



Office de la Propriété

Intellectuelle  
du Canada

Un organisme  
d'Industrie Canada

Canadian  
Intellectual Property  
Office

An agency of  
Industry Canada

CA 2474635 C 2011/03/08

(11)(21) **2 474 635**

(12) **BREVET CANADIEN**  
**CANADIAN PATENT**

(13) **C**

(86) Date de dépôt PCT/PCT Filing Date: 2002/10/31  
(87) Date publication PCT/PCT Publication Date: 2003/08/14  
(45) Date de délivrance/Issue Date: 2011/03/08  
(85) Entrée phase nationale/National Entry: 2004/07/27  
(86) N° demande PCT/PCT Application No.: IB 2002/004537  
(87) N° publication PCT/PCT Publication No.: 2003/066135  
(30) Priorité/Priority: 2002/02/08 (SE0200370-5)

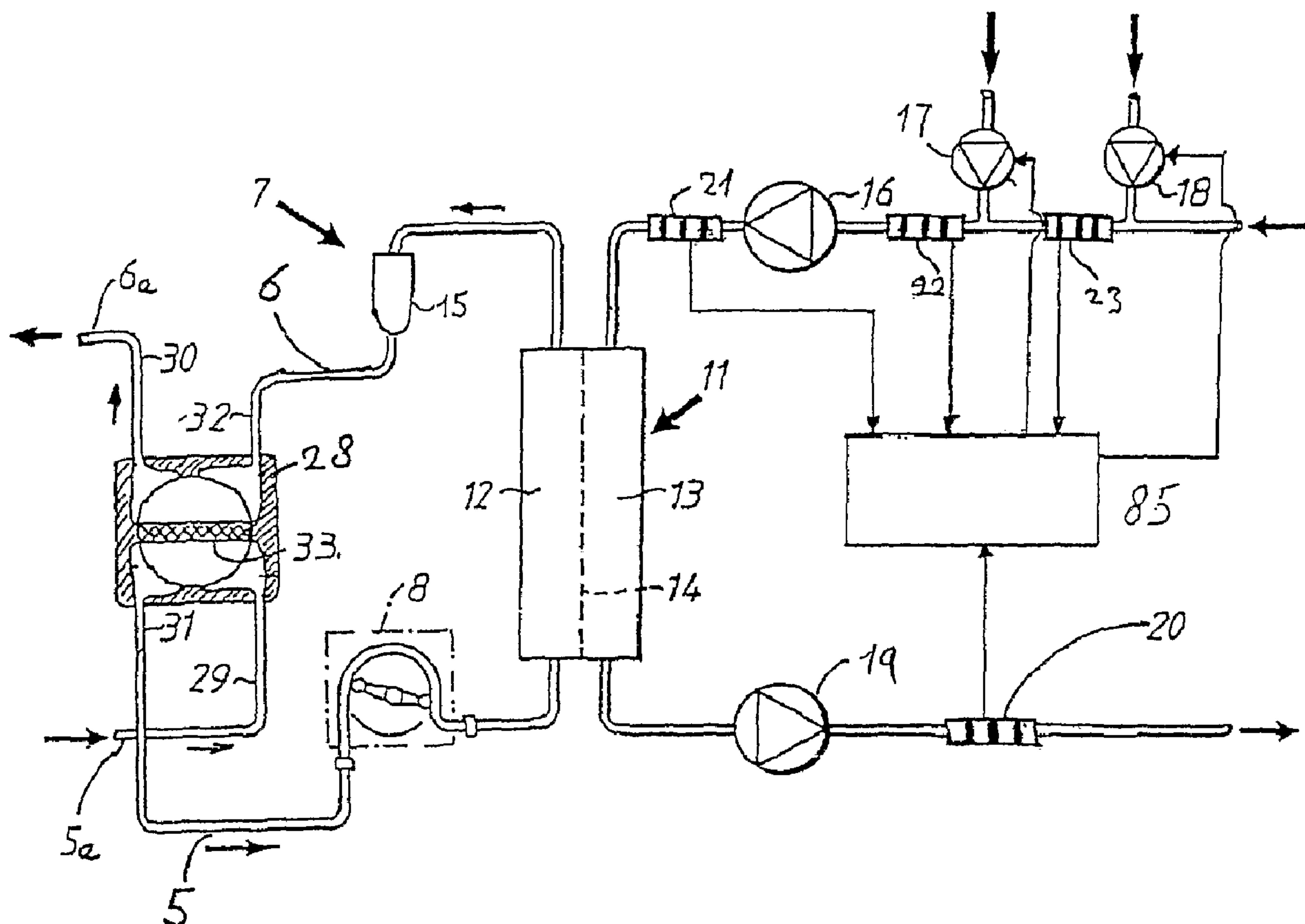
(51) Cl.Int./Int.Cl. *A61M 1/16* (2006.01),  
*A61M 1/34* (2006.01), *A61M 1/36* (2006.01)

(72) Inventeurs/Inventors:  
BENE, BERNARD, FR;  
GOUX, NICOLAS, FR;  
HANSSON, PER, SE;  
HERTZ, THOMAS, SE;  
JANSSON, OLOF, SE;  
PERSSON, ROLAND, SE;  
STERNBY, JAN, SE;  
ASBRINK, PERRY, SE

(73) Propriétaire/Owner:  
GAMBRO LUNDIA AB, SE

(74) Agent: ROBIC

(54) Titre : PROCEDE ET APPAREIL POUR DETERMINER LE DEBIT D'UN FLUIDE VERS UN ACCES  
(54) Title: METHOD AND APPARATUS FOR DETERMINING ACCESS FLOW



(57) Abrégé/Abstract:

A method and apparatus for determining a fluid flow rate in a blood access having an upstream position and a downstream position using a blood treatment apparatus including a blood treatment unit having a semi permeable membrane delimiting a first chamber

## (57) Abrégé(suite)/Abstract(continued):

through which blood removed from said blood access passes and a second chamber through which dialysis liquid passes, an arterial line and a venous line, connected to an inlet and an outlet of the first chamber respectively. The method for determining the fluid flow rate involves measurement of the post dialyzer concentration or conductivity before and after a flow reversal. The invention further involves a method for checking the operating configuration of the arterial and venous lines of the blood treatment apparatus.

**(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)**

**(19) World Intellectual Property Organization  
International Bureau**



(43) International Publication Date  
14 August 2003 (14.08.2003)

PCT

(10) International Publication Number  
**WO 03/066135 A1**

**(51) International Patent Classification<sup>7</sup>:** A61M 1/16, 1/34

(21) International Application Number: PCT/IB02/04537

(22) International Filing Date: 31 October 2002 (31.10.2002)

(25) Filing Language: English

(26) Publication Language: English

**(30) Priority Data:**  
0200370-5 8 February 2002 (08.02.2002) SE

(71) *Applicant (for all designated States except US): GAM-BRO LUNDIA AB [SE/SE]; Magistratsvägen 16, S-226 43 Lund (SE).*

(72) Inventors; and  
(75) Inventors/Applicants (*for US only*): **BENE, Bernard**

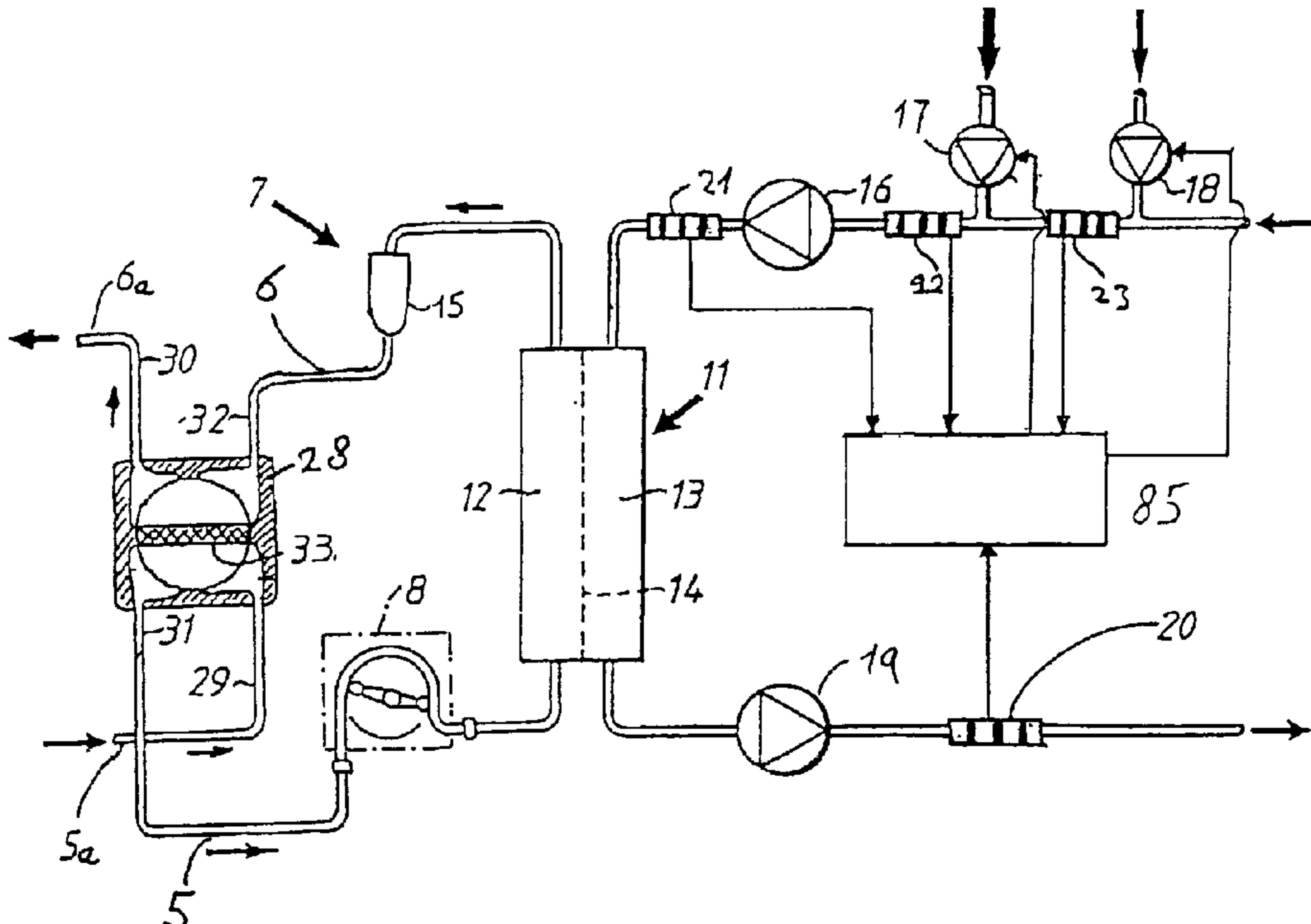
**GOUX, Nicolas** [FR/FR]; 8, impasse des Troignes, F-69290 Craponne (FR). **HANSSON, Per** [SE/SE]; Kvarngatan 4, S-232 34 Arlöv (SE). **HERTZ, Thomas** [SE/SE]; Starvägen 18a, S-227 32 Lund (SE). **JANSSON, Olof** [SE/SE]; Näktergalsgatan 33, S-235 38 Vellinge (SE). **PERSSON, Roland** [SE/SE]; Vasagatan 9b, S-216 11 Limhamn (SE). **STERNBY, Jan** [SE/SE]; Spårsnögatan 45, S-226 52 Lund (SE). **ASBRINK, Perry** [SE/SE]; Lindeborgsgatan 10, S-215 81 Malmö (SE).

(74) Agent: **SUTTO, Luca**; GAMBRO Patent Department, 61, Avenue Tony Garnier, F-69007 Lyon (FR).

**(81) Designated States (national):** AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW.

[Continued on next page]

**(54) Title: METHOD AND APPARATUS FOR DETERMINING ACCESS FLOW**



W003/066135 A1

**(57) Abstract:** A method and apparatus for determining a fluid flow rate in a blood access having an upstream position and a downstream position using a blood treatment apparatus including a blood treatment unit having a semi permeable membrane delimiting a first chamber through which blood removed from said blood access passes and a second chamber through which dialysis liquid passes, an arterial line and a venous line, connected to an inlet and an outlet of the first chamber respectively. The method for determining the fluid flow rate involves measurement of the post dialyzer concentration or conductivity before and after a flow reversal. The invention further involves a method for checking the operating configuration of the arterial and venous lines of the blood treatment apparatus.



**(84) Designated States (regional):** ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

**Published:**

— with international search report

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

TITLE

Method and apparatus for determining access flow.

5 TECHNICAL FIELD

The present invention relates to a method and apparatus for determining fluid flow rate in a patient's blood access. More particularly, the invention relates to the calculation of the fluid flow rate in the blood access based on conductivity measurements of the post dialyzer or 10 other blood treatment unit effluent fluid.

BACKGROUND ART

There are several types of treatments in which blood is taken out in an extracorporeal blood 15 circuit. Such treatments involve, for example, hemodialysis, hemofiltration, hemodiafiltration, plasmapheresis, blood component separation, blood oxygenation, etc. Normally, blood is removed from a blood vessel at a blood access and returned to the same blood vessel.

In hemodialysis and similar treatments, a blood access commonly surgically created in the nature of a arterio-venous shunt, commonly referred to as a fistula. Blood needles are 20 inserted in the fistula. Blood is taken out from the fistula via a needle at an upstream position and blood is returned to the fistula via needle at a downstream position.

