector **52** that is depicted in FIG. **5**. Additionally, the fitting **56** is tightened to its orientation as shown in FIG. **5** to provide a fluid leakage-resistant seal between the fitting **56** and the casing **16**. The cap **78** that is shown in FIG. **6** is also removed and replaced with the connector **62** (FIG. **5**), and the outer end of the connector **62** is coupled to the section of tubing **64**. As will be described in more detail below, the sections of tubing **54**, **64** enable the flow of fluid (such as blood) into and out of the chamber **18** for measuring parameters of the fluid.

The measuring device **14** that is shown in FIGS. **1** and **5** is also illustrated in more detail in FIGS. **8–20**. The device **14** includes a two-part elongated housing **200** that is illustrated as it appears before assembly in FIG. **9**. The two parts could be held by internal barbed connectors (for snap-together assembly) or by screws. Preferably, the housing **200** is made of an impact-resistant plastic material such as a mixture of polycarbonate and acrylonitrile-butadiene-styrene (“ABS”) polymer, and has a smooth outer surface for

facilitating disinfection. Optionally, the inner surface of the housing **200** is coated with an electromagnetically-compatible shielding material.

The measuring device **14** includes a second, female coupling **202** that is optionally made of a metallic material such as anodized aluminum. The coupling **202** has a concave recess with a generally U-shaped configuration in directions perpendicular to the longitudinal axis of the housing **200**. The recess includes two flat, opposed sidewall sections **204** that are interconnected by a central bight section **206** (see, e.g., FIG. **1**). Preferably, the opposed sidewall sections **204** converge as the bight section **206** is approached and extend along respective reference planes that are oriented at an angle in the range of about 28 degrees to about 32 degrees relative to each other. More preferably, the sidewall sections **204** extend along respective reference planes that are oriented at an angle of about 30 degrees relative to each other.

An outer edge portion of each sidewall section **204** has an elongated groove **208** that extends in a direction parallel to the longitudinal axis of the housing **200**. Optionally, one end of the coupling includes a recess **209** (see FIGS. **8–11**) for guiding fluids that may be present from accidental spills and the like away from the area near the sensors **28**, **30**, **32**, **34**.

The measuring device **14** includes a fiber terminal block assembly **210** (see, e.g., FIG. **9**). The fiber block assembly **210** includes a fiber block insert plate **212** and a fiber block end plate **214** that are joined together by a pair of machine screws as depicted in FIG. **10**. Preferably, the insert plate **212** is made of polycarbonate and the end plate **214** is made of aluminum.

The underside of the insert plate **212** is secured to the top of the coupling **202** by four machine screws that are illustrated in FIG. **10**. The underside of the insert plate **212** includes a generally oval-shaped protrusion **216** (see, e.g., FIG. **12**) having a major axis that is parallel to the longitudinal axis of the housing **200**. The protrusion **216** includes a semi-cylindrical keyway **218** with a diameter that is just slightly larger than the diameter of the cassette key **42**. Preferably, the keyway **218** is oriented such that a reference plane that bisects the keyway **218** along its central diametrical plane is also perpendicular to the longitudinal axis of the housing **200**.

The protrusion **216** has a side wall that is matingly received and extends through an oval-shaped opening that is centrally located in the bight section **206** of the coupling **202**. The outer side wall of the protrusion **216** has an oval-shaped configuration in bottom view that is complementary in shape to the oval-shaped configuration of the inner sidewall of the rim **40** of the cassette **12** (see, e.g. FIG. **4**).

FIG. **5** illustrates the cassette **12** and a portion of the measuring device **14** when coupled together. When the measuring device **14** is connected to the cassette **12**, the tabs **50** of the cassette **12** are received in respective grooves **208** of the measuring device **14**. Furthermore, the outer surfaces **47** of the cassette support sections flatly contact the respective flat, facing surfaces of the measuring device sidewall sections **204**. As can be appreciated, the first coupling **44** that is connected to the cassette casing **16** and the second coupling **202** that is connected to the housing **200** together represent a connector for releasably connecting the cassette **12** to the measuring device **14**.

During assembly of the cassette **12** to the measuring device **14**, the casing **16** is pushed in a direction toward the protrusion **216**, and during such pushing motion the outer surfaces of the wedge-shaped tabs **50** function as a cam to deflect the leg portions **46** inwardly and toward each other until such time as the outer edge of each tab **50** is adjacent

the respective groove 208. Once the outer edges of the tabs 50 are adjacent the grooves 208, the inherent bias of the leg portions 46 causes the latter to spread apart and return to their normal configuration in a snap-like manner such that the outer edges of the tabs 50 are received in the grooves 208 and thereby couple the cassette 12 to the measuring device 14.

The protrusion 216 has an outer end portion that matingly and snugly fits within the recess 26 when the measuring device 14 and the cassette 12 are connected together. In addition, when the measuring device 14 and the cassette 12 are connected together, the key 42 of the cassette 12 fits within the mating keyway 218 of the measuring device 14. Advantageously, the key 42 substantially prevents coupling of the cassette 12 to the measuring device 14 when an attempt is made to couple the cassette 12 to the measuring device 14 in an opposite fashion (i.e., in such a manner that the cassette 12 is oriented 180 degrees about a vertical reference axis from its orientation shown in FIG. 5).

The cassette 12 is detached from the measuring device 14 by squeezing the leg portions 46 toward each other; the cassette 12 is then moved away from the device 14 until the tabs 50 clear the grooves 208. Preferably, the outer sides of the coupling 202 are each provided with a fingertip-sized recess 207 to enhance the user's grip on the outer edge of the cassette flanges 48 during detachment of the cassette 12 from the measuring device 12. The recesses 207 also help guide the user's finger toward a position for engaging the center of the flanges 48 at a location close to the tabs 50.

The fiber terminal block assembly 210 includes an upstanding polycarbonate tube 220 that is solvent-welded to the insert plate 212 and receives a thermistor assembly. As shown in FIGS. 5 and 10, the thermistor assembly includes a thermistor support 222 having a central internal passageway and a necked-down lower end section. A thermistor 224 is mounted partially within a cavity of the lower end section of the thermistor support 222 and has a pair of leads that extend through the passageway. An example of a suitable thermistor is part no. SC30BF103A-L8 from Thermometrics.

A cap 226 (see FIG. 5) made of a material such as stainless steel covers the thermistor 224 and is secured by potting compound to a circumscribing side wall of the lower end section of the support 222. As an alternative construction, however, the cap may fit in a recess formed in the lower end section of the support 222. The cap 226 has a thermal conductivity that is high relative to the thermal conductivity of the adjacent plastic material. The potting compound (such as no. H20 or no. 930-4 from Epo-tek) substantially fills the space between the inner surface of the cap 226 and the external surface of the thermistor 224 and facilitates heat transfer between the cap 226 and the thermistor 224.

The support 222 is received in an internal, generally cylindrical passage of the tube 220, and an O-ring 228 located in a peripheral groove of the support 222 provides a fluid-resistant seal. The support 222 is kept in the passage of the tube 220 by a retainer 230 having an upper threaded section that is threaded into a mating threaded section of the tube 220.

The passage of the tube 220 narrows to a somewhat smaller diameter near its lower end and presents an annular shoulder 232 that is depicted in FIG. 5. The support 222 includes an upper, circumscribing cylindrical rib 234 having an outer diameter that is larger than the internal diameter of the tube passage in regions beneath the shoulder 232. In addition, a helical compression spring 236 (see also FIG. 10)

extends between and bears against the lower end of the retainer 230 and an upwardly facing annular wall of the rib 234. The spring 236 yieldably biases the support 222 in a downwardly direction viewing FIG. 5 such that the rib 234 comes to rest against the shoulder 232 when the measuring device 14 is not coupled to the cassette 12.

When the cassette 12 and the measuring device 14 are not coupled together, the lower end of the thermistor cap 226 protrudes beneath the bottom of the plate 212 a distance that is greater than the depth of the well 36 of the cassette 12 relative to the top surface of the recess 26 when viewing FIG. 5. As such, as the cassette 12 is coupled to the measuring device 14, the cap 226 contacts the bottom of the well 36 and urges the support 222 in an upwardly direction against the bias of the spring 236. Once the coupling 44 and the coupling 202 are connected together, the spring 236 tends to retain the bottom of the cap 226 in a position of close contact with the bottom of the well 36 to increase the effective contact area between the cap 226 and the well 36 and reduce thermal resistance. The inner configuration of the well 36 is preferably somewhat complementary and more preferably is closely complementary to the external configuration of the cap 226 in order to improve heat transfer along a path from the fluid in the central portion 24 of the chamber 18, across the joint between the well 36 and the cap 226, and ultimately to the thermistor 224.

With reference to FIG. 12, the insert plate 212 includes four holes 238, 240, 242 and 244 that extend through the protrusion 216 and are arranged in spaced-apart relation along the major axis of the oval-shaped protrusion 216. In addition, the bottom plate includes three holes 246, 248, 250 that are spaced from the protrusion 216 and extend through a raised platform 252 (see FIGS. 10 and 13) located on an upper, outer end section of the insert plate 212. As shown in FIG. 13, a disk 254 made of an optical reference material is placed in the hole 248 and is secured against a shoulder of the hole 248 by a set screw 256 that is threaded into a lower threaded section of the hole 248. The optical reference disk 254 is preferably made of a fluorescing material (such as a 0.002 weight percent fluorescent material in polycarbonate). A suitable fluorescent material is a high quantum efficiency fluorescent dye, such as 'MACROLUX 10GN' brand material from Bayer.

The holes 246, 250 are identical to the hole 248 and each receives a set screw similar to set screw 256 and an optical reference disk 254a, 254b (shown numbered only in FIG. 15). The optical reference disk 254a in the hole 246 is identical to the optical reference disk 254. The optical reference disk 254b in the hole 250 is similar to the optical reference disk 254 but is instead preferably made of a 0.0035 weight percent fluorescent material dissolved in polycarbonate.

The end plate 214 of the fiber terminal block assembly 210 has nine holes 258-274 that are numbered in FIG. 14. The holes 258-274 are arranged in an array of three rows with three holes in each row. The holes 258-274 as well as the holes 238-250 are each adapted to receive a ferrule surrounding an end of a bundle of optical fibers. An exemplary ferrule is designated by the numeral 276 in FIG. 13. An example of a suitable material for the ferrules received in holes 246-250 is brass or stainless steel, an example of a suitable material for the ferrules received in holes 238-244 is stainless steel and an example of a suitable material for the ferrules received in holes 258-274 is brass or aluminum. Optionally, the platform of the bottom plate includes three small threaded openings (see FIGS. 10 and 13), each of which intersects with one of the holes 246, 248, 250. Those

small openings are each adapted to threadably receive a small set screw (not shown) that secures the ferrule in the adjacent hole **246**, **248**, **250** to the platform in order to facilitate assembly.

