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(54) **SYSTEM FOR PERFUSION MANAGEMENT**

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See application file for complete search history.

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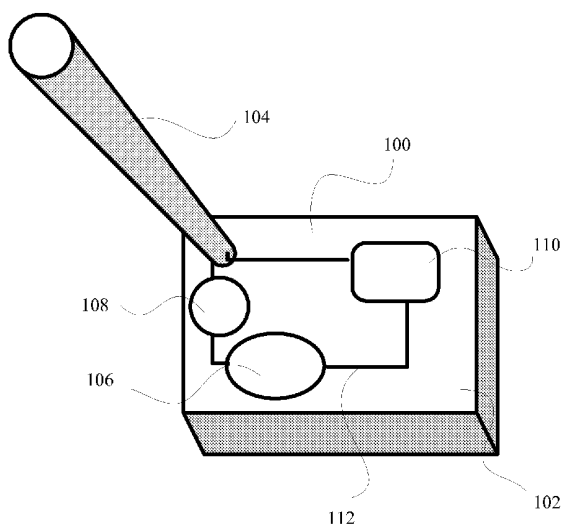
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(57) **ABSTRACT**

A system for perfusion management that monitors, maintains, diagnoses, or treats perfusion deficiencies.

21 Claims, 5 Drawing Sheets



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FIG. 1

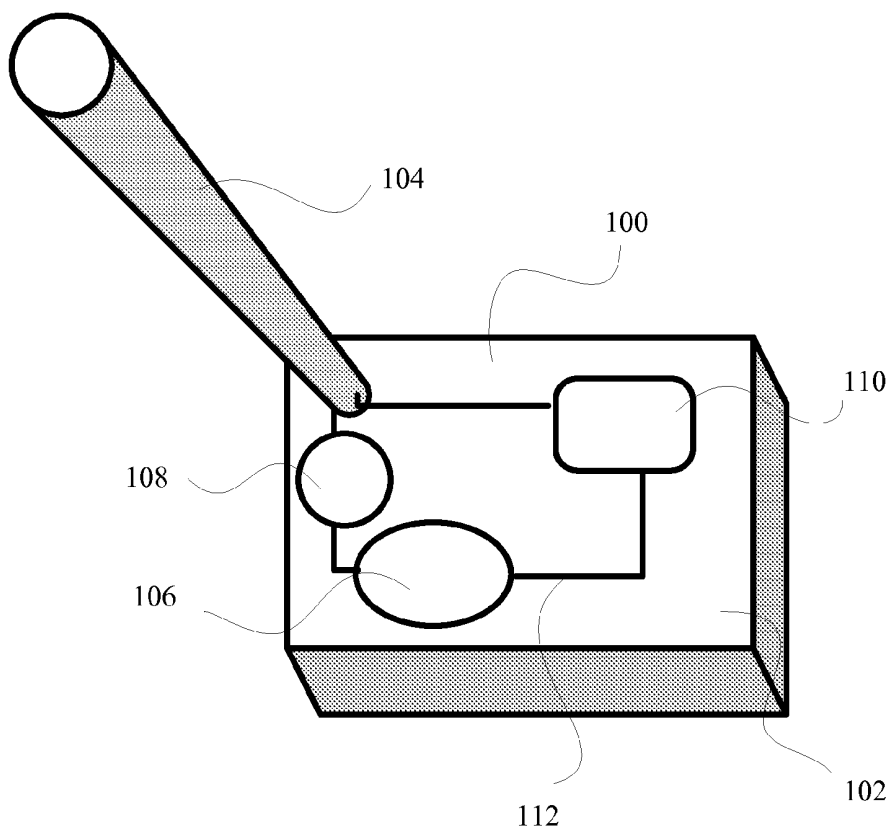
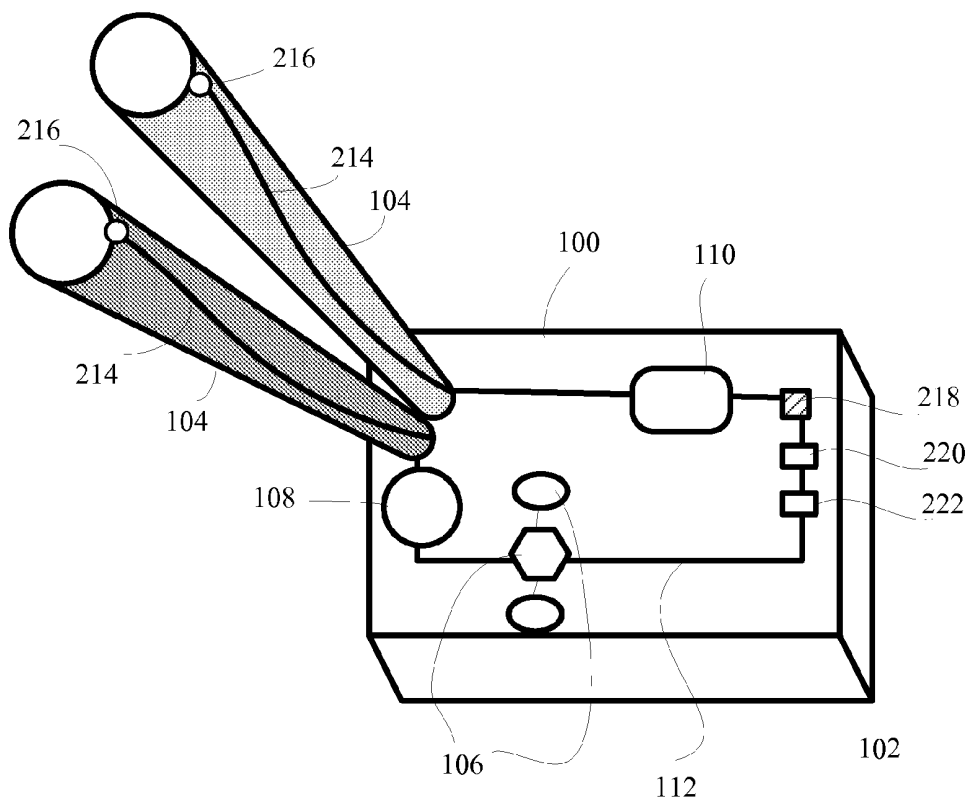


