Product Catalog

VISION BIOMED SERVICES SDN BHD

Quality and Customer Satisfaction is Our Business Best Achievement.
Vision Biomed Services Sdn Bhd wishes to become an
excellent company in medical industry by providing
quality services and customer trust.

HEAMODIALYSIS MACHINE DISINFECTANT

PEROXY PLUS™ RP – COLD STERILANT FOR DIALYZER REPROCESSING

Peroxy Plus RP is used for reprocessing dialyzers machine. Peroxy Plus RP is a mixture of peracetic acid, hydrogen peroxide and acetic acid. Peroxy Plus RP leaves no toxic residues and after reacting with organic material, it decomposes into oxygen and acetic acid. This cold sterilant destroy tuberculocidal, bactericidal and fungicidal.

MDR NO.: GC8658524-181933

COMPOSITION

- Peracetic acid solution (PAA) (>4%)
- Hydrogren peroxide (20%)
- Acetic acid (5>%), stabilizing agents, corrosion inhibitors, RO water.

MICROBIOLOGICAL PROPERTIES

Bactericide, fungicide, virucide and sporicide.

GBL PEROXY PLUS RP

Designed to be used in dialyzer reprocessing systems. Each canister holds five liters of GBL Peroxy Plus RP concentrate. To prevent excessive pressure buildup, all canisters are provided with patented vented caps. These caps should not be altered or replaced. Please store in upright position. GBL Peroxy Plus RP comes with two 5-liter canisters per case.

PACKAGING

Shelf Life	Storage	Packaging	Order information
2 Years	5-25°C	5 liter canister x 2	6127-5000

HEAMODIALYSIS MACHINE DISINFECTANT

CITRO™ PLUS – HEMODIALYSIS MACHINE DISINFECTANT; CITRIC ACID 21%

Used for chemo-thermal disinfectant (combination of chemical and heat disinfectant) of hemodialysis machines with recirculation.
CITRO PLUS is a very strong and effective disinfectant because of synergetic effects of its ingredients.

MDR NO.: GC3506519-36236

COMPOSITION

- Citric acid (21%)
- Malic acid (2.5%)
- Lactic acid (2.5%)

MICROBIOLOGICAL PROPERTIES

A 3% CITRO PLUS solution is bactericidal, fungicidal, virucidal and sporicidal at 84°C

CITRO PLUS

Designed to be used for thermochemical disinfection in hemodialysis machine. Each canister holds five liters of Citro Plus concentrate. To prevent excessive pressure buildup, all canisters are provided with patented vented caps. These caps should not be altered or replaced. Please store in upright position. Citro Plus comes with two 5 liter canisters per case.

PACKAGING

Shelf Life	Storage	Packaging	Order information
2 Years	5-25°C	5 liter canister x 2	6106-5000

MACHINE SURFACE DISINFECTANT

SURFACTO™ NEUTRAL HEMODIALYSIS MACHINE SURFACE DISINFECTANT

Surface disinfectant for hemodialysis machine in between treatments.
Suitable for efficient disinfectant and cleansing of dialysis machine surfaces.

COMPOSITION

- Alkyl dimethyl benzyl ammonium chloride (0.1%), Dodecyl trimethyl ammonium bromide, propylene glycol, benzotriazole, ethanol 96% antioxidants and stabilizing agents.
- Free of glutaraldehyde, formaldehyde, buffers and other phenol derivatives.

MICROBIOLOGICAL PROPERTIES

- Bactericide
- Fungicide

MDR NO.: GB5312319-36237

SPECIFICATION

Shelf Life	Storage	Packaging	Order information
2 Years	5-25°C	1 liter bottle x 20	06009-1000

- Ready to use formula with corrosion inhibitor.
- Formula compatible with acrylic, glass and PVC.
- Aldehyde and phenol free.
- Spray onto the device surface and wait for 1 minute, clean with sterile wipe.
- Ready to use, do not dilute.
- Extremely effective disinfectant with cleansing properties.
- Biodegradable, harmless for environment, safe for humans.
- Does not leave trace or a greasy film.
- Broad antimicrobial spectrum in 1 minute

TEST STRIP

HYDROGEN PEROXIDE (RESIDUAL) TEST STRIP

- To check the level of chemical and must be 0 ppm on dialyzer before using on patients.
- Fast and easy to use.
- Detection range: 0 - 10 ppm (H2O2)

PERACETIC ACID TEST STRIP

- To check the level of chemical and must be 4% ppm on dialyzer before storing.
- Fast and easy to use.
- Detection range: 250 - 2000 ppm
- Detection range (%): 0.5% - 4.0%

MDR NO.: GC6743724-162522

ADVANTAGES

- Results instantly or in less than 15 seconds.
- The dry test strip is stable at room temperature.
- Water resistant label and color blocks.
- Easy to read color blocks.
- More accurate.

SPECIFICATION

Shelf Life	Product name	Range	Packaging	Order information
2 Years	Hydrogen Peroxide (Residual)	0 - 10 ppm	100 strips/btl	HPTS-18-VB
2 years	Peracetic Acid	250 - 2000 ppm	100 strips/btl	PATS-19-VB

TEST STRIP

RESIDUAL CHLORINE TEST STRIP

- To check the level of chlorine.
- Fast and easy to use.
- Detection range: 0 - 1.0 ppm
- Standard : Total Chlorine Negative (<0.1 ppm)

TOTAL HARDNESS TEST STRIP

- To check the level of hardness.
- Fast and easy to use.
- Detection range: 0 - 300 ppm
- Standard : Water Hardness Negative (<17 ppm)

ADVANTAGES

- Results instantly or in less than 15 seconds.
- The dry test strip is stable at room temperature.
- Water resistant label and color blocks.
- Easy to read color blocks.
- More accurate.