The fiber terminal block assembly **210** includes a number of bundles of optical fibers. The optical fiber bundles are omitted from all of the views except that an exemplary network (not accurately depicted) of bundles is shown in FIG. **9** for purposes of explanation. A schematic of the various optical fiber bundles is illustrated in FIG. **15** and shows the true preferred various paths of the optical fiber bundles between the holes **238–250** and the holes **258–274**.

In more detail, and with reference to FIG. **15**, the fiber terminal block assembly **210** includes a first bundle of optical fibers **280** that extends between the hole **240** and the hole **260**, a second bundle of optical fibers **281** that extends between the hole **240** and the hole **270**, a third bundle of optical fibers **282** that extends between the hole **246** and the **270**, a fourth bundle of optical fibers **283** that extends between the hole **242** and the hole **270**, a fifth bundle of optical fibers **284** that extends between the hole **242** and the hole **258**, a sixth bundle of optical fibers **285** that extends between the hole **246** and the hole **264**, a seventh bundle of optical fibers **286** that extends between the hole **244** and the hole **262**, an eighth bundle of optical fibers **287** that extends between the hole **244** and the hole **274**, a ninth bundle of optical fibers **288** that extends between the hole **250** and the hole **274**, a tenth bundle of optical fibers **289** that extends between the hole **250** and the hole **266**, an eleventh bundle of optical fibers **290** that extends between the hole **248** and the hole **226**, a twelfth bundle of optical fibers **291** that extends between the hole **248** and the hole **272**, a thirteenth bundle of optical fibers **292** that extends between the hole **238** and the hole **272** and a fourteenth bundle of optical fibers **293** that extends between the hole **238** and the hole **268**.

A suitable optical fiber for each of the various bundles **280–293** is a fiber having a nominal outer diameter of 0.0022 inch (56 micron) or 0.0020 inch (51 micron), with core glass of Schott LF5 or F2, clad glass of Schott 8250 Corning 7056 or 7052 and a clad thickness of 0.00008 to 0.00012 inch (2–3 microns). Preferably, although not necessarily, all of the optical fibers in the various bundles **280–293** are identical. Of course, other types of fibers and number of fibers in each of the bundles **280–293** may be different from the description set out above.

Preferably, an optical adhesive such as “Epo-tek” brand optical epoxy no. 301 or 301-2 is used to secure bundled ends of the fibers into the ferrules (such as ferrule **276**) to present optical apertures. The bundles **280–293** are preferably coated with a non-fluorescing, optically opaque stiff plastic or rubber material. Ends of some of the bundles **280–293** are intermixed to present bifurcated sections such that the optical fibers received in one ferrule may lead to different ferrules at the opposite ends of the fibers as depicted schematically in FIG. **15**. Moreover, the optical fibers of each optical aperture are randomized (i.e., the fibers of each optical aperture are spatially well-mixed even in instances where the fibers pass through bifurcated sections). For example, the fibers of the optical aperture in hole **240** are randomized such that no circular area having a diameter of 0.020 inch (0.5 mm) or greater exists that has less than three fibers originating from either of the bundles **280**, **281**.

Examples of a suitable number of optical fibers for the middle region of each bundle **280–293** are set out below in Tables Ia and Ib. (For purposes herein, the “middle region” shall mean a central section of the bundle along its length at

a location between its ends and any bifurcated sections.) Tables Ia and Ib identify the target number of optical fibers and overall diameter of the optical aperture (which may include fibers originating from one or more than one bundle) in the ferrules placed in each of the holes **238–250** and **258–274**. The values in Tables Ia and IIa are based on optical fibers having an outer diameter of 56 microns as mentioned above and a packing fraction of 70 percent (i.e. the area occupied by the fibers divided by the overall, circular area of the optical aperture). The values in tables Ib and IIb are based on optical fibers having an outer diameter of 51 microns as mentioned above and a packing fraction of 75 percent. Since small optical fibers are difficult to count during assembly, the manufacturer may instead prefer to measure the diameter of a bundle to more quickly determine by estimation the number of optical fibers in the bundle.

TABLE Ia

Bundle Number	Target Number of Optical Fibers Middle Region of Bundle
280	1150
281	375
282	200
283	375
284	1150
285	600
286	1150
287	375
288	200
289	600
290	600
291	200
292	375
293	1150

TABLE Ib

Bundle Number	Target Number of Optical Fibers Middle Region of Bundle
280	1472
281	560
282	282
283	560
284	1472
285	847
286	1472
287	560
288	282
289	847
290	847
291	282
292	560
293	1472

TABLE IIa

Hole Number	Target Number of Optical Fibers in Optical Aperture	Diameter of Optical Aperture	
		Inch	mm
238–244	1525	0.104	2.64
246–250	800	0.076	1.93
258–262	1150	0.089	2.26
264	600	0.067	1.70
266	1200	0.093	2.36

TABLE IIa-continued

Hole Number	Target Number of Optical Fibers in Optical Aperture	Diameter of Optical Aperture	
		Inch	mm
268	1150	0.089	2.26
270	950	0.081	2.06
272-274	575	0.064	1.62

TABLE IIb

Hole Number	Target Number of Optical Fibers in Optical Aperture	Diameter of Optical Aperture	
		Inch	mm
238-244	2023	0.105	2.66
246-250	1129	0.078	1.98
258-262	1472	0.089	2.26
264	847	0.068	1.73
266	1694	0.096	2.44
268	1472	0.089	2.26
270	1402	0.086	2.18
272-274	842	0.067	1.70

The measuring device 14 also has an optics assembly 300 which includes a block 302 as well as nine optic subassemblies 304-312 (which are numbered in FIG. 16 only). The block 302 has nine cylindrical cavities symmetrically arranged in three rows with three cavities in each row, and one of the optic subassemblies 304-312 is received in each cavity. Preferably, the block 302 is made of a material having a thermal conductivity similar to metal. An example of a suitable material is aluminum. The block 302 could also be made of a ceramic material having an appropriate thermal conductivity.

The optic subassemblies numbered 310, 311 and 312 are excitation optic subassemblies, and the subassembly 312 is illustrated in exploded view in FIG. 16 for exemplary purposes. The subassembly 312 includes a first optical retainer 314 and a second optical retainer 316 which is identical to the first retainer 314. When the retainers 314, 316 are connected together in the manner shown in FIG. 10, internal wall sections of the retainers 314, 316 together present an elongated chamber 318 having a somewhat cylindrical overall configuration with a central, longitudinal reference axis 320.

The first retainer 314 is shown alone in FIGS. 17-19. The retainer 314 has internal wall sections that define a first chamber portion 322 (FIGS. 17 and 18), a second chamber portion 324, a third chamber portion 326 and a fourth chamber portion 328. When the retainers 314, 316 are connected together in the manner shown in FIG. 10, the first chamber portions 322 of the retainers 314, 316 present a generally cylindrical light inlet port having a shape for enshrouding a light emitting diode (as described in more detail below), the second chamber portions 324 present a generally cylindrical subchamber for receiving an optical filter, the third chamber portions 326 present a subchamber for receiving an optical lens, and the fourth chamber portions 328 present a generally cylindrical light outlet port. A small cylindrical opening extends between and communicates the light inlet port with the optical filter subchamber, and a second, small cylindrical opening extends between

and communicates the optical filter subchamber with the lens subchamber.

The wall sections of the retainer 314 defining the chamber portions 324, 326 are each connected to a first set of one or more ribs 330. In the embodiment shown in the drawings, the first set includes two spaced apart ribs 330 that extend in a direction parallel to the central axis 320. Each of the ribs 330 (see, e.g. FIG. 18) preferably extends the entire length of the respective chamber portions 324, 326 and has an outermost deformable tip portion 332 that is shown in more detail in FIGS. 19a. The deformable tip portion 332 may be elastomeric (i.e., it self-returns to its original shape after the force causing the deformation is removed) or non-elastomeric.

The retainer 314 also includes a second set of one or more ribs 334. In the embodiment shown in the drawings, the second set of ribs 334 consists of a single rib that is located along a radial wall section of the third chamber portion 326 directly adjacent the fourth chamber portion 328. The rib 334 has an overall configuration of a triangle in side view as illustrated in FIGS. 17 and 17a, and also has an outermost deformable (elastomeric or non-elastomeric) tip portion 336 that is shown enlarged in FIGS. 17a and 17b.

The second chamber portion 324 and the third chamber portion 326 also include a third set of one or more ribs 338 that extend in reference planes perpendicular to the central axis 320. Six ribs 338 are shown in the embodiment illustrated in FIGS. 17-19. Four of the ribs 338 are located in the second chamber portion 324, and are arranged in opposing fashion such that one pair of ribs 338 extend in a first common reference plane perpendicular to the central axis 320 and the other pair of ribs 338 extend in a second reference plane perpendicular to the central axis 320. The two remaining ribs 338 are located in the third chamber portion 326 and extend in another common reference plane that is perpendicular to the central axis 320. An exemplary rib 338 is shown enlarged in FIG. 17c, and each rib 338 includes an outermost deformable tip portion 340 that may be either elastomeric or non-elastomeric.

The retainer 314 has a symmetrical, diametrical wall section with a pair of pegs 342 and a pair of matching holes 344. One peg 342 and one hole 344 have parallel central reference axes that extend in a common plane perpendicular to the central axis 320. Similarly, the other peg 342 and the other hole 344 also have respective central reference axes that extend in a common plane that is perpendicular to the central axis 320 but spaced from the aforementioned reference plane containing the central axes of the other peg 342 and hole 344.

The wall sections defining the third chamber portion 326 also include an annular, chamfered wall section 346 that extends in a semi-circular path about the central axis 320. The lens subchamber consequently has an overall configuration that includes a generally cylindrical portion and an aligned frustoconical portion having common central axes that are collinear with the central axis 320. Since the retainer 316 is identical to the retainer 314, a detailed description of the retainer 316 need not be provided.

When the retainers 314, 316 are connected together in the manner shown in FIG. 10, the pegs 342 of the retainer 314 are received in the holes 344 of the retainer 316 and the pegs 342 of the retainer 316 are received in the holes 344 of the retainer 314. Preferably, at least one mating pair of the pegs 342 and holes 344 are closer to one end of the subassembly 312 than the other end. As such, if an attempt is made to connect the retainers 314, 316 together in such a manner that the first chamber portions 322 are not opposed from each

other and instead are located at opposite ends, the ends of the retainers **314**, **316** will not be aligned and provide a readily apparent visual indication to the assembler that the retainers **314**, **316** have not been properly oriented with respect to each other.

