

NEW POSSIBILITIES

SIMPLE DELIVERY

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Baxter

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The **Artis Physio Plus** multi-therapy system is Baxter's newest innovation for effective delivery of hemodialysis therapy.

ARTIS PHYSIO PLUS Main performances

TREATMENT MODALITIES

HD double needle (DN) and single needle double pump (SN DP) HDF/HF predilution, HDF postdilution, **UltraControl** HDF modality HDx expanded hemodialysis with the **Theranova*** dialyzer **BiCart Select** system and **BiCart Select Citrate** system **HemoControl** modality: automatic sodium and UF profiling Flexible isolated UF

TREATMENT MONITORING

Programmable patient card for treatment prescription BPM: Blood pressure monitoring Diascan monitoring: measurement of ionic clearance and Kt/V Hemoscan monitoring: measurement of the blood volume variation Smartscan: programmable alert system Dialysis reporting page: treatment summary and event log

ARTIS PHYSIO PLUS Technical specifications

DIMENSIONS AND WEIGHT

| DIMENSIONS AND WEIGHT | |
|---------------------------------------|--|
| Width | 660 mm |
| Depth | 700 mm |
| Height | 1550 mm (without IV pole) |
| Weight | < 135 kg |
| Operating environment | |
| Ambient temperature | +18 to +35°C |
| Relative humidity | 30 to 85% rh. |
| Ambient pressure | 795 to 1060 HPa |
| BLOOD MODULE | |
| Blood flow control | |
| Actual blood flow rate HD DN | 10 to 500 mL/min |
| Mean blood flow rate SN DP | 20 to 270 mL/min |
| Stroke volume SN DP | 25 to 60 mL |
| Blood circuit pressure supervision | |
| Arterial pressure | -300 to +150 mmHg |
| Venous pressure | +10 to +400 mmHg |
| Single needle pressure (SNDP) | +40 to +450 mmHg |
| Air detection | Ultrasonic detector |
| Heparin syringe pump | |
| 1.5 to 10 ml/h (Syringe 30 ml) or 0.5 | ; to 4.0 ml/h (Syringe 10 ml) s, manual bolus and programmable stop time |
| Blood volume monitoring | s, mandat botas and programmable stop time |
| Relative blood volume | -40% to +10% |
| | 40/0 00 110/0 |
| FLUID MODULE | |
| Water supply | |
| Inlet pressure | 150 to 800 kPa |
| Inlet temperature | +5 to +32°C (230V) or +10 to +32°C (115V) |
| Inlet water quality | Must comply with local standards and ISO 13959 standard |
| | |

SYSTEM INTEGRATION

ArtiSet blood tubing system with auto-loading procedure One-button auto-priming procedure Evaclean integrated draining of priming fluid Automated priming, patient connection and rinseback Ultra Prime function for on-line priming, bolus and rinseback

Navpad user interface navigation system

Programmable user timer

Open connection to IT network for clinical data management Integrated service mode

Dialysis fluid preparation and monitoring

| | 5 | |
|---|--|--|
| Flow rate (HD treatment) | 300-800 mL/min (by step of 50 mL/min) | |
| Temperature | +34.0 to +39.5°C | |
| Sodium range | 130 to 160 mmol/L, | |
| Bicarbonate range | 24 to 38 mmol/L | |
| Dry bicarbonate | BiCart cartridge | |
| pH (optional) | 1.0 to 13.0 pH units | |
| Substitution fluid (on-line HDF/HF) | 1.2 to 27 L/h (maximum 150 L) | |
| Concentrate low consumption mode Connection to central concentrate supply (optional) | | |
| Ultrafiltration control | | |
| UF volume | 0 to 24 L, in steps of 0.05L | |
| UF rate | 0 to 3 L/h | |
| Blood leak detection | Infrared light | |
| Disinfection and rinse | | |
| Heat, and chemical disinfection Integrated heat mode with Baxter CWP Heat with CleanCart A or C cartridge (disinfection + deproteneization or decalcification) Chemical Stand-by mode Weekly disinfection program with auto-On/Off Disinfection process history U9000 ultrafilter management | | |
| Power supply | | |
| Voltage | 230/240 VAC (± 10 %) or 115 VAC (± 10 %) | |
| Frequency | 50/60 Hz (± 5 Hz) | |
| | | |

| Ev | tennel commontivity | |
|----|----------------------------|--|
| Ba | attery backup power supply | Up to 15 min |
| Po | wer consumption | Max. 10 A at 230/240 VAC or Max. 16 A at 115 VAC |
| FI | equency | JU/OU HZ (± J HZ) |

External connectivity

Ethernet port 10/100 base T

IT connectivity with HL7 protocol

USB port for Baxter Service Operations

The products meet the applicable provisions of Annex I (Essential Requirements) and Annex II (Full quality assurance system of the Council Directive 93/42/EEC of 14 June 1993, amended by Directive 2007/47/EC)

For safe and proper use of the device mentioned herein, please see the Operators Manual

* Do not use Theranova dialyzers in HDF or HF mode

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