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(54) **Title:** A DEVICE, SYSTEM AND METHOD FOR THE CONTINUOUS PROCESSING AND SEPARATION OF BIOLOGICAL FLUIDS INTO COMPONENTS

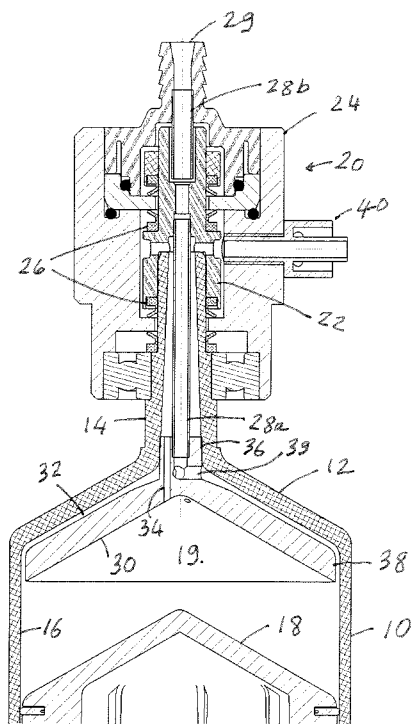


FIG 1B

(57) **Abstract:** A device for the processing and separation of biological fluids into components comprises a hollow centrifugal processing chamber (10) fitted with an inlet/outlet head (20) and preferably with an axially movable piston (18). The inlet/outlet head has two separate inlets/outlets, for instance an axial inlet (29) and a lateral outlet (40). The processing chamber (1) is fitted with an internal flow guide (30) enabling operation of the device in a continuous processing mode wherein biological fluid to be processed is continuously intaken by say the axial inlet (29) and at the same time processed components are continuously removed via say the lateral outlet (40). The continuous processing flow can be driven by an external peristaltic pump (59) and/or by axial displacement of a piston (18) in the chamber (10).

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A Device, System and Method for the Continuous Processing and Separation of Biological
Fluids into Components

Field of the Invention

- 5 This invention relates to a device, a system and a method for the processing and separation of biological fluids into components.

Background of the Invention

The invention relates in particular to a device, a system and a method for the processing and separation of biological fluids into components of the type known from EP-B-0 912 250 and
10 EP-B-1 144 026. Such known devices comprise a hollow centrifugal processing chamber rotatable about an axis of rotation. The processing chamber usually has an inner cylindrical wall enclosing a space for receiving biological fluids to be processed and having an inclined upper wall leading to a neck with an inlet/outlet for biological fluid to be processed and for processed components of the fluid. The space in the hollow processing chamber for receiving
15 biological fluids occupies the entire volume across the hollow processing chamber between facing parts of the inner cylindrical wall and along the length of the cylindrical wall, said space having a given volume delimited by a bottom of the hollow cylindrical chamber, or a variable volume delimited by the position of an axially movable member within the cylindrical wall. The device also comprises a dynamic/static inlet/outlet head or “rotary seal”
20 mounted on the neck of the processing chamber. The inlet/outlet head has a first part rotatable with the processing chamber and a second part that remains stationary. The first and second parts of the inlet/outlet head have a sealing means allowing rotation of the first part relative to the second part. The inlet/outlet head has a central through-passage for the inlet of biological fluid to be processed and the outlet of processed/separated components of the
25 fluid.

Typically the centrifugal processing chamber contains within its cylindrical wall an axially movable member such as a piston that defines a separation space of variable volume wherein the biological fluid to be processed and separated is received. By axially moving this member, biological fluids can be intaken or output.

This known device forms part of a system wherein the processing chamber is part of a disposable set comprising a plurality of containers for receiving on the one hand the biological fluid to be processed and separated and, on the other hand, the separated components, and optionally one or more additional containers for additive solutions.

5 The known device and system are suitable for processing and separating many types of biological fluids including whole blood, apheresis blood, bone marrow blood and expanded cells or stem cells. However, the quantity of biological fluid that can be processed/separated at one time is limited by the maximum volume of the processing chamber's separation space. Therefore, in order to process large volumes of biological fluids with this known device and
10 system, it is necessary to resort to using multiple disposable sets one after another in a single apparatus, or to run several disposable sets in parallel in multiprocessing apparatus.

Other devices for processing and separating biological fluids are known, but their structure is incompatible with that of EP-B-1 144 026 and they do not achieve the same advantages as EP-B-1 144 026. For example WO 2012/137086 describes a blood centrifugal device with a
15 rotatable inner bowl disposed within a rotatable housing defining therebetween a narrow gap for receiving the blood being processed. A stationary structure is disposed in a central recess along the central axis of the device for delivering blood from an inlet at the top to the bottom where it passes into the bottom end of the narrow gap. At the top of the gap a separator separates blood components.

20 Summary of the Invention

An object of the invention is to overcome the mentioned limitation of the known device/system EP-B-1 144 026 by proposing along the following lines an improved portable device working in combination with a disposable centrifugal processing chamber that has the capacity to sediment and separate biological fluid components from a large volume quantity
25 of a biological fluid, including whole blood, apheresis blood, bone marrow blood and expanded cells or stem cells through culturing techniques, down to a very small quantity of volume, by combining a continuous processing flow mechanism advantageously with a variable size processing chamber through displacement of an axially movable element such as a piston.

According to one aspect of the invention there is provided a device of the above-mentioned type which further comprises a flow guide inserted in and rotatable with the processing chamber, the flow guide being located adjacent to, spaced slightly apart from and in shape-matching relationship to the inclined upper wall of the processing chamber to define with the
5 inclined upper wall of the processing chamber an inclined, preferably annular, flow passage, the flow guide being disposed above said space for receiving biological fluids to be processed with its inclined flow passage leading into a top part of the inner cylindrical wall of the processing chamber.

In this improved device, the inlet/outlet head comprises a first inlet/outlet and a second
10 inlet/outlet, the first and second inlets/outlets being disposed on the stationary second part of the inlet/outlet head at a top end and/or at a lateral side thereof, as well as an axial separator in the central through-passage of the inlet/outlet head, the axial separator defining separate first and second axially-directed passages in the inlet/outlet head.

An upper part of one of the first and second axially-directed passages communicates with
15 one of the first and second inlets/outlets, and an upper part of the other of the first and second axially-directed passages communicates with the other of the first and second inlets/outlets. Moreover, a lower part of one of the first and second axially-directed passages communicates with said space in the processing chamber for receiving biological fluids to be processed, and a lower part of the other of the first and second axially-directed passages
20 communicates with the inclined flow passage between the flow guide and the inclined upper wall of the processing chamber.

The inventive device is so arranged that it is operable in a continuous flow mode in which biological fluid to be processed can be inlet via one of the first and second inlets/outlets, while processed biological fluid is simultaneously outlet via the other of the first and second
25 inlets/outlets.

In one embodiment, the axial separator is a central tube extending through the central through-passage of the inlet/outlet head, the central tube being connected (i) to deliver incoming biological fluid through the inside of the central tube and through the space between the flow guide and said inclined upper wall of the processing chamber, to the inner
30 cylindrical wall of the processing chamber, and (ii) to extract processed fluid components

separated at said inclined wall of the processing chamber and that pass through the space between the flow guide and said inclined upper wall of the processing chamber.

In this improved device, an inside part of the processing chamber located beyond (i.e. below) the flow guide can for example be in fluid communication via an aperture in the flow guide to the central through-passage of the inlet/outlet head outside said central tube of the inlet/outlet head; and the stationary second part of the inlet/outlet head has an outlet for the extraction of processed biological fluid components from the inside of the processing chamber, said outlet being in fluid communication with said central through-opening of the inlet/outlet head outside said central tube.

10 In one embodiment, the stationary second part of the inlet/outlet head comprises an external generally cylindrical body, and said outlet is a lateral outlet located in a lateral side of the generally cylindrical body. In this embodiment, the processing chamber can have an upwardly-projecting elongated central neck that extends into the inside of the rotatable first part of the inlet/outlet head up to adjacent the level of the lateral outlet. Also, in this case, the sealing means of the inlet/outlet head can comprise a first seal located axially on one side of the lateral outlet and a second seal located axially on the other side of the lateral outlet.

The axial separator such as a central tube can comprise a stationary axially-outer part extending in the stationary second part of the inlet/outlet head, and a rotatable axially-inner part connected to a central part of the flow guide for rotation therewith. In this embodiment the inner part of the central tube communicates with the space between the flow guide and the inclined upper wall of the processing chamber.

The rotatable first part of the inlet/outlet is typically located inside the stationary second part of the inlet/outlet head. The first inlet /outlet is typically an axial inlet/outlet on the inlet/outlet head and the second inlet /outlet is on a lateral side of the inlet/outlet head.

25 In a particular embodiment of the device: (i) the inclined upper wall of the processing chamber is frusto-conical as is the upper surface of the flow guide or diverter/inverter; (ii) the flow guide has a central sleeve that fits in the neck of the processing chamber leaving a space of several millimeters between the facing frusto-conical surfaces; (iii) the lower end of a central tube forming the axial separator fits in the central sleeve of the flow guide and

communicates with said space between the facing frusto-conical surfaces; and (iv) said aperture in the flow guide is in the form of at least one through passage, preferably three equally-distributed through passages, in its central sleeve that communicate(s) the inside of the processing chamber with the central passage of the inlet/outlet head outside its central tube.

The flow guide or “diverter/inverter” typically comprises a peripheral rim of external cylindrical shape that extends from the periphery of a frusto-conical upper surface of the flow guide. This peripheral cylindrical rim fits in the inner cylindrical wall of the processing chamber below the junction of the inner cylindrical wall and the inclined upper wall of the processing chamber, leaving a small space

As in the known device, the processing chamber usually contains within its cylindrical wall an axially movable member such as a piston that defines a separation space of variable volume wherein the biological fluid to be processed and separated is received. Also as in the known device, the processing chamber of the device according to the invention is part of a disposable set comprising a set of containers for receiving the biological fluid to be processed and separated and the separated components, and optionally one or more additional containers for additive solutions. The device according to the invention also preferably further comprises a cabinet for receiving the processing chamber, the cabinet having drive means for driving the centrifugal processing chamber and preferably also means for controlling the axial position of an axially movable member such as a piston in the processing chamber.

The device according to the invention is also preferably provided with closure members for the inlet/outlet to the processing chamber, said closure members being clips or pinch valves that act on tubing of the disposable set, and/or stopcocks included in the disposable set and/or fitted on a cabinet for receiving the processing chamber.

Another aspect of the invention is a system for the processing and separation of biological fluids into components, comprising the device as discussed previously and further comprising at least one peristaltic pump for pumping incoming biological fluid through the inside of the central tube; and/or for pumping extracted processed biological fluid

components via the central tube; and/or for pumping extracted processed biological fluid components via said outlet on the second part of the inlet/outlet head.

This system optionally comprises a first peristaltic pump operable to pump incoming biological fluid through the inside of the central tube into the processing chamber, and a
5 second main peristaltic pump operable to pump extracted processed biological fluid components via said outlet on the second part of the adaptor head, the first and optional second peristaltic pumps being optionally operable simultaneously to provide continuous flow processing. However, continuous flow can be achieved also and preferably with a single peristaltic pump.

