

theranova⁵⁰⁰

HDx THERAPY, ENABLED BY THE THERANOVA DIALYZER

The THERANOVA dialyzer, featuring an innovative membrane, effectively targets large middle molecules not efficiently removed by currently available dialysis treatments. It provides the opportunity for an expanded hemodialysis therapy, HDx, providing HDF performance and beyond in the removal of middle and larger middle molecules, using regular HD infrastructure.

HDF PERFORMANCE AND BEYOND, AS SIMPLE AS HD

- Markedly greater clearances and intradialytic reduction ratios than regular HD – at ordinary blood flow rates¹
- Equivalent removal of small and conventional middle molecules to high-volume HDF – Greater removal possible for larger middle molecules²
- Albumin removal limited to between 1 and 4 grams^{1,2}
- Compatible with any HD monitor³ and with standard-quality dialysis fluid quality^{4,5}

ACHIEVED THROUGH MEMBRANE INNOVATION

- Higher permeability^{6,7}
- Enhanced selectivity by size exclusion^{6,7}
- A step closer to the natural kidney^{6,7}



TYPICAL PATIENT PROFILE: PATIENTS REQUIRING HIGHER CLEARANCES OF LARGER UREMIC TOXINS, WITHOUT ACCESS TO HDF

The theranova⁵⁰⁰ Dialyzer

INDICATIONS FOR USE	
Indications For Use	THERANOVA dialyzers are indicated for treatment of chronic and acute renal failure by hemodialysis. Do not use for hemodiafiltration, hemofiltration due to higher permeability of larger molecular weight proteins such as albumin

COMPONENTS	MATERIALS	
Membrane	Polyarylethersulfone / Polyvinylpyrrolidone	PAES / PVP – BPA-free
Potting	Polyurethane	PUR
Housing, Header	Polycarbonate	PC
Gasket	Silicon rubber	SIR
Protection Cap	Polypropylene	PP

MEMBRANE	
Membrane design	Asymmetric wall, 3-layer finger structure Medium Cut-Off, narrow pore size distribution For the safe and proper use of device, please refer the Instructions for Use
Effective Membrane Area [m ²]	2.0
Fiber Dimension	
– Inner diameter [µm]	180
– Wall thickness [µm]	35
Sterilizing Agent	Steam
Sterile Barrier	Medical Grade Paper

BLOOD COMPARTMENT	
Blood Compartment Volume [ml]	105
Residual Blood Volume [ml]	<1

DIALYSIS FLUID QUALITY REQUIREMENTS ^{4,5}	
Minimum Requirements	Standard Dialysis Fluid Quality ISO 11663:2014 or ANSI/AAMI RD62 standard

PERFORMANCES*	
UF-coefficient [ml/(h·mmHg)]	59
Pressure Drop – Blood Compartment [mmHg]	
Qb=200	≤80
Qb=300	≤120
Qb=400	≤160
Qb=500	≤200
Qb=600	≤240
Pressure Drop – Dialysate Compartment [mmHg]	
Qd=300	≤15
Qd=500	≤25
Qd=800	≤40

LIMITS FOR USE	
Maximum TMP [mmHg]	600
Operating blood flow range [ml/min]	200-600
Operating dialysate flow range [ml/min]	300-800

STORAGE CONDITIONS	
Storage conditions	<30 °C; <86 °F

IN-VITRO CLEARANCES (at UF = 0 ml/min)

	Qb / Qd	ml/min
Urea (60 Da)	200/500	199
	300/500	285
	400/500	351
	400/800	381
	500/800	454
Phosphate (95 Da)	200/500	194
	300/500	267
	400/500	320
	400/800	354
Creatinine (113 Da)	200/500	196
	300/500	274
	400/500	331
	400/800	365
Vitamin B12 (1.4 kDa)	500/800	428
	200/500	169
	300/500	215
	400/500	249
	400/800	280
	500/800	317

	Qb / Qd	ml/min
Inulin (5.2 kDa)	200/500	139
	300/500	170
	400/500	193
	400/800	216
	500/800	241
	200/500	128
Cytochrome C (12 kDa)	300/500	155
	400/500	175
	400/800	196
	500/800	217
Myoglobin (17 kDa)	200/500	110
	300/500	130
	400/500	147
	400/800	163
	500/800	180

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, please refer to the Instructions for Use

CE 0086

* According to ISO 8637:

UF-coefficient: measured with bovine blood, Hct 32%, Pct 60g/l, 37°C

Pressure drop blood: measured with bovine blood, Hct 32%, Pct 60g/l, 37°C, UF = 0 ml/min

Pressure drop dialysate: measured with dialysate

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- Baxter. Data on file. *Theranova Limited Controlled Distribution Report 2016*
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- Boschetti-de-Fierro A, et al. *MCO membranes: Enhanced Selectivity in High-Flux Class*. Scientific Reports (2015); 5: 18448
- Krause B, et al. *Highly selective membranes for blood purification*. Euromembrane Congress 2015, Abstract E139

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