SPECIFICATION

Shelf Life	Product name	Range	Packaging	Order information
2 years	Residual Chlorine	0 - 1.0 ppm	100 strips/btl	CTS-20-VB
2 years	Total Hardness	0 - 300 ppm	100 strips/btl	THTS-21-VB

HAEMODIALYSIS BICARBONATE BAG (BIBAG)

- Is a polypropylene bag filled with sodium bicarbonate powder which enables ready to use saturated liquid bicarbonate concentrate by mixing with water based on the dilution factors set on the HD machine.
- It is non-sterile and the powder is stored in enclosed bag which minimize the risk of contamination.
- Indicated for use in patients undergoing hemodialysis for acute and chronic renal failure and intended to be used as one component in the preparation of dialysate in a 3 - stream proportioning HD machine equipped with bag module.
- The connector is designed to readily gets plugged in to the slot provided in the HD machine.
- Available for Fresenius dialysis machine model 4008S and 4008S NG/5008.

MDR NO.: GC9752722-84818

ADVANTAGES

- Handling - easy to handle, easy to transport, occupies less space and reduces labour.
- Hygiene - Promotes hygienic delivery of Sodium Bicarbonate and Prevents Microbial growth and ensures safe dialysis treatment.
- Environment - Minimum wastage, environment friendly, no spillage.

SPECIFICATION

Shelf Life	Size	Packaging	Order information	Consumption
2 Years	555 grams	18 bags/box	SBB555 4008S NG/5008	5 hours/treatment (single treatment)
2 Years	650 grams	18 bags/box	SBB650 4008S NG/5008	5 hours/treatment (single treatment)
2 years	900 grams	16 bags/box	SBB900 4008S	8 hours/treatment (2 treatment)
2 years	900 grams	12 bags/box	SBB900 4008S NG/5008	8 hours/treatment (2 treatment)

HAEMODIALYSIS BICARBONATE CARTRIDGE (BICART)

- The haemodialysis bicarbonate cartridge is filled with sodium bicarbonate and housed in a polypropylene container.
- When placed in the designated holder, water flows through the cartridge to generate a saturated bicarbonate solution ready for use. This solution is then proportioned with haemodialysis concentrates (35X or 45X) and purified water in the dialysis machine to produce bicarbonate-based dialysis fluid.
- Can be used for single treatment (standard) and double treatment (high volume)
- Suitable for Nipro, Gambro and BBraun dialysis machines.

ADVANTAGES

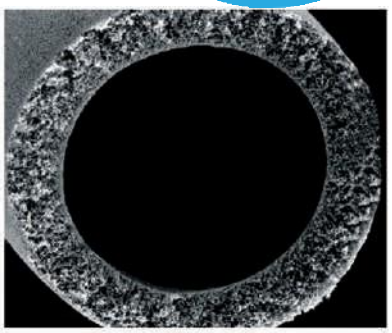
- Handling - easy to handle, easy to transport, occupies less space and reduces labour.
- Hygiene - Promotes hygienic delivery of Sodium Bicarbonate and Prevents Microbial growth and ensures safe dialysis treatment.
- Minimum wastage, environment friendly, no spillage and save labour.

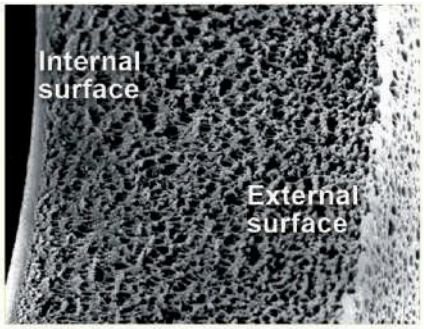
SPECIFICATION

Shelf Life	Size	Packaging	Order information	Consumption
2 Years	500 grams	10 cartridges/box	SL2-HBC-500	5 hours/treatment (single treatment)
2 years	650 grams	10 cartridges/box	SL2-HBC-650	6 hours/treatment (single treatment)
2 years	720 grams	10 cartridges/box	SL2-HBC-720	7 hours/treatment (2 treatment)
2 years	760 grams	10 cartridges/box	SL2-HBC-760	7 hours/treatment (2 treatment)
2 years	900 grams	10 cartridges/box	SL2-HBC-900	8 hours/treatment (2 treatment)
2 years	1100 grams	10 cartridges/box	SL2-HBC-1100	10 hours/treatment (2 treatment)

DIALYSIS FLUID FILTER

DF SERIES





Core Features

- **First** Chinese Manufacturer of Endotoxin Filters
- Significant **Biocompatibility**
- **Compatible** to a Wide Range of Machines
- Open to **Multiple Disinfection** Methods
- **Longer** usage time

MDR NO.: GB4794825-194066

Model	DF210, DF210-T	DF220	DF230
Effective Membrane Area	2.1 m²	2.2 m²	0.9 m²
Fiber Membrane Diameter	200 μm	200 μm	250 μm
Fiber Membrane Thickness	40 μm	40 μm	50 μm
Maximum Service Life	900 hours or 160 treatments	3 months or 100 treatments	3 months or 150 treatments
Connector	Hansen connector	/	Hansen connector
Filtration Rate	≥1000 mL/min	≥1000 mL/min	≥1000 mL/min
Particle Contents after Filtration	Per each square millimeter, there are less than 25 particles with a diameter larger than 10μm and less than 3 particles with a diameter larger than 25μm.		
Bacterial Contents after Filtration	≤0.1CFU/mL	≤0.1CFU/mL	≤0.1CFU/mL
Endotoxin Contents after Filtration	<0.03EU/mL	<0.03EU/mL	<0.03EU/mL
Maximum TMP	600 mmHg	600 mmHg	600 mmHg
Material of Shell and End Cover	Polycarbonate	Polypropylene	Polycarbonate
Material of Fiber Membrane	Polyethersulfone	Polyethersulfone	Polyethersulfone
Material of Sealing Ring	Silicone	Silicone	Silicone
Material of Sealing Glue	Polyurethane	Polyurethane	Polyurethane

DIALYSATE FLUID FILTER

HEMONOVA SERIES

DIALYSATE FILTER

PRODUCT ADVANTAGES

- Independent intellectual property rights of the film making process
- Enhanced retention of bacterial endotoxins
- BPA free, full security
- Multiple models can be adapted to different clinical models

TECHNICAL SPECIFICATIONS

Model number	HEMONOVA-F	HEMONOVA-B	HEMONOVA-N	HEMONOVA-G
Surface area	2.2 m²	2.4m²	2.4m²	2.4m²
Membrane inner diameter	200±20（μm）			
Membrane wall thickness	40±10（μm）			
Direction of filtration	Medial membrane → Outside of membrane			
Maximum operating pressure	200KPa			
Period of use/Filter exchange	160 times/3 months			
Method of sterilization	Sterilization by electron beam radiation			
Filtration rate	≥1000ml/min			
Particle filtration performance	The content of particles of 10um and more than 10um in the filtrate was ≤25/mL The content of particles of 25um and more than 25um in the filtrate was ≤3 /mL			
Bacterial filtration performance	The amount of bacteria in the filtrate was≤1CFU/10mL			
Endotoxin filtration performance	The endotoxin content in the filtrate was <0.03EU/mL			

HOLLOW FIBER HEMODIALYZER

- PES is simpler and it has better stable physical and chemical properties than PS.