10 Another aspect of the invention is a method of processing and separating biological fluids into components using the device or the system as discussed above, the method comprising the following steps:

- 15 (a) filling the processing chamber with a biological fluid via the first inlet/outlet in the inlet/outlet head, for example an axial inlet via a central tube, with the second inlet/outlet of the inlet/outlet head closed, for example a lateral outlet on the second part of the adaptor head closed;
- (b) rotating the processing chamber to centrifuge biological fluid in the chamber in order to separate different components of the biological fluid onto the inner cylindrical wall of the processing chamber or lighter or waste components into the inside of the
20 processing chamber;
- (c) operating in a continuous processing mode wherein filling of the processing chamber with a biological fluid via a first inlet/outlet, for example via a central tube, is carried out while at the same time extracting, via a second inlet/outlet, for example a lateral outlet on the second part of the inlet/outlet head, waste or light components that have
25 separated into the inside of the processing chamber;
- (d) closing of said first inlet/outlet whereby the processing chamber is no longer being filled with biological fluid, and continuing to extract components , for example via a lateral outlet; and

(e) closing of said second inlet/outlet, for example a lateral outlet whereby processed components that have separated into the inside of the processing chamber are no longer extracted via said outlet, and extracting through the first inlet/outlet, for example via said central tube, components that have separated onto the inner cylindrical wall of the processing chamber.

In this method, filling of the processing chamber with a biological fluid and/or extracting components is preferably produced by or is assisted by pumping, preferably by an external peristaltic pump, and/or by displacing in the processing chamber an axially movable member such as a piston to alter the volume of a separation space of variable volume defined by the axially movable member.

Components are preferably extracted via said second outlet by pumping, preferably with a peristaltic pump. In a preferred, embodiment during the continuous processing of step (c) biological fluid is continuously supplied to the processing chamber by pumping with a peristaltic pump, and separated components are continuously extracted by pumping with a peristaltic pump.

An advantage of the described method is that continuous processing in step (c) can be continued to process/separate a volume of biological fluid that exceeds the maximum separation volume of the processing chamber.

Just prior to and/or during step (e), the centrifugal processing chamber can be accelerated/decelerated and/or rotated in opposite directions to loosen and/or mix separated components that have adhered on the inner cylindrical wall of the processing chamber.

The centrifuging during step (b) is typically continued during steps (c) and (d), continuously or discontinuously.

Brief Description of the Drawings

The invention will now be described by way of example with reference to the accompanying drawings, wherein:

- FIG. 1A is a cross section through the upper part of a processing chamber with its inlet/outlet head, according to the prior art:

- FIG. 1B is a cross section through the upper part of a processing chamber according to the present invention with its inlet/outlet head;
- FIG. 1C is a view of the inventive processing chamber, like FIG. 1B, schematically showing the flow of incoming biological fluid and of outgoing processed components;
- 5 - FIG. 1D is a view of the inlet/outlet head of the inventive processing chamber, like FIG. 1B, schematically showing the stationary and rotating parts;
- FIG. 2 shows three views of a flow guide or diverter/inverter in plan view, side view and in perspective;
- 10 - FIG. 3a, FIG. 3b and FIG. 3c show variants of the continuous flow processing chamber;
- FIG. 4A and FIG. 4B show two examples of disposable sets including the processing chamber of the present invention;
- FIG. 5 is a perspective view of a cabinet for receiving a processing chamber of the present invention;
- 15 - FIG. 6 is a perspective view of another cabinet for receiving a processing chamber of the present invention;
- FIG. 7A and FIG. 7B show, in two parts, a flow diagram of the main processing sequence;
- 20 - FIG. 8A, FIG. 8B, FIG. 8C, FIG. 8D and FIG. 8E schematically show a processing chamber according to the present invention during different phases of operation; and
- FIG. 9 shows a variant of the processing chamber without a piston.

Detailed Description

FIG. 1 A shows the top part of a prior art device of the type known from EP-B-0 912 250. Such known devices comprise a hollow centrifugal processing chamber 10 rotatable about an axis of rotation. The processing chamber 10 has an inner cylindrical wall 16 enclosing a space 19 for receiving biological fluids to be processed and an inclined upper wall 12 leading to a neck 14 having an inlet/outlet 29 for biological fluid to be processed and for processed components of the fluid. The space 19 in the hollow processing chamber 10 for receiving biological fluids occupies the entire uninterrupted volume across the hollow processing chamber 10 between facing parts of the inner cylindrical wall 16 along the length of the

cylindrical wall. Said space has a given volume delimited by a bottom (10B, Fig. 9) of the hollow cylindrical chamber 10, or a variable volume delimited by the position of an axially movable member such as piston 18 within the cylindrical wall. A dynamic/static inlet/outlet head 20 or “rotary seal” is mounted on the neck 14. The inlet/outlet head 20 has an inner first part 22 rotatable with the processing chamber’s neck 14 and an outer/upper second part 24 that remains stationary. The first and second parts 22,24 of the inlet/outlet head 20 have two double seals 26 allowing rotation of the first part 22 with the neck 14 relative to the stationary second part 24. The inlet/outlet head has a central through-passage ending with the inlet/outlet 29 for the inlet of biological fluid to be processed and the outlet of processed/separated components of the fluid. A central stationary tube 28 is held by the part 24.

The centrifugal processing chamber 10 contains within its inner cylindrical wall 16 an axially movable member namely a piston 18 that defines a separation space 19 of variable volume wherein the biological fluid to be processed and separated is received. By axially moving this member/piston 18, biological fluids can be intaken into the separation space 19 or output, via the central through-passage 29.

It can be seen that with the known device the biological fluid to be processed is intaken via the central through-passage (axial inlet/outlet 29) and the processed components are extracted also via the central through-passage (the same axial outlet/inlet 29). As mentioned previously, the quantity of biological fluid that can be processed/separated at one time is limited by the maximum volume of the separation space 19 of the known device’s processing chamber 10. Consequently, a given processing chamber of the prior art cannot be used to process large volumes of biological liquid.

This limitation is overcome by the present invention. FIG. 1B shows a processing chamber of the present invention, using the same reference numerals as in FIG. 1A to designate like parts.

As shown in the example of FIG. 1B, a device according to the invention further comprises a flow guide (“diverter/inverter”) 30 inserted in and rotatable with the processing chamber 10. The flow diverter/inverter 30 is located adjacent to, spaced slightly apart from and in shape-matching relationship to the inclined upper wall 12 of the processing chamber 10, leaving an

inclined frusto-conical annular space 32 of several mm. A central tube in two parts, an inner part 28a and an outer part 28b, extends through the central through-passage of the inlet/outlet head 20. The central tube 28a, 28b is connected, on the one hand, to deliver incoming biological fluid through the inside of the central tube 28b, 28a and through the space 32
5 between the flow diverter/inverter 30 and said inclined wall 12, to the inner cylindrical wall 16 of processing chamber 10 and, on the other hand, to extract processed fluid components separated at the inclined wall 16 of processing chamber 10 and that pass through the space 32 between the flow diverter/inverter 30 and said inclined upper wall 12.

In this improved device, the inside part 19 of the processing chamber located beyond (i.e.
10 below) the flow diverter/inverter 30 is in fluid communication via an aperture 34 in the central part of the flow diverter/inverter 30 to the central through-passage of the inlet/outlet head 20 outside said central tube 28a.

Also, the outer stationary second part 24 of the inlet/outlet head 20 has a lateral outlet 40 for the extraction of processed biological fluid components from the inside 19 of the processing
15 chamber 10, said lateral outlet 40 being in fluid communication with said central through-opening 29 of the inlet/outlet head 20 outside said central tube 28a.

In this embodiment, the processing chamber's upwardly-projecting elongated central neck 14 extends into the inside of the rotatable first part 22 of the inlet/outlet head 20 up to adjacent the level of the lateral outlet 40. Also, in this case, the sealing means of the inlet/outlet head
20 20 comprise two first seals 26 located axially on one side of the lateral outlet 40 and two second seals 26 located axially on the other side of the lateral outlet 40.

The central tube 28a, 28b comprises a stationary axially-outer part 28b extending in the stationary second part 24 of the inlet/outlet head 20, and a rotatable axially-inner part 28a connected to a central part of the flow diverter/inverter 30. The inner part of the central tube
25 28a communicates with the space 32 between the flow diverter/inverter 30 and the inclined upper wall 12 via openings 39.

The rotatable first part 22 of the inlet/outlet head 20 is located laterally inside the stationary second part 24 of the inlet/outlet head, and inside the stationary outer end.

As usual, the inclined upper wall 12 of the processing chamber 10 is frusto-conical as is the upper surface of the flow diverter/inverter 30. As shown, the flow diverter/inverter 30 has an upwardly-protruding central sleeve 36 that fits in the neck 14 of the processing chamber 10 leaving the space 32 of several millimeters between the facing frusto-conical surfaces. The lower end of the central tube 28a fits in this central sleeve 36 and communicates with said space 32 via openings 39 in sleeve 36. In this example, the flow diverter/inverter 30 has an aperture in the form of at least one axially-directed through passage 34, preferably three equally-distributed axial through passages 34 (Fig. 2), in its central sleeve 36 that communicate(s) the inside 19 of the processing chamber 10 with the central passage of the inlet/outlet head 20 outside its central tube 28a.

The flow diverter/inverter 30 typically comprises a peripheral rim 38 of external cylindrical shape that extends from the periphery of a frusto-conical upper surface of the flow diverter/inverter 30. This peripheral cylindrical rim 38 fits in the inner cylindrical wall 16 of the processing chamber 10, below the junction of the inner cylindrical wall 16 and the inclined upper wall 12, leaving a space.

Figure 1C shows the flow of biological fluid entering and exiting the processing chamber 10. First the biological fluid enters the chamber through the inlet port 29 situated in the middle-top part. The fluid falls down by passing first through the static part 24 of the inlet/outlet head 20, then continuing through the dynamic part 22 potentially in rotation, and then reaching the flow diverter/inverter 30. The flow diverter/inverter 30 forces the fluid to continue its way from the central axial to the external side of the conical chamber 10 in rotation, all as shown by the dark line "IN". This ensures a smooth transition for cells that switch from a static to a spinning mode and potentially subject to a high centrifugation force.

The biological fluid inside the processing chamber 10 is then centrifuged, and biological cells are separated from supernatant or other medium, by a density physical principle after being exposed to a horizontal sedimentation force. Cells remain compacted at the external side of the chamber 10 at the inner cylindrical wall 16, while supernatant or medium can be removed via the central axis. As indicated by the gray arrow "OUT", the fluid to be discarded is then pumped from the bottom-central axis of the chamber (centre of space 19), passes through the apertures 34 in the centre of the diverter/inverter 30, and goes up outside

the tube 28a in rotation. Then, after being aspirated up to the height of the lateral outlet 40, the separated biological fluid to be discarded is extracted through the lateral outlet 40 in the static part 24 of the inlet/outlet head. The discarded biological fluid thus switches from a high spinning mode to a static mode. The working of this mechanical mechanism might potentially be damaging for cells. By damaging, is meant creating a pressure, friction, sheer force or even cutting cells by the spinning mechanical part that will either potentially differentiate stem cells making them losing their totipotent properties, or even kill them by apoptosis, necrosis or other destructive mechanisms. For this reason, it is ensured that only discarded product with no cells or with only a low cell concentration are going through the described pathway.