FIG. 2



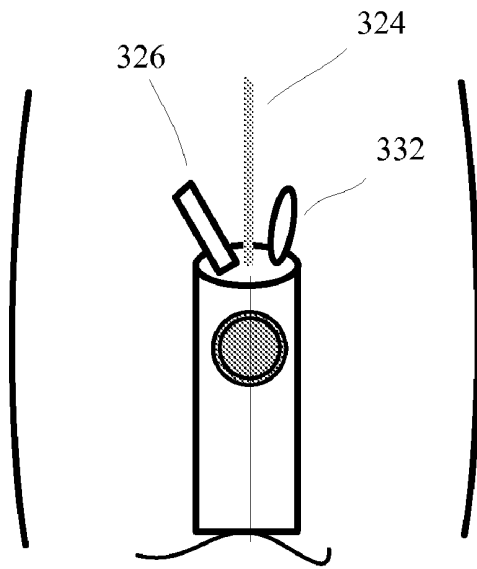


FIG. 3A

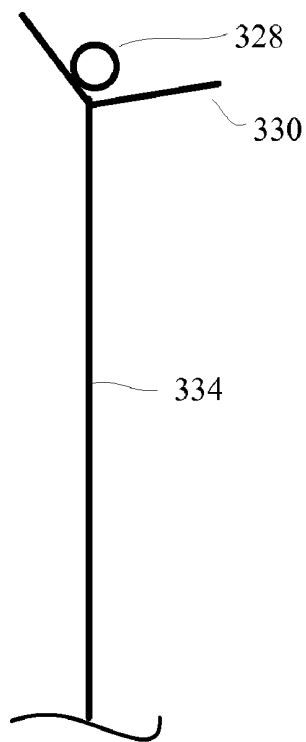


FIG. 3B