A plano-convex lens **348** (see FIGS. **16** and **16a**) is received in the third chamber portions **326**, and includes a cylindrical portion having an outer peripheral cylindrical wall **350** and a convex, dome-shaped portion having an outer wall **352** with a configuration of a partial sphere. The central diametrical axis of the dome-shaped portion and the central axis of the cylindrical portion lie along a common reference axis known as the optical axis of the lens **348**. An example of a suitable lens has a 6 mm focal length and a diameter of 6 mm such as part no. 45078 from Edmund Scientific. Preferably, the orientation of the chamfered wall section **346** of the third chamber portions **326** (relative to the central axis **320**) is somewhat similar to the orientation of the engaged region of the outer wall **352** (relative to the central axis of the dome-shaped portion) in areas where the sections **346** contact the outer wall **352** as the retainers **314**, **316** are assembled together.

During assembly, and as the retainers **314**, **316** are closed around the lens **348**, the cylindrical wall **350** comes into contact with the ribs **330**, **334**, **338** in interference fit relation. As the retainers **314**, **316** approach a fully closed, connected together orientation, the tip portions **332**, **336**, **340** in the lens subchamber crush and deform while exerting respective forces on the lens **348** in certain directions. More specifically, as the retainers **314**, **316** are closed, the tip portions **332**, **340** of the first and third set of ribs **330**, **338** direct a force on the lens **348** in a radially inwardly direction toward the central axis **320**, while the tip portions **336** of the second set of ribs **334** direct a force on the lens **348** having vector components extending in radially inward directions toward the central axis **320** as well as vector components extending parallel to the central axis **320** in directions toward the chamfered wall sections **346**.

The orientation of the ribs **330**, **334**, **338** that are located in the third chamber portions **326** is selected to guide the lens **348** into proper concentric relationship so that the optical axis of the lens **348** is aligned in parallel with and preferably precisely collinear with the central axis **320**. As the retainers **314**, **316** are closed together, the optical axis of the lens **348** is shifted as necessary by the ribs **330**, **334**, **338** and brought into a position that coincides with the central reference axis **320**. The ribs **334** urge the dome-shaped outer wall **352** into a position of snug contact with the chamfered wall sections **346**. The ribs **334** also exert a force on the lens **348** that is sufficient to pivot the lens **348** upon the wall sections **346** and shift the rear, flat face of the cylindrical lens portion next to the light outlet port **328** into an orientation such that the rear face is precisely perpendicular to the central axis **320** once the retainers **314**, **316** are closed. The ribs **330**, **338** function to shift the lens **348** in a lateral direction as the retainers **314**, **316** are closed so that the optical axis of the lens **346** is collinear with the central axis **320**.

As the retainers **314**, **316** are closed, the deformable tip portions **332**, **336**, **340** cold flow to accommodate the configuration of the lens **348** (and any surface irregularities) and secure the lens **348** in place. Such deformable tip portions **332**, **336**, **340** (along with other regions of the plastic retainers **314**, **316**) also tend to protect the lens **348** from damage when the optics assembly **300** or the measuring device **14** is subjected to impact or vibration.

The second chamber portions **324** of the retainers **314**, **316** are adapted to receive an optical filter **354**. The filter **354**

that is shown in FIG. **10** for exemplary purposes has a cylindrical configuration with a central axis. As the retainers **314**, **316** are brought toward a closed, connected-together orientation, the tip portions **332** of the ribs **330** located in the second chamber portions **324** and the tip portions **340** of the ribs **338** located in the second chamber portions **324** crush and deform, and urge the filter **354** to such an orientation that the central axis of the filter is collinear with the central axis **320** of the subassembly **312**. Once the retainers **314**, **316** are fully closed, the ribs **330**, **338** snugly engage the cylindrical side wall of the filter **354** and securely retain it in place.

The retainers **314**, **316** when closed together present a smooth cylindrical outer surface having a diameter that is preferably closely similar or identical to the internal diameter of the hole of the block **302** receiving the subassembly **312**. More preferably, the cylindrical outer surface presented by the retainers **314**, **316** when assembled together is slightly larger than the hole in the block **302** receiving the subassembly **312** in order to establish a slight interference fit between the block **302** and the subassembly **312**. Once the subassembly **312** is received in the hole, the retainers **314**, **316** remain assembled together without the need for adhesive, fasteners or the like. Alternatively, however, an adhesive or mechanical fastener could be provided to hold the retainers **314**, **316** in the block.

Each of the retainers **314**, **316** is preferably integrally-molded and made of a relatively soft, deformable plastic material such as an ABS polymer or a blend of ABS polymer and polycarbonate. An example of a suitable ABS polymer is no. 8540H from TAITALAC. Preferably, the material has a limited cold flow and yet is sufficient elastic to securely hold the optical component in place.

Optionally, the tip portions **332**, **336**, **340** or alternatively the entire ribs **330**, **334**, **338** are made of a plastic material that has a higher modulus of elasticity than the modulus of elasticity of the plastic material of the remainder of the retainers **314**, **316**. Insert molding, for example, could be used to provide tip portions **332**, **336**, **340** that are made of a material different than the remainder of the material of the retainers **314**, **316**. In any case, the material of the tip portions **332**, **336**, **340** and/or the entire ribs **330**, **334**, **338** is selected such that each tip portion **332**, **336**, **340** snugly engages and retains a force on the lens **348** and/or the filter **354** even after repeated cycles of temperature fluctuations. Preferably, the retainers **314**, **316** are made of a material that has a smaller thermal conductivity than the thermal conductivity of the material of the block **302**, in order to help insure that the temperature of the lens **348** and the filters **354** of the various subassemblies **304**–**312** is substantially identical even during periods of temperature fluctuation of the block **302**.

The excitation optic subassemblies **310**, **311** are identical to the excitation optic subassembly **312** with the exception of the optical filters. In particular, the optical filter **354** of the subassembly **312** has an overall diameter of 0.25 inch (6 mm) and a length of 0.145 inch to 0.175 inch (3.6 to 4.4 mm), has a pass band with a center wavelength of 398 nm and transmits 50% of peak transmission at wavelengths of 385 nm and 410 nm. The optical filter for the subassembly **311** is identical to the filter **354** except that the optical filter for the subassembly **311** has a pass band with a center wavelength of 413 nm and transmits 50% of peak transmission at wavelengths of 400 nm and 425 nm. The optical filter for the subassembly **310** is identical to the filter **354** except that the optical filter for the subassembly **310** has a pass band with a center wavelength of 475 nm and transmits 50% of peak transmission at wavelengths of 460 nm and 490 nm.

Other aspects of the subassemblies **310, 311** are identical to the subassembly **312** and as such a detailed description of the subassemblies **310, 311** need not be provided.

However, the optic subassemblies **304–309** are emission optic subassemblies and are somewhat different than the excitation optic subassemblies **310–312** as can be appreciated by reference to the subassembly **304** that is shown in exploded view in FIG. **16**. The subassembly **304** includes two retainers **314a, 316a** that are essentially similar to the retainers **314, 316** except for the differences set out below. The retainer **314a** alone is illustrated in FIGS. **20** and **21**. The retainers **314a, 316a** are identical and as such a detailed description of the retainer **314a** will suffice to describe the retainer **316a** as well. The retainer **314a** has internal wall sections that define a first chamber portion **322a**, a second chamber portion **324a**, a third chamber portion **326a**, a fourth chamber portion **328a** and a fifth chamber portion **329a**. When the retainers **314a, 316a** are connected together, the first chamber portions **322a** present a generally cylindrical light inlet port and the fourth chamber portions **328a** present a generally cylindrical light outlet port. Moreover, when the retainers **314a, 316a** are closed, the second chamber portions **324a** present a generally cylindrical filter sub-chamber for receiving an optical filter **354a** similar to the filter sub-chamber of subassembly **312**, and the third chamber portions **326a** present a first lens sub-chamber for receiving an optical lens **348a** similar to the lens sub-chamber of the subassembly **312**. The fifth chamber portions **329a** when the retainers **314a, 316a** are closed present a second lens sub-chamber for receiving an optical lens **349a** that is similar to the first lens sub-chamber of the subassembly **304**, but the second lens sub-chamber is oriented 180 degrees opposite from the orientation of the first lens sub-chamber with respect to a central reference axis of the subassembly **304**.

The retainer **314a** has first, second and third sets of ribs **330a, 334a, 338a** respectively that are preferably identical to the first rib set **330**, the second rib set **334** and the third rib set **338** of the subassembly **312**. As a consequence, as the retainers **314a, 316a** are closed, the ribs **330a, 334a, 338a** function to shift the lenses **348a, 349a** as well as the filter **354a** as necessary to bring the optical axes of the lens **348a, 349a** and the central axis of the filter **354a** into parallel alignment and preferably collinear alignment with the central axis of the subassembly **304** once assembled. The retainers **314a, 316a** also each have two pegs and two mating holes similar to the pegs **342** and holes **344** of the retainers **314, 316**. Preferably, however, the peg and hole on each side of each retainer **314a, 316a** are spaced apart from each other a distance that is somewhat different than the spacing between the pegs **342** and holes **344** of the retainers **314, 316** in directions parallel to the central axis **320**, in order to reduce the likelihood that one of the retainers **314, 316** is accidentally connected to one of the retainers **314a, 316a**.

The emission optic subassemblies **305–309** are each identical to the emission optic subassembly **304** except for the optical filters. Specifically, the optical filter **354a** of the subassembly **304** as well as the optical filters for the subassemblies **305, 307** are each identical to the filter **354** described above except that the filters for the subassemblies **304, 305, 307** have a pass band with a center wavelength of 550 nm and transmit 50% of peak transmission at wavelengths of 515 nm and 585 nm. The optical subassembly **306** has an optical filter that is identical to the filter **354** except that the optical filter for the subassembly **306** has a pass band with a center wavelength of 485 nm and transmits 50% of

peak transmission at wavelengths of 450 nm and 520 nm. The optical filters for the subassemblies **308, 309** are identical to the filter **354** except that the optical filters for the subassemblies **308, 309** have a pass band with a center wavelength of 500 nm and transmit 50% of peak transmission at wavelengths of 465 nm and 535 nm. The lenses **348a, 349a** of the subassembly **304** as well as the corresponding lenses of the subassemblies **305–309** are each identical to the lens **348** described above.

The retainers **314, 314a, 316, 316a** provide important advantages, inasmuch as they not only secure the lenses **348, 348a, 349a** and the filters **354, 354a** in proper alignment but also present built-in apertures for receiving light emitting diodes and photodiodes as described in further detail below. In addition, the retainers **314, 314a, 316, 316a** provide molded, built-in light inlet ports and light outlet ports of a suitable size for adjacent optical apertures. The retainers **314, 314a, 316, 316a** provide an inexpensive yet simple and efficient means for manufacture and assembly of the various components without the need for gaskets or washers as is common with other optical assemblies (such as, for example, conventional optical assemblies wherein lenses, filters and washers are dropped in sequence into a cylindrical chamber of a one-piece housing).