The arterio-venous shunt or fistula is blood access having capability of providing a high 25 blood flow and being operative during several years and even tens of years. It is produced by operatively connecting, for example, the radial artery to the cephalic vein at the level of the forearm. The venous limb of the fistula thickens during the course of several months, permitting repeated insertion of dialysis needles.

An alternative blood access to the fistula is the arterio-venous graft, in which a connection is generated from, for example, the radial artery at the wrist to the basilic vein. The connection is made with a tube graft made from e.g. autogenous saphenous vein or from 30 polytetrafluoroethylene (PTFE, Teflon). The needles are inserted in the graft.

A further example of a blood access is a silicon, dual-lumen catheter surgically implanted into one of the large veins.

Further type of blood access find use in specific situations, like a no-needle arterio-venous 35 graft consisting of a T-tube linked to a standard PTFE graft. The T-tube is implanted in the skin. Vascular access is obtained either by unscrewing a plastic plug or by puncturing a septum of said T-tube with a needle. Other methods and devices are also known.

During the above blood treatment therapies, hemodialysis for instance, it is desirable to obtain a constant blood flow rate of 150 - 500 ml/min or even higher, and the access site

must be prepared for delivering such flow rates. The blood flow in an AV fistula is often 800 ml/min or larger, permitting delivery of a blood flow rate in the desired range.

In the absence of a sufficient forward blood flow, the extracorporeal circuit blood pump will take up some of the already treated blood entering the fistula via the venous needle, so

5 called access or fistula recirculation, leading to poor treatment results and progressive reduction of treatment efficiency.

A common cause of poor flow with AV fistulas is partial obstruction of the venous limb due to fibrosis secondary to multiple venipunctures. Moreover, stenosis causes a reduction of access flow.

10 It has been found that access flow rate often exhibit a long plateau time period with sufficient access flow, followed by a short period of a few weeks with markedly reduced access flow leading to recirculation and ultimately access failure. By constantly monitoring the evolution of the access flow during consecutive treatment sessions, it is possible to detect imminent access flow problems. Proper detection of access flow reduction may help in carrying out a 15 maintenance procedure on the access thereby avoiding any access failure.

A non-invasive technique that allows measurement of flow through AV fistulas and grafts is colour Doppler ultrasound. Magnetic Resonance Imaging (MRI) has also been used. However, these techniques require expensive equipment and are not easily used in the dialysis clinic environment.

20 Several methods have been suggested for monitoring recirculation and access flow. Many of these methods involve injection of a marker substance in blood, and the resultant recirculation is detected. The methods normally involve measurement of a property in the extracorporeal blood circuit. Examples of such methods can be found in US 5,685,989, US 5,595,182, US 5,453,576, US 5,510,716, US 5,510,717, US 5,312,550, etc.

25 Such methods have the disadvantage that they require the injection of the marker substance and external equipment for the measurements.

More recently, EP 928 614 and WO 00/24440, suggest to measure a post dialyzer concentration of a substance, in particular urea in the effluent fluid before and after a flow reversal, i.e. before the flow reversal the arterial line carries blood from an upstream position 30 of the blood access, and the venous line carries blood towards a downstream position of the blood access, whereas the arterial line carries blood from an downstream position of the blood access, and the venous line carries blood towards a upstream position of the blood access after the flow reversal. A valve for such reversal is shown in i.e. US 5,605,630 and US 5,894,011. A disadvantage in these methods is the requirement for special equipment 35 for measuring the urea concentration. Urea sensors are as such available but they are not standard equipment for most of the dialysis monitors and they have also a considerable maintenance costs.

SUMMARY OF THE INVENTION

The present invention covers a method for determining a fluid flow rate ( $Q_a$ ) in a blood access having an upstream position and a downstream position using a blood treatment apparatus, the blood treatment apparatus including:

a blood treatment unit having a semi permeable membrane delimiting a first chamber through which blood removed from said blood access passes and a second chamber through which dialysis liquid passes,

an arterial line connected to an inlet of the first chamber, and

a venous line connected to an outlet of the first chamber,

10 said arterial and venous lines being able to be switched between at least a normal configuration, in which said arterial line carries blood from said upstream position of said blood access, and said venous line carries blood towards said downstream position of said blood access, and at least a reversed configuration, in which said arterial line carries blood from said downstream position of said blood access, and said venous line carries blood towards said upstream portion of said blood access,

said method comprising the steps of:

- passing a dialysis liquid through the second chamber of said treatment unit, at least for a time interval  $T$  said dialysis liquid upstream the treatment unit comprising at least a substance having a concentration ( $C_i$ ) different from the concentration of the same substance in blood,

20 during said interval  $T$  a switching occurring between one and the other of said normal and reversed configurations;

- keeping the concentration ( $C_i$ ) of said at least a substance in the dialysis liquid at the treatment unit inlet substantially constant during said interval  $T$ ; obtaining, downstream the treatment unit, a first post-treatment unit conductivity of the dialysis liquid or a first post treatment unit concentration ( $C_R$ ;  $C_N$ ) of said substance in the dialysis liquid, said first

3a

conductivity or concentration relating to the venous and arterial lines configured according to one of said normal or reversed configuration, said first post treatment unity conductivity or concentration referring to the dialysis liquid before switching the venous and arterial lines and during said time interval T;

- obtaining, downstream the treatment unit, a second post treatment unit conductivity of the dialysis liquid or post treatment unit concentration ( $C_R$ ;  $C_N$ ) of said substance in the dialysis liquid, said second conductivity or concentration relating to the venous and arterial lines configured according to the other of said normal or reversed configuration, said second post treatment unity conductivity or concentration referring to the dialysis liquid after switching of the venous and arterial lines and during said time interval T; and
- calculating the fluid flow rate ( $Q_a$ ) in said blood access as a function of:
  - o said first post treatment unit concentration or conductivity and of
  - o said second post treatment unit concentration or conductivity.

The present invention also covers an apparatus for determining the fluid flow rate ( $Q_a$ ) in a blood access having a downstream position and an upstream position, the apparatus comprising:

- 20 a. a dialysis liquid source,
- b. a treatment unit, having a semi permeable membrane delimiting a first chamber through which blood removed from said blood access passes and a second chamber through which dialysis liquid passes,
- c. a dialysis liquid line for circulating dialysis liquid in the second chamber;
- d. an arterial line connected to an inlet of the first chamber,
- e. a venous line connected to an outlet of the first chamber,
- f. said arterial and venous lines being able to be configured according to at least a normal configuration, in which said arterial line carries blood from said upstream position of said blood access, and said venous line carries

3b

blood towards said downstream position of said blood access, and to at least a reversed configuration, in which said arterial line carries blood from said downstream position of said blood access, and said venous line carries blood towards said upstream portion of said blood access,

- g. means for switching the venous and arterial lines, during said time interval T, between one of said normal and reversed configurations to the other of said normal and reversed configurations;
- h. means for varying a concentration (Ci) of at least a substance of the dialysis liquid upstream the treatment unit;
- 10 i. a sensor operating downstream the treatment unit for detecting a post-treatment unit conductivity of the dialysis liquid or a post treatment unit concentration of said substance in the dialysis liquid, and
- j. a control unit capable of performing the following steps:
  - o operating said varying means in such a way that, at least for a time interval T, said dialysis liquid circulating upstream the treatment unit comprises at least a substance having a concentration (Ci) different from the concentration of the same substance in blood, the control unit acting on the varying means to keep substantially constant the concentration Ci of said at least a substance during said time interval T;
  - o obtaining from said sensor a first post-treatment unit conductivity of the dialysis liquid or a first post treatment unit concentration of said substance in the dialysis liquid, for the venous and arterial lines being configured according to one of said normal or reversed configuration, said first conductivity or concentration relating to the dialysis liquid before switching the venous and arterial lines and during said time interval T,
  - o obtaining, from said sensor a second post treatment unit conductivity of the dialysis liquid or post treatment unit

20

3c

concentration of said substance in the dialysis liquid, for the venous and arterial lines being configured according to the other of said normal or reversed configuration, said second conductivity or concentration relating to the dialysis liquid after switching of the venous and arterial lines and during said time interval T; and

- o calculating the fluid flow rate (Qa) in said blood access as a function of said first post treatment unit concentration or conductivity and of said second post treatment unit concentration or conductivity.

10 The present invention further covers a computer program product comprising a computer readable memory storing apparatus executable instructions thereon, which when executed by the control unit of a blood treatment apparatus as defined above render the control unit able of performing the steps of the control unit as defined above.

The present invention also covers the use of an apparatus as defined above, for determining a fluid flow rate in a blood access of a patient.

By providing means for creating a difference in conductivity between the dialysis fluid and blood and by providing a post treatment unity conductivity cell, the apparatus can determine the blood access flow, with relatively inexpensive modifications to conventional dialysis apparatuses.

According to a preferred embodiment, a first and second concentration or conductivity are measured on the post treatment unit fluid flowing downstream the treatment unit, or so called effluent fluid.

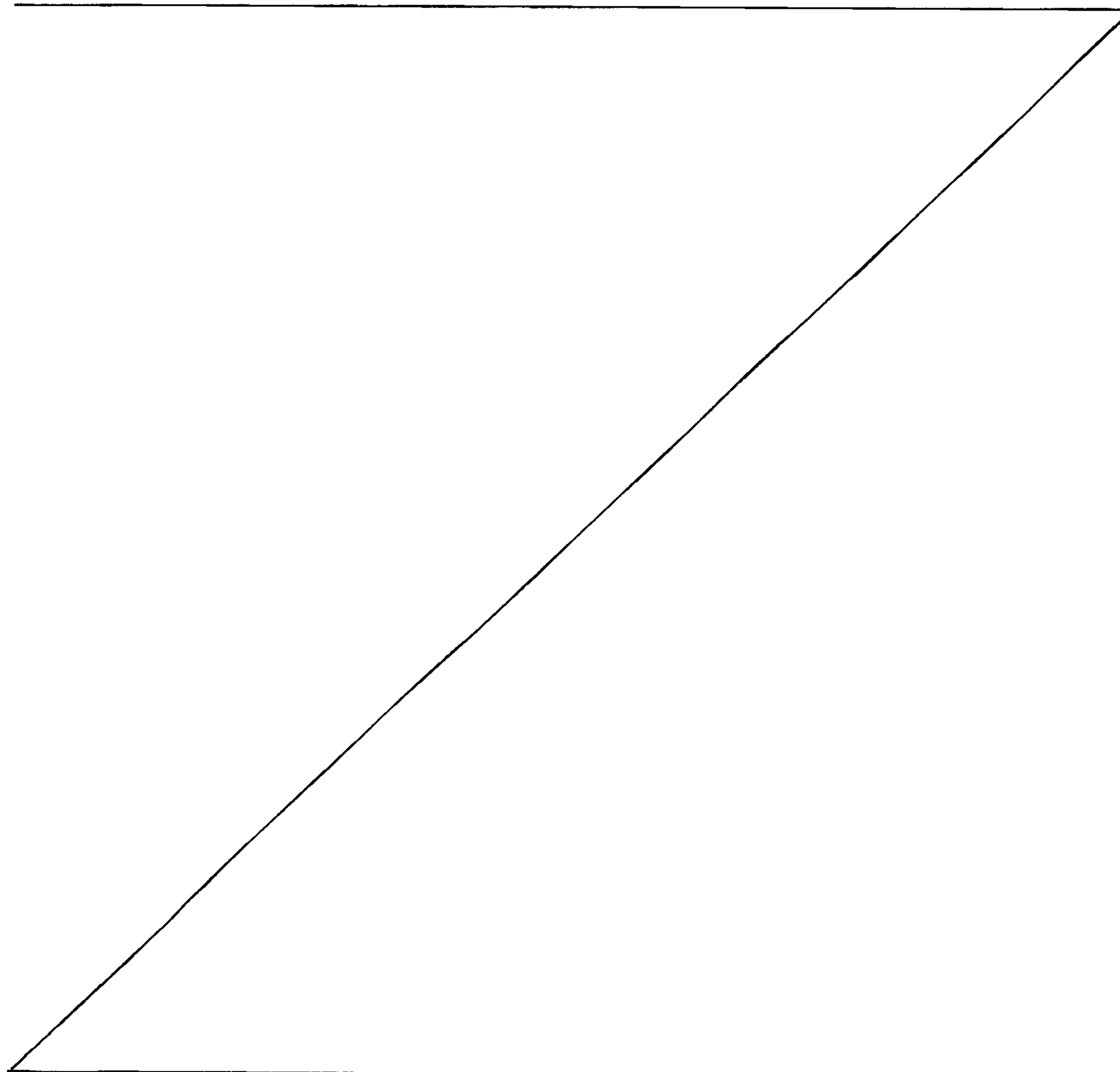
During normal dialysis a blood flow in a first direction is created by operating a blood pump, in which the arterial line carries blood from said upstream position of said blood access, and the venous line carries blood towards said downstream position of said blood access (normal configuration of the lines).

3d

A blood flow in a second direction, in which said arterial line carries blood from said downstream position of said blood access, and said venous line carries blood towards said upstream portion of said blood access (reversed configuration of the lines), may be created by

- manually connecting the arterial line to the downstream position of the blood access and the venous line to an upstream position of the blood access, or by
- connecting the arterial line to both the upstream and the downstream position of the blood access and connecting the venous line to both the upstream and the downstream position of the blood access, closing one of the connections between

---



the arterial line with the blood access and opening the other and closing one of the connections between the venous line with the blood access and opening the other, or by

providing a valve able to connect the arterial line with the upstream position of the access point and the venous line with the downstream position of the access point in a first position of said valve and able to connect the arterial line with the downstream position of the access point and the venous line with the upstream position of the access point in a second position of said valve.