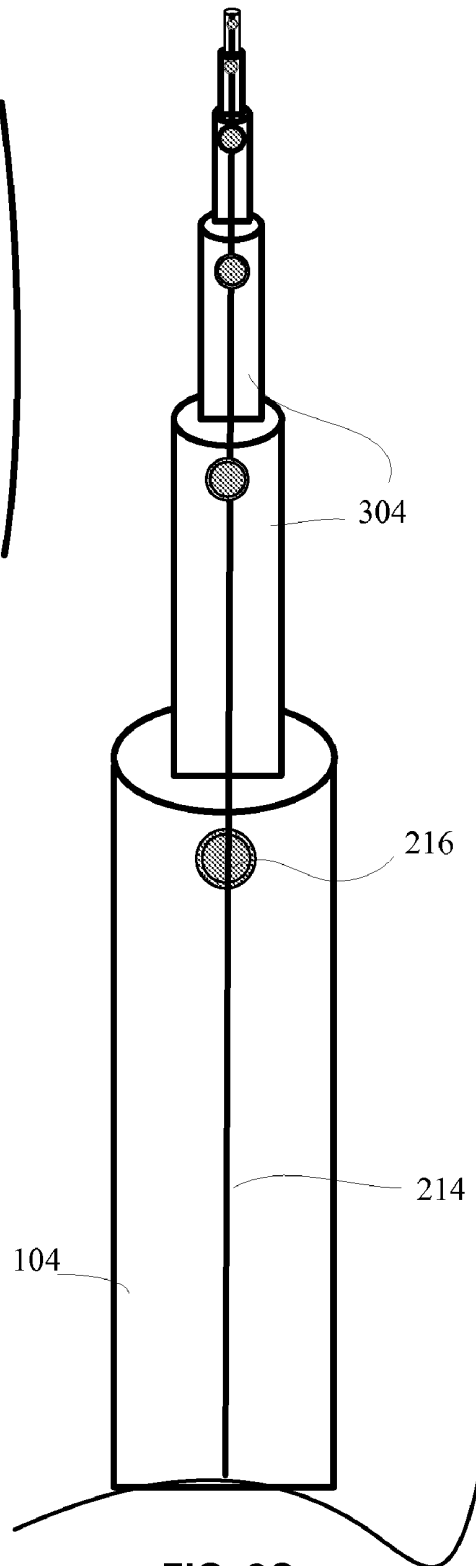


FIG. 3C

FIG. 4

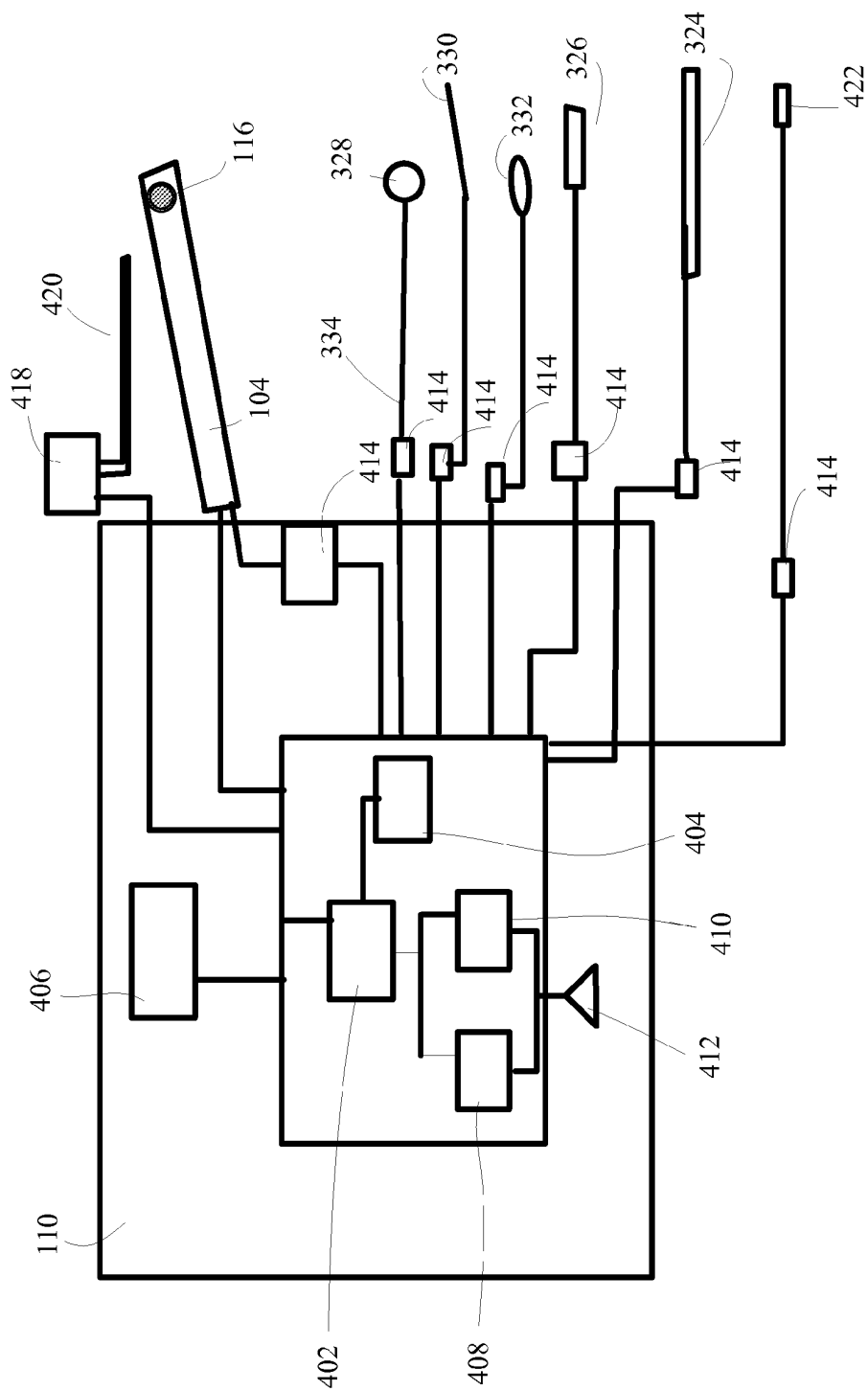
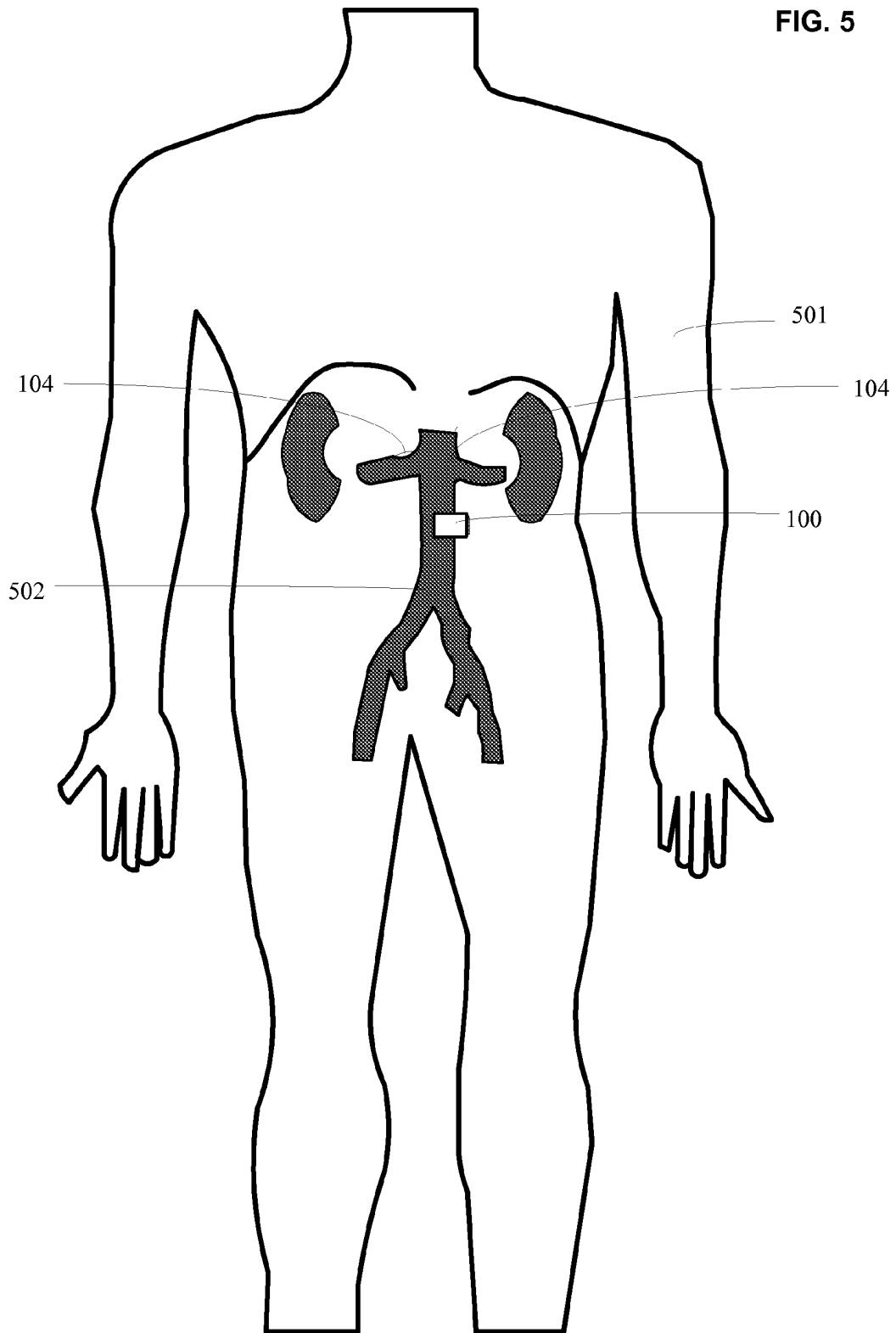