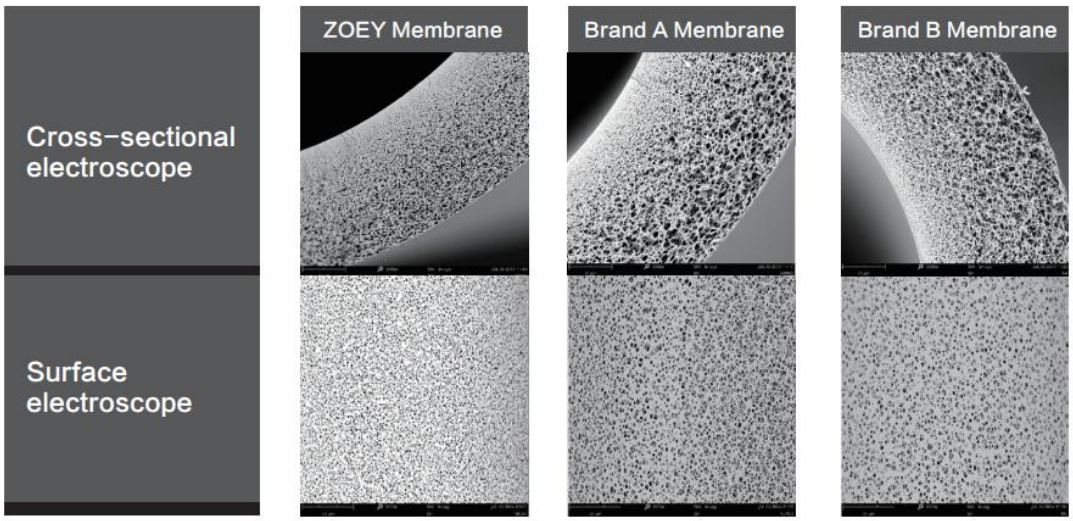
ADVANTAGE

- PP shell, PES membrane, BPA free
- Better bio-compatibility
- Excellent toxin clearance
- Optimized product design
- Smaller blood volume

CONTRAST

The section microstructure shows that ZOEY's hollow fiber membrane has the tightest dense layer, the smallest aperture change, and more uniform surface distribution compared with other 2 types membranes.

MDR NO.: GC9530122-97957



HOLLOW FIBER HEMODIALYZER

PRODUCT PARAMETERS

Product specification		14L	16L	18L	20L	22L	24L
A (Effective surface area) m^2		1.4	1.6	1.8	2.0	2.2	2.4
V (blood priming volume) mL		78	91	101	110	121	134
UF-coefficient mL/h/mmHg		21	24	27	30	33	36
TMP max		500 mmHg					
Max. dialysate flow		800 mL/min					
△P (pressure drop blood)		≤13KPa (QB=300 mL/min)					
Clearance rate	Urea	182	189	193	197	200	200
Q_B=200mL/min	Creatinine	170	180	187	193	198	200
Q_D=500mL/min	Phosphate	170	174	181	193	196	199
Q_F=10mL/min	Vitamin B_{12}	110	119	130	145	161	173
Clearance rate	Urea	245	251	259	266	274	286
Q_B=300mL/min	Creatinine	234	238	246	253	260	267
Q_D=500mL/min	Phosphate	217	224	239	251	256	265
Q_F=10mL/min	Vitamin B_{12}	134	140	154	167	178	188
Clearance rate	Urea	283	292	302	313	325	338
Q_B=400mL/min	Creatinine	260	269	285	294	303	312
Q_D=500mL/min	Phosphate	252	259	269	281	291	301
Q_F=10mL/min	Vitamin B_{12}	147	159	170	184	196	207
Membrane		PES hollow fiber membrane					
Sterilization method		Radiation sterilization					

Product specification		14H	16H	18H	20H	22H	24H
A (Effective surface area) m^2		1.4	1.6	1.8	2.0	2.2	2.4
V (blood priming volume) mL		78	91	101	110	121	134
UF-coefficient mL/h/mmHg		51	58	65	72	80	87
TMP max		500 mmHg					
Max. dialysate flow		800 mL/min					
△P (pressure drop blood)		≤13KPa (QB=300 mL/min)					
Clearance rate	Urea	197	198	199	200	200	200
Q_B=200mL/min	Creatinine	193	195	199	199	200	200
Q_D=500mL/min	Phosphate	181	186	191	194	196	196
Q_F=10mL/min	Vitamin B_{12}	138	150	159	165	169	175
	β2-MG β2	48	53	59	65	72	80
Clearance rate	Urea	275	283	288	292	292	297
Q_B=300mL/min	Creatinine	261	269	275	274	278	283
Q_D=500mL/min	Phosphate	236	248	255	262	268	274
Q_F=10mL/min	Vitamin B_{12}	168	184	195	203	211	223
	β2-MG β2	/	/	/	/	/	/
Clearance rate	Urea	318	332	344	350	352	365
Q_B=400mL/min	Creatinine	290	300	311	320	330	338
Q_D=500mL/min	Phosphate	268	284	296	310	318	326
Q_F=10mL/min	Vitamin B_{12}	186	202	218	225	233	245
	β2-MG β2	/	/	/	/	/	/
S.C.	Inulin	0.9±10%					
	β2-MG β2	≥0.7					
	Myohemoglobin	≥0.55					
	Albumin	≤0.01					
Membrane		PES hollow fiber membrane					
Sterilization method		Radiation sterilization					

In vitro testing; According to EN1283, ISO 8637 and YY0053; The above data represent in vitro test performance only, and may vary depending on the patient's blood composition and clinical setting for in vivo use.