FIG. 1D illustrates the static and dynamic part of the inlet/outlet head 20 of the processing chamber, with airtightness or water tightness provided via seals 26. The central light part 22 indicates the dynamic mechanical pieces that will spin during a normal procedure. The external dark part 24 indicates the static mechanical pieces that will not spin during a procedure. Tightness is ensured through four seals 26 represented in black.

FIG. 2 illustrates the design of the flow diverter/inverter 30. The upper conical surface has the same shape and dimensions as the internal surface of the inclined upper wall 12 of the processing chamber. When assembled, a small canal or space 32 of few millimetres remains for allowing the fluid to pass through. In the centre, the central aperture 35 in the central sleeve 36 is connected by laterally-oriented apertures 39 leading in to the space 32 between the frusto-conical surfaces. This diverts the fluid and forces the biological fluid to flow down on the conical upper part. The three apertures 39 at 120 degrees each and at a distance of few millimetres ensure that central fluid in the chamber is pumped up through the external side of the dynamic part 22, and thus reaching the lateral outlet port 40.

FIG. 3 illustrates design variants of the continuous flow processing chamber.

FIG. 3a is a variant of the flow diverter/inverter 30 that could be used in case the flow does not need to be inverted. In this design, the inlet port 29 will downwardly deliver the biological fluid entering via the middle of the centrifugal chamber, while the outlet port 40 is connected with the path situated in the side of the internal chamber, i.e. via the inclined

annular space 32. This drawing removes the inverted flow path. The inlet port 29 is connected with the central axis of the chamber, and the outlet port 40 is connected with side channels or space 32 of the chamber. Continuous flow processing is the same, only the inlet/outlet port connections are inverted.

- 5 FIG. 3b simplifies the design by using only two seals 26 for ensuring tightness instead of four seals. Each seal 26 has two radially-directed lips that will ensure tightness when pressure or vacuum is generated by the intermediary of a peristaltic pump or up and down movement of the piston.

FIG. 3c is design variant of the two seals 26 each having two axially-directed lips aligned as
10 shown for ensuring tightness as described in relation with FIG. 3b.

Disposable Kits

FIG. 4A shows a typical disposable kit that will be used with the inventive processing chamber 10/20/40. On the upper-left, a bag 50 containing a volume to be processed is connected to the disposable kit. On the bottom-left, a collection bag 52 is attached to the kit
15 for collecting concentrated cell solutions. On the top-right side, a bag 54 is optionally connected to the disposable kit and may contain washing solution for rinsing the chamber at the very end of the procedure, or eventually for suspending concentrated cell solution in a nutritive solution. On the bottom-right, a waste bag 56 is used for discarding solution that has to be eliminated from cells.

20 A buffer 51 containing a filter is inserted in the input line, having the aim to filter aggregates or other unwanted material from the volume to process, and also preventing air bubbles that could flow inside the processing chamber. In addition, a stopcock ramp 58 is used for switching between the multiple bags, or eventually for priming air if necessary through an air filter for ensuring sterility. The disposable set can also include pinch valves located on the
25 tubing for controlling the opening/closing of the tubes.

On the waste line, an air filter 55 for connecting to a pressure sensor is used for monitoring constantly the maximum pressure or vacuum applied inside the disposable set. Also a peristaltic pump 59 is connected on the waste line, and will generate the flow by minimally potentially damaging cells that will not circulate through the pump mechanism.

An air buffer 57 is preferably connected between the lateral outlet 40 of the processing chamber and the peristaltic pump 59. This air buffer 57 has a role of limiting pressure jerks when the peristaltic pump 59 is operated. The air buffer can be a drip chamber.

As shown, the waste line leading to wastebag 56 is connected to the lateral outlet 40 of the processing chamber's inlet/outlet head 20.

FIG. 4B illustrates a similar disposable kit that has multiple input-output and is able to process and switch between more biological solutions. Bag selection is done through eight stopcocks 58 ensuring up to eight input or output bags.

FIG.5 shows a modified version of the cabinet of a known system that integrates a peristaltic pump. As shown the system comprises a cabinet 60 including profiled front part 62 for receiving a centrifugal processing chamber via swing-opening plates 64. The cabinet top also incorporates a spectrometric sensor 66 including bubble detection working with ultrasound technology for measuring biological elements and air bubbles flowing to the processing chamber. A control 67 for a peristaltic pump is located on the cabinet side and a stopcock motor control 68 on its top. On its sides the cabinet also has hooks 69 for attaching tubing of an installed disposable set.

This system works in accordance with the disposable kit illustrated in FIG. 4A. In addition, for a proper calculation of air bubbles entering the processing chamber, an incorporated optical line sensor can integrate an additional ultrasound emitter and receptor that is able to calculate real-time volume of air passing through the tube.

FIG. 6 shows a modified version of the same system that integrates a peristaltic pump and four stopcock motors with individual controls 68. This system works in accordance with the disposable kit illustrated in FIG. 4B.

The continuous processing flow function achievable with the device according to the invention is mainly ensured via one or more external peristaltic pumps generating the flow, on the one hand through the inlet port 29 pumping the biological fluid to the external rotating side 16 of the processing chamber where the sedimentation G force is at maximum and, on the other hand, through the lateral outlet port 40 situated in the centre of the processing

chamber's inlet/outlet head 20 for removing plasma, supernatant or other unwanted biological fluid of lower densities where there is no or little sedimentation force applied. By combining this functionality with a variable processing volume for example as described in EP-B-0 912 250, this invention solves existing limitations of small volume collection of concentrated cells starting from large volume biological fluids.

The main processing sequence

A main application of the invention relates to the fast volume reduction of large biological fluid into a small collection of concentrated cells, as illustrated in FIG. 8A to FIG. 8E. The flow-diagrams of FIG. 7A and FIG. 7B set out the processing steps corresponding to each of FIG. 8A to FIG. 8E.

The first step consists of inserting the centrifugal processing chamber 10 inside the equipment with inlet and outlet ports 29, 40 closed, as shown in FIG. 8A. Then, filling of the centrifugal chamber 10 by pumping into it the volume to be processed can start, FIG. 8B. After connecting the inlet port 29 with the biological fluid to be processed, the mechanism of aspirating down the piston 18 by the intermediary of a pneumatic system that creates a vacuum is done as described in EP-B-0 912 250. During filling, there is potentially a need to remove remaining air that may have stayed trapped in the processing chamber 10. For performing this action, the piston 18 can be pushed slightly up by generating compressed air pressure below the moveable piston. Remaining air and a small portion of filled volume will be re-extracted in the volume to be processed. Then filling can start again until the piston 18 reaches the bottom side of the processing chamber 10.

As indicated in FIG. 8B, during filling the axial inlet 29 is open and the lateral outlet 40 is closed.

At this point, the separation space of processing chamber 10 is filled with a biological solution and sedimentation can start. Sedimentation process is done through centrifugation of the disposable set, as described for example in EP-B-1 144 026.

After some time of centrifugation at a certain rotating speed, blood or biological cells are separated from supernatants or medium solution and the continuous flow for large volume separation can start, as shown in FIG. 8C. In this state, the lateral outlet port 40 is connected

with a waste bag and an external peristaltic pump connected on the waste line is activated. There is the possibility that trapped air still remains in the channel of the outlet port 40, but it will be expelled in the waste bag as soon as peristaltic pumps will be activated. Now, the biological fluid is continuously pumped and separated from the initial volume connected to the inlet port 29 of the chamber, by being centrifuged in the separation chamber in rotation, and finally with supernatant being removed from the central axis of the processing chamber and redirected to the waste bag through the outlet port. During the entire separation process, biological cells will remain in the processing chamber 10, and the speed flow is calculated in order to ensure a sufficient time for the cells to be exposed to a separation force that will guarantee a good separation.

As indicated in FIG. 8C, during continuous processing, the axial inlet 29 is open for the continuous intake of biological fluid, and the lateral outlet port 40 is open for the continuous removal of separated waste.

Once the desired volume has been entirely processed, the peristaltic pump(s) stop. At the beginning of the procedure, the user has set a requested final volume in the software user interface of the system, so at this stage the inlet port 29 is closed and the piston 18 starts moving up by applying the same air-pressure mechanism as described previously, as shown in FIG 8D (axial inlet 29 closed; lateral outlet port 40 open). During this stage, rotating speed is maintained to ensure proper cell separation, and piston 18 moves up at a validated rotation speed and discards the remaining volume into the waste bag until the final volume desired corresponds to the remaining volume in the processing chamber 10.

The last stage is the recovery procedure of the concentrated cells, shown in FIG. 8E. The remaining cell solution in the disposable chamber 10 is firstly mixed through several motor acceleration/decelerations on both directions in order to detach/mix cells potentially stuck on the inner plastic walls 16 of the disposable chamber 10. Then, the inlet port 29 is opened while the outlet port 40 is closed, and piston 18 starts extracting concentrated cells via the axial inlet/outlet port 29. By doing so, cells never pass through the peristaltic pump mechanism that could cause some mechanical stress on the biological cells or stem cells. In addition, for ensuring a proper extraction of cells out of the processing chamber, a dilution phase can also be performed by pumping a small amount of washing solution or suspension

buffer in the centrifugal chamber 10, by mixing the solution again and then extracting the solution in the final desired solution. By doing so, cell losses are minimized.

Variant of the processing chamber without a piston

FIG 9 shows a variant of the processing chamber 10 whose cylindrical wall extends to a
5 bottom wall 10B providing an inner space of fixed volume for receiving biological fluids to be processed. The upper part (inlet/outlet head 20) has the same design as before 10 but the processing chamber 10 does not contain a moveable piston for providing a variable volume.

This chamber 10 of given volume is able to process large volumes through a continuous flow where the final volume to be collected is equal to the processing chamber volume. The
10 chamber 10 can be used with same cabinet and disposable kits as described previously. This variant of the invention greatly simplifies the technology as no pneumatic system is needed anymore for moving a piston up and down, and no infrared detection system is needed for detecting real-time volume remaining in the processing chamber. With this variant, peristaltic pumps are used to intake biological fluid and output separated components.