FIG. 5



SYSTEM FOR PERFUSION MANAGEMENT

CROSS-REFERENCE TO RELATED APPLICATIONS

The present application is related to, claims the earliest available effective filing date(s) from (e.g., claims earliest available priority dates for other than provisional patent applications; claims benefits under 35 USC §119(e) for provisional patent applications), and incorporates by reference in its entirety all subject matter of the following listed applications; the present application also claims the earliest available effective filing date(s) from, and also incorporates by reference in its entirety all subject matter of any and all parent, grandparent, great-grandparent, etc. applications of the following listed applications:

1. U.S. patent application Ser. No 10/827,576 entitled A SYSTEM WITH A RESERVOIR FOR PERFUSION MANAGEMENT, naming Lowell L. Wood, Jr. as inventor, filed on Apr. 19, 2004.
2. U.S. patent application Ser. No. 10/827,578 entitled A SYSTEM WITH A SENSOR FOR PERFUSION MANAGEMENT, naming Lowell L. Wood, Jr. as inventor, filed on Apr. 19, 2004.
3. U.S. patent application Ser. No. 11/891,356 entitled A SYSTEM FOR PERFUSION MANAGEMENT, naming Lowell L. Wood Jr. as inventor, filed on Aug. 9, 2007.
4. U.S. patent application Ser. No. 11/891,333 entitled A SYSTEM WITH A RESERVOIR FOR PERFUSION MANAGEMENT, naming Lowell L. Wood Jr. as inventor, filed on Aug. 9, 2007.
5. U.S. patent application Ser. No. 11/891,355 entitled A SYSTEM WITH A RESERVOIR FOR PERFUSION MANAGEMENT, naming Lowell L. Wood Jr. as inventor, filed on Aug. 9, 2007.
6. U.S. patent application Ser. No. 11/891,573 entitled A TELESCOPING PERFUSION MANAGEMENT SYSTEM, naming Lowell L. Wood, Jr. as inventor, filed on Aug. 9, 2007.
7. U.S. patent application Ser. No. 11/891,371 entitled A TELESCOPING PERFUSION MANAGEMENT SYSTEM, naming Lowell L. Wood, Jr. as inventor, filed on Aug. 9, 2007.

TECHNICAL FIELD

The present application relates, in general, to detection and/or treatment.

SUMMARY

In one aspect, a system includes, but is not limited to at least one arm; a control circuit coupled to said at least one arm; at least one receptacle coupled to said at least one arm; and at least one sensor coupled to said at least one arm, said system configured for implantation in an animal. In addition to the foregoing, other system aspects are described in the claims, drawings, and text forming a part of the present application.

In one aspect, a method includes but is not limited to: making a device, by providing a cavity for storing a receivable; and including at least one transmitting channel coupled to said cavity for delivering said receivable from said cavity to a predetermined location in an animal, cooperative with a monitoring system and a detecting system including logic or software operative for delivering said receivable. In addition

to the foregoing, other method aspects are described in the claims, drawings, and text forming a part of the present application.

In another aspect, a method includes but is not limited to: placing a device comprising a receiving unit for storing a deliverable and in communication with a pliable duct; directing said pliable duct to a location; and administering said deliverable from said receiving unit to said location cooperative with a guiding system and a detecting system. In addition to the foregoing, other method aspects are described in the claims, drawings, and text forming a part of the present application.

In one or more various aspects, related systems include but are not limited to circuitry and/or programming for effecting the herein-referenced method aspects; the circuitry and/or programming can be virtually any combination of hardware, software, and/or firmware configured to effect the herein-referenced method aspects depending upon the design choices of the system designer.

In addition to the foregoing, various other method and/or system aspects are set forth and described in the text (e.g., claims and/or detailed description) and/or drawings of the present application.

The foregoing is a summary and thus contains, by necessity; simplifications, generalizations and omissions of detail; consequently, those skilled in the art will appreciate that the summary is illustrative only and is NOT intended to be in any way limiting. Other aspects, inventive features, and advantages of the devices and/or processes described herein, as defined solely by the claims, will become apparent in the non-limiting detailed description set forth herein.

BRIEF DESCRIPTION OF THE FIGURES

FIG. 1 is a front-plan view of a device for perfusion management **100**.

FIG. 2 is a front-plan view of another aspect of the device for perfusion management **100**.

FIGS. 3A, 3B, and 3C show exploded views of various aspects of the device of FIG. 1, including an extensible finger **104**.

FIG. 4 is a schematic view of the control circuit **110** and devices in communication with the control circuit **110**.

FIG. 5 illustrates an example wherein the device for perfusion management **100** is placed in a selected location in a human body **501**.

The use of the same symbols in different drawings typically indicates similar or identical items.

DETAILED DESCRIPTION

The present application uses formal outline headings for clarity of presentation. However, it is to be understood that the outline headings are for presentation purposes, and that different types of subject matter may be discussed throughout the application (e.g., device(s)/structure(s) may be described under the process(es)/operations heading(s) and/or process(es)/operations may be discussed under structure(s)/process(es) headings). Hence, the use of the formal outline headings is not intended to be in any way limiting.