PHARMACEUTICAL (SODIUM CHLORIDE)

WATER SOFTENING SALT

Our pharmaceutical grade salt is the ideal choice for numerous pharmaceutical applications including dialysis solutions, enteral and parenteral infusion solutions. Its versatility makes it an indispensable component in pharmaceutical manufacturing. It is used in medical, pharmaceutical and cosmetic industries for various applications such as drug manufacturing, medical solutions and cosmetics manufacturing.

PACKAGING

- 10kg
- 25kg

SPECIFICATION

Pharmaceutical Sodium Chloride		
Products Name	**Items**	**Standard Value**
Quality Standard	NaCL	99.5% min
	Moisture	1% max
	Water Insolubles	2% max
	Dimension	16x16mm, 20x8mm
	Weight per tablet	5.0 ~ 11.0g

GUARD FILTER

MELT BLOWN FILTER CARTRIDGES

FEATURES AND BENEFITS

- Designed for use in breathing and anesthetic systems for the protection of the patient, hospital personnel and the equipment from potential microbial contamination.
- 100% high purity polypropylene (melt blown), no adhesives.
- High dirt holding capacity and low-pressure loss.
- To remove unwanted particles, pollutants and chemicals from liquids.

TYPICAL APPLICATIONS

- Seawater RO pre-filtration
- Portable water filtration
- Chemical filtration - wide chemical compatibility
- Plating baths
- Amine Filtration
- Meets FDA requirement for food and beverage contact

SPECIFICATION

Filter Media	Melt Blown Polypropylene	**Micron Rating**	1.0,5.0,10.0, 20,50,70,100um
Inner Core	Polypropylene	**Filtration Efficiency**	≥80%
Outer Diameter	63mm (2.5")/ 114mm (4.5")	**Length (based on Cut Ends)**	10", 20"
Inner Outer	28mm (1.10")	**Filtration Area (m2)**	0.4 m2 per 10"

AUTOMATIC DIALYZER REPROCESSOR MACHINE

ROXYPLUS

Equipped with the advance technology with inbuild report generation with dialyzer consumption record.

DESIGN

ROXYPLUS very unique but yet simple system, makes it easy to use at a same time robust. Stainless Steel frame keeps the machine rust free for years together. Capacitive touchscreen are very much easy to operate even with wet gloves in hands. Large 7” Display modules enable user to take a reading on a system in just a glance, even from distance with ease.

MDR NO.: GC6743724-162522

QUALITY

ROXYPLUS stands apart due to its quality. Even a smallest component used in system is tested for its quality. Tubes are made from medical grade PVC. High quality quick release coupling and connectors are used for ease of service. All the connectors and valves used in system are complies with strict ISO and CE quality system regulation.

SAFETY

ROXYPLUS offers world class safety to both patients and users. With non-return valves on RO water inlet and drain outlet also on sterilant intake tube prevents from any contaminations from either side. No one can bypass the step or any step of machine, this gives a full safety to patients, that the dialyzer reprocessed on ROXYPLUS are safe to reuse as per AAMI standards. The Peroxy plus RP sterilant and ROXYPLUS system combination requires no premixing, eliminating the possibility of human error during the dilution process. All the care has been taken to prevent any chance of cross contamination.

FAST (TIME SAVING)

It takes only 8-9 minutes to reprocess the one dialyzer with ROXYPLUS. Also, it takes only 3 minutes to disinfect the system after it is used.

ECONOMICAL

ROXYPLUS uses only 27ml of Peroxy Plus RP cold sterilant for each Dialyzer. With the proper processing, one can easily increase the number of reuses with all the protocols followed to make the treatment more economical for the patients. Also, this gives a chance to Nephrologists to use the dialyzer with the reverse ultra-filtration to achieve maximum clearance even with reused dialyzers.

USER FRIENDLY

Graphical User Interface of the system is designed in a such a way that every user can operate system effectively. All the basic parameter is set in the software. So, user has just to start the process, automatically it will process the dialyzers hence all the possible human error is avoided. If any problem is there with dialyzers machine will notify by audible voice acknowledgment and Multicolour LED status indication on front panel.

AUTOMATIC DIALYZER REPROCESSOR MACHINE

REALIBILITY

Innovative hydraulics design with not a single electro-magnetic motor inside the machine makes machine more reliable. No moving parts hence no ware-n-tear. Also, system is designed in such a way that the start of each reprocessing, system will perform System Check cycle of all the parameters before each reprocessing. Most Advanced microprocessor and precise sensors provide most accurate readings.

FUTURE READY

System has very unique features like:

- In-built report generation to generate all vital reports directly from system
- User can generate month wise reprocessing record with dialyzer consumption
- Voice Assistance throughout the process
- Easy data transfer via USB port & front panel
- System has dedicated programs for the machine Rinse and Sanitization, Hypo Disinfection cycles
- Label printer and can be directly connected to machine, can be used to print patient data on a shelf adhesive label
- Precleaning option for effective cleaning of clotted dialyzer
- It has all the safety and compliance which industry expect from the product like ROXYPLUS

TECHNICAL SPECIFICATION

MODEL :

Single Station (ADR – 1) & Double Station (ADR – 2)

TECHNICAL PARAMETERS :

Dimensions	:	13” (L) x 16” (W) x 25” (H)	Weight	:	Approx. 30KG
Voltage	:	110V / 230V AC – 50/60Hz – 2.5A,	Power	:	150W
Ambient Temp:		10° to 45° C	Realtive Humidity	:	10% to 80% RH

DISPLAY

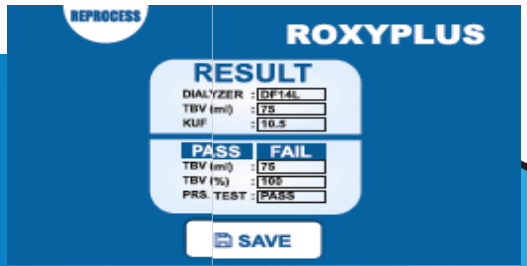
Bigger 7”,65K Color TFT LCD Display with Capacitive Touchscreen and Multi-touch Support. Simple yet distinctive Graphical User interface.