Those skilled in the art will recognize that the concepts of the mating retainers and an optical block may also be used for other optical assemblies such as binoculars, telescopes and the like. Moreover, a useful assembly may be provided by molding lenses and/or optical filters in place and integral with one of the retainers. As another alternative, the retainers could be constructed to present folded or offset optic axes such as is often provided for, e.g. binoculars. If the assembly is to be used in applications where rapid heat transfer between the block and the retainers is desired (for example, in assemblies where the optic components are heated), the retainers could be made of a material other than plastic and a thermal potting compound could be placed between the retainers and the block.

Although the optics assembly **300** as described in detail above is presently preferred, an alternative optics assembly may be constructed by eliminating the block **302** and providing other structure to hold the retainers together and to also hold the subassemblies together as a group if desired. For example, rings or snap-action clips may be constructed to hold each pair of retainers together, and a framework, grid, strapping or other structure may be provided to hold assembled pairs of the retainers to other assembled pairs of retainers. Optionally, the configuration of each assembled pair of retainers may be a shape other than cylindrical such as hexagonal, square, rectangular or triangular in instances where it is desired to establish flat, face-to-face contact between adjacent retainer pairs and avoid air gaps (as may be desired, for example, to facilitate heat transfer). As another option, each retainer may be provided with structure (such as snap-action pegs or tabs or a mortise and tenon arrangement) to interlock with the other retainer of that subassembly, or to interlock with retainers of other subassemblies.

As an additional option, a number of retainers for a number of different subassemblies may be integrally molded together in side-by-side relation as an array that is then connected to a similar array after the optical components are in place in each subassembly chamber, in order to reduce the number of parts that might otherwise be separately handled. For example, three retainers similar to retainers **316** in FIG. **16** could be integrally molded and then assembled to three retainers similar to the retainers **314** in FIG. **16** (with the

option, as explained above, that each retainer **314** is identical to each retainer **316**). Moreover, the backs of the retainers of one row could be integrally molded to the backs of the retainers of the adjacent row to further reduce the number of parts in any one optics assembly. Further, the assembled pairs of such retainers could have shapes other than cylindrical such as hexagonal, square, rectangular, triangular or the like as mentioned above.

The measuring device **14** also includes an electrical assembly **400** (see, e.g., FIG. 9) that includes an electro-optics subassembly **402** and a printed circuit board subassembly **404** having a printed circuit board substrate and various electronic components mounted on the substrate. The electro-optics subassembly **402** includes a coupling plate **406** (see also FIG. 22) having six somewhat oval-shaped recesses **408** that are arranged in two rows with three recesses **408** in each row. The coupling plate **406** also includes three circular recesses **410** that are arranged along a single row and in columnar alignment with the recesses **408**, such that the recesses **408**, **410** present an array of nine recesses with three recesses in each of three rows.

Preferably, at least a portion of the coupling plate **406** is made of an elastomeric material. As one example, the coupling plate **406** may include an aluminum substrate that is covered or partially covered with an elastic material such as polyurethane or silicone. Optionally, the substrate presents recesses corresponding to the recesses **408**, **410**, and a layer of elastomeric material is received in each recess. Optionally, the elastomeric material extends past the four edges of the metal substrate and covers the entire extent of a flat rear surface of the metal substrate that faces away from the optics assembly **300**. Preferably, the elastomeric material is electrically non-conductive and has holes that are aligned with but somewhat smaller than the holes in the metal provided for electrical leads. As such, the leads are kept out of contact with the aluminum substrate.

The electro-optical subassembly **402** includes three light emitting diodes or "LEDs" **412** (see FIGS. 8 and 10), each of which have a circular base that is snugly received in a respective one of the recesses **410**. Each LED **412** also has a dome-shaped portion that is snugly, complementally received in the light inlet port presented by the first chamber portions of the respective retainers of subassemblies **310**–**312**. The retainers of subassemblies **310**–**312** also have a circular cavity (see, e.g., cavity **323** in FIG. 17 and 18) for receiving the remainder of the circular base of the corresponding LED. An example of a suitable LED is a gallium nitride LED such as part no. NLPB-500 from Nichia or alternatively a laser diode.

The electro-optics subassembly **402** also includes six solid state light detectors or photodiodes **414**, each of which includes a body that is partially and snugly received in a respective one of the recesses **408**. The photodiodes **414** may be either silicone photodiodes or avalanche photodiodes. An example of a suitable silicon photodiode **414** is part no. S1133-14 from Hamamatsu. The LEDs **412** as well as the photodiodes **414** each include a pair of leads that extend through respective holes of the coupling plate **406**, and in particular through the holes of the metal substrate and elastomeric material as described above.

A pair of machine screws **416** (see FIG. 10) extend through a folded-over portion of the printed circuit board substrate, through respective holes **418** (see FIG. 22) of the coupling plate **406**, through respective holes **420** (see FIG. 16) of the block **302** and are threaded into matingly-threaded holes **422** (see FIG. 14) of the fiber block end plate **214**. Preferably, the block **302** is also provided with a pair of

protruding, parallel alignment posts **424** that are received in respective holes **426** (see FIG. 14) of the insert plate **214** to facilitate assembly. Once the screws **416** are tightened into the insert plate **214**, the elastic material extending over the substrate of the coupling plate **406** provides a yieldable layer that helps to prevent damage to the electrical assembly **400** whenever the measuring device **14** is subject to physical shock forces as may occur, for example, whenever the measuring device **14** is accidentally dropped.

As shown for example in FIG. 9, a peripheral edge portion of the coupling plate **406** protrudes outwardly from adjacent areas of the optic block **302** as well as the printed circuit board subassembly **404**. The peripheral edge portions of the coupling plate **406** are received in channels **428** of the two parts of the housing **200**. The protruding, peripheral edge portions of the coupling **406** in combination with the channels **428** provide a stable mounting support for the optics assembly **300** and the electrical assembly **400**, and also serve to reduce the amount of convective heat that would otherwise flow from the area within the housing **200** surrounding the printed circuit board subassembly **404** to the area within the housing **200** surrounding the remainder of the components including the optics assembly **300** and the fiber block assembly **210**.

The printed circuit board subassembly **404** includes a controller **430** (see FIG. 23) that receives power, clock timing and instructional signals from a remote monitor **432** (see FIG. 24). A flexible electrical cable **434** (FIG. 8) operatively interconnects the controller **430** and the monitor **432**. As another option, however, the electrical assembly **400** may be powered by batteries located in or adjacent to the housing **200** and the electrical cable **434** replaced by a bundle of optical fibers or by a telemetric communication device such as a device providing radio frequency or optical frequency signals.

The printed circuit board substrate of the subassembly **404** is folded into a rectangular, box-like configuration as shown in FIGS. 8–10. The box-like configuration includes four side portions and two end portions that surrounds the space where the electronic components on the substrate are mounted. Such construction advantageously provides electrical shielding for the electronic components and also serves to somewhat thermally isolate the components from the LEDs.

The controller **430** is electrically connected (connection not shown in drawings) to three drivers **436** that, in turn, are each electrically connected to a corresponding one of the LEDs **412**. The drivers **436** energize the respective LEDs **412** in sequence in accordance.

The printed circuit board subassembly **404** includes three analog-to-digital converters **438**, **439**, **440** that are shown in FIG. 23 as well as in FIG. 15. A suitable analog-to-digital converter is catalog no. DDC101 from BURR-BROWN. As schematically depicted in FIG. 15, the converter **438** is electrically connected to the photodiodes **414** associated with the optic subassemblies **305** and **306**, the converter **439** is electrically connected to the photodiodes **414** associated with the optic subassemblies **307** and **308** and the converter **440** is electrically connected the diode **414** associated with the optic subassemblies **304**, **309**. Additionally, a signal amplifier or opamp **442** (such as catalog no. AD795 from Analog Devices or catalog no. OPA 124U from Burr-Brown) is interposed in the electrical lead between the photodiode **414** of the subassembly **309** and the converter **439**.

In use, the controller **430**, upon receiving a certain signal from the monitor **432**, directs a signal to one of the drivers **436** which, in turn, energizes the corresponding one of the

LEDs 412. Light then travels from that LED 412 through the adjacent optical fibers of the fiber block assembly 210 and to the corresponding sensor 28–34 where it is absorbed. The sensor then emits light at a different wavelength. The amount of emitted light is determined by the analyte concentration (such as calibration fluid or blood) in the fluid chamber 18 of the cassette 12. Light emitted from such sensor 28–34 is directed through other optical fibers of the fiber block assembly 210 and to one of the photodiodes 414, and the converter 438–440 that is electrically connected to that diode 414 provides a digital output signal that is representative of the amount of light flux detected. The controller 430, in accordance with a preselected time delay interval, reads data received from the appropriate converter 438–440 and passes such data to the monitor 432 via cable 434.

The optical filter 354 within each of the excitation optic subassemblies 310–312 serves to pass substantially all of the light having wavelengths within its pass band and block substantially all of the light having wavelengths that are not within the pass band. The lens 348 of each excitation optic subassembly 310–312 focuses light emitted from the optical filter 354 onto the optical aperture of the adjacent fiber optic bundle.

Light that is emitted from the sensors 28–34 and directed to the emission optic subassemblies 304–309 passes from the optical aperture of the adjacent fiber optic bundle to the corresponding lens 348a. The lens 348a serves to collimate and direct the light along the optical axis to the adjacent optical filter 354a. The second lens 349a then focuses the light onto the corresponding, adjacent photodiode 414. The filter 354a passes most of the light having wavelengths within the pass band and blocks substantially all of the light having wavelengths outside of the pass band. For any given optical pathway, the filter 354 selects the wavelengths of light that excite the fluorescent dye in the corresponding sensor 28–34, and blocks substantially all other light. The filter 354a selects the wavelengths of light that have been emitted by the fluorescent dye and block substantially all remaining light (including any reflected excitation light).

The converters 438–440 convert the analog signal received from the photodiodes 414 into a digital data stream that is representative of the amount of light flux sensed by the photodiodes 414. The digital data is forwarded to the controller 430 which, in turn, sends the data to the monitor 432.

FIG. 15 shows that the light emitted from the LED 412 that is received in the subassembly 310 is simultaneously directed to the sensor 30, the sensor 32 and the reference disk 254 adjacent the hole 246. Light emitted from the sensor 30 is detected by the photodiode 414 next to the subassembly 305 and converted to a digital signal by the converter 438. Light emitted from the reference disk 254 next to the hole 246 is detected by the photodiode 414 of subassembly 307 and is converted to a digital signal by the converter 439. Light that is detected from the sensor 32 by the photodiode 414 adjacent the subassembly 304 is converted to a digital data stream by the converter 440. The digital data stream from the three converters 438–440 is received by the controller 430 and forwarded to the monitor 432.