1. Perfusion Management Device(s) and/or Process(es)

With reference now to FIG. 1, shown is a front plan view illustrative of various exemplary perfusion management device(s) and/or process(es). Accordingly, the present application first describes certain specific exemplary structures of FIG. 1; thereafter, the present application illustrates certain specific exemplary processes. Those having skill in the art

will appreciate that the specific devices and processes described herein are intended as merely illustrative of their more general counterparts.

A. Structure(s) and or Device(s)

With reference to the figures, and with reference now to FIG. 1, shown is a front-plan view of a device for perfusion management **100**. The device for perfusion management **100** includes a body portion **102** from which at least one arm **104** projects. A receptacle **106** within the body portion **102** contains a fluid, for example, a fluid for treatment. A controllable valve **108** provides a path through which the fluid may travel to the at least one arm **104**.

A control circuit **110** provides a control signal that may open or close the control valve **108**. The control circuit **110** may be connected to a sense line **112** that allows it to monitor the fluid levels within the receptacle **106**. Additionally, a sense line **114** connects the control circuit **110** to a sensor **116** at the distal end of the at least one arm **104**.

Referring now to FIG. 2, depicted is an aspect of the device for perfusion management **100** which includes the body portion **102** from which a set of at least one arm **104** projects. In one aspect, each of the at least one arm **104** is in fluid communication with a respective receptacle **106**. Each receptacle **106** may be filled with a different fluid for delivery. In another approach, the one or more receptacles **106** may be coupled to a mixing chamber **224** where the fluid contents of each receptacle **106** are present for mixing and the mixed contents enter the arm for delivery to a selected location. The choice of fluid in the receptacle **106** may depend, for example, on the purpose of the device, for example, treatment of colon cancer, treatment of breast cancer, or treatment of an arterial disease. The choice of fluid in the receptacle **106** includes, but is not limited to, for example, a chemical, a chemical compound, a protein, a lipoprotein, a glycoprotein, a sugar, a lipid, an antigen, an antibody, a cytokine, a peptide, a neurotransmitter, a hormone, an ion, a messenger molecule, a nucleic acid, an engineered nucleic acid, a nucleic acid vector, a drug, a cell, a cell fragment, a cell organelle, a liposome, a pharmaceutical agent, a biological material, or a biological fraction. The receptacle **106** may also be utilized for storage and disposal of operational fluids. Also, although the exemplary embodiment described herein focuses primarily on fluid delivery, one skilled in the art will understand that fluid-like substances, such as gels, and fluidizable substances or non-fluid type substances, such as small solid particles, may be delivered in accordance with the invention. It will also be appreciated by those having skill in the art that the nature of the fluid in the receptacle **106** includes, for example, and is not limited to, a liquid, a solution, a mixture, a gel, a colloid, a colloid of a suitable viscosity, a suspension, an emulsion, or any material of low shear-strength for delivery to a site.

In one aspect one or more fluids are delivered to one or more of selected locations by the device for perfusion management **100**. The selected location may be, for example, in proximity to or within a tumor, a circulatory system, an aorta, a vena cava, a site of therapy, or a site of investigation in an animal.

Continuing to refer to FIG. 2, a pump **218** provides fluid at a controlled flow rate for delivery to a site from the receptacle **106**. It will be appreciated by those skilled in the art that the type of pump is not critical to the invention and may include, for example, a mechanical pump, a piezoelectric pump, an osmotic pump, a source of pressure, or a device for maintaining a positive flow of fluid through the device. Additionally, fluid flow may be further modulated with micro valves and self-pressurizing fluidic reservoirs. Moreover, in some applications, the fluid may be delivered without a pump. For

example, fluid delivery may be controlled using a pressurized bladder, controlled dissolution or dilution of a material, a drip or gravity type of approach, or any other suitable approach to deliver an appropriate amount or an appropriate delivery-rate of the fluid.

With reference now to FIG. 3, depicted is an exploded view of the arm **104** showing a plurality of extended parts **304** with the sensor **116** at the distal end of each of the extended parts **304**. In one aspect of the invention, the sensor **116** is an array of sensors, deployed from one or more portholes, at the distal end of each of the extended parts **304**. In one approach, the portholes are sized and shaped to provide access through which the sensors **116** may be deployed. The portholes may include seals, stress relief or other features appropriate for proper mechanical deployment. In one approach, one or more of the portholes can be controllably opened or closed to provide communication exterior to the arm or main body. The sensor **116** may be retracted within the porthole and deployed through the porthole. Where the porthole can be opened and closed, the porthole can close to limit communication and can be opened for deployment. The array of sensors may include, but is not limited to, for example, sensors for detecting pressure, temperature, chemical, gas, electrolyte, flow, volume, composition, or concentration. In an alternate aspect of the invention, microelectrodes, such as, for example, solid-state microelectrodes, are sensitized with an agent for detecting a relevant interacter. Examples of the agent include, but are not limited to, for example, agonists of angiogenesis. The choice of sensor **116** depends on the physiological variable being monitored, treated, or controlled. The term "physiological variable" refers to any and all measurements relating to the functioning of a living organism in normal, sub-normal, or abnormal states.

Continuing to refer to FIG. 3 and referring now to FIG. 4, an operative tool **324** is coupled to the distal most extended part **304**, or deployed from the porthole, or carried by the arm **104**, further including a carrying line **334** in communication with the control circuit **110**. The operative tool **324** includes, but is not limited to, for example, one or more of a combination of, a tool positioner, an ablation device, a laser, a vacuum, a siphon **326**, an evacuation device, a fluid dispenser **328**, a cauterizer **330**, a stent **332**, a tissue-liquefying device, or a source of an electric charge or electromagnetic radiation **422**. The vacuum or the siphon **326** is employed for removing a cell, a mass of cells, a tissue, a fluid, a gel, a sample, a debris, a contaminant, or other material for which removal is desired or appropriate. The ablation device operates for perturbing or reducing the structural integrity or viability of a cell, a mass of cells, an assembly of biological materials exhibiting shear strength, or a tissue. The assembly of biological materials includes, for example, blood clots, cartilage, or bone. The source of an electric charge or electromagnetic radiation **422** includes, but is not limited to, for example, steady state electric currents, time-varying electric currents, alternating electric currents, radio waves, microwaves, ultraviolet rays, infrared rays, optical rays, terahertz beams, and the like.