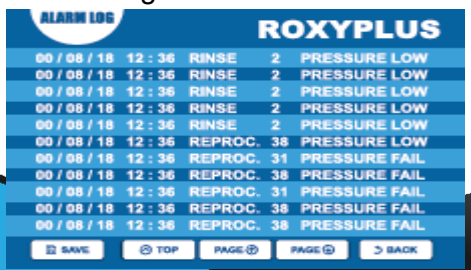
HARDWARE

- 1.4 GHz ARM Processor with 1 GB of RAM and 32 GB of storage
- Built-in Bluetooth and Wi-Fi Functionality
- USB and ethernet for file sharing
- Status Notification Indicator LED
- Voice assistance throughout the operation machine
- Inbuilt Reports Generation
- EMI and RF Filter with Twin Fuse for extra safety

BASIC REQUIREMENT

Water Requirement:	Pressure	:	20 PSI to 50 PSI
	Purity	:	RO water as per AAMI and ANSI standard
	Flow (min)	:	3L/Min.
	Water Consumption	:	Approx. 14L for 1 dialyzer
Drainage	:	Capacity to Drain 6L/ min, should not located 1 feet above the ground	





REVERSE OSMOSIS SYSTEM

PURECARE

- User Friendly with Complete Alarm Indication
- Low Maintenance Cost
- Compact Equipment
- Great Water Filtration System

PURECARE Reverse Osmosis System is a system that produces high quality ultrapure water for hemodialysis use that complied with AAMI standards. This system is designed to meet all current standards for hemodialysis water quality.

REVERSE OSMOSIS SYSTEM

FEATURES AND BENEFITS

- Consistent and safe
- Complies with AAMI water quality
- 7 alarm indicators
- Touch Screen Control Panel
- 316 grade stainless steel frame
- Easy operating system
- Automatic on and shutdown timer

PRESSURE GAUGES

- Raw Water - Guard Filter In
- Sediment - Guard Filter Out
- Carbon 1 - System In
- Carbon 2 - System Out
- Softener - Permeate

TECHNICAL SPECIFICATIONS

MODEL	600LPH	900LPH	1200LPH	1500LPH	1800LPH
PRODUCT WATER	10 L/min	15L/min	20L/min	25L/min	30L/min
REJECT WATER	10 L/min	15L/min	20L/min	25L/min	30L/min
MEMBRANE	2 Units	3 Units	4 Units	5 Units	6 Units
FLOW METER	Product/Reject Meter Digital				
INPUT VOLTAGE	Single Phase 220V – 240V (50Hz)				
CURRENT (AMPS)	16.0A				
HP PUMP	1x220-230 / 240V 4.0Kw 50Hz				
DIGITAL AUTOTROL	273/278				
RO MEMBRANE	Filmtec				
MEMBRANE SIZE	4040				

MEDICAL EQUIPMENT

DIALYSIS CHAIR

- Dialysis is a method of removing excess waste products and water from the blood. It is very time consuming and usually takes several hours.
- This model has been specifically designed for dialysis patients also can use for oncology treatment and blood donation.
- This dialysis chair features a foot board for relaxation (calf cramps) and increased upholstery thickness, thereby improving patient comfort as well.

PRODUCT FEATURES

- Padded seat, reclining backrest, armrests and footrest control by hand lever located at side of the chair.
- Four swivel casters with lock, facilitate transport and maneuverability in confined areas.
- Available in variety of colors.
- Keeps patients comfortable and may better accommodate
- patients who may faint during a procedure.

SPECIFICATION

Model	Material	Dimension (LxWxH)
DC 13	PU Polyurethane Leather / PVC Upholstery	860mm x 960mm x 1005mm

MEDICAL EQUIPMENT

VITAL SIGN MONITOR
COMPLETE WITH STAND

PORTABLE SUCTION PUMP

ELECTROCARDIOGRAPHY MACHINE
(ECG) 3 CHANNEL & 12 CHANNEL

AED CARDIOLIFE

AED HEARTSINE
SAMARITAN

EMERGENCY CART

DIALYZER RACK

OXYGEN CYLINDER
WITH REGULATOR

MEDICAL EQUIPMENT

WARD SCREEN 4 PANEL

DIGITAL CHAIR WEIGHING SCALES

DRESSING TABLE STAINLESS STEEL

TROLLEY PLANE

EXAMINATION COUCH WITH STEPSTOOL

CARDIAC TABLE
Type: Standard

OTHER DIALYSIS PRODUCT

BRINE TANK

BACTERIA FILTER

ACTIVATED CARBON FILTER

PP MELT BLOWN CARTRIDGE FILTER
PORE : 1 MICRON

TEST STRIP

OUR CERTIFICATE

No. Siri: 014658
Serial No.:

ASAL
ORIGINAL

PIHAK BERKUASA PERANTI PERUBATAN

MEDICAL DEVICE AUTHORITY

PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY
AKTA PERANTI PERUBATAN 2012 (AKTA 737)
MEDICAL DEVICE ACT 2012 (ACT 737)
LESEN ESTABLISMEN
ESTABLISHMENT LICENCE
Seksyen 24(1) Akta 737
Section 24(1) of Act 737

No. Lesen / Licence No. : **MDA-B611-W125**
Tarikh Sah Lesen / Licence Validity Date : **28/05/2025 - 27/05/2028**

Lesen adalah dengan ini diberi kepada: / *Licence is hereby granted to:* **VISION BIOMED SERVICES SDN.BHD**

yang beralamat di: / *of:* **NO 85 JALAN PENGKALAN INDAH 2 BANDAR PENGKALAN INDAH 31650 IPOH PERAK**

Sebagai: / *as:* **WAKIL DIBERI KUASA** / ***AUTHORIZED REPRESENTATIVE***

Orang yang bertanggungjawab: / *Person Responsible:* **NARENDRAN A/L BALAKRISHNAN(I/C NO.: 750911-08-6339)**

Lesen ini diberikan tertakluk kepada peruntukan-peruntukan di bawah Akta 737 dan peraturan-peraturan dibawahnya serta syarat-syarat seperti di Lampiran 1.
This licence is granted subject to the provisions under Act 737 and its subsidiary legislations and the conditions as in Attachment 1.