CLAIMS

1. A device for the processing and separation of biological fluids into components, comprising:

- 5 - a hollow centrifugal processing chamber (10) rotatable about an axis of rotation, the processing chamber having an inner cylindrical wall (16) enclosing a space (19) for receiving biological fluids to be processed and having an inclined upper wall (12) leading to a neck (14) having an inlet/outlet (29) for biological fluid to be processed and for processed components of the fluid, wherein the space (19) in the hollow processing chamber for
10 receiving biological fluids occupies the entire volume across the hollow processing chamber (10) between facing parts of the inner cylindrical wall (16) and along the length of the cylindrical wall, said space (19) having a given volume delimited by a bottom (10B) of the hollow cylindrical chamber, or a variable volume delimited by the position of an axially movable member (18) within the cylindrical wall (16); and
- 15 - a dynamic/static inlet/outlet head (20) mounted on the neck (14) of the processing chamber (10), the inlet/outlet head having a first part (22) rotatable with the processing chamber and a second part (24) that remains stationary, the first and second parts of the inlet/outlet head being connected by a sealing means (26) allowing rotation of the first part relative to the second part, the inlet/outlet head having a central through-passage for the inlet of biological
20 fluid to be processed and the outlet of processed/separated components of the fluid;

Characterized in that the device further comprises a flow guide (30) inserted in and rotatable with the processing chamber (10), the flow guide being located adjacent to, spaced slightly apart from and in shape-matching relationship to the inclined upper wall (12) of the processing chamber to define with the inclined upper wall of the processing chamber an
25 inclined flow passage (32), the flow guide (30) being disposed above said space (19) for receiving biological fluids to be processed with its inclined flow passage (32) leading into a top part of the inner cylindrical wall (12) of the processing chamber; and wherein the inlet/outlet head (20) comprises:

- a first inlet/outlet (29) and a second inlet/outlet (40), the first and second inlets/outlets being disposed on the stationary second part (24) of the inlet/outlet head (20) at a top end and/or at a lateral side thereof, and
- an axial separator (28a, 28b) in the central through-passage of the inlet/outlet head (20), the axial separator defining separate first and second axially-directed passages in the inlet/outlet head,
- an upper part of one of the first and second axially-directed passages communicating with one of the first and second inlets/outlets (29, 40), and an upper part of the other of the first and second axially-directed passages communicating with the other of the first and second inlets/outlets (29, 40),
- a lower part of one of the first and second axially-directed passages communicating with said space (19) in the processing chamber for receiving biological fluids to be processed, and a lower part of the other of the first and second axially-directed passages communicating with the inclined flow passage (32) between the flow guide (30) and the inclined upper wall (12) of the processing chamber,

arranged so the device is operable in a continuous flow mode in which biological fluid to be processed can be inlet via one of the first and second inlets/outlets (29, 40), while processed biological fluid is simultaneously outlet via the other of the first and second inlets/outlets (29, 40).

- 20 2. The device according to claim 1, wherein the stationary second part (24) of the inlet/outlet head (20) comprises an external generally cylindrical body said first inlet/outlet is an axial inlet/outlet (29) in the inlet/outlet head (20), and said second inlet/outlet is a lateral inlet/outlet(40) located in a lateral side of the generally cylindrical body.
- 25 3. The device according to claim 2 wherein the processing chamber has an upwardly-projecting elongated central neck (14) that extends into the inside of the rotatable first part (22) of the inlet/outlet head (20) up to adjacent the level of the lateral inlet/outlet (40)..

4. The device according to claim 2 or 3, wherein the sealing means (26) of the inlet/outlet head comprise a first seal located axially on one side of the lateral inlet/outlet (40) and a second seal located axially on the other side of the lateral inlet/outlet (40).
- 5 5. The device according to any preceding claim wherein said axial separator is a central tube (28a, 28b) that comprises a stationary axially-outer part (28b) extending in the stationary second part (24) of the inlet/outlet head, and a rotatable axially-inner part (28a) connected to a central part of the flow guide (30), said rotatable axial-inner part of the central tube communicating with the space (32) between the flow guide (30) and said inclined upper wall (12) of the processing chamber.
- 10 6. The device according to any preceding claim wherein the rotatable first part (22) of the inlet/outlet head (20) is located inside the stationary second part (24) of the inlet/outlet head (20).
7. The device according to claim 5 wherein :
- 15 (i) the inclined upper wall (12) of the processing chamber (10) is frusto-conical as is the upper surface of the flow guide (30);
- (ii) the flow guide (30) has a central sleeve (36) that fits in the neck (14) of the processing chamber leaving a space of several millimeters between the facing frusto-conical surfaces;
- 20 (iii) the lower end of the central tube (28a) fits in the central sleeve (36) of the flow guide (30) and communicates with said space (32) between the facing frusto-conical surfaces; and
- (iv) the flow guide (30) comprises an aperture in the form of at least one through passage (34), preferably three equally-distributed through passages, in its central sleeve (36) that communicate(s) the inside of the processing chamber (10) with the
- 25 central passage of the inlet/outlet head outside its central tube (28a, 28b).

8. The device according to any preceding claim wherein the flow guide (30) comprises a peripheral rim (38) of external cylindrical shape that extends from the periphery of a frusto-conical upper surface of the flow guide (30), said peripheral cylindrical rim (38) fitting in the inner cylindrical wall (16) of the processing chamber below the junction of the inner cylindrical wall and the inclined upper wall (12) of the processing chamber.
9. The device according to any preceding claim wherein the processing chamber (10) contains within its cylindrical wall (16) an axially movable member (18) such as a piston that defines a separation space (19) of variable volume wherein the biological fluid to be processed and separated is received.
10. The device according to any preceding claim wherein the processing chamber (10) is part of a disposable set comprising a set of containers (52, 54, 56) for receiving the biological fluid to be processed and separated and the separated components, and optionally one or more additional containers for additive solutions.
11. The device according to any preceding claim further comprising a cabinet (60) for receiving the processing chamber, the cabinet having drive means for driving the centrifugal processing chamber and preferably also means for controlling the axial position of an axially movable member such as a piston in the processing chamber.
12. The device according to claim 11 or claim 12 when it depends on claim 11, further comprising closure members for the inlet/outlet to the processing chamber, said closure members being clips or pinch valves that act on tubing of the disposable set, and/or stopcocks (58) included in the disposable set and/or fitted on a cabinet for receiving the processing chamber.
13. A system for the processing and separation of biological fluids into components, comprising the device of any one of claims 1 to 12 and further comprising at least one peristaltic pump (59) for pumping incoming biological fluid through one of the first inlets/outlets; and/or for pumping extracted processed biological fluid components through the other of the inlets/outlets.

14. A method of processing and separating biological fluids into components using the device according to any one of claims 1 - 12 or the system of claim 13, the method comprising the following steps:
- 5 (a) filling the processing chamber (10) with a biological fluid via one of said inlets/outlets (29, 40) with the second inlet/outlet closed (29, 40);
- (b) rotating the processing chamber (10) to centrifuge biological fluid in the chamber in order to separate different components of the biological fluid onto the inner cylindrical wall (16) of the processing chamber or into the inside (19) of the processing chamber;
- 10 (c) operating in a continuous processing mode wherein filling of the processing chamber (10) with a biological fluid is carried out while at the same time extracting, via the second inlet/outlet components that have separated into the inside of the processing chamber;
- (d) closing the first inlet/outlet (29, 40) whereby the processing chamber is no longer
15 being filled with biological fluid, and continuing to extract components that have separated into the inside of the processing chamber via said second inlet/outlet (29, 40); and
- (e) closing said second inlet/outlet (29, 40) whereby processed components that have
20 separated into the inside of the processing chamber are no longer extracted via said second inlet/outlet (29, 40), and extracting via said first inlet/outlet (29, 40) components that have separated onto the inner cylindrical wall of the processing chamber.
15. The method according to claim 14 wherein filling of the processing chamber with a biological fluid is produced by or is assisted by pumping, preferably by an external
25 peristaltic pump (59), and/or by displacing in the processing chamber an axially movable member such as a piston (18) to alter the volume of a separation space of variable volume defined by the axially movable member.
16. The method according to claim 14 or 15 wherein components are extracted from the processing chamber by pumping, preferably with a peristaltic pump (59).

17. The method according to any one of claims 14 to 16 wherein continuous processing in step (c) is continued to process/separate a volume of biological fluid that exceeds the maximum separation volume of the processing chamber (10).
18. The method according to any one of claims 14 to 17 wherein just prior to and/or during step (e) the centrifugal processing chamber (10) is accelerated/decelerated and/or rotated in opposite directions to loosen and/or mix separated components that have adhered on the inner cylindrical wall of the processing chamber.
19. The method according to any one of claims 14 to 18 wherein centrifuging during step (b) is continued during steps (c) and (d), continuously or discontinuously.

AMENDED CLAIMS

received by the International Bureau on 09 August 2016 (09.08.2016)

1. A device for the processing and separation of biological fluids into components, comprising:
- 5 - a hollow centrifugal processing chamber (10) rotatable about an axis of rotation, the processing chamber having an inner cylindrical wall (16) enclosing a space (19) for receiving biological fluids to be processed and having an inclined upper wall (12) leading to a narrower neck (14) that extends axially from the cylindrical wall (16) and its inclined upper wall (12) and that has an inlet/outlet (29) for biological fluid to be processed and for
10 processed components of the fluid, the cylindrical wall (16), its inclined upper wall (12) and the narrower neck (14) all being in one piece, wherein the space (19) in the hollow processing chamber for receiving biological fluids occupies the entire volume across the hollow processing chamber (10) between facing parts of the inner cylindrical wall (16) and along the length of the cylindrical wall, said space (19) having a given volume delimited by a
15 bottom (10B) of the hollow cylindrical chamber, or a variable volume delimited by the position of an axially movable member (18) within the cylindrical wall (16) ; and
- a dynamic/static inlet/outlet head (20) mounted on and around the neck (14) of the processing chamber (10), the inlet/outlet head having a first part (22) rotatable with the processing chamber and a second part (24) that remains stationary, the first and second parts
20 of the inlet/outlet head being connected by a sealing means (26) allowing rotation of the first part relative to the second part, the inlet/outlet head having a central through-passage for the inlet of biological fluid to be processed and the outlet of processed/separated components of the fluid;
- Characterized in that** the device further comprises a flow guide (30) inserted in and
25 rotatable with the processing chamber (10), the flow guide being located adjacent to, spaced slightly apart from and in shape-matching relationship to the inclined upper wall (12) of the processing chamber to define with the inclined upper wall of the processing chamber an inclined flow passage (32), the flow guide (30) being disposed above said space (19) for receiving biological fluids to be processed with its inclined flow passage (32) leading into a

top part of the inner cylindrical wall (12) of the processing chamber; and wherein the inlet/outlet head (20) comprises:

- a first inlet/outlet (29) and a second inlet/outlet (40), the first and second inlets/outlets being disposed on the stationary second part (24) of the inlet/outlet head (20) at a top end and/or at a lateral side thereof, and
- an axial separator (28a, 28b) in the central through-passage of the inlet/outlet head (20), the axial separator defining separate first and second axially-directed passages in the inlet/outlet head,
- an upper part of one of the first and second axially-directed passages communicating with one of the first and second inlets/outlets (29, 40), and an upper part of the other of the first and second axially-directed passages communicating with the other of the first and second inlets/outlets (29, 40),
- a lower part of one of the first and second axially-directed passages communicating with said space (19) in the processing chamber for receiving biological fluids to be processed, and a lower part of the other of the first and second axially-directed passages communicating with the inclined flow passage (32) between the flow guide (30) and the inclined upper wall (12) of the processing chamber,

arranged so the device is operable in a continuous flow mode in which biological fluid to be processed can be inlet via one of the first and second inlets/outlets (29, 40), while processed biological fluid is simultaneously outlet via the other of the first and second inlets/outlets (29, 40).