Continuing to refer to FIG. 3, it will be appreciated by those having skill in the art that the operative tool **324** may include a set of devices having general or "multi-purpose" utility. The operative tool **324** may include, but is not limited to, for example, a combination of the fluid dispenser **328**, the siphon **326**, and the ablation device. In this example the operative tool combination, for example, delivers the fluid or gel, ablates cells, and removes debris.

Continuing to refer to FIG. 3, the plurality of extended parts **304** may themselves be hollow forming a conduit for delivery of the fluid to a site, or for housing a circuitry cou-

pling the control circuit 110 to the operative tool 324, or for housing a mechanism that guides the arm 104 or the plurality of extended parts 304.

With reference now to FIG. 4, illustrated is a schematic view of the control circuit 110 and devices in communication with the control circuit 110. The device for perfusion management 100 shows a data transmitter 410, and a data receiver 408 coupled to the control circuit 110. An antenna 412 may be used for transmitting data to the exterior wirelessly. The antenna 412 is shown diagrammatically, but may be a structure, such as a strip antenna, that may be integrated in a manner that does not impair or significantly perturb system performance. The control circuit 110 is depicted as having a processor 402 coupled to a memory 404 that provides data storage and retrieval capability, and a power source 406. Feedback circuitry or logic circuitry provides communication between the control circuit 110 and devices in communication with it. In some applications, a software program providing instructions may be stored in the memory 404 to control operation of the control circuitry or to store data gathered under control of the control circuitry. Additionally, the control circuit 110 may have components for system integrated digital data gathering, processing, storage, compression and transmission. These can provide data control capabilities and operation control capabilities. For example, the transmission components may communicate through the antenna 412 to a person, system, computer, or device exterior to the body. This communication can allow data gathered by the sensors to be displayed, stored or otherwise processed in the external environment. Additionally, this communication may allow for the processed data or a plurality of new data to be received from the exterior by the device for perfusion management 100. Data compression can allow the control circuitry to store data representing larger amounts of data to be stored in the memory 404 or to be transmitted to the exterior environment in a more efficient manner.

Continuing to refer to FIG. 4, one or more of the operative tools 324 are mounted on an actuator 414 which allows for the independent movement of each tool. Alternatively, one or more operative tools 324 may be mounted as a unit on one actuator 414 and moved as a group, for example, forming an aspirating-dispensing unit. For example, the fluid dispenser 328 and the siphon 326 may be mounted together as a group. The actuator 414 may be a motor, a piezo electrically driven actuator, a micromechanical or electrical effector, or the like.

Continuing to refer to FIG. 4, the arm 104 may include an imaging device deployed from the porthole or from the distal end of the arm 104 or carried by a carrying line 334. The term "imaging device" being used herein to designate in general those components, circuits, assemblies and sub-assemblies comprising electrical, optical, acoustic, or opto-electronic components. In one aspect, the control circuit 110 is coupled to the imaging device that includes a laser 418, or a source of light or scene-illuminating radiation, coupled to an optical feed line 420 to illuminate an area. A charge coupled device is positioned to capture data from the illuminated area and provides an electronic signal indicative of the area imaged. Conventional circuitry then produces a digital representation that may be displayed, stored in the memory 404, or otherwise processed. The displayed image may serve, for example, for guiding the arm 104 to the selected location or for determining the efficacy of a treatment or a procedure. One skilled in the art will recognize that the imaging device described herein is exemplary of imaging devices and that other imaging devices, including for example, raster and line-scanning imagers, nonvisible spectral imagers, and fluorescence imagers, may be included.

With reference now to FIG. 5, the device for perfusion management 100 is depicted implanted in an aorta 502 with the arm 104 traveling a blood vessel in a human body 501. Additionally, the device for perfusion management 100 is configured for full or partial placement in the human body 501. The configuration may incorporate a combination of the following criteria, including but not limited to, dimensions, composition, shape, power dissipation level, or texture. In one aspect, the body portion 102 is sized for implantation in proximity to the aorta 502 or the vena cava and the arm 104 is sized for traveling a blood vessel in an animal, for example, the human body 501. In this aspect, if the vasculature decreases two-fold, each of the extending parts 304 has about a two-fold decrease in diameter. The length of the arm 104, for example, depends upon the distance between the selected location and the location of the body portion 102, and the route traveled by the arm 104 to arrive at the selected location. It will be appreciated by those having skill in the art that the arm 104 including the one or more of the operative tools 324 is of a size, dimension or shape operable for traveling one or more blood vessel of decreasing or increasing luminal diameter. It will also be appreciated by those having skill in the art that the arm 104 and the one or more operative tool 324 may pass through the wall of a lumen, or trans-luminally, to the surrounding tissue for detecting, delivery of a treatment, or for sampling. It will also be appreciated by those having skill in the art that the trans-luminal mode described is not limited to blood vessels and includes the space or cavity of an organ or structure.