MUNALITHARAN PARAMASUA
KETUA EKSEKUTIF
CHIEF EXECUTIVE
PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY

ASAL
ORIGINAL

PIHAK BERKUASA PERANTI PERUBATAN

MEDICAL DEVICE AUTHORITY

PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY
AKTA PERANTI PERUBATAN 2012 (AKTA 737)
MEDICAL DEVICE ACT 2012 (ACT 737)
LESEN ESTABLISMEN
ESTABLISHMENT LICENCE
Seksyen 24(1) Akta 737
Section 24(1) of Act 737

No. Lesen / Licence No. : **MDA-5771-K124**
Tarikh Sah Lesen / Licence Validity Date : **06/04/2024 - 05/04/2027**

Lesen adalah dengan ini diberi kepada: / *Licence is hereby granted to:* **VISION BIOMED SERVICES SDN.BHD**

yang beralamat di: / *of:* **NO.85, JALAN PENGKALAN INDAH 2, BANDAR PENGKALAN INDAH, 31650 IPOH PERAK**

Sebagai: / *as:* **PEMBUAT** / ***MANUFACTURER***

Orang yang bertanggungjawab: / *Person Responsible:* **NARENDRAN A/L BALAKRISHNAN(I/C NO.: 750911-08-6339)**

Lesen ini diberikan tertakluk kepada peruntukan-peruntukan di bawah Akta 737 dan peraturan-peraturan dibawahnya serta syarat-syarat seperti di Lampiran 1.
This licence is granted subject to the provisions under Act 737 and its subsidiary legislations and the conditions as in Attachment 1.

MUNALITHARAN PARAMASUA
KETUA EKSEKUTIF
CHIEF EXECUTIVE
PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY

ASAL
ORIGINAL

PIHAK BERKUASA PERANTI PERUBATAN

MEDICAL DEVICE AUTHORITY

PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY
AKTA PERANTI PERUBATAN 2012 (AKTA 737)
MEDICAL DEVICE ACT 2012 (ACT 737)
LESEN ESTABLISMEN
ESTABLISHMENT LICENCE
Seksyen 24(1) Akta 737
Section 24(1) of Act 737

No. Lesen / Licence No. : **MDA-5697-D124**
Tarikh Sah Lesen / Licence Validity Date : **06/04/2024 - 05/04/2027**

Lesen adalah dengan ini diberi kepada: / *Licence is hereby granted to:* **VISION BIOMED SERVICES SDN.BHD**

yang beralamat di: / *of:* **NO.85, JALAN PENGKALAN INDAH 2, BANDAR PENGKALAN INDAH, 31650 IPOH PERAK**

Sebagai: / *as:* **PENGEDAR** / ***DISTRIBUTOR***

Orang yang bertanggungjawab: / *Person Responsible:* **NARENDRAN A/L BALAKRISHNAN(I/C NO.: 750911-08-6339)**

Lesen ini diberikan tertakluk kepada peruntukan-peruntukan di bawah Akta 737 dan peraturan-peraturan dibawahnya serta syarat-syarat seperti di Lampiran 1.
This licence is granted subject to the provisions under Act 737 and its subsidiary legislations and the conditions as in Attachment 1.

MUNALITHARAN PARAMASUA
KETUA EKSEKUTIF
CHIEF EXECUTIVE
PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY

ASAL
ORIGINAL

PIHAK BERKUASA PERANTI PERUBATAN

MEDICAL DEVICE AUTHORITY

PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY
AKTA PERANTI PERUBATAN 2012 (AKTA 737)
MEDICAL DEVICE ACT 2012 (ACT 737)
SIJIL PENDAFTARAN PERANTI PERUBATAN
MEDICAL DEVICE REGISTRATION CERTIFICATE
Seksyen 5(1) Akta 737
Section 5(1) of Act 737

No. Pendaftaran / Registration No.: **GC3506519-36236**
Tarikh Sah Pendaftaran / Registration Validity Date: **10/11/2024 - 18/11/2029**

Sijil ini adalah dengan ini diberi kepada: / *This certificate is hereby issued to:* **VISION BIOMED SERVICES SDN.BHD**

yang beralamat di: / *which is located at:* **NO 85 JALAN PENGKALAN INDAH 2 BANDAR PENGKALAN INDAH, 31650 IPOH PERAK DARUL RIDZUAN**

bagi mengesahkan peranti perubatan seperti yang dinyatakan dalam Lampiran 1 adalah berdaftar di bawah Seksyen 5(1) Akta 737.
to confirm that the medical device as detailed out in Attachment 1 is registered under Section 5(1) of Act 737.

Pendaftaran ini diberikan tertakluk kepada peruntukan-peruntukan di bawah Akta 737 dan peraturan-peraturan yang dibuat dibawahnya serta syarat-syarat seperti di Lampiran 2.
This registration is granted subject to the provisions under Act 737 and its subsidiary legislations and the conditions as in Attachment 2.