Preferably, the calculation of the fluid flow rate in the blood access is carried out by  
10 using the formula:

$$Q_a = f(C_r, C_i, C_n, Q_{uf}, T_r);$$

According to an embodiment the following formula can be used:

$$Q_a = (T_r - Q_{uf}) * (C_r - C_i) / (C_n - C_r),$$

In which  $Q_a$  is the fluid flow rate in the blood access,  $T_r$  transport rate of substances over the semi permeable membrane of the treatment unit referred to the venous and arterial lines in normal condition,  $Q_{uf}$  is the ultrafiltration flow rate,  $C_r$  is the post treatment unit conductivity after flow reversal,  $C_i$  is pre treatment unit conductivity, and  $C_n$  is the post treatment unit conductivity before the flow reversal.

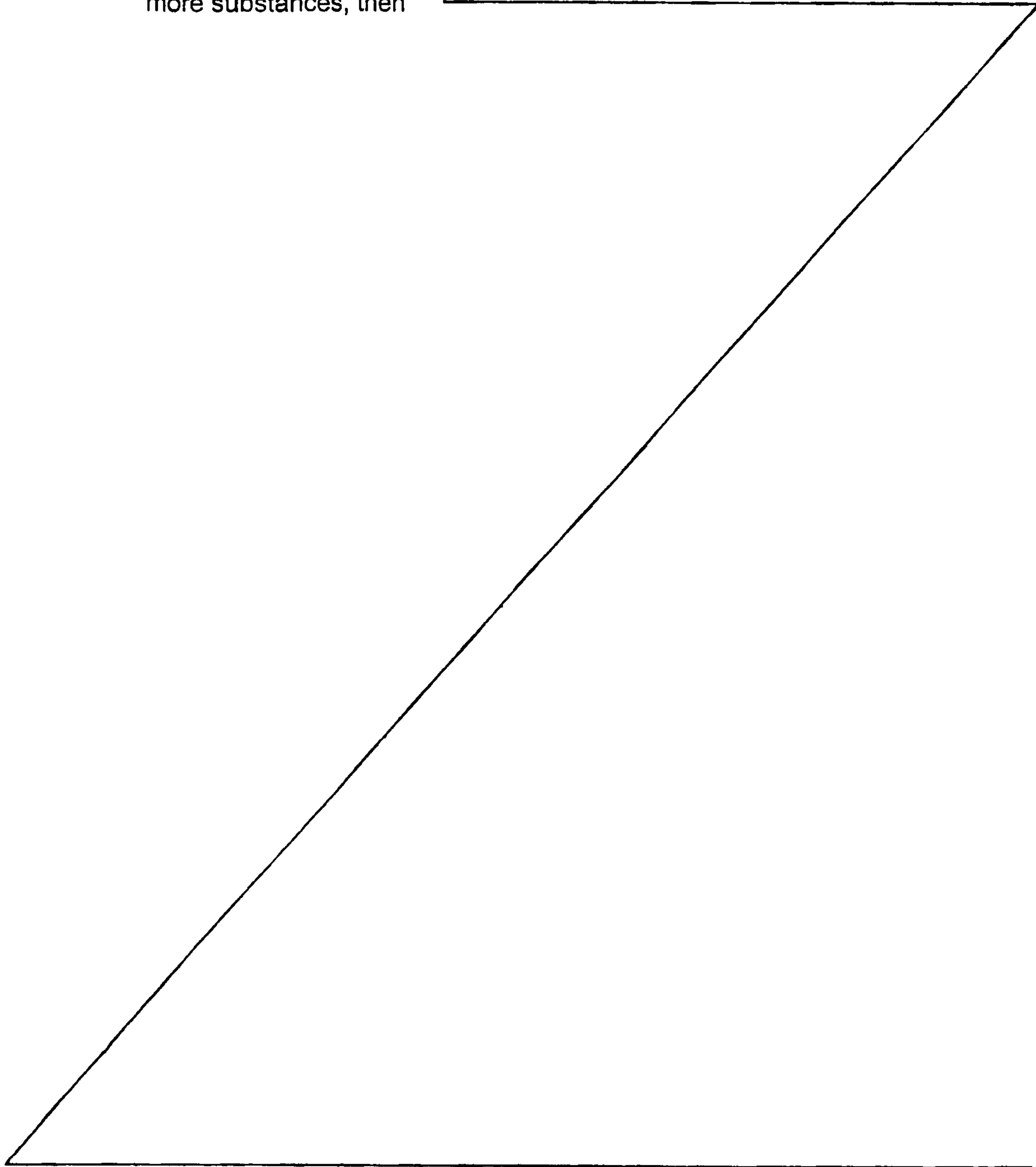
For determination of the transport rate  $T_r$ , the effective ionic dialysance  $D$  can be used. The  
20 effective ionic dialysance  $D$  determined for example as described in EP 658 352. Alternatively, the transport rate can be derived from experience values of a particular dialyzer.

The effective urea clearance, determined by other methods known in the art, can also be used for the transport rate  $T_r$ , since it has been found to be very similar to effective ionic dialysance.

4a

Preferably, to an embodiment of the invention the method (and corresponding blood treatment apparatus) for determining Qa comprises the following steps:

- a. circulating a first dialysis liquid into the second chamber inlet of said treatment unit, said first dialysis liquid presenting a treatment concentration for one or more substances, then



- b. increasing or decreasing at a time  $T_i$  the concentration of the substance in the dialysis liquid for circulating to the second chamber inlet, during a time interval  $T$ , a second liquid having a concentration  $C_i$  for said one or more substances different from the concentration of the same substances in blood,
- 5 c. switching the venous and arterial lines between one to the other of said normal and reversed configurations during the time interval  $T$ ,
- d. obtaining the first post treatment unit conductivity of the dialysis liquid or first concentration of said substance in the dialysis liquid, relating to the dialysis liquid before switching the venous and arterial lines and during said time interval  $T$ ,
- 10 e. obtaining the second post treatment unit conductivity of the dialysis liquid or second post treatment unit concentration of said substance in the dialysis liquid, relating to the dialysis liquid after switching of the venous and arterial lines and during said time interval  $T$ , during said time interval  $T$ , the concentration  $C_i$  of said substance(s) being kept substantially constant.
- 15

According to another feature of the invention it may be provided to that, during said time interval  $T$ , the following consecutive sub-steps are executed:

- a. First configuring the said arterial and venous lines according to the normal configuration for obtaining said first concentration or first conductivity, and then
- 20 b. configuring the arterial and venous lines according to the reversed configuration for obtaining said second concentration or conductivity.

25 Alternatively, during said time interval  $T$ , the following consecutive sub-steps may be provided with:

- a. First, configuring the said arterial and venous lines according to the reversed configuration for obtaining said first post treatment unit concentration or conductivity, and then
- 30 b. configuring the arterial and venous lines according to the normal configuration for obtaining said second concentration or conductivity.

Thanks to this alternative option it is possible to first configure the lines in the reversed configuration for the execution of the  $Q_a$  determination. As for  $Q_a$  calculation, a measurement in the normal configuration is also necessary, by starting in reversed configuration and then passing to normal configuration there is no risk to leave the lines in reverse configuration which would lead to a reduced treatment efficiency.

Another advantage with this modified procedure is that we have an automatic indication that the lines have actually been returned to normal for the rest of the treatment, otherwise there will be no access flow measurement. With the original procedure it is much more difficult for the machine to detect if the lines are left in the reversed position for the rest of the treatment.

5

In term of fistula flow determination, notice that two things will happen if we go from reversed lines back to normal instead of the other way around. First of all, the clearance measured at the conductivity change will be a clearance with reversed lines. This clearance is lower than the normal clearance, how much is determined by the access flow rate. Secondly, the 10 conductivity change caused by returning the lines to normal will go in the opposite direction to normal. The sign of the conductivity change can be handled just by using the absolute value of the change, but the lower clearance value needs to be handled by a change in the formula. As the access flow rate (A) depends on normal configuration clearance ( $K_n$ ), ultrafiltration rate (UF) and reversed flow configuration clearance ( $K_r$ ) according to

$$15 \quad A = (K_n - UF) \cdot \left( \frac{K_r}{K_n - K_r} \right) \quad (1)$$

then

$$A = (K_n - UF) \cdot R \quad (2)$$

with R determined from the inlet conductivity ( $C_i$ ), and the outlet conductivities in normal ( $C_n$ ) and reversed ( $C_r$ ) positions according to

$$20 \quad R = \left( \frac{C_r - C_i}{C_n - C_r} \right) \quad (3)$$

Combining (1) and (2) we see that

$$K_n \cdot R = K_r \cdot R + K_r \quad (4)$$

Access flow rate can therefore be calculated as

$$A = (K_n - UF) \cdot R = K_r \cdot R + K_r - UF \cdot R = (K_r - UF) \cdot R + K_r \quad (5)$$

25 Since  $K_r$  is the measured clearance when the lines are reversed, the only modification to the formula for access flow that has to be made if the lines are reversed from the beginning is that we must add the measured clearance. Note however for the calculation of R that  $C_n$  and  $C_r$  will switch positions time wise if the lines are reversed from the start (i.e.  $C_r$  will be measured before  $C_n$ ).

30

Note that in the present description and in the claims  $C_n$  refers always to conductivity-concentration of the effluent dialysis fluid in normal configurations of the lines while  $C_r$  refers always to conductivity-concentration of the effluent dialysis fluid in reversed configuration of the lines. If the time sequence adopted is first reversed than normal configuration: the first 35 post treatment unit conductivity-concentration of the dialysis liquid is  $C_r$  while the second post treatment unit conductivity-concentration is  $C_n$ . If the time sequence adopted is first

normal than reversed configuration: the first post treatment unit conductivity-concentration of the dialysis liquid is  $C_n$  while the second post treatment unit conductivity-concentration is  $C_r$ .

During execution of the above-disclosed method, the post treatment unit conductivities (first and second) are measured after a delay allowing equilibrium to establish.

Preferably, according to a feature of the invention, the post treatment unit conductivity after the flow reversal is measured at various intervals or continuously so that the value of the conductivity at the time of the flow reversal can be determined by extrapolating the measured values backwards to the moment of the flow reversal. In this way the method can compensate for drift of parameters between the time when the flow is reversed until the time where a substantial equilibrium is reached.

#### BRIEF DESCRIPTION OF THE DRAWINGS

In the following detailed portion of the present description, the invention will be explained in more detail with reference to the exemplary embodiments shown in the drawings, in which Fig. 1 is a partially schematic view of a forearm of a patient provided with an AV fistula.

Fig. 2 is a schematic diagram of an extracorporeal circuit and part of the fluid path of a dialysis machine.

Fig. 3 is a schematic diagram of an extracorporeal circuit including a flow reversal valve.

Fig. 4 is the schematic diagram of Fig. 3, with the valve turned for reversed blood flow

Fig. 5 is a graph showing the conductivities before and after flow reversal, and

Fig. 6 is another graph showing the conductivities before and after flow reversal.

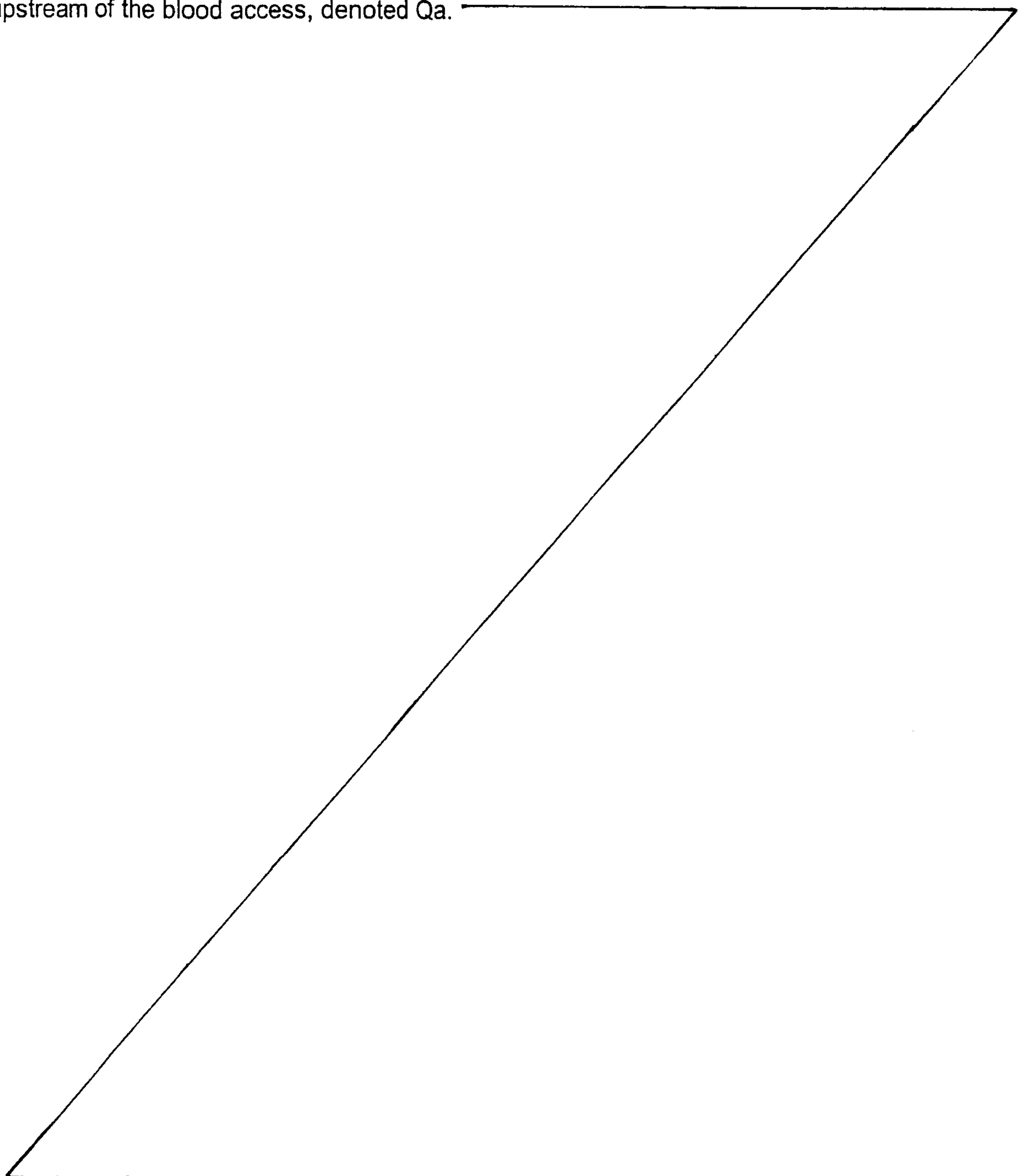
#### DESCRIPTION OF DETAILED EMBODIMENTS OF THE INVENTION

For the purpose of this description, a blood access is a site in which a fluid in a tube can be accessed and removed from and/or returned to the tube. The tube may be a blood vessel of a mammal, or any other tube in which a fluid is flowing. The general term blood access as used here includes arterio-venous fistulas, arterio-venous grafts, and dual-lumen catheters amongst other similar types of blood access that allow for an upstream access position and a downstream access position.