It will also be appreciated by those having skill in the art that the device for perfusion management 100 and its components, such as, for example, the arm 104, the plurality of extended parts 304, or one or more operative tools 324, has a size, dimension, shape, material, and properties of flexion, retraction, and extension to allow for the steering, guiding, or positioning of the components of the device for perfusion management 100. For example, the arm 104 may need to be steered around an occlusion or a fork in the vasculature. In this example, the arm 104 may need to be retracted, repositioned and then extended in a new direction. Extending, retracting or repositioning of the arm 104 may be accomplished by techniques known in the art, for example, by using a guide wire or a by employing a shape polymer. In another aspect, the arm may be retracted and then "punched through" an occlusion to dislodge, penetrate, or grapple it. In this example, a laser beam, a shearing tool, a cutting tool, or a drug, for example, a lytic drug, may be employed to degrade or reduce the occlusion. In this example, subsequent to the dislodgement and degradation of the occlusion, the siphon 326 or an evacuation device is employed to evacuate any debris, before the arm 104 continues traveling the circulatory system. It will also be appreciated by those skilled in the art that the device for perfusion management 100 is not restricted to traveling the circulatory system but may be implanted in any tissue, such as, for example, nerve, epithelial, dermal, sub-dermal, connective, or muscle tissue. Additionally, the device for perfusion management 100 may be implanted in inter-tissue spaces, or inter-organ spaces, for example, those found within a body cavity.

In one aspect the device for perfusion management 100 includes an array of sensors 116 positioned across the plurality of extended parts 304 for monitoring, tracking, or mapping a gradient of temperature, pressure, flow, or material concentration in one or more locations. The one or more locations may be, for example, a tissue, an artery, or a vein. In another aspect the device for perfusion management 100 has an auto-correct feature for correcting a sub-normal or abnormal gra-

cient of temperature, pressure, flow, or material concentration. In yet another aspect the device for perfusion management **100** has an auto-correct feature for detecting and correcting a sub-normal or abnormal gradient of temperature, pressure, flow, or material concentration.

The device for perfusion management **100** may be composed of materials known in the art, for example, a metal, a ceramic, a glass, a plastic, a polymer, a biologically compatible material, or a combination. For example, the device for perfusion management **100** may be made of helically-coiled stainless steel wire and coated with a polymer, such as, Teflon™. In another example; the device for perfusion management **100** may be made of helically-coiled stainless steel wire and coated with a polymer and impregnated with one or more of a biological material, for example, including but not limited to, anti coagulants, or inhibitors.

B. Operation(s) and/or Process(es)

Those having skill in the art will appreciate that some or all of the components of the device for perfusion management **100** may be present ex vivo. In one implementation, the device for perfusion management **100** is placed in proximity to the location on the animal, for example, the human body **501**, and the at least one arm **104** directed to the selected location and an effective agent delivered in proximity to the selected location. The arm **104** may be retracted after such a delivery, leaving the device for perfusion management **100** in place at the location, until time for a future delivery of the effective agent or another procedure or operation. In this implementation, the majority of the device for perfusion management **100** is ex vivo while the arm **104** alternates between ex vivo and in vivo states.

In another aspect, some or all the components of the device for perfusion management **100** are present in vivo. In one implementation, the device for perfusion management **100** is placed in proximity to the location within the animal, for example, the human body **501**, and the arm **104** directed to a selected location and an effective agent delivered in proximity to the selected location. The arm **104** may be retracted after such a delivery, leaving the device for perfusion management **100** in place at the location, until time for a future delivery or another operation. In this implementation, the majority of the device for perfusion management **100** is in vivo while the arm **104** alternates between retracted, partially retracted or unretracted states.

In one implementation, the device for perfusion management **100** is operable by a person. The person monitors, guides, positions, and performs other actions/operations or manages a response consistent with the device for perfusion management **100** being managed by the person. In such an implementation a separate display device can present imagery to aid the person. The imagery may be captured as described above with reference to FIG. **4**, may be computer generated or may be captured by a separate imaging device internal to or external to the animal, for example, the human body **501**. Actions may be performed under control of the person who may be on site or may be linked from a remote location, or the device for perfusion management **100** may be programmed to perform some or all functions automatically. For example, the device for perfusion management **100** may be programmed to perform functions, such as, lumen clearance, lumen maintenance, monitoring of concentrations, sending of alerts, delivery of one or more of the effective agent at timed intervals or locations, self-check, or self-diagnosis. It will be appreciated by those of skill in the art that the device for perfusion management **100** may be programmed for complete automatic operation of one or more functions.

C. Variation(s), and/or Implementation(s)

Those having skill in the art will recognize that the present application teaches modifications of the devices, structures, and/or processes within the spirit of the teaching herein. For example, the device for perfusion management **100** need not be limited to managing perfusion. The device provides a mechanism for exploring one or more regions and/or reaching a location within an animal, obtaining information, communicating this information, performing operations, performing procedures, or providing treatment. The treatment includes but is not limited to, for example, treatment of a subnormal, abnormal or pathological condition. In another example, the device for perfusion management **100** may find utility in the management of physiological functions, the detection or elimination of pathological functions or conditions, and/or treatment of a disease or a pathological state of non-human animals. Other modifications of the subject matter herein will be appreciated by one of skill in the art in light of the teachings herein.

The foregoing described aspects depict different components contained within, or connected with, different other components. It is to be understood that such depicted architectures are merely exemplary, and that in fact many other architectures can be implemented which achieve the same functionality. In a conceptual sense, any arrangement of components to achieve the same functionality is effectively “associated” such that the desired functionality is achieved. Hence, any two components herein combined to achieve a particular functionality can be seen as “associated with” each other such that the desired functionality is achieved, irrespective of architectures or intermedial components. Likewise, any two components so associated can also be viewed as being “operably connected”, or “operably coupled”, to each other to achieve the desired functionality.