MUNALITHARAN PARAMASUA
KETUA EKSEKUTIF
CHIEF EXECUTIVE
PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY

ASAL
ORIGINAL

PIHAK BERKUASA PERANTI PERUBATAN

MEDICAL DEVICE AUTHORITY

PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY
AKTA PERANTI PERUBATAN 2012 (AKTA 737)
MEDICAL DEVICE ACT 2012 (ACT 737)
SIJIL PENDAFTARAN PERANTI PERUBATAN
MEDICAL DEVICE REGISTRATION CERTIFICATE
Seksyen 5(1) Akta 737
Section 5(1) of Act 737

No. Pendaftaran / Registration No.: **GB5312319-36237**
Tarikh Sah Pendaftaran / Registration Validity Date: **01/10/2024 - 30/09/2029**

Sijil ini adalah dengan ini diberi kepada: / *This certificate is hereby issued to:* **VISION BIOMED SERVICES SDN.BHD**

yang beralamat di: / *which is located at:* **NO 85 JALAN PENGKALAN INDAH 2 BANDAR PENGKALAN INDAH, 31650 IPOH PERAK DARUL RIDZUAN**

bagi mengesahkan peranti perubatan seperti yang dinyatakan dalam Lampiran 1 adalah berdaftar di bawah Seksyen 5(1) Akta 737.
to confirm that the medical device as detailed out in Attachment 1 is registered under Section 5(1) of Act 737.

Pendaftaran ini diberikan tertakluk kepada peruntukan-peruntukan di bawah Akta 737 dan peraturan-peraturan yang dibuat dibawahnya serta syarat-syarat seperti di Lampiran 2.
This registration is granted subject to the provisions under Act 737 and its subsidiary legislations and the conditions as in Attachment 2.

MUNALITHARAN PARAMASUA
KETUA EKSEKUTIF
CHIEF EXECUTIVE
PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY

ASAL
ORIGINAL

PIHAK BERKUASA PERANTI PERUBATAN

MEDICAL DEVICE AUTHORITY

PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY
AKTA PERANTI PERUBATAN 2012 (AKTA 737)
MEDICAL DEVICE ACT 2012 (ACT 737)
SIJIL PENDAFTARAN PERANTI PERUBATAN
MEDICAL DEVICE REGISTRATION CERTIFICATE
Seksyen 5(1) Akta 737
Section 5(1) of Act 737

No. Pendaftaran / Registration No.: **GC8658524-181933**
Tarikh Sah Pendaftaran / Registration Validity Date: **02/09/2024 - 01/09/2029**

Sijil ini adalah dengan ini diberi kepada: / *This certificate is hereby issued to:* **VISION BIOMED SERVICES SDN.BHD**

yang beralamat di: / *which is located at:* **NO 85 JALAN PENGKALAN INDAH 2 BANDAR PENGKALAN INDAH, 31650 IPOH PERAK DARUL RIDZUAN**

bagi mengesahkan peranti perubatan seperti yang dinyatakan dalam Lampiran 1 adalah berdaftar di bawah Seksyen 5(1) Akta 737.
to confirm that the medical device as detailed out in Attachment 1 is registered under Section 5(1) of Act 737.

Pendaftaran ini diberikan tertakluk kepada peruntukan-peruntukan di bawah Akta 737 dan peraturan-peraturan yang dibuat dibawahnya serta syarat-syarat seperti di Lampiran 2.
This registration is granted subject to the provisions under Act 737 and its subsidiary legislations and the conditions as in Attachment 2.

MUNALITHARAN PARAMASUA
KETUA EKSEKUTIF
CHIEF EXECUTIVE
PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY

OUR CERTIFICATE

ASAL
ORIGINAL

PIHAK BERKUASA PERANTI PERUBATAN — MEDICAL DEVICE AUTHORITY

PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY
AKTA PERANTI PERUBATAN 2012 (AKTA 737)
MEDICAL DEVICE ACT 2012 (ACT 737)
SIJIL PENDAFTARAN PERANTI PERUBATAN
MEDICAL DEVICE REGISTRATION CERTIFICATE
Seksyen 5(1) Akta 737
Section 5(1) of Act 737

No. Pendaftaran: **GC6743724-162522** *Registration No.:*
Tarikh Sah Pendaftaran: **14/02/2024 - 13/02/2029** *Registration Validity Date:*

Sijil ini adalah dengan ini diberi kepada: **VISION BIOMED SERVICES SDN.BHD**
This certificate is hereby issued to:

yang beralamat di: **NO.85, JALAN PENGKALAN INDAH 2, BANDAR PENGKALAN INDAH, , 31650 IPOH PERAK DARUL RIDZUAN**
which is located at:

bagi mengesahkan peranti perubatan seperti yang dinyatakan dalam Lampiran 1 adalah berdaftar di bawah Seksyen 5(1) Akta 737.
to confirm that the medical device as detailed out in Attachment 1 is registered under Section 5(1) of Act 737.

Pendaftaran ini diberikan tertakluk kepada peruntukan-peruntukan di bawah Akta 737 dan peraturan-peraturan yang dibuat dibawahnya serta syarat-syarat seperti di Lampiran 2.
This registration is granted subject to the provisions under Act 737 and its subsidiary legislations and the conditions as in Attachment 2.

MUHALITHARAN PARAMASUA
KETUA EKSEKUTIF
CHIEF EXECUTIVE
PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY

No. Siri: **045386** *Serial No.:*

ASAL
ORIGINAL

PIHAK BERKUASA PERANTI PERUBATAN — MEDICAL DEVICE AUTHORITY

PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY
AKTA PERANTI PERUBATAN 2012 (AKTA 737)
MEDICAL DEVICE ACT 2012 (ACT 737)
SIJIL PENDAFTARAN PERANTI PERUBATAN
MEDICAL DEVICE REGISTRATION CERTIFICATE
Seksyen 5(1) Akta 737
Section 5(1) of Act 737

No. Pendaftaran: **GC9752722-84818** *Registration No.:*
Tarikh Sah Pendaftaran: **27/01/2022 - 26/01/2027** *Registration Validity Date:*

Sijil ini adalah dengan ini diberi kepada: **VISION BIOMED SERVICES SDN.BHD**
This certificate is hereby issued to:

yang beralamat di: **NO.85, JALAN PENGKALAN INDAH 2, BANDAR PENGKALAN INDAH, , 31650 IPOH PERAK DARUL RIDZUAN**
which is located at:

bagi mengesahkan peranti perubatan seperti yang dinyatakan dalam Lampiran 1 adalah berdaftar di bawah Seksyen 5(1) Akta 737.
to confirm that the medical device as detailed out in Attachment 1 is registered under Section 5(1) of Act 737.