7a

The general terms dialyzer or blood treatment unit as used here include filters for hemodialysis, hemofilters, hemodiafilters, plasmafilters and ultrafilters.

The fluid flow rate is the flow rate of the fluid in the tube or blood vessel immediately upstream of the blood access, denoted  $Q_a$ .



The general term dialysis as used here includes hemodialysis, hemofiltration, hemodiafiltration and therapeutic plasma exchange (TPE), among other similar treatment procedures.

5 The general term effluent fluid as used here refers to the dialysis fluid downstream of the dialyzer or blood treatment unit.

The general term "transport of substances or ions though the semi permeable membrane" includes any parameter that is indicative of the rate at which substances or ions pass through the dialyzer membrane. Examples of such parameters are, clearance, urea clearance, dialysance, ionic dialysance and effective ionic dialysance.

10 The general term ionic dialysance as used here refers to a variable that expresses the transport of ions through the dialyzer membrane. The ionic dialysance is ion dependent, i.e. different ions have different dialysance values. It is also dependent on blood flow, dialysate flow and Quf, so during measurements when determining the access flow these must preferably be held constant. The effective ionic dialysance, herein denoted D, further 15 depends on recirculation effects in the fistula and the cardiopulmonary circuit, and is obtained for example as described by EP 658 352. The major ions determining the conductivity of dialysate liquid are sodium and chloride

20 Fig. 1 discloses a forearm 1 of a human patient. The forearm 1 comprises an artery 2, in this case the radial artery, and a vein 3, in this case the cephalic vein. Openings are surgically created in the artery 2 and the vein 3 and the openings are connected to form a fistula 4, in which the arterial blood flow is cross-circuited to the vein. Due to the fistula, the blood flow through the artery and vein is increased and the vein forms a thickened area downstream of the connecting openings. When the fistula has matured after a few months, the vein is 25 thicker and may be punctured repeatedly. Normally, the thickened vein area is called a fistula.

An arterial needle 5a, to which is connected a piece of tube, is placed in an upstream position in the fistula, in the enlarged vein close to the connected openings and a venous needle 6a, to which is connected a piece of tube, is placed in a position downstream of the 30 arterial needle, normally at least five centimeters downstream thereof.

As described above, the blood access can also be an arterio-venous graft, a double lumen catheter or other similar arrangements.

35 The needles 5a and 6a are connected to a tube system, shown in Fig. 2, forming an extracorporeal circuit 7 comprising a blood pump 8, such as a peristaltic pump. The blood pump propels blood from the fistula, through the arterial needle, the extracorporeal circuit, the venous needle, and back into the fistula.

The extracorporeal blood circuit 7 shown in Fig. 2 further comprises an arterial clamp 9 and a venous clamp 10 for isolating the patient from the extracorporeal circuit should an error occur.

5 Downstream of pump 8 is a dialyzer 11, comprising a first, so called blood chamber 12 and a second, so called dialysis fluid chamber 13 separated by a semi permeable membrane 14. Further downstream of the dialyzer is a drip chamber 15, separating air from the blood therein.

10 The bloodline upstream of the dialyzer 11 is referred to as the arterial line 5, whereas the bloodline downstream from the dialyzer 11 is known as the venous line 6. The arterial and venous lines 5 and 6 are able to be configured according to at least a normal configuration, in which said arterial line carries blood from said upstream position of said blood access and said venous line carries blood towards said downstream position of said blood access, and

15 to at least a reversed configuration, in which said arterial line carries blood from said downstream position of said blood access and said venous line carries blood towards said upstream portion of said blood access.

20 In the normal configuration, blood passes from the arterial needle past the arterial clamp 9 to the blood pump 8. The blood pump drives the blood through the dialyzer 11 and further via the drip chamber 15 and past the venous clamp 10 back to the patient via the venous needle. The drip chamber may comprise an air detector, adapted to trigger an alarm should the blood emitted from the drip chamber comprise air or air bubbles. The blood circuit may comprise further components, such as pressure sensors etc.

25 The dialysis fluid chamber 14 of the dialyzer 11 is provided with dialysis fluid via a first pump 16, which obtains dialysis fluid from a source of pure water, normally RO-water, mixed with one or several concentrates of ions, varying means including metering pumps 17 and 18 being shown for metering such concentrates. Sensors comprising a conductivity cell 22 and a conductivity cell 23 are provided downstream of the points where the concentrates are mixed into the main fluid stream. The signal of the respective conductivity cell 22,23 is in a closed loop manner compared with the desired conductivity and the speed of the pumps 17 and 18 are controlled in response. A further conductivity cell 21, connected to the protective system of the dialysis machine, is provided downstream from all concentrate mixing steps

30 measuring the final total conductivity. The protective system compares the measured final conductivity with a calculated final conductivity and puts the dialysis machine in a safe state, if anything should have gone wrong in the mixing steps.

35

A control unit 85 operates said varying means for circulating a dialysis liquid in the second chamber of said treatment unit in such a way that, at least for a time interval T, said dialysis liquid upstream the treatment unit has a concentration (Ci) of one or more substances different from the concentration of the same substance(s) in blood.

5

According to an embodiment of the invention the difference in concentration is measured as a difference in the conductivity, because most of the components in the dialysis liquid are electrolytes and thus a change in their concentration will inherently lead to a change in the conductivity of the dialysis liquid. It will be understood though, that the invention can also be carried out using the concentration of substances that have no or little effect on the conductivity of the liquid that they are dissolved in, such as urea or glucose.

10

A preferable range for the dialysate conductivity during the blood access flow measurement is 14,5 to 17,5 mS/cm, preferably about 15 to 16 mS/cm. Thus a conductivity difference between the blood and the dialysate of about 1 to 2 mS/cm is created.

15

20

In the specific embodiment shown in figures 5 and 6 an increase in conductivity (concentration of one or more electrolytes) is applied to the fluid upstream the second chamber 13. Said increase starts at time Ti in order to bring the second chamber inlet conductivity to a substantially constant value Ci for a certain time interval T.

According to a first alternative, the invention can work even if instead of an increase a decrease in conductivity or concentration is applied to the fluid at the inlet of the second chamber.

25

According to a second alternative, if the dialysis liquid inherently has the required difference in conductivity with respect to the blood, then no change in conductivity shall be created for performing the method according to the invention.

30

35

A major contribution to the conductivity of the dialysis liquid is sodium chloride. From a physiological standpoint and for best control, the preferred way to adjust the final total conductivity is therefore to change the concentration of sodium chloride. The control unit 85 changes the setting of sodium chloride and in response the speed of metering pump 17 and/or 18 is adjusted as described above. In many types of dialysis apparatus however, the sodium chloride is in a concentrate container together with all the minor amounts of other electrolytes e.g. potassium, magnesium, calcium and peracetic acid, the so called "A concentrate". This concentrate contributes about 12 mS/cm of the usual final 14 mS/cm conductivity. The remainder of the conductivity comes from the bicarbonate concentrate. In

such a dialysis machine (not shown) the conductivity is set by changing the amount of A concentrate in the same way as described above for sodium chloride alone.

Though less attractive from a physiological point of view, it is also possible to change the 5 concentration of all electrolytes, i.e. inclusive bicarbonate simultaneously. It is also possible to change the concentration of any other electrolytes or other components such as glucose.

An exchange of substances between the blood and the dialysis fluid takes place in the .0 dialyzer 11 through the semi permeable membrane 14. The exchange may take place by diffusion under the influence of a concentration gradient, so called hemodialysis, and/or by convection due to a flow of liquid from the blood to the dialysis fluid, so called ultrafiltration.

From the dialysis fluid chamber 14 of the dialyzer is emitted a fluid called the effluent fluid, which is driven by a second pump 19 via a conductivity cell 20 to drain. The conductivity cell 15 measures continuously or at various intervals, the conductivity of the effluent fluid emitted from the dialyzer, to provide an effluent fluid conductivity.

As described above, the present invention provides a method of non-invasively measuring the fluid flow in the fistula immediately before the arterial needle, using the conductivity cell 20 and the dialysis circuit as shown in Fig. 2.

By measuring the first post dialyzer liquid conductivity-concentration during normal dialysis (or normal configuration of the venous and arterial lines) and then reversing the positions of the needles (reversed configuration) and measuring the second post dialyzer conductivity-concentration with the needles in the reversed position, the control unit is able to calculate 25 the blood flow in the blood access, without the addition of any substance to the blood or the dialysis fluid solely for the sake of the measurement.

Note that in order to pass from the normal configuration of the lines to the reversed 30 configuration of the lines the following alternative options can be used.

One way of achieving flow reversal in the needles is by manually disconnecting the needles from the bloodlines and reconnecting the arterial needle to the venous bloodline and the venous needle to the arterial bloodline (not shown). Various other ways for achieving the flow reversal are known to the skilled person.

35 Another embodiment usable for switching the lines between the normal and the reversed condition and vice-versa is shown in Figs. 3 and 4. These figures relate to a schematic diagram of the dialysis circuit according to Fig. 2 with the addition of a valve 28 to perform the flow reversal. The arterial needle 5a is connected to an arterial inlet line 29 of the valve

and the venous needle 6a is connected to a venous inlet line 30 of the valve. The blood pump is connected via arterial line 5 to a first outlet line 31 of the valve and the blood returning from the dialyzer 11 is connected via the venous line 6 to a second outlet line 32 of the valve. The valve 28 comprises a valve housing and a pivotable valve member 33, which 5 is pivotable from the normal position shown on the drawing to a reverse position pivoted 90° in relation to the normal position. In the normal position shown in Fig. 3, the arterial needle 5a is connected to the blood pump 8 and the venous needle 6a is connected to the outlet of the dialyzer, via the drip chamber 15. In the reversed position shown in Fig. 4, the arterial needle 5a is connected to the outlet of the dialyzer and the venous needle 6a is connected to the blood pump 8, as required. Thus the flow is "reversed", and the arterial line 5 carries 10 blood from a downstream position of the blood access, and the venous line 6 carries blood towards an upstream position of the blood access. According to an embodiment, the dialysis machine automatically controls the change of the valve position.

15 As mentioned before other systems may be used to pass form a configuration to the other; for instance manually changeable connections in the arterial line to the downstream position of the blood access and in the venous line to an upstream position of the blood access. Alternatively the lines may be designed to present first conduits connecting the arterial line to both the upstream and the downstream position of the blood access and second conduits 20 connecting the venous line to both the upstream and the downstream position of the blood access. In order to operate the configuration, means for selectively closing one of the first conduits between the arterial line and the blood access and means for selectively closing one of the conduits between the venous line and the blood access can be provided. Such closing means can be manually operable valves or valves controlled by the blood treatment 25 apparatus. Pinch valves, cam valves or clamps having portions active on respective tube portions can be used.

As a further alternative flow distribution means can be used able of connecting the arterial line with the upstream position of the access point and the venous line with the downstream position of the access point, in a first state of said flow distribution means, and able to 30 connect the arterial line with the downstream position of the access point and the venous line with the upstream position of the access point, in a second state of said flow distribution means.

35 Figures 5 and 6 are graphs of measured pre and post dialyzer conductivities. The horizontal axis represent the lapsed times and the vertical axis represent the measured conductivity in mS/cm. In figures 5,6 it is assumed to start with the venous and arterial lines in normal condition and to switch the lines into the reversed condition during the time interval T of

change of the conductivity of the dialysis fluid. As already mentioned it is possible to execute the method according to the invention starting with the reversed condition.