In somewhat similar fashion, the light emitting from the LED 412 of the subassembly 312 is directed to the sensor 34 as well as the reference disk 254 that is adjacent the hole 250. Light emitted from the sensor 34 is detected by the photodiode 414 next to the subassembly 306, and the analog output of the photodiode 414 is converted to a digital data

stream by the converter 438. At the same time, light emitted from the reference disk 254 adjacent the hole 250 is detected by the photodiode 414 next to the subassembly 308, and the analog signal from that photodiode 414 is converted to a digital data stream by the converter 439.

When the LED 412 of the subassembly 311 is energized, light is directed simultaneously to the sensor 28 and to the reference disk 254 that is adjacent the hole 248. Light emitted from the reference disk 254 adjacent the hole 248 is detected by the photodiode 414 next to the subassembly 308, and the analog output of the photodiode 414 is changed to a digital data stream by the converter 439. Light that is emitted from the sensor 28 is detected by the photodiode 414 next to the subassembly 309, and the analog output from that photodiode 414 is amplified by amplifier 442 and directed to the converter 440 which changes the analog signal to a digital data stream.

The schematic shown in FIG. 15 enables the use of only three converters 438–440 even though four sensors (i.e., sensors 28–34) and three reference disks 254 are in use, and also enables the use of only three LEDs 412. Such a time sharing or “multiplexing” arrangement conserves the number of components needed for the electrical assembly 400 and also conserves space so that the housing 200 may be relatively small. In addition, such an arrangement reduces the amount of heat that might otherwise be generated within the housing 200.

A schematic block diagram of the monitor 432 is shown in FIG. 24. The monitor 432 includes an interface card 450 that is connected to the cable 434 leading from the electrical subassembly 404. The interface card 450 is electrically coupled to an auxiliary board 452 that, in turn, is connected to a primary controller or “SBC” (single board computer) 454. The interface card 450 also provides power to the LEDs 412.

The SBC controls the amplitude and duration of the pulse drive to the LEDs 412. The SBC 454 is connected to a memory card 456 as well as a display 458 such as an LED display. A power inverter 460 for the display 458 is also connected to the SBC 454.

The monitor 432 also includes a power supply 462 that is connected to a source of line voltage. The power supply 462 is electrically coupled to a charger 464 that maintains a battery 466 in charged condition. The power supply 462 as well as the battery 466 are connected to a power selector 468. The monitor 432 includes a switch 470 that, when closed, sends power to three DC/DC converters 472. The DC/DC converters 472 as well as a printer 474 are both electrically coupled to the auxiliary board 452.

FIG. 25 is a schematic illustration of a cardiopulmonary by-pass circuit 500 that advantageously employs the measuring device 14 and the cassette 12 described above. The circuit 500 includes a first length of tubing 502 having an inlet that is placed in communication with a venous blood vessel (preferably the vena cava) of a patient 504 undergoing surgery such as open heart surgery. The tubing 502 also includes an outlet that is connected to a venous bag reservoir 506. A cardiotomy reservoir 508 is also connected by means of a second length of tubing 510 to the venous bag reservoir 506.

The circuit 500 also includes a third length of tubing 512 having an inlet that is connected to an outlet port of the venous bag reservoir 506. The third length of tubing 512 has an outlet that is connected to a peristaltic or centrifugal pump 514.

The pump 514 has an outlet that is coupled to an inlet of a fourth length of tubing 516. The fourth length of tubing

516 has an outlet that is coupled to an inlet port of an oxygenator 518. The oxygenator 518 has an outlet port that is connected to an inlet of a fifth length of tubing 520.

The fifth length of tubing 520 has an outlet that is in communication with an inlet port of an arterial filter 522. An outlet port of the arterial filter 522 is connected to an inlet of a sixth length of tubing 523. The sixth length of tubing 523 also has an outlet that is connected to an arterial blood vessel (preferably the aorta) of the patient 504.

The cardiopulmonary by-pass circuit 500 also includes a shunt passageway 524. In the embodiment illustrated in FIG. 25, the shunt passageway 524 includes a sixth length of tubing 526 having an inlet that is connected to an outlet port of the arterial filter 522. The sixth length of tubing 526 also has an outlet that is connected to an inlet port of a blood parameter measurement cassette such as the cassette 12 shown in FIGS. 1-7. As an example, the sixth length of tubing 526 may be the same as the tubing 54 shown in FIG. 5.

The shunt passageway 524 also includes a seventh length of tubing 528 having an inlet that is connected to an outlet port of the cassette 12. The seventh length of tubing 528 is optionally the same as the tubing 84 that is shown in FIG. 5. The seventh length of tubing 528 also has an outlet port that is connected to an inlet port of a sampling port 530.

The shunt passageway 524 further includes an eighth length of tubing 532 having an inlet that is coupled to an outlet port of the sampling port 530. The eighth length of tubing 532 has an outlet that is connected to the inlet port of the venous bag reservoir 506 in the embodiment shown in FIG. 25. Although not shown in the drawing, the shunt passageway 524 may optionally include a valve to limit or interrupt the passage of blood through the tubing 526, 528 and 532.

The passages within the flexible tubing 502, 510, 512 and 516 as well as within the cardiomy reservoir 508, the venous bag reservoir 506 and the pump 514 comprise a venous passageway. The passages within the flexible tubing 520, 523 as well as within the arterial filter 522 represent an arterial passageway. The outlet of the shunt passageway 524 (i.e., the outlet of the eighth length of tubing 532 in the embodiment shown in FIG. 25) may be placed at any one of a number of different locations along the venous passageway, and is shown as connected to the inlet port of the venous bag reservoir 506 for exemplary purposes only. Preferably, the outlet of the shunt passageway 524 is placed in the venous passageway upstream of the pump 514 in order to enable blood to flow through the shunt passageway 524 without the need for an auxiliary pump.

The inlet of the shunt passageway 524 (i.e., the inlet of the sixth length of tubing 526 in the embodiment shown in FIG. 25) is preferably placed in communication with the arterial passageway at any one of a number of different locations along the latter, and more preferably is connected to an outlet port of the arterial filter 522 as depicted in FIG. 25. As another option, however, the inlet of the shunt passageway 524 may be located along the venous passageway, preferably at a location downstream of the pump 514.

In the circuit 500 shown in FIG. 25, the shunt passageway 524 diverts a portion of the patient's blood flowing through the arterial passageway and directs the diverted portion back to the venous passageway. Advantageously, placement of the cassette 12 with its sensors 28-34 along the length of the shunt passageway 524 avoids the necessity of placing the cassette 12 in series relation with either the venous passageway or the arterial passageway, and yet allows for blood to move continuously past the sensors 28-34 at all times if

desired. Moreover, the shunt passageway 524 can be connected to or disconnected from the venous and arterial passageways during a surgical procedure if desired without interrupting the flow of blood through the venous passageway and arterial passageway to the patient 504.

By contrast, conventional cardiopulmonary by-pass circuits with blood gas monitoring devices have typically included a flow-through cassette or cell that is interposed in the venous or arterial passageway and has one or more sensors for determining parameters of blood flowing through the cell. However, since the flow-through cell is in series with the venous and arterial passageways, the cell must be connected to the venous or arterial passageway before blood is directed through the by-pass circuit, a disadvantage in certain instances. Such an arrangement also effectively precludes the possibility of changing the cell if, for example, one or more of the sensors is defective.

Preferably, the shunt passageway 524 has an average cross-sectional area in reference planes perpendicular to the flow of blood that is smaller than the average cross-sectional area of the venous and/or the arterial passageways in reference planes perpendicular to the flow of blood. For example, the shunt passageway 524 including the flexible tubing 526, 528 and 532 may have an internal diameter of 0.125 inch (3.2 mm), while the venous and arterial passageways including the flexible tubing 502, 512, 516, 520 and 523 may have an average internal diameter in the range of from about 0.25 inch (6.3 mm) to about 0.5 inch (12.7 mm). The smaller internal area of the shunt passageway 524 allows the use of relatively small tubing sizes which can be easily attached to or disconnected from the venous and arterial passageways at any time without undue interruption to the flow of blood through the venous and arterial passageways. For example, the shunt passageway 524 may be connected to the venous and arterial passageways some time after blood has begun to flow through the latter, as may be desired in the surgical suite after the patient has arrived from the pre-surgical suite with the arterial and venous passageways installed and in use.

Furthermore, the shunt passageway 524 is an advantage in that the sampling port 530 can be placed in close proximity to the cassette 12 and help insure that the blood samples that are taken from the sampling port 530 better correlate with the measurement of the blood characteristics determined by the cassette 12 and the measuring device 14. This arrangement also avoids the need for a syringe or other device to withdraw blood samples from the sampling port as is the case with many conventional cardiopulmonary by-pass circuits.

The cassette 12 and the device 14 may also be used in a non-cardiopulmonary bypass shunt application. For example, in an intensive care unit or other critical care setting, an arterial-to-venous shunt passageway could be constructed by connecting relatively small diameter flexible tubing (e.g., 0.125 inch (3.1 mm) to 0.19 inch (4.8 mm) outer diameter) to an arterial blood vessel and a venous blood vessel using conventional cannulation techniques. Blood would then flow through the tubing due to the arterial-venous pressure differential. A cassette such as cassette 12 connected to the tubing would be used to monitor one or more parameters of the blood.

Advantageously, the longitudinal axis of the measuring device 14, and particularly the longitudinal axis of the housing 200, is generally parallel, and preferably is parallel to the direction of blood flowing through the fluid chamber 18 of the cassette 12. Such construction provides a compact assembly and further reduces the possibility of eddy currents or other disruptions in the flow of blood that might otherwise

tend to facilitate clotting of the blood. The parallel arrangement is particularly useful when the cassette **12** and the measuring device **14** are placed in the vicinity of the patient's body.

FIG. **26** is a schematic illustration of a cardiopulmonary by-pass circuit **500a** in accordance with another embodiment of the invention. In FIGS. **25** and **26**, elements that are identified by the same numerals are identical and as a consequence a detailed description of those elements need not be repeated. However, the cardiopulmonary by-pass circuit **500a** of FIG. **26** has a shunt passageway **524a** that is somewhat different than the shunt passageway **524**.

More particularly, the shunt passageway **524a** includes a sixth length of tubing **526a** having an inlet that is connected to the arterial filter **522** and an outlet that is connected to a sampling port **530a**. The shunt passageway **524a** also includes a seventh length of tubing **528a** having an inlet that is connected to the sampling port **530a** and an outlet that is connected to the cassette **12**. An eighth length of tubing **532a** of the shunt passageway **524a** has an inlet that is connected to the cassette **12** and an outlet that is connected to the cardiomy reservoir **508**.