2. The device according to claim 1, wherein the stationary second part (24) of the inlet/outlet head (20) comprises an external generally cylindrical body said first inlet/outlet is an axial inlet/outlet (29) in the inlet/outlet head (20), and said second inlet/outlet is a lateral inlet/outlet(40) located in a lateral side of the generally cylindrical body.
3. The device according to claim 2 wherein the processing chamber's upwardly-projecting elongated central neck (14) extends into the inside of the rotatable first part (22) of the inlet/outlet head (20) up to adjacent the level of the lateral inlet/outlet (40).

4. The device according to claim 2 or 3, wherein the sealing means (26) of the inlet/outlet head comprise a first seal located axially on one side of the lateral inlet/outlet (40) and a second seal located axially on the other side of the lateral inlet/outlet (40).
5. The device according to any preceding claim wherein said axial separator is a central tube (28a, 28b) that comprises a stationary axially-outer part (28b) extending in the stationary second part (24) of the inlet/outlet head, and a rotatable axially-inner part (28a) connected to a central part of the flow guide (30), said rotatable axial-inner part of the central tube communicating with the space (32) between the flow guide (30) and said inclined upper wall (12) of the processing chamber.
6. The device according to any preceding claim wherein the rotatable first part (22) of the inlet/outlet head (20) is located inside the stationary second part (24) of the inlet/outlet head (20).
7. The device according to claim 5 wherein :
- (i) the inclined upper wall (12) of the processing chamber (10) is frusto-conical as is the upper surface of the flow guide (30);
- (ii) the flow guide (30) has a central sleeve (36) that fits in the neck (14) of the processing chamber leaving a space of several millimeters between the facing frusto-conical surfaces;
- (iii) the lower end of the central tube (28a) fits in the central sleeve (36) of the flow guide (30) and communicates with said space (32) between the facing frusto-conical surfaces; and
- (iv) the flow guide (30) comprises an aperture in the form of at least one through passage (34), preferably three equally-distributed through passages, in its central sleeve (36) that communicate(s) the inside of the processing chamber (10) with the central passage of the inlet/outlet head outside its central tube (28a, 28b).
8. The device according to any preceding claim wherein the flow guide (30) comprises a peripheral rim (38) of external cylindrical shape that extends from the periphery of a

frusto-conical upper surface of the flow guide (30), said peripheral cylindrical rim (38) fitting in the inner cylindrical wall (16) of the processing chamber below the junction of the inner cylindrical wall and the inclined upper wall (12) of the processing chamber.

- 5 9. The device according to any preceding claim wherein the processing chamber (10) contains within its cylindrical wall (16) an axially movable member (18) such as a piston that defines a separation space (19) of variable volume wherein the biological fluid to be processed and separated is received.
- 10 10. The device according to any preceding claim wherein the processing chamber (10) is part of a disposable set comprising a set of containers (52, 54, 56) for receiving the biological fluid to be processed and separated and the separated components, and optionally one or more additional containers for additive solutions.
- 15 11. The device according to any preceding claim further comprising a cabinet (60) for receiving the processing chamber, the cabinet having drive means for driving the centrifugal processing chamber and preferably also means for controlling the axial position of an axially movable member such as a piston in the processing chamber.
- 20 12. The device according to claim 11, further comprising closure members for the inlet/outlet to the processing chamber, said closure members being clips or pinch valves that act on tubing of the disposable set, and/or stopcocks (58) included in the disposable set and/or fitted on a cabinet for receiving the processing chamber.
- 25 13. A system for the processing and separation of biological fluids into components, comprising the device of any one of claims 1 to 12 and further comprising at least one peristaltic pump (59) for pumping incoming biological fluid through one of the first and second inlets/outlets (29, 40); and/or for pumping extracted processed biological fluid components through the other of the first and second inlets/outlets (29, 40).
14. A method of processing and separating biological fluids into components using the device according to any one of claims 1 - 12 or the system of claim 13, the method comprising the following steps:

- (a) filling the processing chamber (10) with a biological fluid via one of said inlets/outlets (29, 40) with the second inlet/outlet closed (29, 40);
- (b) rotating the processing chamber (10) to centrifuge biological fluid in the chamber in order to separate different components of the biological fluid onto the inner cylindrical wall (16) of the processing chamber or into the inside (19) of the processing chamber;
- (c) operating in a continuous processing mode wherein filling of the processing chamber (10) with a biological fluid is carried out while at the same time extracting, via the second inlet/outlet components that have separated into the inside of the processing chamber;
- (d) closing the first inlet/outlet (29, 40) whereby the processing chamber is no longer being filled with biological fluid, and continuing to extract components that have separated into the inside of the processing chamber via said second inlet/outlet (29, 40); and
- (e) closing said second inlet/outlet (29, 40) whereby processed components that have separated into the inside of the processing chamber are no longer extracted via said second inlet/outlet (29, 40), and extracting via said first inlet/outlet (29, 40) components that have separated onto the inner cylindrical wall of the processing chamber.
15. The method according to claim 14 wherein filling of the processing chamber with a biological fluid is produced by or is assisted by pumping, preferably by an external peristaltic pump (59), and/or by displacing in the processing chamber an axially movable member such as a piston (18) to alter the volume of a separation space of variable volume defined by the axially movable member.
16. The method according to claim 14 or 15 wherein components are extracted from the processing chamber by pumping, preferably with a peristaltic pump (59).
17. The method according to any one of claims 14 to 16 wherein continuous processing in step (c) is continued to process/separate a volume of biological fluid that exceeds the maximum separation volume of the processing chamber (10).

18. The method according to any one of claims 14 to 17 wherein just prior to and/or during step (e) the centrifugal processing chamber (10) is accelerated/decelerated and/or rotated in opposite directions to loosen and/or mix separated components that have adhered on the inner cylindrical wall of the processing chamber.
- 5 19. The method according to any one of claims 14 to 18 wherein centrifuging during step (b) is continued during steps (c) and (d), continuously or discontinuously.