For determining the fluid flow rate in the blood access, a gradient between the conductivity of the dialysis fluid ( $C_i$ ) at the dialyzer inlet and the blood ( $C_b$ ) is created (Fig. 5). Hereto the conductivity of the dialysis liquid is increased from the conventional value of 14 mS/cm (first dialysis liquid having conductivity which corresponds roughly to the conductivity of blood) to 16 mS/cm (second dialysis liquid). The difference may be of another magnitude and, as already mentioned, can also be created by reducing the conductivity of the dialysis fluid. The conductivity of the second liquid is at least 2mS/cm (2 milli-Siemens / centimeter) higher than the conductivity of the first liquid if the conductivity of the first liquid is less or equal to 15mS/cm.

The conductivity gradient is preferably obtained by changing the sodium chloride concentration, but may also be obtained by varying the concentrations of any of the other electrolytes present in dialysis fluid. The change in electrolyte concentration can in advanced dialysis machines such as the Gambro AK 200 S® be executed by changing the settings or programming a step through the user interface. Use of conductivities instead of concentrations is simpler, more reliable, cheaper to implement as it employs the conventional sensors of the treatment apparatus, does not need determination of D or K in two different conditions.

In Figs. 5 and 6 the conductivity of the dialysis fluid  $C_i$  prepared by the dialysis monitor is increased from 14 to 16 mS/cm at time  $T_i$ . The conductivity  $C_n$  of the post dialyzer fluid, the effluent fluid, will begin to increase at time  $T_0$  with a delay  $T_0-T_i$  caused by the volume of the tubes and the dialyzer.  $C_n$  will reach a semi stable value only after some time. Because the increased conductivity of the dialysis liquid causes a transport of ions from the dialysis liquid to the blood, which therefore also slowly increases in conductivity, there will be a slow drift in of the post dialyzer conductivity. The value of  $C_n$  may be determined after the respective value has become substantially stable, as shown in Figure 5. In order to further improve the precision of the method the value of  $C_n$  may be extrapolated forward to the point in time of the flow reversal  $T_{rev}$ . Alternatively, the value of  $C_n$  may be determined while it is still increasing by estimating which substantially stable value  $C_n$  would have reached after an equilibrium has been established by using numerical methods such as curve fitting or and/or extrapolation, in order to determine the value of  $C_n$  at  $T_{rev}$ , shown in Figure 6. The latter approach will allow the method to be carried out in a shorter time span.

The next step is to reverse the flow at  $T_{rev}$  (cf. Figs. 5 and 6) as described above, i.e. a blood flow in a second direction is created in which the venous line 6 carries treated blood from the dialyzer 11 via arterial needle 5a to the upstream position of the blood access. The arterial line 5 draws in blood from the downstream position via venous needle 6a towards the dialyzer 11.

The effect of this measure is a further increase in the effluent conductivity, which after the flow reversal is referred to as  $Cr$ .  $Cr$  will reach a semi stable value only asymptotically. The value of  $Cr$  may be determined after it has become substantially stable, as shown in Figure 5. The value of  $Cr$  may be extrapolated backwards to the point in time of the flow reversal  $T_{rev}$ . Alternatively, the value may be determined while the conductivity is still increasing by estimating which substantially stable value  $Cr$  would have reached at  $T_{rev}$  after an equilibrium has been established by using numerical methods such as curve fitting or extrapolation, as shown in figure 6.

15 The volumes in the dialyzer and connecting tubes that need to be exchanged cause the delay. During the delay period, changes in other parameters may occur and could influence the measurement negatively. The preferred method uses therefore the values extrapolated, to the point in time where the flow reversal took place. The above techniques allow 20 estimating the value of  $Cn$  and of  $Cr$  at the same time  $Tr$ , thereby increasing the accuracy in  $Qa$  calculation.

Unit 85 may then calculate the fluid flow rate in the blood access in accordance with the formula:

25 
$$Qa = (Tr - Quf) * (Cr - Ci) / (Cn - Cr),$$

wherein:

$Qa$  = fluid flow rate in the blood access

$Tr$  = transport rate of substances through the semipermeable membrane

$Ci$  = dialysis liquid conductivity upstream the treatment unit or dialyzer 11

30  $Cn$  = effluent conductivity referring to the dialysis liquid before flow reversal

$Cr$  = effluent conductivity referring to the dialysis liquid after flow reversal

$Quf$  = ultrafiltration flow rate (Quf).

The transport rate may be based on experience values of a particular dialyzer, such 35 as the clearance, calculated from dialyzer capacity and flow rates or measured by comparing a pre-dialysis blood sample with an initial dialysis liquid urea concentration. Alternatively the transport rate ( $Tr$ ) corresponds to measured effective ionic dialysance ( $D$ ) or to measured

clearance K of the dialyzer, preferably the urea clearance value. The ultrafiltration flow rate Quf is on conventional dialysis machines continuously measured and monitored. The equation can therefore be solved and the fluid flow rate in the blood access is determined.

5 Alternatively to what described above with reference to figures 5,6, the measurement of Qa may be obtained by first configuring the lines in the reversed configuration. Then a change in conductivity or concentration (for instance by means of a step increase or decrease in the concentration of defined solutes in the dialysis liquid) is created and finally the concentration or conductivity of the dialysis liquid downstream the dialyzer is measured both for the liquid 10 in reversed condition and for the liquid in normal condition. This second approach is convenient if the Qa measurement is carried out at the beginning of the dialysis session. Indeed the patient can be first connected to the treatment apparatus with the lines in reversed configuration; then when necessary the lines are reversed, the Qa calculated and the treatment can prosecute normally at high efficiency with no need of further line switching 15 as the line are already in normal configuration.

In case the method is performed starting from the reversed configuration, then the Qa is still calculated as a function of the above-identified parameters.

If Tr is determined from the measured clearance K or the measured effective ionic 20 dialysance D in vivo values obtained when said venous and arterial lines are in the normal configuration, the fluid flow rate (Qa) in said blood access is calculated by the formula  $Qa = (Tr - Quf) * (Cr - Ci) / (Cn - Cr)$ , where Tr is the transport rate when the lines are in the normal configuration.

25 If Tr is obtained from the measured clearance K or the measured effective ionic dialysance D in vivo values obtained when said venous and arterial lines are in the reversed configuration, the fluid flow rate (Qa) in said blood access is calculated by the formula  $Qa = (Tr_r - Quf) * (C_r - Ci) / (C_n - C_r) + Tr_r$ , where  $Tr_r$  is the transport rate when the lines are in the reversed configuration.

30

The measured clearance K or the measured effective ionic dialysance D in vivo values can obtained by the following steps:

- a. passing a third dialysis liquid through the second chamber of said treatment unit, said dialysis liquid presenting a concentration for at least one substance, then
- 35 b. obtaining a third post treatment unit conductivity of the dialysis liquid or third post treatment unit concentration of said substance for the third dialysis liquid,

c. at least for a second time interval, increasing or decreasing the concentration of the substance in the third dialysis liquid for passing a fourth liquid through the second chamber inlet, said fourth liquid having a concentration of at least said substance different from the concentration of the same substance in the third liquid,

5 d. obtaining a fourth post treatment unit conductivity of the dialysis liquid or fourth post treatment unit concentration of said substance for the fourth dialysis liquid, calculating the in vivo value of K or D as a function of said third post treatment unit concentration or conductivity and of said fourth post treatment unit concentration or conductivity.

10 In particular the measured clearance K or the measured ionic dialysance D can be determined during the time interval T so as to use the change in conductivity necessary for the implementation of the present invention. In this case a separate modification of the liquid arriving at the second chamber 13 is not necessary and the third liquid corresponds to the 15 first liquid (before the step in figures 5,6) and the fourth liquid corresponds to the second liquid (after the step in figures 5,6).

Practically if only ions concentration is altered, and again referring to the example of figure 5,

20  $Tr=K$

$$K = (D + U) \cdot \left(1 - \frac{\Delta C_o}{\Delta C_i}\right) \quad \frac{\Delta C_o}{\Delta C_i} \text{ being the inverse of the rate between the step in conductivity of the dialysis fluid at the dialyser inlet and the corresponding step of the dialysis liquid at the outlet of the dialyzer}$$

$$A = (K - U) \cdot \left(\frac{C_i - C_r}{C_r - C_n}\right)$$

25 According to another feature of the invention a method and corresponding apparatus is provided for checking if the arterial and venous lines are in said normal or in said reversed configuration is provided for. This check can be executed at any time during treatment. If the check is carried out after the lines switching it can serve to provide an alert signal in case the operator (manual switching) or the apparatus (automatic switching) failed to return the lines 30 in the normal configuration.

The step of checking if the arterial and venous lines are in the normal or in the reversed configuration comprises the following steps:

Determining the in vivo value of a parameter selected in the group comprising:

- Effective ionic dialysance D or

- b. Effective clearance K or
- c. a parameter proportional to effective ionic dialysance or
- d. a parameter proportional to effective clearance,

Comparing the in vivo value of said parameter with a corresponding threshold value for determining if the venous and arterial lines are in said normal or in said reversed configuration.

In case effective ionic dialysance D is used, any known method for in vivo determination of D can be used, such as the one described in EP 658 352.

10 A simple way of determining D comprises the steps of:

passing a third dialysis liquid through the second chamber inlet of said treatment unit, said dialysis liquid presenting a concentration for at least one substance, then

obtaining a third post treatment unit conductivity of the dialysis liquid or third post treatment unit concentration of said substance for the third dialysis liquid, at least for a second time interval, increasing or decreasing the concentration of the substance in the third dialysis liquid for passing a fourth liquid through the second chamber inlet, said fourth liquid having a concentration of at least said substance different from the concentration of the same substance in the third liquid,

20 obtaining a fourth post treatment unit conductivity of the dialysis liquid or fourth post treatment unit concentration of said substance for the fourth dialysis liquid, calculating the in vivo value of D as a function of said third post treatment unit concentration or conductivity and of said fourth post treatment unit concentration or conductivity.

Once obtained the effective ionic dialysance value D, than D can be compared with a threshold value, which can be a set value or a calculated value or a measured value.

In vivo determination of D can of course be carried out during the time interval T.

30 In case the step of checking if the arterial and venous lines are in said normal or in said reversed configuration is carried out during the time interval T, then the following alternative procedure can be used:

- Comparing said obtained first post-treatment unit conductivity of the dialysis liquid or first post treatment unit concentration of said substance in the dialysis liquid with said obtained second post treatment unit conductivity of the dialysis liquid or second post treatment unit concentration of said substance in the dialysis liquid,

- Determining if said conductivity or concentration are increasing after the switching step. Indeed as can be seen in figure 5, if the conductivity of blood is lower than that of the dialysis liquid, after switching into reversed condition, a sudden increase in conductivity of the dialysis liquid downstream the dialyzer is registered.

5 The upstream conductivity cell should preferably calibrated relative to the downstream conductivity cell 20 for improved accuracy. Preferably temperature compensated conductivity cells are used to improve the accuracy of the method.

10

The value for  $C_i$  may be determined by measuring the conductivity of the dialysis fluid before it enters the dialyzer. Alternatively the set value for the dialysis fluid conductivity may be used, since the actual conductivity will only differ marginally from the set value as dialysis monitors control the conductivity of the dialysis fluid very accurately.

15

**WHAT IS CLAIMED IS:**

1. A method for determining a fluid flow rate (Qa) in a blood access having an upstream position and a downstream position using a blood treatment apparatus, the blood treatment apparatus including:

a blood treatment unit having a semi permeable membrane delimiting a first chamber through which blood removed from said blood access passes and a second chamber through which dialysis liquid passes,

an arterial line connected to an inlet of the first chamber, and

a venous line connected to an outlet of the first chamber,

said arterial and venous lines being able to be switched between at least a normal configuration, in which said arterial line carries blood from said upstream position of said blood access, and said venous line carries blood towards said downstream position of said blood access, and at least a reversed configuration, in which said arterial line carries blood from said downstream position of said blood access, and said venous line carries blood towards said upstream portion of said blood access,

said method comprising the steps of:

- passing a dialysis liquid through the second chamber of said treatment unit, at least for a time interval T said dialysis liquid upstream the treatment unit comprising at least a substance having a concentration (Ci) different from the concentration of the same substance in blood, during said interval T a switching occurring between one and the other of said normal and reversed configurations;

- keeping the concentration (Ci) of said at least a substance in the dialysis liquid at the treatment unit inlet substantially constant during said interval T; obtaining, downstream the treatment unit, a first post-treatment unit conductivity of the dialysis liquid or a first post treatment unit concentration (C<sub>R</sub>; C<sub>N</sub>) of said substance in the dialysis liquid, said first

10

20

conductivity or concentration relating to the venous and arterial lines configured according to one of said normal or reversed configuration, said first post treatment unity conductivity or concentration referring to the dialysis liquid before switching the venous and arterial lines and during said time interval T;

- obtaining, downstream the treatment unit, a second post treatment unit conductivity of the dialysis liquid or post treatment unit concentration ( $C_R$ ;  $C_N$ ) of said substance in the dialysis liquid, said second conductivity or concentration relating to the venous and arterial lines configured according to the other of said normal or reversed configuration, said second post treatment unity conductivity or concentration referring to the dialysis liquid after switching of the venous and arterial lines and during said time interval T; and
- calculating the fluid flow rate ( $Q_a$ ) in said blood access as a function of:
  - o said first post treatment unit concentration or conductivity and of
  - o said second post treatment unit concentration or conductivity.

2. Method according to claim 1, wherein the step of passing a dialysis liquid through the second chamber comprises the following sub-steps:

- a. passing a first dialysis liquid through the second chamber inlet of said treatment unit, said first dialysis liquid presenting a treatment concentration for said substance, then
- b. increasing or decreasing at a time  $T_i$  the concentration of the substance in the dialysis liquid for passing through the second chamber inlet a second liquid which, during the time interval T, has a concentration of at least said substance different from the concentration of the same substance in blood.