The cardiopulmonary by-pass circuit **500a** also includes a ninth length of tubing **534a** having an inlet that is connected to an inlet of the venous bag reservoir **506** and an outlet that is connected to the sampling port **530a**. A pump **536a** is interposed in the ninth length of tubing **534a** for directing blood from the venous bag reservoir **506** through the ninth length of tubing **534a** and to the sampling port **530a**.

The sampling port **530a** includes valves to selectively interrupt the flow of blood from either the sixth length of tubing **526a** or the ninth length of tubing **534a**. For example, the valves of the sampling port **530a** may be adjusted to enable the flow of blood through the sixth length of tubing **526a** and to the cassette **12** and interrupt the flow of blood in the ninth length of tubing **534a**, so that the sensors **28-34** of the cassette **12** may be used to monitor the parameters of arterial blood. As another option, the valves of the sampling port **530a** may be adjusted to interrupt the flow of blood through the sixth length of tubing **526a** and enable the flow of blood through the ninth length of tubing **534a** and to the cassette **12** so that the sensors **28-34** may be used for monitoring the parameters of venous blood. Such an arrangement enables the user to alternatively monitor both venous and arterial blood parameters without the need for two cassettes and measuring devices.

Another alternative embodiment of the invention is schematically illustrated in FIG. **27**, wherein is shown a cardiopulmonary by-pass circuit **500b**. The components shown in FIGS. **25** and **27** that bear the same identifying numerals are identical and a detailed description of such components need not be repeated. However, the cardiopulmonary by-pass circuit **500b** includes a shunt passageway **524b** that is somewhat different than the shunt passageways **524**, **524a** described hereinabove.

More particularly, the shunt passageway **524b** includes a sixth length of tubing **526b** having an inlet that is connected to an outlet of the arterial filter **522** and an outlet that is connected to the inlet of the cassette **12**. A seventh length of tubing **528b** of the shunt passageway **524b** has an inlet that is connected to the outlet connector **62** of the cassette **12** and an outlet that is connected to an inlet port of a sampling port **530b**. An eighth length of tubing **532b** of the shunt passageway **524b** has an inlet that is connected to an outlet port of the sampling port **530b** and an outlet that is connected to the cardiomy reservoir **508**.

Additionally, the shunt passageway **524b** includes a ninth length of tubing **534b** having an inlet that is connected to an

inlet port of the venous bag reservoir **506** and an outlet that is connected to an inlet fitting of a second cassette **12a**. Optionally, the second cassette **12a** is identical to the first cassette **12**. The second cassette **12a** has an outlet connector (such as the Luer connector **62** shown in FIG. **5**) that connects to an inlet of a tenth length of tubing **538b**. The tenth length of tubing **538b** has an outlet that is connected to an inlet of the sampling port **532b**. A pump **536b** is interposed in the tenth length of tubing **538b** to direct blood from the inlet of the venous bag reservoir **506** to the sampling port **530b**.

The cardiopulmonary by-pass circuit **500b** that is shown in FIG. **27** enables the user to monitor characteristics of blood simultaneously in both the arterial and venous passageways. As an alternative to the circuits **500a**, **500b** shown respectively in FIGS. **26** and **27**, the inlet of the ninth length of tubing **534a**, **534b** may be connected to the fourth length of tubing **516**, and in such cases the pressure provided by the pump **514** is preferably sufficient to direct the flow of blood through the tubing **534a**, **534b** without the need for the pumps **536a**, **536b**. Additionally, the outlet of the ninth length of tubing **534a**, **534b** as described in the alternative embodiments of the preceding sentence may optionally be connected to the inlet of the venous bag reservoir **506** instead of the sampling port **530a**, **530b** (which in these options can now be omitted).

The shunt passageways **524a**, **524b**, like the shunt passageway **524**, preferably have an average cross-sectional area in reference planes perpendicular to the flow of blood that is smaller than the average cross-sectional area of either of the venous or arterial passageways when considered in reference planes perpendicular to the flow of blood. The shunt passageways **524a**, **524b** can be conveniently attached or detached from the circuits **500a**, **500b** respectively without interrupting the flow of blood passing through the arterial and venous passageways.

Cassette **12a** according to another embodiment of the invention is illustrated in FIGS. **28-31** and broadly includes a cassette body **15a** that is shown alone in FIGS. **28** and **29** and a cassette casing **16a** that is shown alone in FIG. **30**. The body **15a** includes a central support member **17a** for mounting one or more sensors that are adapted to sense one or more parameters in a fluid such as blood. In the embodiment shown in FIGS. **28** and **29**, the support member **17a** carries four sensors **28-34** that are identical to the sensors **28-34** described in connection with the cassette **12** above, although it should be understood in this regard that different sensors or a smaller or larger number of sensors may alternatively be provided.

The sensors **28-34** of the cassette body **15a** are located in an oval-shaped recess **26a** that is surrounded by a rim **40a**. A key **42a** is integrally connected to a side wall of the rim **40a** as shown in FIG. **28**. The recess **26a**, the rim **40a** and the key **42a** are similar in configuration to the recess **26**, the rim **40** and the key **42** described above in order to matingly and releasably couple to the protrusion **216** of the measuring device **14** when desired.

The cassette body **15a** also includes a male coupling **44a** for detachable connection to the female coupling **202** of the measuring device **14**. The coupling **44a** has a convex, generally U-shaped configuration in reference planes perpendicular to an axis extending through the center of the four sensors **28-34**. The coupling **44a** includes opposed leg portions **46a** that extend outwardly from the body **15a** in a direction away from the outward extension of the rim **40a**. Each leg portion **46a** includes three support sections having flat, coplanar outer surfaces **47a** adapted for contact with the

coupling 202. Preferably, the outer surfaces 47a of the opposed leg portions 46a converge as the body 15a is approached and extend along reference planes that are oriented in an angle in the range of about 28 degrees to about 32 degrees relative to each other. More preferably, the outer surfaces 47a extend along respective reference planes that are oriented at an angle of about 30 degrees relative to each other.

A flange 48a is integrally connected to the outer end of each leg portion 46a. Flanges 48a lie in a common plane that is parallel to the aforementioned axis extending through the center of the four sensors 28-34. The leg portions 46a are somewhat flexible and can be moved slightly toward each other under the influence of finger pressure, but also have sufficient memory to quickly and repeatedly return to their original, normal orientation as shown in the drawings once finger pressure is released.

An outer, central region of each leg portion 46a is integrally connected to a wedge-shaped tab 50a. The tabs 50a extend away from each other and outwardly from the respective leg portions 46a along respective reference planes that are oriented at an angle of about 80 degrees relative to each other. Additionally, a distal edge of each tab 50a extends in a reference plane that is oriented at an angle of 25 degrees relative to the direction of extension of the flanges 48a. Outermost edges of the tabs 50a are spaced outwardly relative to adjacent regions of the respective leg portions 46a and lie in a common reference plane that is between the body 15a and the aforementioned reference plane containing the flanges 48a.

Additionally, the leg portions 46a each have two rectangular openings 49a that are located between the body 15a and the tabs 50a, and preferably are located closely adjacent the body 15a. Additionally, the cassette body 15a has a groove 51a as illustrated in FIG. 29 for receiving an O-ring 51a that is depicted only in FIG. 31.

The cassette casing 16a that is shown alone in FIG. 30 has walls defining a chamber 18a, an inlet 53a for admitting a fluid such as blood to the chamber 18a and an outlet 55a for discharging fluid from the chamber 18a. The casing 16a has a side opening 19a that is surrounded by a wall 61a and extends into the chamber 18a. The casing 16a also includes a pair of elongated wings 57a that extend outwardly in a direction away from the chamber 18a. Two snaps or ears 59a, each having a protruding, wedge-shaped configuration, are connected to each wing 57a at a location next to the chamber 18a.

The wings 57a are somewhat flexible and can be moved slightly toward each other under the influence of finger pressure, but also have sufficient memory to quickly and repeatedly return to their original, normal orientation as shown in the drawings once finger pressure is released. Preferably, the cassette body 15a and the cassette casing 16a are each injection-molded as initially separate pieces of a plastic material such as relatively clear medical grade polycarbonate.

The ears 59a of the casing wings 57a and the openings 49a of the cassette body 15a constitute a connector for detachably coupling the casing 16a to the body 15a. The body 15a and the casing 16a are shown coupled together in FIG. 31, where it can be observed that each of the ears 59a are received in a respective one of the openings 49a to securely couple the casing 16a to the body 15a. When the casing 16a is so connected to the body 15a, the O-ring 51a (FIG. 31) engages the wall 61a (FIG. 30) of the casing 16a in order to close the opening 19a and provide a leak-resistant fluid seal between the body 15a and the casing 16a.

Advantageously, the wedge-shaped configuration of the ears 59a functions as a cam to deflect the respective wings 57a inwardly and in a direction toward each other as the ears 59a slide across the inner surfaces of the leg portions 46a when the casing 16a is coupled to the body 15a. As such, the user can securely couple the casing 16a to the body 15a by pressing on the casing 16a adjacent the fluid chamber 18a, and the user need not squeeze the wings 57a together. Once the outer edges of the ears 59a are moved past the outer sides of the openings 49a, the inherent resiliency of the wings 57a causes the ears 59a to snap in place in the openings 49a. However, the casing 16a may be detached from the body 15a when desired by pressing the wings 57a toward each other until such time as the outer edges of the ears 59a clear the openings 49a, and then moving the casing 16a away from the body 15a.

FIG. 31 also illustrates connection of the cassette 12a to the female coupling 202 of the measuring device 14 described above. The tabs 50a snap into respective grooves 208 of the coupling 202 as the cassette 12a is moved toward the measuring device 14. In this regard, the tabs 50a are similar in function and operation to the tabs 50, and provide a snap-action connection to releasably couple the cassette 12a to the measuring device 14. When it is desired to detach the cassette 12a from the measuring device 14, the flanges 48a are moved inwardly by finger pressure toward each other until such time as the tabs 50a clear the grooves 208, and the cassette 12a may then be removed from the female coupling 202.

In other respects, the cassette 12a is similar in construction and function to the cassette 12. For example, the cassette 12a is used for calibration purposes and may be also used if desired for monitoring parameters of blood. As another example, internal walls of the cassette 12a defining the chamber 18a are preferably provided with a hydrophilic surface such as a coating of heparin. The chamber 18a also preferably includes chamber portions similar to the portions 20, 22, 24 and 25 described above.