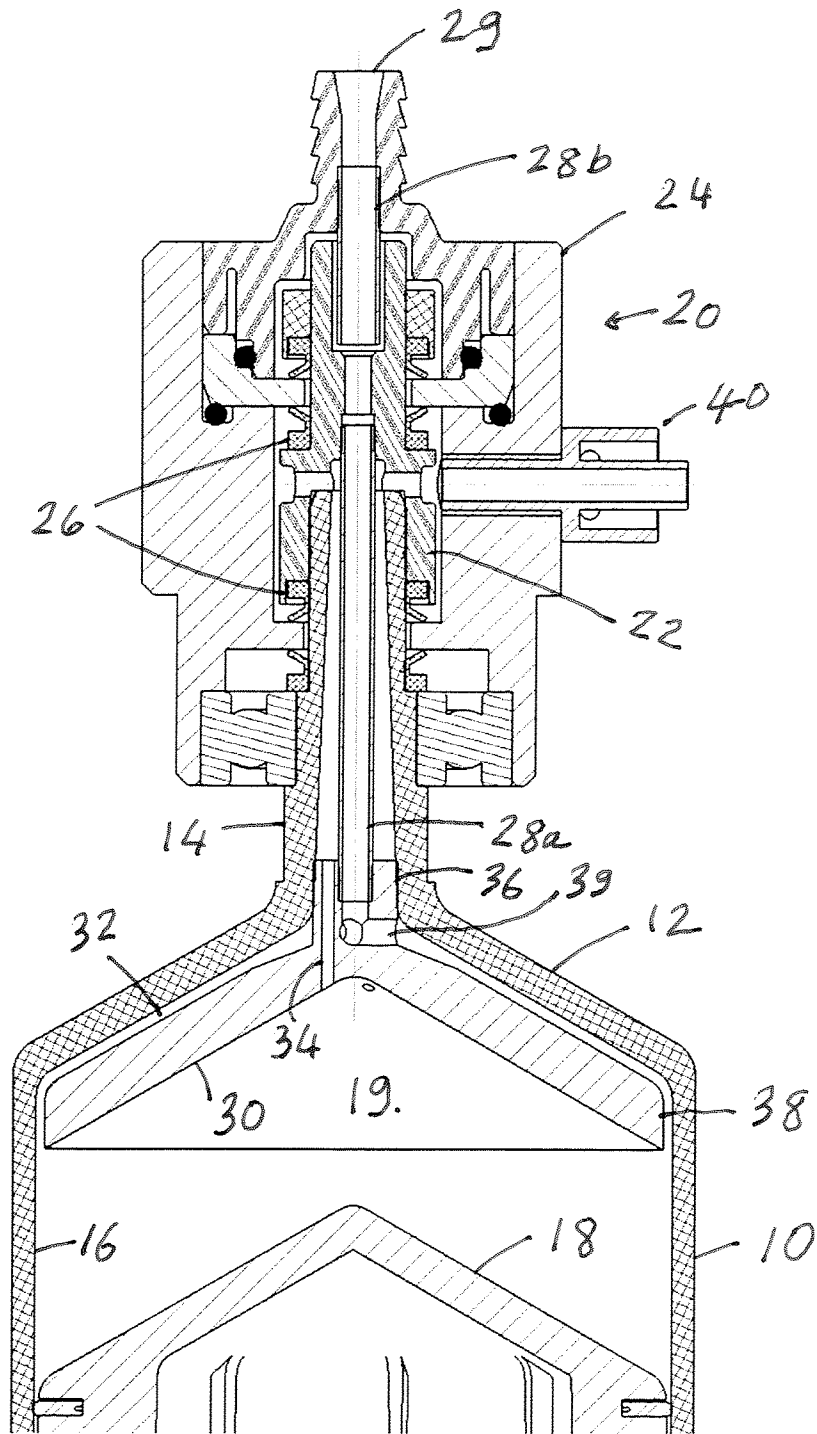


FIG. 1B

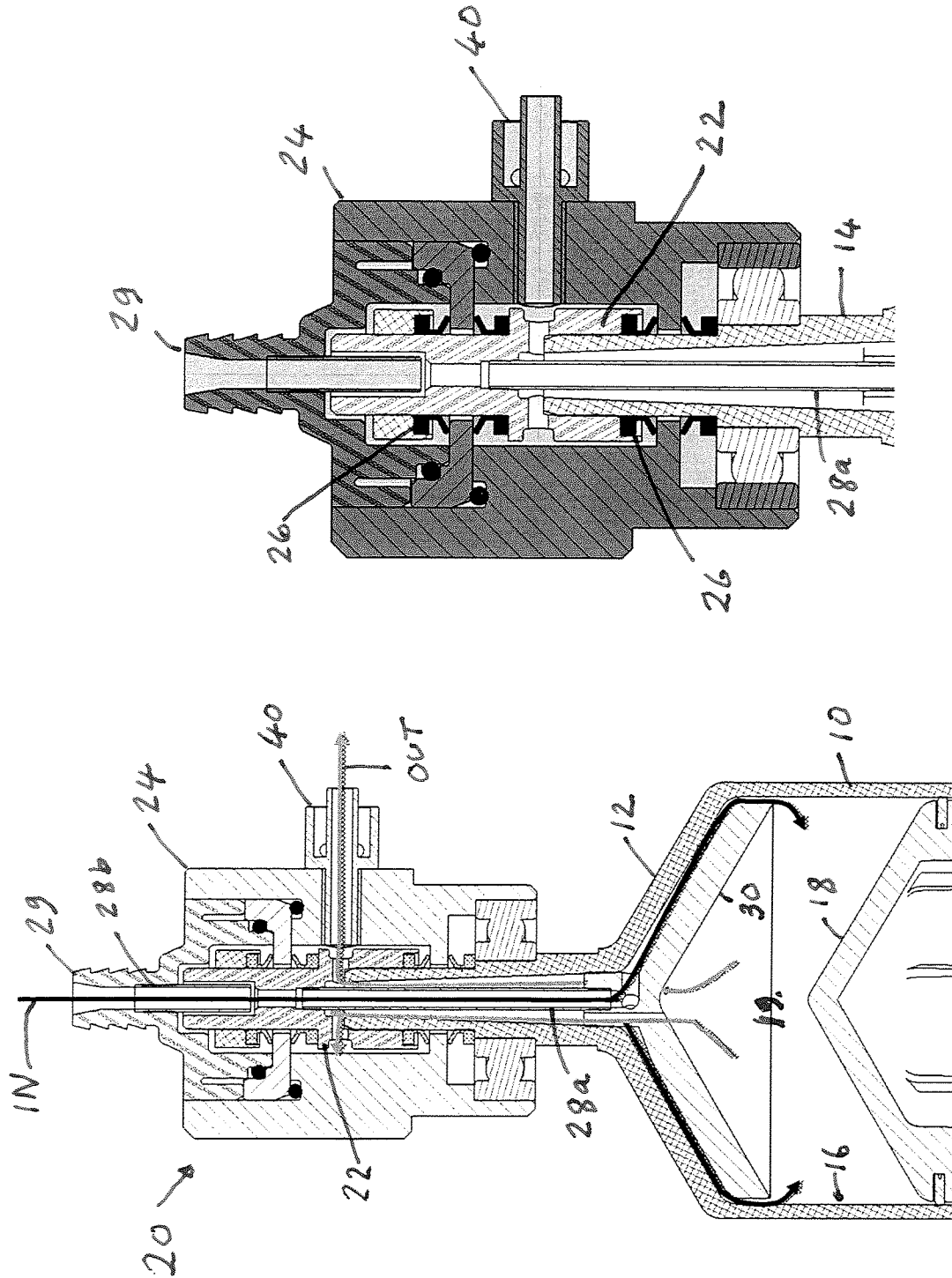


FIG. 1D

FIG. 1C

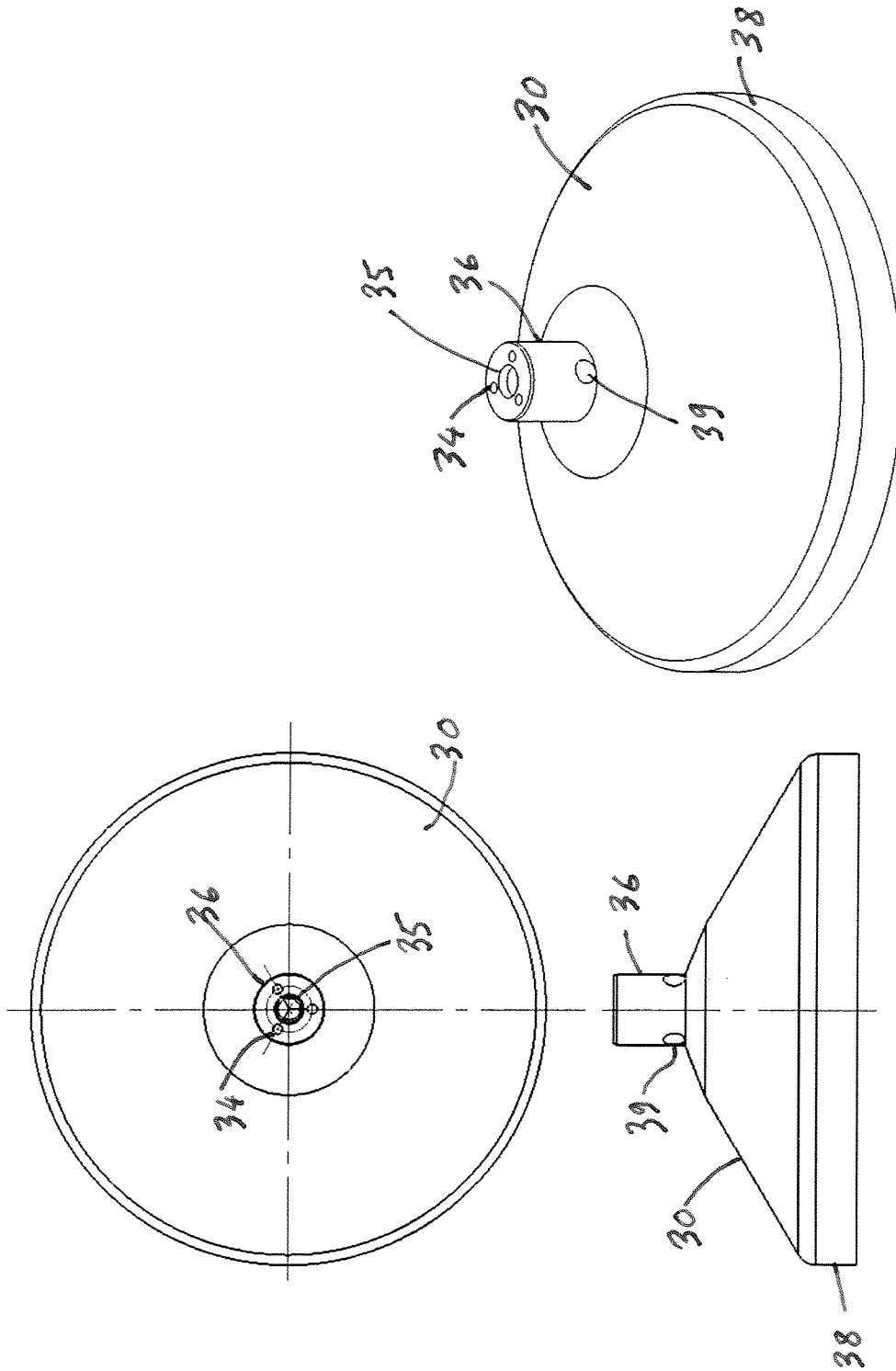


FIG. 2

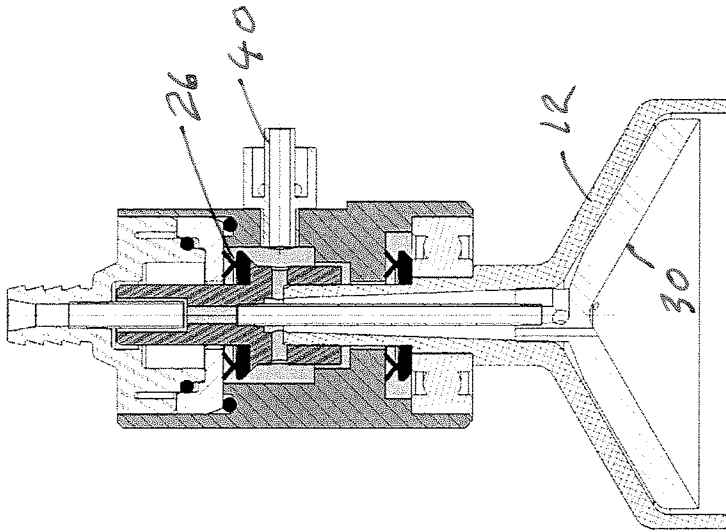


FIG. 3c

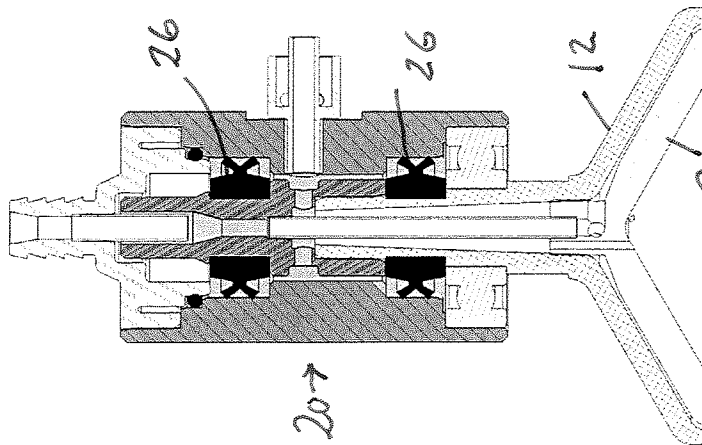


FIG. 3b

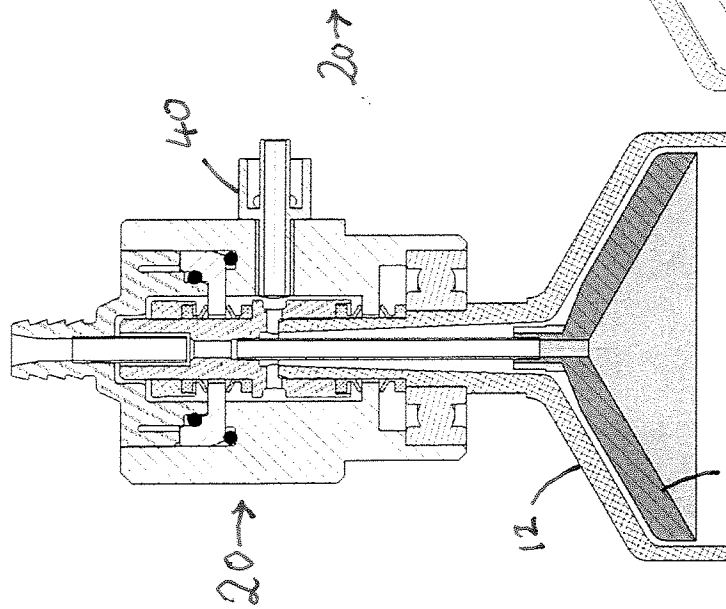


FIG. 3a

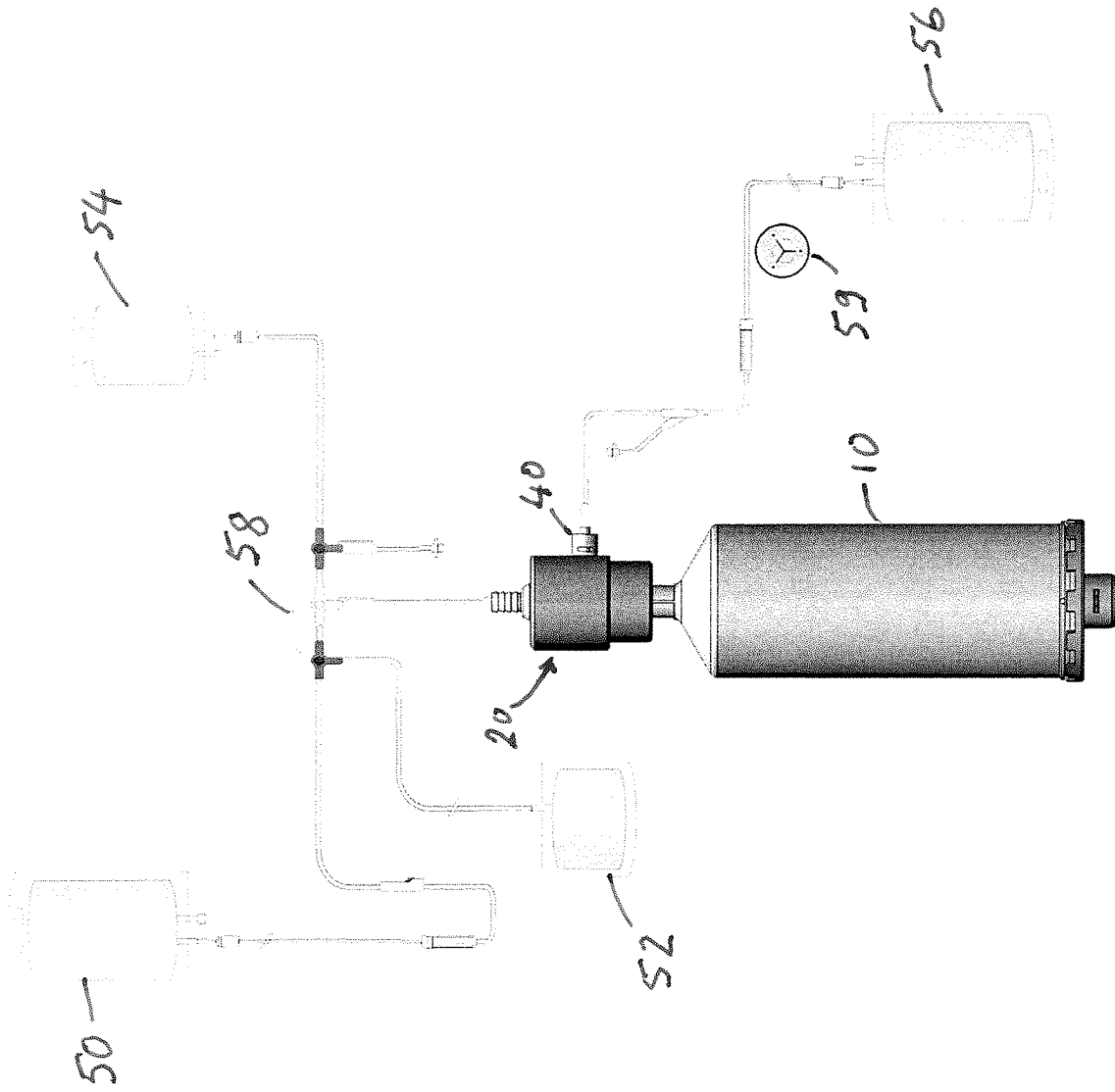


FIG. 4A

Drawing 4B: variant of the disposable kit

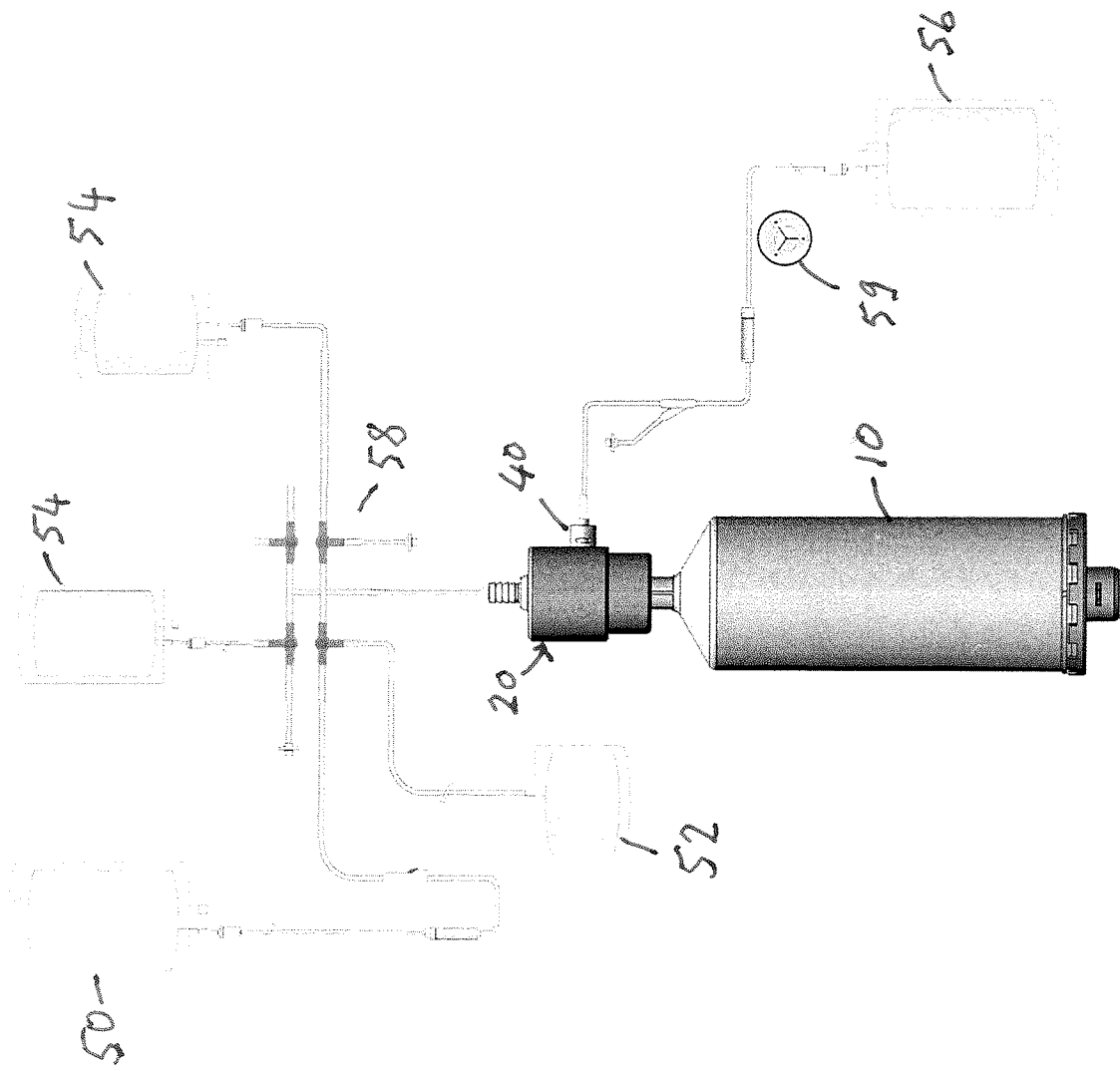


FIG. 4B

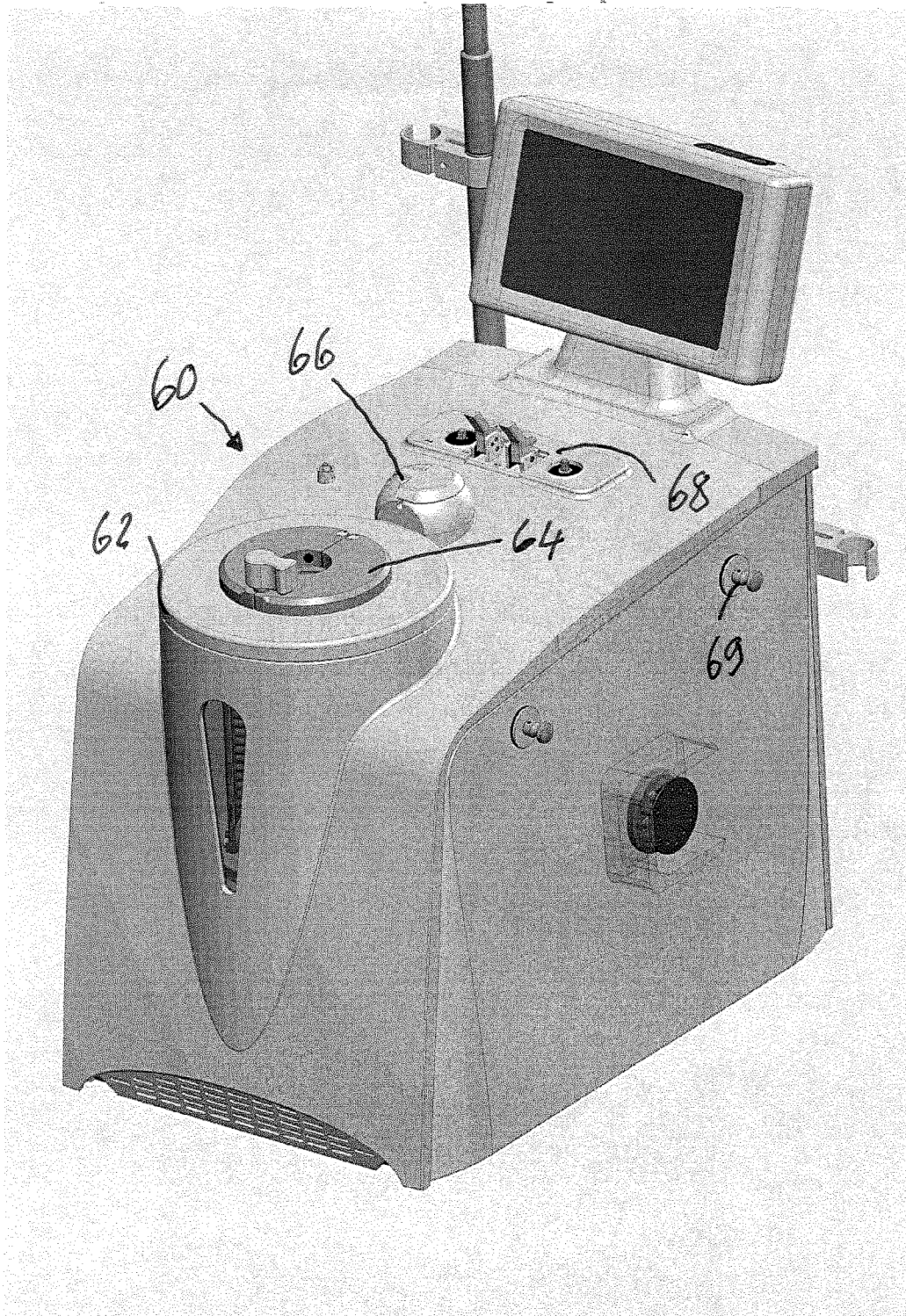


FIG. 5

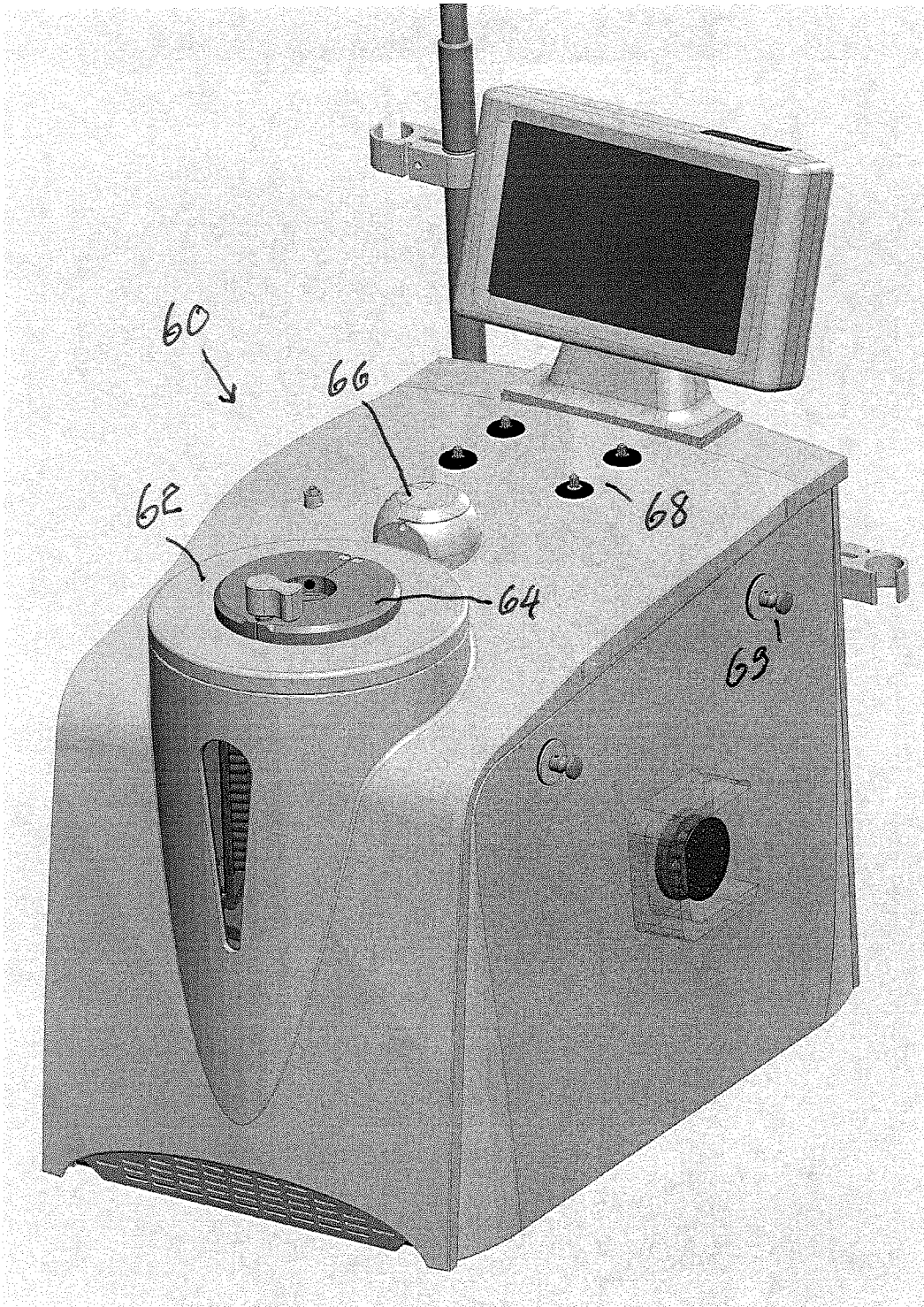


FIG. 6

Drawing 7: flow chart-main application



FIG. 7

Drawing 8: functional view of the new chamber at different processing stages

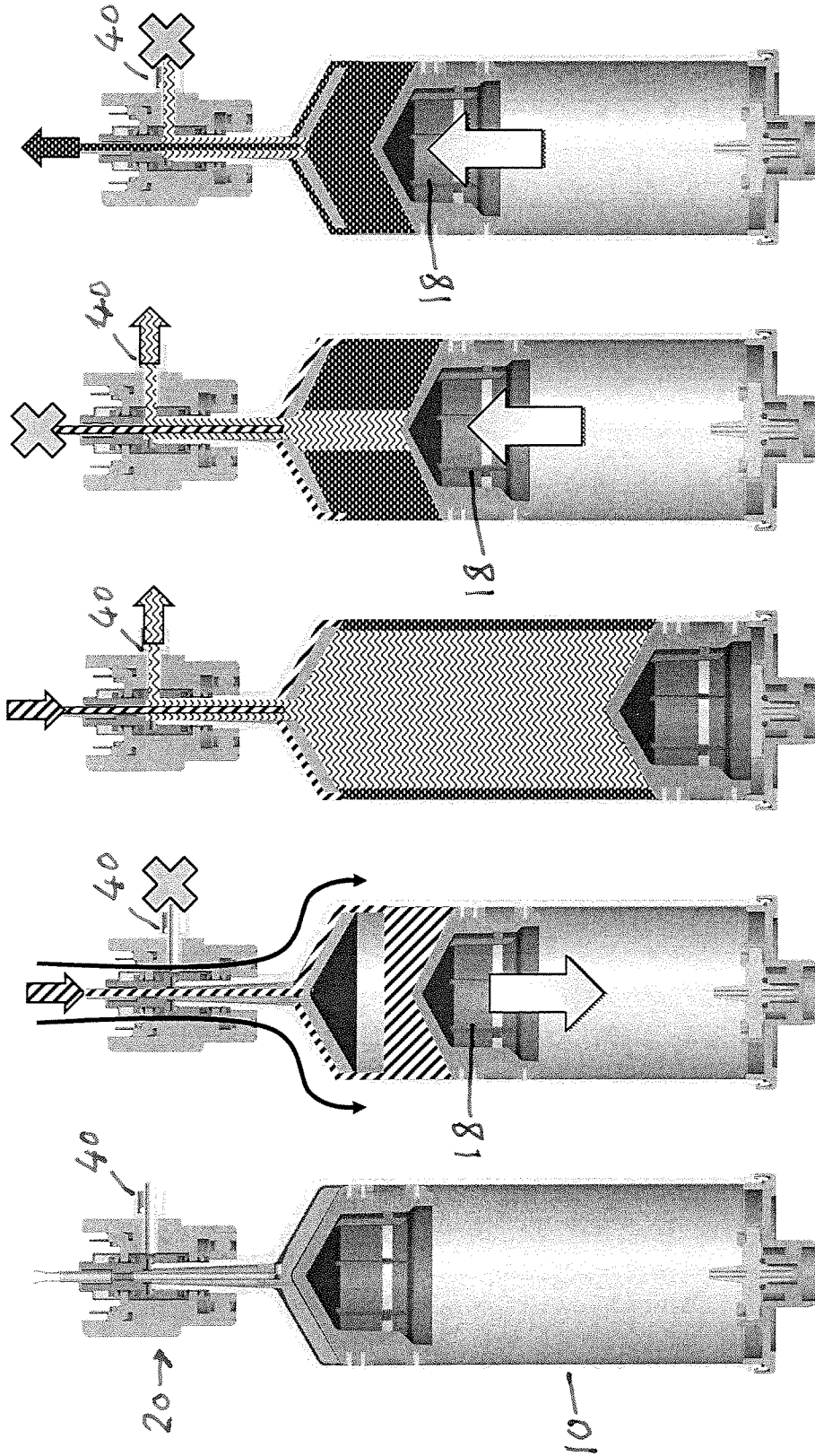


FIG. 8E

FIG. 8D

FIG. 8C

FIG. 8B

FIG. 8A

FIG. 8

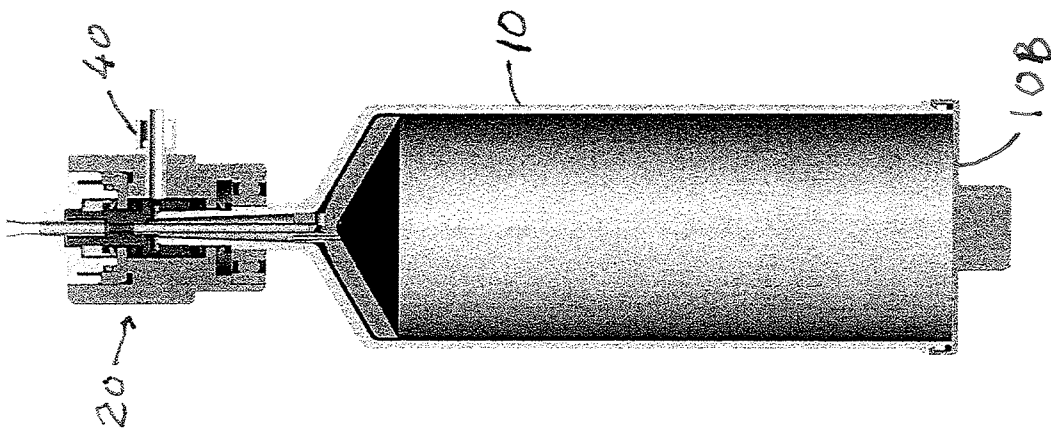


FIG. 9

INTERNATIONAL SEARCH REPORT

International application No
PCT/IB2016/050107

A. CLASSIFICATION OF SUBJECT MATTER
INV. B04B5/04 A61M1/36 A61M1/02
ADD.
According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED
Minimum documentation searched (classification system followed by classification symbols)