3. Method according to claim 1, wherein the step of passing a dialysis liquid through the second chamber comprises the following sub-steps:

- a. passing a first dialysis liquid through the inlet of the second chamber of said treatment unit, said first dialysis liquid presenting a treatment concentration for prefixed substances; and
- b. during the time interval  $T_i$ , increasing or decreasing at a time  $T_i$  the concentration of more than one of said prefixed substances in the dialysis liquid for passing through the second chamber inlet a second liquid having a concentration of said substances different from the concentration of the same substances in blood.

4. Method according to claim 2, wherein said substance is an ion.

10 5. Method according to claim 3, wherein said substances are ions.

6. Method according to claims 3, wherein the fluid flow rate ( $Q_a$ ) in said blood access is calculated as a function of said first post treatment unit concentrations or conductivity and of said second post treatment unit concentrations or conductivity.

7. Method according to claim 1, wherein a step of checking if the arterial and venous lines are in said normal or in said reversed configuration is provided for.

8. Method according to claim 7, wherein the step of checking if the arterial and venous lines are in the normal or in the reversed configuration comprises the following steps:

a. determining the in vivo value of a parameter selected in the group consisting

20 of:

- i. effective ionic dialysance  $D$ ,
- ii. effective clearance  $K$ ,
- iii. a parameter proportional to effective ionic dialysance, and
- iv. a parameter proportional to effective clearance; and

b. comparing the in vivo value of said parameter with a corresponding threshold value for determining if the venous and arterial lines are in said normal or in said reversed configuration.

9. Method according to claim 8, wherein the step of determining the in vivo value of said parameter comprises the steps of:

a. passing a third dialysis liquid through the second chamber inlet of said treatment unit, said dialysis liquid presenting a concentration for at least one substance, then

b. obtaining a third post treatment unit conductivity of the dialysis liquid or third post treatment unit concentration of said substance for the third dialysis liquid,

c. at least for a second time interval, increasing or decreasing the concentration of the substance in the third dialysis liquid for passing a fourth liquid through the second chamber inlet, said fourth liquid having a concentration of at least said substance different from the concentration of the same substance in the third liquid,

d. obtaining a fourth post treatment unit conductivity of the dialysis liquid or fourth post treatment unit concentration of said substance for the fourth dialysis liquid, and

e. calculating the in vivo value of said parameter as a function of said third post treatment unit concentration or conductivity and of said fourth post treatment unit concentration or conductivity.

10. Method according to claim 8, wherein said threshold value is a set value or a calculated value or a measured value.

11. Method according to claim 8, wherein the step of in vivo determination of said parameter is carried out during the time interval T.

12. Method according to claim 8, wherein a further step of sending an alert signal is provided for in case the comparing step determines that the venous and arterial lines are in said reversed configuration.

13. Method according to claim 7, wherein the step of checking if the arterial and venous lines are in said normal or in said reversed configuration is carried out during said first time interval T.

14. Method according to claim 13, wherein the step of checking if the arterial and venous lines are in said normal or in said reversed configuration comprises the following sub-steps:

10        - comparing said obtained first post-treatment unit conductivity of the dialysis liquid or first post treatment unit concentration of said substance in the dialysis liquid with said obtained second post treatment unit conductivity of the dialysis liquid or second post treatment unit concentration of said substance in the dialysis liquid; and

      - determining if said conductivity or concentration are increasing after the switching step.

15. Method according to claim 1, further comprising the steps of:

20        a. determining the transport rate (Tr) of ions through the semi permeable membrane,

      b. obtaining the first post treatment unit conductivity (Cn; Cr) for the dialysis liquid before switching the venous and arterial lines,

      c. obtaining the second post treatment unit conductivity (Cr; Cn) for the dialysis liquid after switching the venous and arterial lines, and

      d. calculating the fluid flow rate (Qa) in said blood access as a function of said first and second post treatment unit conductivities and of said transport rate.

16. Method according to claim 15, wherein the fluid flow rate (Qa) is calculated from the values of said transport rate (Tr), said first post treatment unit conductivity

(Cn), said second post treatment unit conductivity (Cr), and the conductivity of the dialysis liquid (Ci) upstream the treatment unit.

17. Method according to claim 15 or 16, wherein said first and second post treatment unit conductivities (Cn, Cr) are obtained by measuring the conductivity of the effluent fluid exiting from the second chamber of said treatment unit.

18. Method according to any one of claims 1 to 17, further comprising the step of obtaining the ultrafiltration flow rate (Quf).

19. Method according to claim 16, wherein the fluid flow rate (Qa) in said blood access is calculated by the formula  $Qa = (Tr) * (Cr - Ci) / (Cn - Cr)$ .

10 20. Method according to claim 18, wherein the fluid flow rate (Qa) in said blood access is calculated by the formula  $Qa = (Tr - Quf) * (Cr - Ci) / (Cn - Cr)$ .

21. Method according to claim 18, wherein Tr is determined from the measured clearance K or the measured effective ionic dialysance D in vivo values obtained when said venous and arterial lines are in the normal configuration, the fluid flow rate (Qa) in said blood access being calculated by the formula  $Qa = (Tr - Quf) * (Cr - Ci) / (Cn - Cr)$ , where Tr is the transport rate when the lines are in the normal configuration.

22. Method according to claim 18, wherein  $Tr_r$  is transport rate of ions though the semi permeable membrane determined from the measured clearance K or the measured effective ionic dialysance D in vivo values obtained when said venous and arterial lines are in the reversed configuration, the fluid flow rate (Qa) in said blood access being calculated by the formula  $Qa = (Tr_r - Quf) * (Cr - Ci) / (Cn - Cr) + Tr_r$ .

23. Method according to claim 17 or 18, wherein the measured clearance K or the measured effective ionic dialysance D in vivo values are obtained by the following steps of:

- a. passing a third dialysis liquid through the second chamber of said treatment unit, said dialysis liquid presenting a concentration for at least one substance, then
- b. obtaining a third post treatment unit conductivity of the dialysis liquid or third post treatment unit concentration of said substance for the third dialysis liquid,
- c. at least for a second time interval, increasing or decreasing the concentration of the substance in the third dialysis liquid for passing a fourth liquid through the second chamber inlet, said fourth liquid having a concentration of at least said substance different from the concentration of the same substance in the third liquid,
- d. obtaining a fourth post treatment unit conductivity of the dialysis liquid or fourth post treatment unit concentration of said substance for the fourth dialysis liquid, and
- e. calculating the in vivo value of K or D as a function of said third post treatment unit concentration or conductivity and of said fourth post treatment unit concentration or conductivity.

24. Method according to claim 15 or 20, wherein the transport rate (Tr) corresponds to:

- experience values of a particular dialyser,
- calculated values,
- measured effective ionic dialysance (D),
- using a predialysis blood sample together with the initial dialysis liquid urea concentration, or
- measured clearance K of the dialyser.

25. Method according to claim 21 or 22, wherein the measured clearance K or the measured ionic dialysance D are in vivo values determined during the time interval T.

26. Method according to claim 2 or 25, wherein the third liquid corresponds to the first liquid and the fourth liquid corresponds to the second liquid.

27. Method according to claim 3 or 25, wherein the third liquid corresponds to the first liquid and the fourth liquid corresponds to the second liquid.

28. Method according to claim 2 or 3, wherein during said time interval T the conductivity of the second liquid is higher than the conductivity of the first liquid.

29. Method according to claim 28, wherein during said time interval T the conductivity of the second liquid is at least 1mS/cm higher than the conductivity of the first liquid.

10 30. Method according to claim 29, wherein during said time interval T the conductivity of the second liquid is at least 2mS/cm higher than the conductivity of the first liquid if the conductivity of the first liquid is less or equal to 15mS/cm.

31. Method according to any one of claims 28 to 30 comprising the following steps:

- changing the conductivity in the first liquid upstream the treatment unit to define the second liquid,
- keeping substantially constant during said time interval T the conductivity of the second liquid upstream the treatment unit,
- waiting a delay after start of said conductivity change and then determine the time T0 when a prefixed change in conductivity occurs in the liquid downstream of the dialyzer,
- measuring a plurality of first values of the conductivity of the liquid downstream the treatment unit after said time T0,
- calculating the first post-treatment unit conductivity of said liquid from said plurality of values;

20

- f. switching the lines from one of said configurations to the other of said configurations;
- g. measuring a plurality of second values of the conductivity of the liquid downstream the treatment unit after said time switching, and
- h. calculating the second post-treatment unit conductivity of said liquid from said plurality of values.

32. Method according to claim 31, wherein the further step of changing the conductivity in the second liquid upstream the treatment unit is provided for.

33. Method according to claim 31, wherein the measurement of the first values is  
10 carried out after a delay from time T0.

34. Method according to claim 31, wherein the moment Trev when the switching occurs is determined, the measurement of the second values being carried out after a delay from time Trev.

35. Method according to claim 34, wherein the plurality of second values of concentration or conductivity after the switching step is continuously or intermittently measured and the concentration or conductivity (Cr) at the time of the switching  $T_{rev}$  is determined by extrapolating the measured values backwards to the moment ( $T_{rev}$ ) of the switching.

36. Method according to any one of claims 1 to 35, wherein the conductivity of  
20 the dialysis fluid is adjusted by varying the sodium chloride concentration, or by varying the concentration of all A-concentrate electrolytes simultaneously, or by varying the concentration of all electrolytes in the dialysis liquid.

37. An apparatus for determining the fluid flow rate (Qa) in a blood access having a downstream position and an upstream position, the apparatus comprising:  
a. a dialysis liquid source,

b. a treatment unit, having a semi permeable membrane delimiting a first chamber through which blood removed from said blood access passes and a second chamber through which dialysis liquid passes,

c. a dialysis liquid line for circulating dialysis liquid in the second chamber;

d. an arterial line connected to an inlet of the first chamber,

e. a venous line connected to an outlet of the first chamber,

f. said arterial and venous lines being able to be configured according to at least a normal configuration, in which said arterial line carries blood from said upstream position of said blood access, and said venous line carries blood towards said downstream position of said blood access, and to at least a reversed configuration, in which said arterial line carries blood from said downstream position of said blood access, and said venous line carries blood towards said upstream portion of said blood access,

10 g. means for switching the venous and arterial lines, during said time interval T, between one of said normal and reversed configurations to the other of said normal and reversed configurations;

h. means for varying a concentration (Ci) of at least a substance of the dialysis liquid upstream the treatment unit;

i. a sensor operating downstream the treatment unit for detecting a post-20 treatment unit conductivity of the dialysis liquid or a post treatment unit concentration of said substance in the dialysis liquid, and

j. a control unit capable of performing the following steps:

- o operating said varying means in such a way that, at least for a time interval T, said dialysis liquid circulating upstream the treatment unit comprises at least a substance having a concentration (Ci) different from the concentration of the same substance in blood, the control unit acting on the varying means to keep substantially constant the concentration Ci of said at least a substance during said time interval T;

10

- obtaining from said sensor a first post-treatment unit conductivity of the dialysis liquid or a first post treatment unit concentration of said substance in the dialysis liquid, for the venous and arterial lines being configured according to one of said normal or reversed configuration, said first conductivity or concentration relating to the dialysis liquid before switching the venous and arterial lines and during said time interval T,
- obtaining, from said sensor a second post treatment unit conductivity of the dialysis liquid or post treatment unit concentration of said substance in the dialysis liquid, for the venous and arterial lines being configured according to the other of said normal or reversed configuration, said second conductivity or concentration relating to the dialysis liquid after switching of the venous and arterial lines and during said time interval T; and
- calculating the fluid flow rate (Qa) in said blood access as a function of said first post treatment unit concentration or conductivity and of said second post treatment unit concentration or conductivity.

38. Apparatus according to claim 37, wherein said sensor comprises a post

20 treatment unit conductivity (Cn,Cr) cell.

39. Apparatus according to claim 37, wherein the varying means are designed for increasing or decreasing the concentration of one or more substances in the dialysis liquid.

40. Apparatus according to claim 37, wherein the control unit is programmed to calculates the fluid flow rate (Qa) in said blood access as a function of said first post treatment unit concentrations or conductivity and of said second post treatment unit concentrations or conductivity.

41. Apparatus according to claim 37, wherein during said time interval T the control unit is programmed to act on said switching means to carry out the following consecutive sub-steps:

- a. first configuring the said arterial and venous lines according to the normal configuration for obtaining said first concentration or first conductivity (Cn); then
- b. configuring the arterial and venous lines according to the reversed configuration for obtaining said second concentration or conductivity (Cr); and
- 10 c. returning the arterial and venous lines to the normal configuration for prosecution of the blood treatment.

42. Apparatus according to claim 37, wherein during said time interval T the control unit acts on said switching means to carry out the following consecutive sub-steps:

- first, configuring the said arterial and venous lines according to the reversed configuration for obtaining said first post treatment unit concentration or conductivity (Cr), and then
- configuring the arterial and venous lines according to the normal configuration for obtaining said second concentration or conductivity (Cn)

20 and then prosecuting the blood treatment.

43. Apparatus according to claim 37, wherein the control unit is able to perform a step of checking if the arterial and venous lines are in said normal or in said reversed configuration.

44. Apparatus according to claim 41, wherein after having configured the arterial and venous lines according to the reversed configuration the control unit is able to perform a further step of checking if the arterial and venous lines are in said first or in said reversed configuration.