Additionally, various fittings and connectors may be coupled to the inlet 53a and the outlet 55a for connection with flexible tubing. Alternatively, a cap such as cap 56 may be connected to the outlet port 55a, and a gas filter assembly similar to the filter assembly 66 may be coupled to the inlet 53a. Such an assembly enables the user to calibrate the sensors 28-34 in the manner similar to the procedure described above for the cassette 12.

The cassette casing 16a has a relatively small internal diameter and is used during calibration. Optionally, the casing 16a may also be used when the cassette 12a forms part of a shunt passageway such as the shunt passageway 524, 524a, 524b in FIGS. 25-27. An alternative cassette casing 16a is illustrated in FIGS. 32 and 33, and is useful where the cassette 12a is to be used as a flow-through cell for sensing in an arterial or a venous passageway. For example, the casing 16b may have an inlet 53b and an outlet 55b that is adapted to receive flexible tubing having a nominal inner diameter of 0.5 inch (12.5 mm).

The casing 16b includes four wedge-shaped snaps or ears 59b which are located in pairs along opposite sides of an oval-shaped opening 19b. The ears 59b have the same spatial relationship to each other as the ears 59a described above, and are adapted to be snapped into the openings 49a of the cassette body 15a when the casing 16b is connected to the body 15a. However, since the ears 59b are not coupled to wings such as the wings 59a, the ears 59b serve to connect the casing 16b to the body 15a in substantially permanent fashion and as a consequence the casing 16b cannot readily be detached from the body 15a once coupled together.

The casing **16b** has an internal fluid chamber **18b** with a side opening **19b**. An oval-shaped membrane **61b** extends across the opening **19b** and is joined to an oval-shaped land **63b**. The membrane **61b** is connected to the land **63b** by ultrasonic welding or adhesive bonding. The membrane **61b** is made of material having a series of small perforations such as 0.005 inch (0.12 mm) track-etched polycarbonate.

Aside from the membrane **61b**, the cassette casing **16b** is made using two initially separate pieces: a first piece **65b** and a second piece **67b** as shown in FIG. 33. The first and second pieces **65b**, **67b** are joined together using, for example, an ultrasonic welding or an adhesive bonding procedure.

FIG. 34 is an end cross-sectional view through the casing **16b**, the body **15a** and the female coupling **202** of the measuring device **14**. As can be observed in FIG. 34, the configuration of the fluid chamber **18b** in the vicinity of the sensors **28-34** is generally oval-shaped, an advantage in that a somewhat larger chamber area can be provided within the confines of the leg portions **46a**. The cross-sectional area of the generally oval-shaped portion of the fluid chamber **18b** shown in FIG. 34 is approximately equal, and preferably equal, to the circular cross-sectional area of the inlet **53b** and the outlet **55b**. Moreover, wall sections defining the fluid chamber **18b** also preferably provide a smooth transition between the circular cross-sectional areas and the middle, generally oval-shaped area to avoid undue disruption of the flow of blood through the chamber **18b**.

FIGS. 35 and 36 depict a cap **69b** adapted for use with the casing **16b**. The cap **69b** includes a central, generally oval-shaped member **71b** that is adapted to extend over and protect the membrane **61b** before such time as the casing **16b** is connected to the body **15a**. Preferably, the member **71b** has a shape complementary in configuration to the membrane **61b** when the membrane **61b** is fixed to the casing **16b**. The cap **69b** also has a groove **73b** to carry an O-ring (not shown) for sealing engagement with the casing **16b** in order to establish a hermetic seal over the membrane **61b**.

The cap **69b** includes a pair of flexible wings **75b**, each of which has a pair of rectangular openings **77b** as shown in FIG. 35. As the cap **69b** is pressed onto the casing **16b**, the wings **75b** engage the wedge-shaped ears **59b** and deflect outwardly in a direction away from each other. Once the ears **59b** are moved to a position adjacent respective openings **77b**, the inherent resiliency of the wings **75b** causes the wings **75b** to self-return to their normal orientation as shown in FIGS. 35 and 36 to securely couple the cap **69b** to the casing **16b**. The cap **69b** may be easily detached from the casing **16b** when desired by pressing the wings **75b** in a direction toward each other until the ears **59b** clear the openings **77b** and then moving the cap **69b** away from the casing **16b**.

FIGS. 37 and 38 illustrate for exemplary purposes two other casings for use with the body **15a**. In FIG. 37, for example, a casing **16c** includes a first piece **65c** and a second piece **67c** that together with the cassette body **15a** present a fluid chamber **18c**. The fluid chamber **18c** has a generally circular cross-sectional area along its entire length. Since the chamber **18c** is smaller in area than, for example, the cross-sectional area of the chamber **18b** shown in FIG. 34, the need for an oval-shaped middle portion of the chamber **18c** is avoided. In other respects, the casing **16c** is similar to the casing **16b**. For example, the casing **16c** has a membrane similar to membrane **61b**.

As an example of use of the cassette **12a**, the body **15a** is preferably coupled to the casing **16a** and then shipped to the user in such coupled-together relation. Additionally, a gas

filter assembly such as assembly **66** (see, e.g., FIG. 5) is connected to the cassette **12a** along with a fitting and cap such as fitting **56** and cap **78** (see, e.g., FIG. 6). The user then connects the gas filter assembly to a source of calibration gas, and loosens the cap. The calibration gas is then directed through the fluid chamber **18a** in the manner described above with regard to calibration of the cassette **12**.

Once calibration is complete, the casing **16a** is removed from the body **15a** by squeezing the wings **57a** toward each other. During such time, the body **15a** preferably remains connected to the coupling **202** of the measuring device **14**. Next, the cap **69b** is removed from the selected casing **16b** or **16c**. If, for example, the cassette **12a** is intended to be used in an arterial or a venous passageway having lengths of tubing with an internal nominal diameter of 0.5 inch (12.5 mm), the casing **16b** is connected to the body **15a** in the manner shown in FIG. 34. Once the procedure is completed, the body **15a** with the casing **16b** is removed from the coupling **202** and discarded. On the other hand, if the cassette **12a** is intended for use in a smaller passageway, the casing **16c** is coupled to the body **15a** in the manner illustrated in FIG. 37.

Advantageously, the membrane **61b** enables the user to connect the casing **16b** to an arterial or venous passageway before such time as the body **15a** is coupled to the casing **16b**. As a consequence, the casing **16b** may be used to pass fluid along the arterial and venous passageways without interruption of the blood flow to begin monitoring.

The casing **16c** has an elongated, external rib **82** that is also shown in FIG. 39. The rib **82** extends along the length of the fluid chamber **18c** and is located on the casing **16c** at a position remote and opposite the cassette body **15a**. Preferably, the rib **82** extends away from the chamber **18c** along a reference plane that bisects the angle between the flexible leg portions.

The rib **82** is a distinct advantage when coupling the casing **16c** to the body **15a**. Once a pair of adjacent ears **59c** are received in openings **49a**, finger pressure can be exerted on the rib **82** to rock the casing **16c** in a lateral direction about its longitudinal axis in order to enable the remaining ears **59c** to snap in the remaining, respective openings. In addition to facilitating such rocking motion, the rib **82** also serves to stiffen the casing **16c**. A similar rib can be used with casings of other sizes as well, such as the casings **16**, **16a** and **16d** described above. Moreover, and as shown in FIG. 39, the rib **82** can optionally include one or more post sections that serve to stiffen the rib **82** and help the rib **82** resist deflection due to lateral forces.

The cassette assembly that is shown in FIG. 38 includes the body **15a** as mentioned above, but in this instance the body **15a** is coupled to a casing **16d** that is advantageously a unitary, single component with the exception of a membrane that is similar to membrane **61b**. The cross-sectional area of a fluid chamber **18d** of the cassette **12a** shown in FIG. 38 is substantially the same as the area of the fluid chamber **18a** depicted in FIG. 31, and as such the cassette **12a** as depicted with the casing **16d** in FIG. 38 is conveniently adapted for use with a shunt passageway such as passageway **524** in FIGS. 25-27. The casing **16d** is useful in instances where, for example, the manufacturer wishes to ship a single unit that will be used for both calibration and blood monitoring, and lacks wings (such as wings **57a** in FIG. 31) to reduce the likelihood that the body **15a** will detach from the casing **16d** in use. In other respects, the casing **16d** is similar to the casings **16b**, **16c** and has, among other things, ears **59d** similar to ears **59b** for coupling the casing **16d** to the body **15a**. In addition, the shipping cap **69b** may be used with the casings **16c**, **16d** as desired.

What is claimed is:

1. A system for measuring one or more parameters of blood comprising:
 - a casing having walls defining a chamber for receiving a quantity of blood;
 - at least one sensor connected to said casing adjacent said chamber for sensing at least one parameter of the blood in said chamber;
 - a measuring device including a housing, at least one light source connected to the housing and at least one light detector connected to the housing; and
 - a connector for releasably coupling said casing to said housing, said connector including a first coupling connected to said casing and a second coupling connected to said housing, said first coupling having a convex configuration and said second coupling having a concave configuration.
2. The system of claim 1 wherein said first coupling and said second coupling have matching, generally U-shaped configurations.
3. The system of claim 1 wherein said first coupling includes at least one flexible leg portion with an outwardly extending tab, and wherein said second coupling includes at least one groove for receiving said tab.
4. The system of claim 1 wherein said first coupling has a generally U-shaped configuration with a central section and two leg portions that extend away from said central section, said leg portions each including at least one outer surface, said at least one outer surface of one of said leg portions and said at least one outer surface of the other of said leg portions extending in respective reference planes that are oriented at an angle in the range of about 28 degrees to about 32 degrees relative to each other.
5. The system of claim 4 wherein each of said leg portions includes an outwardly extending tab.
6. The system of claim 4 wherein said central section includes a recess and a generally oval-shaped rim surrounding said recess, and wherein said at least one sensor is located in said recess.
7. The system of claim 4 wherein said leg portions are movable toward each other under the influence of finger pressure.
8. The system of claim 7 wherein said leg portions each include at least one outwardly extending tab.
9. The system of claim 1 wherein said first coupling includes a central section having a recess and a generally oval-shaped rim surrounding said recess, and wherein said at least one sensor is located in said recess.
10. The system of claim 1 wherein said at least one sensor includes at least two sensors, wherein said first coupling includes a rim surrounding said at least two sensors, and wherein said first coupling includes an alignment key connected to said rim and extending between an adjacent pair of said at least two sensors.
11. The system of claim 1 wherein said at least one sensor includes a potassium sensor.
12. A cassette for measuring one or more parameters of blood comprising:
 - a casing having walls defining a chamber for receiving a quantity of blood;
 - at least one sensor connected to said casing adjacent said chamber for sensing at least one parameter of the blood in said chamber; and
 - a coupling for connecting said casing to a blood parameter measurement device, said coupling having a convex configuration with a central section, wherein said at

least one sensor is located in said central section, and wherein said coupling includes at least one flexible leg portion with a tab for releasable connection to the blood parameter measurement device, said at least one leg portion extending in a direction away from said central section, said tab extending outwardly in a lateral direction from said at least one leg portion.