Pendaftaran ini diberikan tertakluk kepada peruntukan-peruntukan di bawah Akta 737 dan peraturan-peraturan yang dibuat dibawahnya serta syarat-syarat seperti di Lampiran 2.
This registration is granted subject to the provisions under Act 737 and its subsidiary legislations and the conditions as in Attachment 2.

AHMAD SHARIFF BIN HAMBALI
KETUA EKSEKUTIF
CHIEF EXECUTIVE
PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY

ASAL
ORIGINAL

PIHAK BERKUASA PERANTI PERUBATAN — MEDICAL DEVICE AUTHORITY

PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY
AKTA PERANTI PERUBATAN 2012 (AKTA 737)
MEDICAL DEVICE ACT 2012 (ACT 737)
SIJIL PENDAFTARAN PERANTI PERUBATAN
MEDICAL DEVICE REGISTRATION CERTIFICATE
Seksyen 5(1) Akta 737
Section 5(1) of Act 737

No. Pendaftaran: **GB4794825-194066** *Registration No.:*
Tarikh Sah Pendaftaran: **11/01/2025 - 10/01/2030** *Registration Validity Date:*

Sijil ini adalah dengan ini diberi kepada: **VISION BIOMED SERVICES SDN.BHD**
This certificate is hereby issued to:

yang beralamat di: **NO 85 JALAN PENGKALAN INDAH 2 BANDAR PENGKALAN INDAH, 31650 IPOH PERAK DARUL RIDZUAN**
which is located at:

Peranan establismen: **Wakil Diberi Kuasa**
Role of establishment: ***Authorized Representative***

bagi mengesahkan peranti perubatan seperti yang dinyatakan dalam Lampiran 1 adalah berdaftar di bawah Seksyen 5(1) Akta 737.
to confirm that the medical device as detailed out in Attachment 1 is registered under Section 5(1) of Act 737.

Pendaftaran ini diberikan tertakluk kepada peruntukan-peruntukan di bawah Akta 737 dan peraturan-peraturan yang dibuat dibawahnya serta syarat-syarat seperti di Lampiran 2.
This registration is granted subject to the provisions under Act 737 and its subsidiary legislations and the conditions as in Attachment 2.

MUHALITHARAN PARAMASUA
KETUA EKSEKUTIF
CHIEF EXECUTIVE
PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY

TÜVNORD

Certificate

Management system as per

Good Distribution Practice for Medical Devices (GDPMD)

MDA/RR No.1, November 2015 pursuant to Appendix 4, Third Schedule of Medical Device Regulation 2012
This certification was conducted in accordance with the TUV NORD (MALAYSIA) SDN BHD auditing and certification procedures and is subject to regular surveillance audits

Vision Biomed Services Sdn.Bhd
No 85, Jalan Pengkalan Indah 2,
Bandar Pengkalan Indah, 31650, Ipoh, Perak,
Malaysia

applies a management system in line with the above standard for the following scope

Scope
Authorized Representative, Import, Storage, Handling and Distribution (including Transportation) of Disinfectant, Single Use, and Active Devices for Dialysis.

Special Storage and Handling condition : None
Outsource activities : Transportation/Courier, Pest Control

Certificate Registration No. 222 16001
Valid from 2025-03-11
Valid until 2028-03-10
Initial certification 2016

2025-03-09
Issuance Date
Ir. Bill Kong, Managing Director
Certification Body at TUV NORD (MALAYSIA) SDN BHD

TUV NORD (MALAYSIA) SDN BHD
No. 9F-1A, 9th Floor, Tower 2 @ PFCC, Jalan Puteri 1/2, Bandar Puteri Puchong, 47100 Puchong, Selangor, Malaysia

GDPMD — Medical Device AUTHORITY MALAYSIA

TÜVNORDGROUP

TÜVNORD

Certificate

Management system as per

ISO 13485 : 2016 / MS ISO 13485 : 2017

In accordance with Appendix 4, Third Schedule of Malaysia Medical Devices Regulations 2012 and in accordance with the TUV NORD (MALAYSIA) SDN BHD auditing and certification procedure, it is hereby certified that

VISION BIOMED SERVICES SDN BHD (874930-A)
No. 85, Jalan Pengkalan Indah 2,
Bandar Pengkalan Indah,
31650 Ipoh, Perak,
Malaysia

applies a management system in line with the above standard for the following scope

Scope

Legal Manufacturer, Sales, Distribution and Storage of Automatic Dialyzer Reprocessor, Hemodialysis Bicarbonate Bag, Water Test Strips, Bacteria Filter, Carbon Filter, Guard Filter and Vacuum Salt

Manufacture of Reverse Osmosis System

Installation, Testing and Commissioning, Servicing of Automatic Dialyzer Reprocessor and Reverse Osmosis System

Certificate Registration No. 221 20001
Valid from 2023-04-06
Valid until 2026-04-05
Initial certification 2020

2024-03-14
Issuance Date
Ir. Bill Kong, General Manager
Certification Body at TUV NORD (MALAYSIA) SDN BHD

TUV NORD (MALAYSIA) SDN BHD
MDA Registration No: MDA/CAB - 008
No. 9F-1A, 9th Floor, Tower 2 @ PFCC
Jalan Puteri 1/2,
Bandar Puteri Puchong
47100 Puchong, Selangor, Malaysia

MS ISO13485 — Medical Device AUTHORITY MALAYSIA

TÜVNORDGROUP

CONTACT US

VISION BIOMED SERVICES SDN BHD

HQ Office :
Unit A-3A-15, Kompleks Perindustrian Em Hub, Persiaran Surian,
Seksyen 3, Taman Sains Selangor Kota Damansara, 47810 Petaling Jaya Selangor.
Tel : 03-8679 7679

Warehouse :
No.85 Jalan Pengkalan Indah 2, Bandar Pengkalan Indah, 31650 Ipoh, Perak.
Tel : 05-321 9422

visionbiomedservices@gmail.com
Website : www.visionbiomed.com.my

GDPMD - 222 16001
MDA - 8611-W125
MDA - 5771-K124
MDA - 5697-D124
ISO 13485 : 2016 / MS ISO 13485 : 2017 NO : 221 20001