45. Apparatus according to claim 37, wherein the control unit is able to perform the following steps:

- a. determining the transport rate (Tr) of ions through the semi permeable membrane,
- b. obtaining the first post treatment unit conductivity (Cn, Cr) relating to the dialysis liquid before switching the venous and arterial lines,
- c. obtaining the second post treatment unit conductivity (Cr, Cn) relating to the dialysis liquid after switching the venous and arterial lines, and
- d. calculating the fluid flow rate (Qa) in said blood access as a function of said first and second post treatment unit conductivities and of said transport rate.

10

46. Apparatus according to claim 45, wherein the fluid flow rate (Qa) is calculated from the values of said transport rate (Tr), said first post treatment unit conductivity (Cn,Cr), said second post treatment unit conductivity (Cr,Cn), and the conductivity of the dialysis liquid (Ci) upstream the treatment unit.

47. Apparatus according to claim 45, wherein said first and second post treatment unit conductivities (Cn, Cr) are obtained by said sensor.

48. Apparatus according to any one of claims 37 to 47, further comprising means acting on the dialysis line for causing an ultrafiltration flow rate (Quf).

20

49. Apparatus according to claim 46 or 48, wherein the fluid flow rate (Qa) in said blood access is calculated by the formula  $Qa=(Tr)*(Cr-Ci)/(Cn-Cr)$  or by the formula  $Qa=(Tr-Quf)*(Cr-Ci)/(Cn-Cr)$ , wherein Tr is determined from the measured clearance K or the measured ionic dialysance D in vivo values obtained when said venous and arterial lines are in the normal configuration.

50. Apparatus according to claim 46 or 48, wherein Tr is the transport rate obtained from the measured clearance K or the measured ionic dialysance D in

vivo values obtained when said venous and arterial lines are in the reversed configuration, the fluid flow rate (Qa) in said blood access being calculated by the formula  $Qa = (Trr - Quf) * (Cr - Ci) / (Cn - Cr) + Trr$ .

51. Apparatus according to any one of claims 37 to 50, further comprising means for preparing dialysis liquid with a conductivity different from said blood, preferably said means comprises means for controlled mixing of electrolyte concentrates with water.

52. Apparatus according to any one of claims 37 to 51, wherein said switching means comprises:

- 10      - manually changeable connections in the arterial line to the downstream position of the blood access and in the venous line to an upstream position of the blood access, or
- first conduits connecting the arterial line to both the upstream and the downstream position of the blood access and second conduits connecting the venous line to both the upstream and the downstream position of the blood access, means for selectively closing one of the first conduits between the arterial line and the blood access and means for selectively closing one of the conduits between the venous line and the blood access, or
- 20      - a valve able to connect the arterial line with the upstream position of the access point and the venous line with the downstream position of the access point in a first position of said valve and able to connect the arterial line with the downstream position of the access point and the venous line with the upstream position of the access point in a second position of said valve, or
- flow distribution means for connecting the arterial line with the upstream position of the access point and the venous line with the downstream position of the access point in a first state of said flow

distribution means and able to connect the arterial line with the downstream position of the access point and the venous line with the upstream position of the access point in a second state of said flow distribution means.

53. Apparatus according to any one of claims from 37 to 52, wherein said treatment unit comprises one selected in the group consisting of:

- i. a dialyzer;
- ii. an hemofilter;
- iii. a plasmafilter;
- iv. an hemodiafilter; and
- v. an ultrafilter.

10 54. Computer program product comprising a computer readable memory storing apparatus executable instructions thereon, which when executed by the control unit of a blood treatment apparatus as defined in any one of claims 37 to 53 render the control unit able of performing the steps of the control unit as defined in any one of claims 37 to 53.

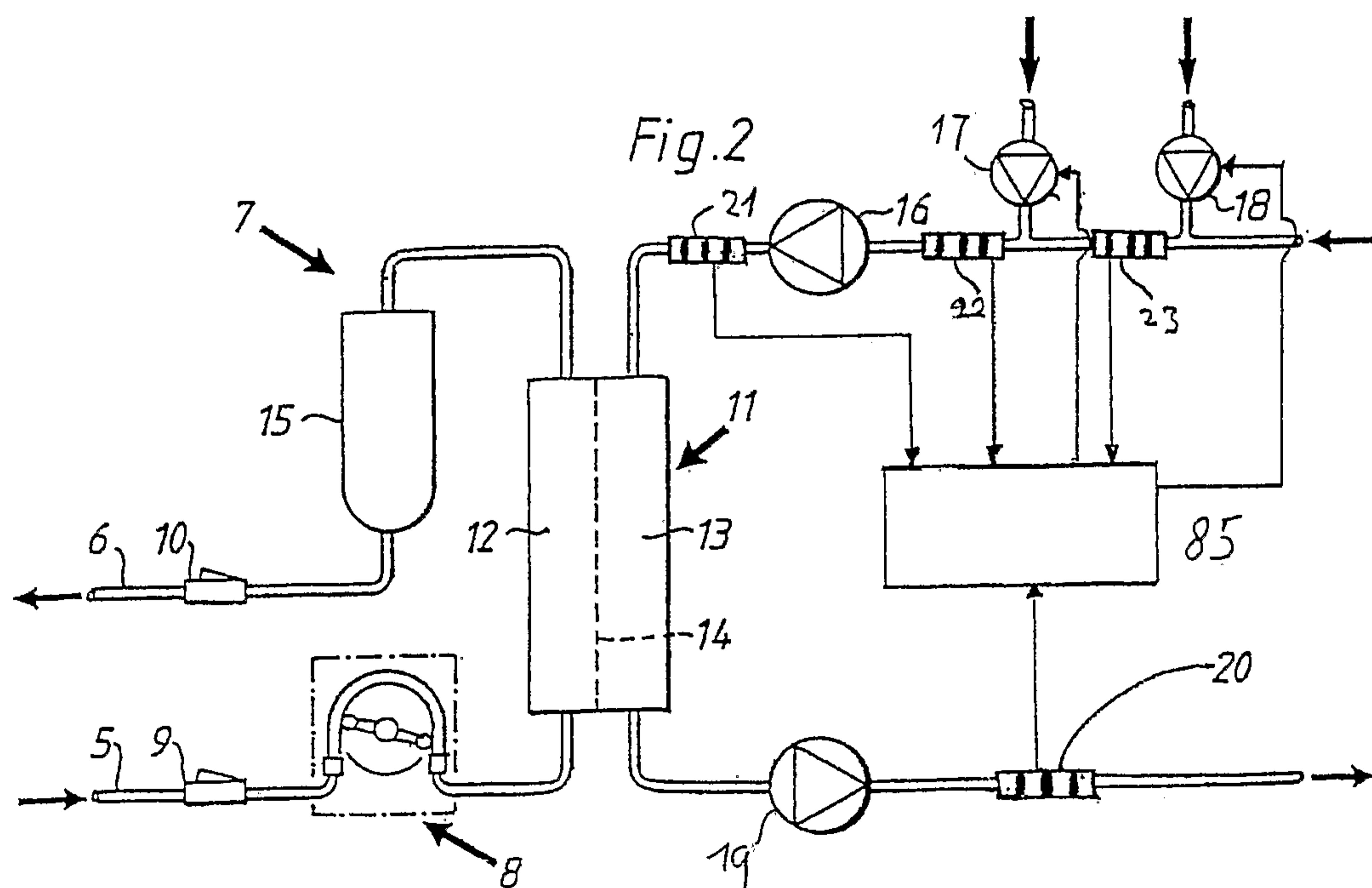
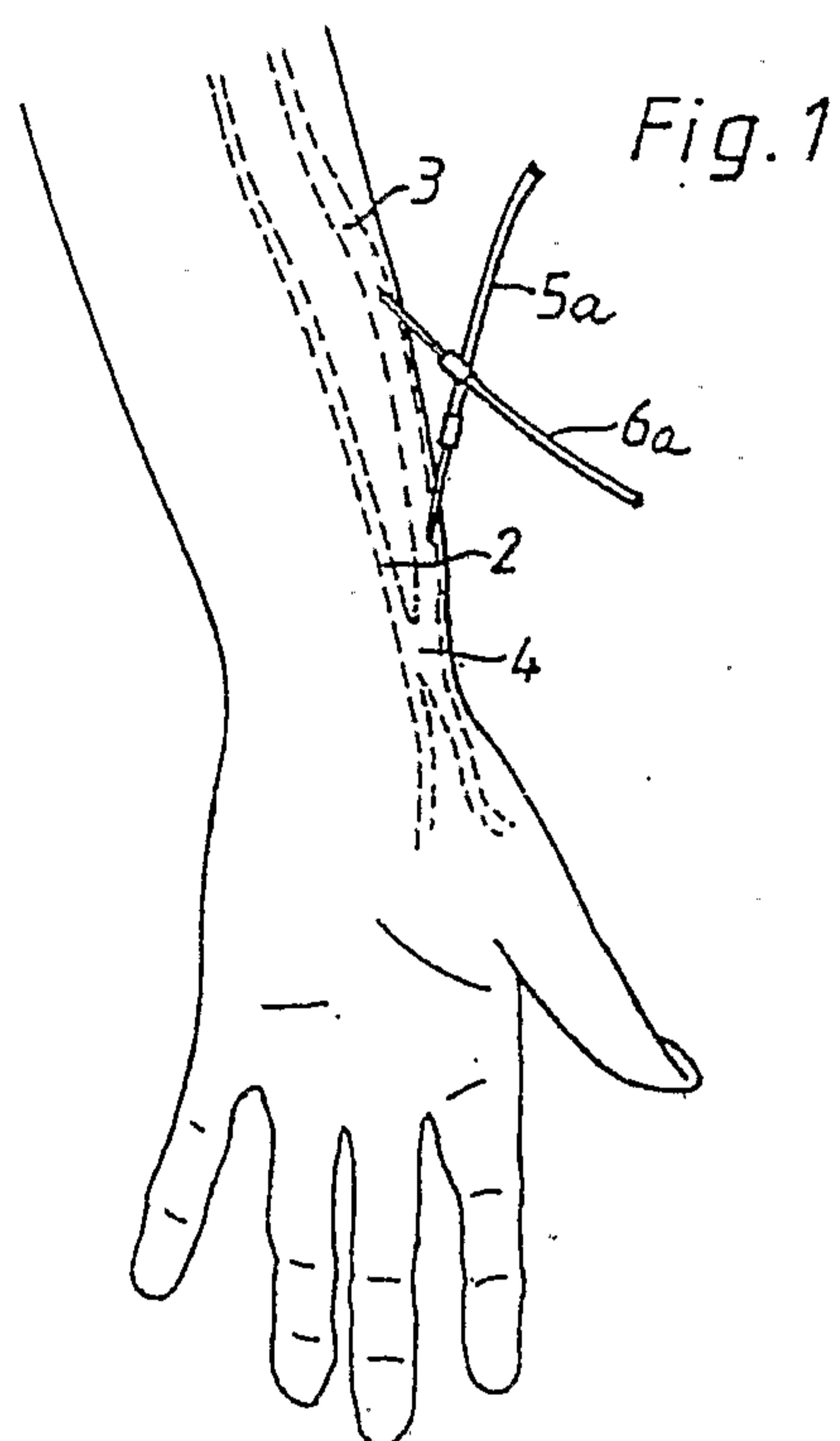
55. The computer program product according to claim 54, wherein it is stored on a magnetic or optic data carrier.