13. The cassette of claim 12 wherein said at least one leg portion includes two flexible leg portions that extend away from said central section, said two leg portions each including at least one outer surface, said at least one outer surface of one of said two leg portions and said at least one outer surface of the other of said two leg portions extending in respective reference planes that are oriented at an angle in the range of about 28 degrees to about 32 degrees relative to each other.

14. The cassette of claim 13 wherein each of said two leg portions includes an outwardly extending tab.

15. The cassette of claim 13 wherein said central section includes a recess and a generally oval-shaped rim surrounding said recess, and wherein said at least one sensor is located in said recess.

16. The cassette of claim 13 wherein said two leg portions are movable toward each other under the influence of finger pressure.

17. The cassette of claim 16 wherein said two leg portions are integrally connected to said central section.

18. The cassette of claim 12 wherein said at least one sensor includes at least two sensors, wherein said coupling includes a rim surrounding said at least two sensors, and wherein said coupling includes an alignment key connected to said rim and extending between an adjacent pair of said at least two sensors.

19. The cassette of claim 12 wherein said at least one sensor includes a potassium sensor.

20. The cassette of claim 12 wherein said chamber has a longitudinal axis and includes an inlet portion, an outlet portion and a middle portion between said inlet portion and said outlet portion, wherein said inlet portion and said outlet portion each have a generally circular configuration in reference planes perpendicular to said longitudinal axis, wherein said middle portion has a non-circular configuration in reference planes perpendicular to said longitudinal axis, said circular configurations of said inlet portion and said outlet portion each having respective areas that are each approximately equal to the area of said non-circular configuration of said middle portion.

21. The cassette of claim 12 wherein said coupling is releasably connected to said casing.

22. The cassette of claim 12 wherein said coupling is integrally connected to said casing.

23. A cassette for measuring one or more parameters of blood comprising:

- a casing having walls defining a chamber for receiving a quantity of blood;
- at least one sensor connected to said casing adjacent said chamber for sensing at least one parameter of the blood in said chamber; and
- a coupling for connecting said casing to a blood parameter measurement device, said coupling having a convex configuration and having two outer surfaces extending in respective reference planes that are oriented at an angle in the range of about 28 degrees to about 32 degrees relative to each other.

24. The cassette of claim 23, wherein said reference planes are oriented at an angle of about 30 degrees relative to each other.

25. The cassette of claim 23, wherein said coupling includes two leg portions and at least one tab connected to at least one of said leg portions.

26. The cassette of claim 25, wherein said leg portions each have an outer end region, and wherein said at least one tab includes a pair of tabs extending away from each other, each of said tabs being coupled to a respective one of said outer end regions.

27. The cassette of claim 23, wherein said coupling includes a central section having a recess and a generally oval-shaped rim surrounding said recess, and wherein said at least one sensor is located in said recess.

28. The cassette of claim 23, wherein said at least one sensor includes at least two sensors, wherein said coupling includes a rim surrounding said at least two sensors, and wherein said coupling includes an alignment key connected to said rim and extending between an adjacent pair of said at least two sensors.

29. The cassette of claim 23, wherein said chamber has a longitudinal axis and includes an inlet portion, an outlet portion and a middle portion between said inlet portion and said outlet portion, wherein said inlet portion and said outlet portion each have a generally circular configuration in reference planes perpendicular to said longitudinal axis, wherein said middle portion has a non-circular configuration in reference planes perpendicular to said longitudinal axis, said circular configurations of said outlet portions having respective areas that are each approximately equal to the area of said non-circular configuration of said middle portion.

30. The cassette of claim 23, wherein said coupling is releasably connected to said casing.

31. The cassette of claim 23, wherein said coupling is integrally connected to said casing.

32. The cassette of claim 23, wherein said coupling includes two leg portions that are flexible for movement toward each other under the influence of finger pressure.

33. The cassette of claim 32, wherein said leg portions each include at least one outwardly extending tab.

34. The cassette of claim 23, wherein said at least one sensor includes a potassium sensor.

35. A cassette for measuring one or more parameters of blood comprising:

a casing having walls defining a chamber for receiving a quantity of blood;

at least one sensor connected to said casing adjacent said chamber for sensing at least one parameter of the blood in said chamber; and

a coupling having a convex configuration for connecting said casing to a blood parameter measurement device, wherein said chamber has a longitudinal axis and includes an inlet portion, an outlet portion and a middle portion between said inlet portion and said outlet portion, wherein said inlet portion and said outlet portion each have a generally circular configuration in reference planes perpendicular to said longitudinal axis, wherein said middle portion has a non-circular configuration in reference planes perpendicular to said longitudinal axis, said circular configurations of said inlet portion and said outlet portion having respective areas that are each approximately equal to the area of said non-circular configuration of said middle portion.

36. The cassette of claim 35, wherein said non-circular configuration is an oval.

37. A cassette adapted for releasable coupling to a blood parameter measuring device for measuring one or more parameters of blood, said cassette comprising:

a body;

at least one sensor connected to said body for sensing at least one parameter of the blood;

a casing having walls sections at least partially defining a chamber, an inlet for admitting blood to said chamber and an outlet for discharging blood from said chamber; and

a connector for detachably coupling said body to said casing in such a manner that said at least one sensor is located next to said chamber for sampling one or more parameter of blood flowing through said chamber.

38. The cassette of claim 37, wherein said cassette includes a pair of leg portions connected to said body, and wherein said connector detachably couples said casing to said body at a location between said leg portions.

39. The cassette of claim 38, wherein said casing is elongated and has an external rib extending along its length.

40. The cassette of claim 38, wherein said leg portions each include at least one opening, and wherein said casing includes a pair of wings each having at least one protruding ear that is received in a respective one of said at least one openings when said casing is coupled to said body.

41. The cassette of claim 40, wherein said wings are flexible and movable toward each other under the influence of finger pressure.

42. The cassette of claim 38, including a membrane secured to said casing for communicating blood in said chamber with said at least one sensor.

43. The cassette of claim 42, wherein said casing has a side opening, wherein said membrane extends across said side opening, and wherein said casing includes a pair of flexible wings extending away from said chamber in a direction away from said side opening.

44. The cassette of claim 43, wherein said at least one sensor includes at least two sensors arranged along a reference axis, and wherein said side opening has a longitudinal axis that is parallel with said reference axis when said casing is connected to said body.

45. The cassette of claim 37, wherein said body includes a coupling having a convex configuration for connection to a blood parameter measuring device.

46. The cassette of claim 45, wherein said coupling includes at least one flexible leg portion with a tab for releasable connection to a blood parameter measurement device, said at least one leg portion extending in a direction away from said central section, said tab extending outwardly in a lateral direction from said at least one leg portion.

47. The cassette of claim 46, wherein said coupling includes a central section and wherein said at least one leg portion includes two flexible leg portions that extend away from said central section, said two leg portions each including at least one outer surface, said at least one outer surface of one of said two leg portions and said at least one outer surface of the other of said two leg portions extending in respective reference planes that are oriented at an angle in the range of about 28 degrees to about 32 degrees relative to each other.

48. The cassette of claim 47, wherein each of said two leg portions includes an outwardly extending tab.

49. The cassette of claim 37, wherein said coupling includes two outer surfaces extending in respective reference planes that are oriented at an angle in the range of about 28 degrees to about 32 degrees relative to each other.

50. The cassette of claim 37, wherein said body includes a central section with a recess and a generally oval-shaped rim surrounding said recess, and wherein said at least one sensor is located in said recess.

51. The cassette of claim 37, wherein said at least one sensor includes at least two sensors, wherein said coupling includes a rim surrounding said at least two sensors, and wherein said coupling includes an alignment key connected to said rim and extending between an adjacent pair of said at least two sensors.

52. The cassette of claim 37, wherein said at least one sensor includes a potassium sensor.

53. The cassette of claim 37, wherein said chamber has a longitudinal axis and includes an inlet portion, an outlet portion and a middle portion between said inlet portion and said outlet portion, wherein said inlet portion and said outlet portion each have a generally circular configuration in reference planes perpendicular to said longitudinal axis, wherein said middle portion has a non-circular configuration in reference planes perpendicular to said longitudinal axis, said circular configurations of said inlet portion and said outlet portion each having respective areas that are each approximately equal to the area of said non-circular configuration of said middle portion.

54. A cassette adapted for releasable coupling to a blood parameter measuring device for measuring one or more parameters of blood, said cassette comprising:

a body;

at least one sensor connected to said body for sensing at least one parameter of the blood; and

a casing detachably coupled to said body, said casing having walls defining an elongated chamber for receiving a quantity of blood, said casing also having an external rib extending along its length at a location remote from said body.

55. The cassette of claim 54, wherein said cassette includes two leg portions with outer surfaces that extend in respective reference planes that are oriented at an angle in the range of about 28 degrees to about 32 degrees relative to each other.

56. The cassette of claim 55, wherein said body includes an oval-shaped recess and wherein said at least one sensor is located in said recess.

57. The cassette of claim 55, wherein each of said two leg portions includes at least one outwardly extending tab.

58. The cassette of claim 57, wherein said at least one sensor includes at least two sensors, wherein said body

includes a rim surrounding said at least two sensors, and wherein said body includes an alignment key connected to said rim and extending between an adjacent pair of said at least two sensors.

59. The cassette of claim 54, wherein said at least one sensor includes a potassium sensor.

60. The cassette of claim 54 wherein said body includes a coupling having a convex configuration for connection to a blood parameter measuring device.

61. The cassette of claim 54 wherein said casing includes a side opening, and wherein said cassette includes a membrane extending across said side opening.

62. A cassette for measuring one or more parameters of blood comprising:

a casing having walls defining a chamber for receiving a quantity of blood;

at least one sensor connected to said casing adjacent said chamber for sensing at least one parameter of blood in said chamber; and

a coupling for connecting said casing to a blood parameter measurement device, said coupling including a generally oval-shaped recess and wherein said at least one sensor is located in said recess.

63. The cassette of claim 62 wherein said at least one sensor includes at least two sensors and wherein said coupling includes an alignment key extending between an adjacent pair of sensors.

64. The cassette of claim 62 wherein said at least one sensor includes at least two sensors, wherein said coupling includes a rim surrounding said at least two sensors, and wherein said coupling includes an alignment key connected to said rim and extending between an adjacent pair of said at least two sensors.

65. The cassette of claim 62 wherein said at least one sensor includes a potassium sensor.

66. The cassette of claim 62 wherein said coupling is releasably connected to said casing.

67. The cassette of claim 62 wherein said coupling is integrally connected to said casing.

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