While particular aspects of the present subject matter described herein have been shown and described, it will be obvious to those skilled in the art that, based upon the teachings herein, changes and modifications may be made without departing from this subject matter described herein and its broader aspects and, therefore, the appended claims are to encompass within their scope all such changes and modifications as are within the true spirit and scope of this subject matter described herein. Furthermore, it is to be understood that the invention is defined solely by the appended claims. It will be understood by those within the art that, in general, terms used herein, and especially in the appended claims (e.g., bodies of the appended claims) are generally intended as “open” terms (e.g., the term “including” should be interpreted as “including but not limited to,” the term “having” should be interpreted as “having at least,” the term “includes” should be interpreted as “includes but is not limited to,” etc.). It will be further understood by those within the art that if a specific number of an introduced claim recitation is intended, such an intent will be explicitly recited in the claim, and in the absence of such recitation no such intent is present. For example, as an aid to understanding, the following appended claims may contain usage of the introductory phrases “at least one” and “one or more” to introduce claim recitations. However, the use of such phrases should not be construed to imply that the introduction of a claim recitation by the indefinite articles “a” or “an” limits any particular claim containing such introduced claim recitation to inventions containing only one such recitation, even when the same claim includes the introductory phrases “one or more” or “at least one” and indefinite articles such as “a” or “an” (e.g., “a” and/or “an” should typically be interpreted to mean “at least one” or “one or more”); the same holds true for the use of definite articles

used to introduce claim recitations. In addition, even if a specific number of an introduced claim recitation is explicitly recited, those skilled in the art will recognize that such recitation should typically be interpreted to mean at least the recited number (e.g., the bare recitation of “two recitations,” without other modifiers, typically means at least two recitations, or two or more recitations), etc.

Furthermore, in those instances where a convention analogous to “at least one of A, B, and C, etc.” is used, in general such a construction is intended in the sense one having skill in the art would understand the convention (e.g., “a system having at least one of A, B, and C” would include but not be limited to systems that have A alone, B alone, C alone, A and B together, A and C together, B and C together, and/or A, B, and C together, etc.). In those instances where a convention analogous to “at least one of A, B, or C, etc.” is used, in general such a construction is intended in the sense one having skill in the art would understand the convention (e.g., “a system having at least one of A, B, or C” would include but not be limited to systems that have A alone, B alone, C alone, A and B together, A and C together, B and C together, and/or A, B, and C together, etc.). It will be further understood by those within the art that virtually any disjunctive word and/or phrase presenting two or more alternative terms, whether in the description, claims, or drawings, should be understood to contemplate the possibilities of including one of the terms, either of the terms, or both terms. For example, the phrase “A or B” will be understood to include the possibilities of “A” or “B” or “A and B.”

What is claimed is:

1. A method for making a device, comprising: providing a cavity for storing a receivable; including at least one transmitting channel coupled to said cavity for delivering said receivable from said cavity to a predetermined location in an animal, cooperative with a monitoring system and a detecting system including logic and/or software operative for delivering said receivable; and coupling one or more operative tools deployable via one or more openable and/or closeable portholes to the transmitting channel.
2. The method according to claim 1, further comprising transferring a fluid to said cavity.
3. The method according to claim 2, wherein said device includes a pump, and/or a source of pressure, coupled to said cavity for transferring said fluid.
4. The method according to claim 1, further comprising coupling a motor and/or an actuator to said at least one transmitting channel.
5. The method according to claim 1, further comprising configuring said device to be monitored and/or controlled by a person.
6. The method according to claim 1, further comprising configuring said device for implantation in said predetermined location in said animal.
7. The method according to claim 1, further comprising coupling said cavity to a mixing cavity.

8. The method according to claim 1, further comprising coupling a source of a chemical, a chemical compound, a protein, a lipoprotein, a glycoprotein, a sugar, a lipid, an antigen, an antibody, a cytokine, a peptide, a neurotransmitter, a hormone, an ion, a messenger molecule, a nucleic acid, an engineered nucleic acid, a nucleic acid vector, a drug, a cell, a cell fragment, a cell organelle, a liposome, a pharmaceutical agent, a biological material, and/or a biological fraction internal and/or external to said cavity.

9. The method according to claim 1, further comprising coupling a source of two or more of a chemical, a chemical compound, a protein, a lipoprotein, a glycoprotein, a sugar, a lipid, an antigen, an antibody, a cytokine, a peptide, a neurotransmitter, a hormone, an ion, a messenger molecule, a nucleic acid, an engineered nucleic acid, a nucleic acid vector, a drug, a cell, a cell fragment, a cell organelle, a liposome, a pharmaceutical agent, a biological material, and/or a biological fraction internal and/or external to said cavity.

10. The method according to claim 1, further comprising coupling a device for data gathering, processing, storage and/or transmission to said monitoring system and/or said detecting system.

11. The method according to claim 1, wherein said monitoring system communicates wirelessly.

12. The method according to claim 1, wherein said monitoring system is operative for guiding and/or extending and/or moving said transmitting channel.

13. The method according to claim 1, wherein said monitoring system comprises a processor, a feedback circuit, and/or a logic circuit.

14. The method according to claim 1, wherein said monitoring system is a processor further comprising a stored software and/or firmware program cooperative with said processor.

15. The method according to claim 1, wherein said transmitting channel is formed from a plurality of extending parts.

16. The method according to claim 1, wherein said plurality of extending parts includes portions that are hollow.

17. The method according to claim 1, further comprising coupling a flow regulator and/or a flow monitor to said transmitting channel and/or said cavity.

18. The method according to claim 1, further comprising coupling a device for partially blocking and/or shunting a liquid flow to said transmitting channel.

19. The method according to claim 1, further comprising coupling a dispenser to said transmitting channel, said dispenser including a fluid controller operative to provide a fluid at a controlled rate.

20. The method according to claim 1, further comprising coating said at least one transmitting channel with a polymer and/or a biocompatible material.

21. The method according to claim 1, wherein said detecting system includes coupling an imager, a pressure sensor, a temperature sensor, a chemical sensor, a gas sensor, an electrolyte sensor, a concentration sensor, a composition sensor, and/or a flow sensor coupled to said at least one transmitting channel.

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摘要(译)

用于灌注管理的系统，其监测，维持，诊断或治疗灌注缺陷。