B04B A61M
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
EPO-Internal, EMBASE

C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 2007/213191 A1 (CHAMMAS JACQUES [US]) 13 September 2007 (2007-09-13) the whole document paragraphs [0020], [0047] - [0050], [0054] - [0058], [0070] - [0073]; figures 1, 2, 4, 11	1-13
Y	WO 96/11747 A2 (TRANSFUSION TECHNOLOGIES CORP [US]; HEADLEY THOMAS D [US]; POWERS EDWA) 25 April 1996 (1996-04-25) the whole document page 28 - page 30; figures 3, 4, 11, 40 page 61, line 20 - line 22 page 6, line 33 - page 7, line 11 page 41, line 3 - line 7 ----- -/--	1-13

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search 20 April 2016	Date of mailing of the international search report 28/04/2016
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Authorized officer Pinta, Violaine

INTERNATIONAL SEARCH REPORT

International application No
PCT/IB2016/050107

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	EP 1 144 026 B1 (BIOSAFE S A [CH]) 28 July 2004 (2004-07-28) the whole document	1-13
A	----- WO 2014/070209 A1 (HAEMONETICS CORP [US]) 8 May 2014 (2014-05-08) the whole document paragraph 0046 - column 5, paragraph 0064; figures 3-5 paragraph [0075] - paragraph [0089]; figures 6A, 6B	1-13
A	----- GB 2 181 371 A (CARDIVASCULAR SYSTEMS) 23 April 1987 (1987-04-23) the whole document figures 1-2	1-13
A	----- WO 2006/079238 A1 (ROCHAT JEAN-DENIS [CH]) 3 August 2006 (2006-08-03) the whole document figures 2, 4, 5	1-13
A	----- US 3 104 225 A (DI BENEDETTO CHARLES) 17 September 1963 (1963-09-17) the whole document	1-13
A	----- US 3 145 713 A (LATHAM JR ALLEN) 25 August 1964 (1964-08-25) the whole document column 8; figures 8-9	1-13
A	----- JP H07 256150 A (HITACHI KOKI KK) 9 October 1995 (1995-10-09) figure 1	1-13

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Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: 14-19 (partially)
because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy or surgery. Claims 14-19 were partially searched on the basis of the device of claims 1-12 or system of claim 13.
2. Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of additional fees.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

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