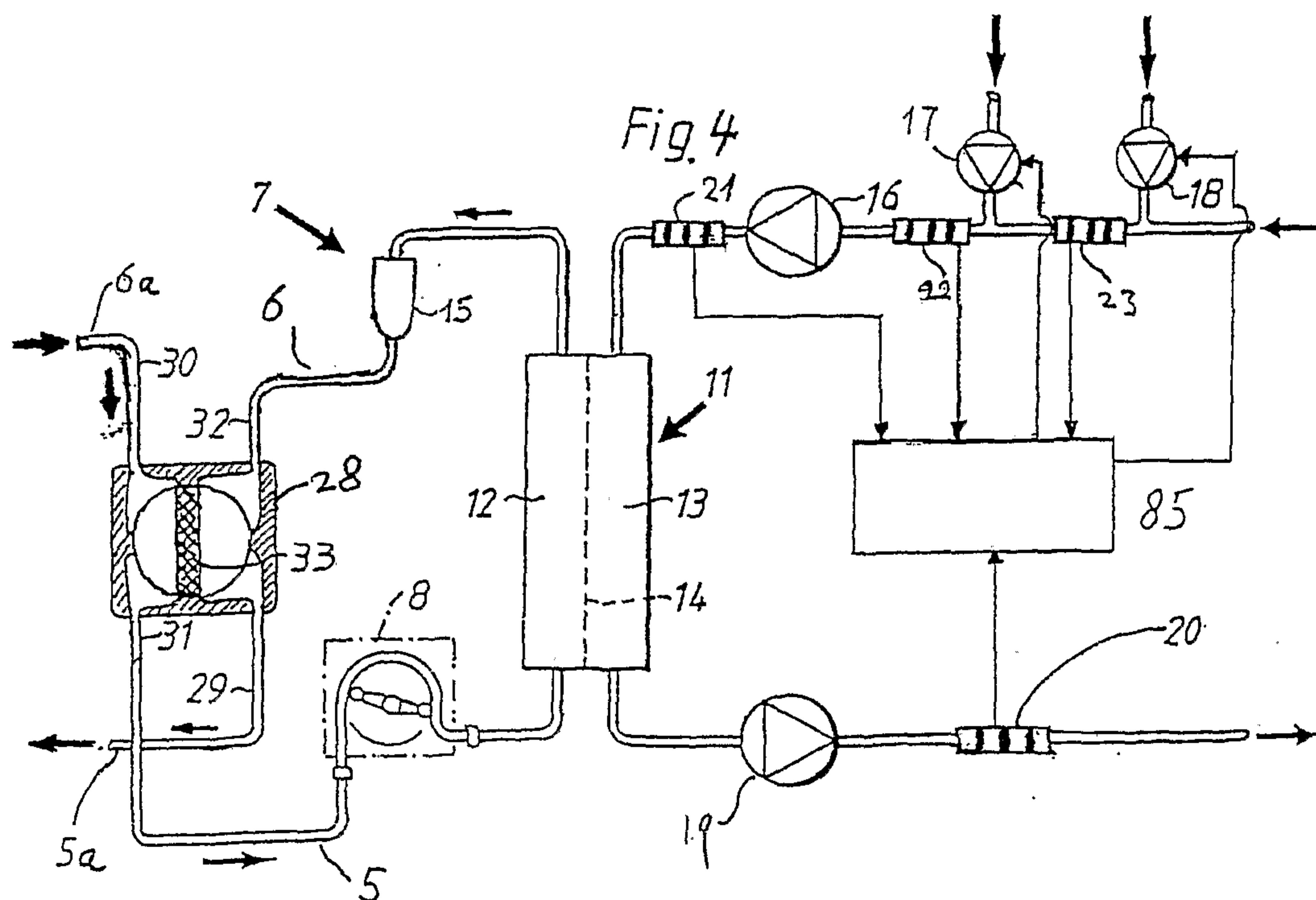
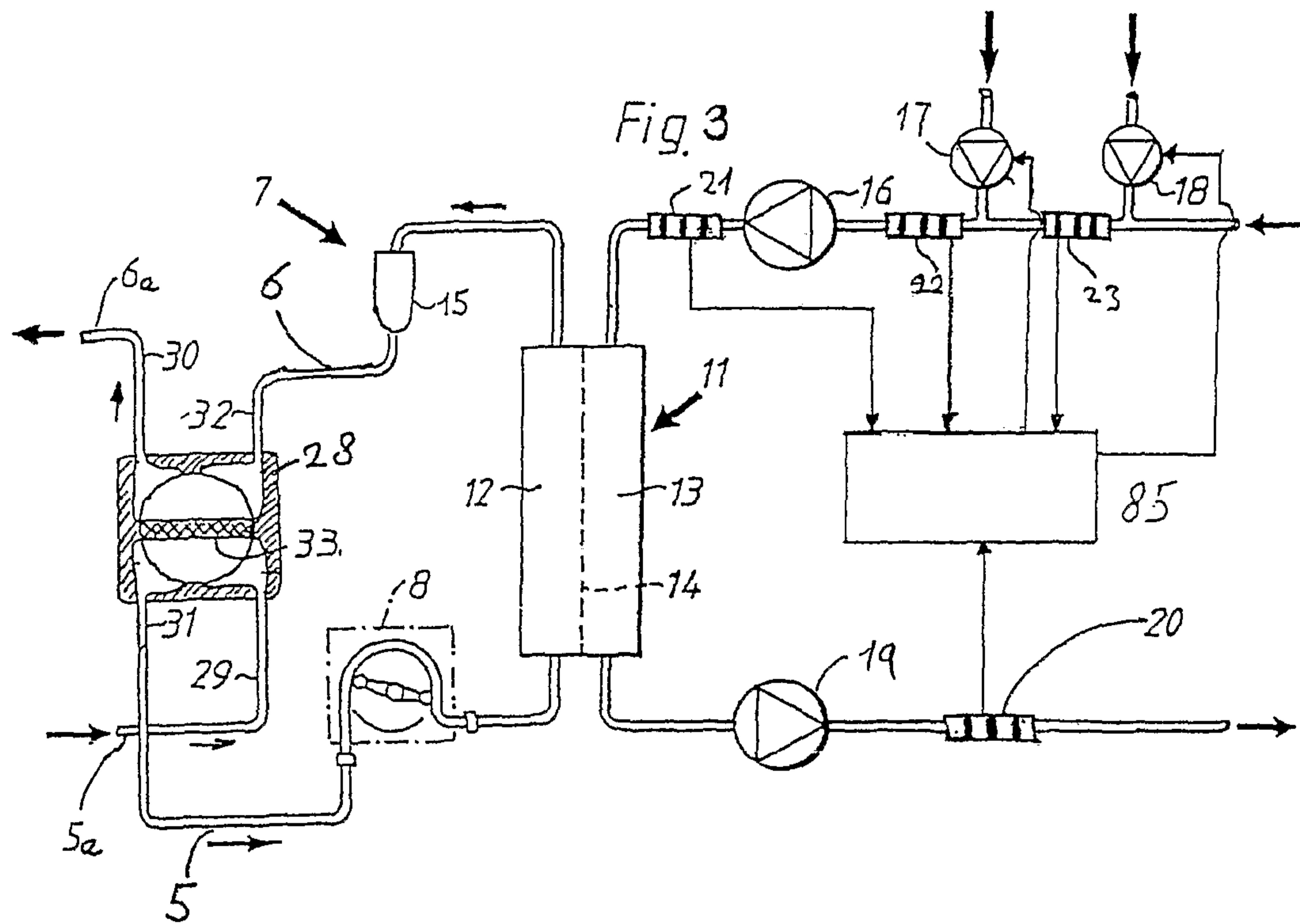
20 56. The computer program product according to claim 54, wherein it is stored on a computer memory.

57. The computer program product according to any one of claims 54 to 56, wherein it is stored on a read only memory.

58. The computer program product according to claim 54, wherein it is stored on a computer remote from the blood treatment apparatus and is able to be transmitted on an electric or electromagnetic signal.

59. Use of a blood treatment apparatus as defined in any one of claims 37 to 53, for determining a fluid flow rate in a blood access of a patient.





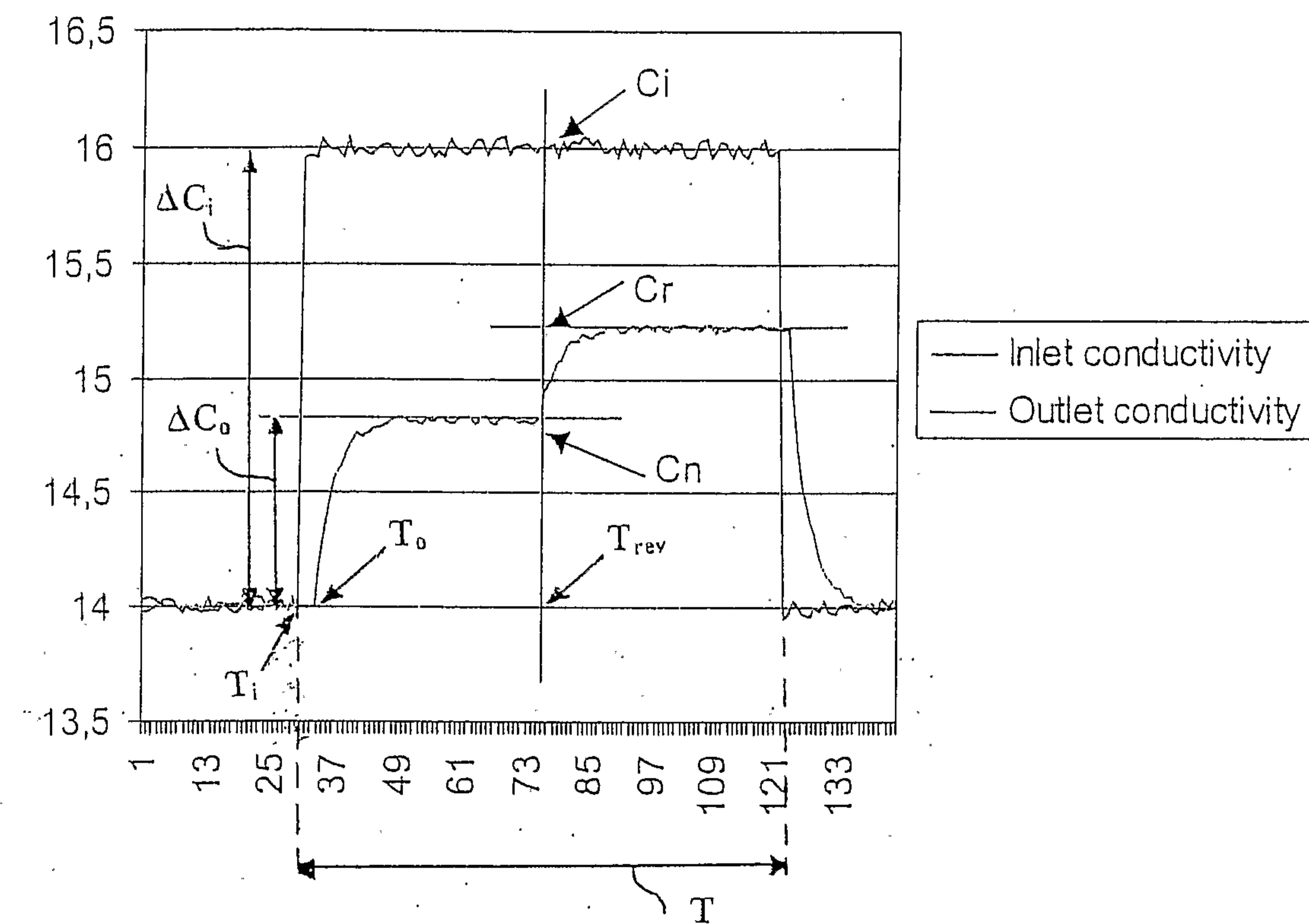


Fig. 5

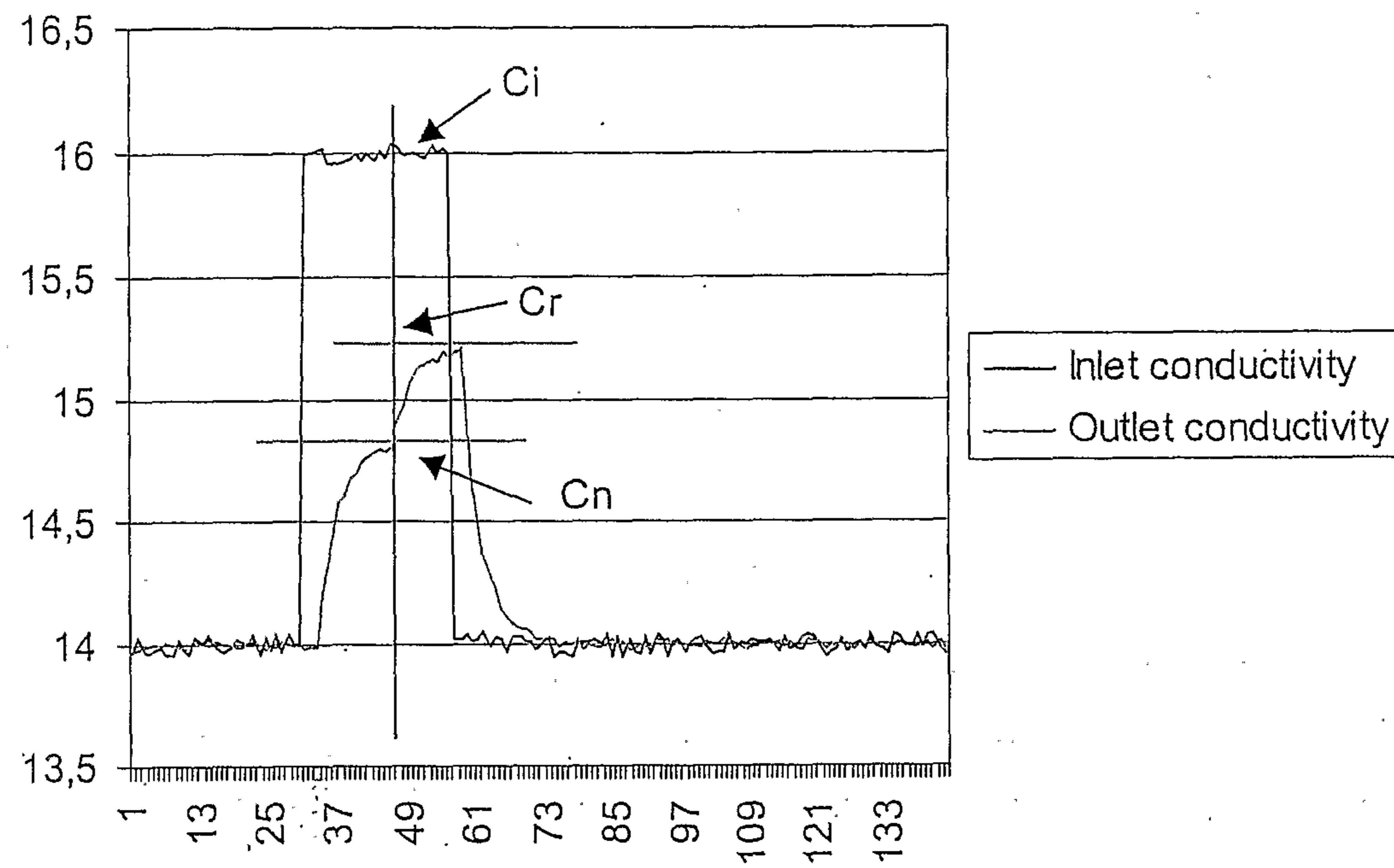


Fig.6

