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ฉลากถุงยาที่ขอแก้ไขเปลี่ยนแปลง ตำรับยาชื่อ Prismasol B0 เลขทะเบียนที่ 2C 15083/61 (N)

> ขอรับรองว่าข้อความบนฉลากของ ภาชนะบรรจุทุกชนิดและหีบห่อ ถูกต้องตรงตามมาตรา ๒๕ (๓) หรือ ๒๗ (๒) และ (๓) แล้วแต่กรณี



PRISMASOL BO

Solution for haemofiltration and haemodialysis. For intravenous use and/or continuous haemodialysis. Potassium free.





Each bag contains			Before rec	After	After reconstitution	
		Α	В		A+B	
Calcium chloride dihydrate		1.29 g		0.258 g/L		
Magnesium chloride hexahydrate		0.51 g		0.102 g/L		
(S)-Lactic acid		1.35 g	0.270 g/L		0.270 g/L	
Sodium chloride			30.64 g	6.13 g/L		
Sodium bicarbonate			14.68 g	2.94 g/L		
Excipients: Water for injections		QS 250 mL	QS 4750 mL	QS 5000 mL		
Carbon dioxid	e for pH adjustme	ent				
After recons	stitution, A + B					
	Na ⁺	Ca ²⁺	Mg ²⁺	CI-	HCO ₃ -	C ₃ H ₅ O ₃ -
mmol/L	140	1.75	0.5	109.5	32	3
mEq/L	140	3.5	1	109.5	32	3
Theoretical osmolality: 287 mOsm/l						
ISOTONIC						

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Keep out of the sight and reach of children. Dosage as directed by a physician. Use only with appropriate extra-renal replacement equipment. Not for direct infusion: mix both compartments before use. Mix additives before connecting the bag to the extracorporeal circuit. Use only if solution is clear and free from visible particles. Sterile and free from bacterial endotoxins. Check for leaks. Do not use with a haemodialysis monitor. Use in one patient on one occasion only. Contains no antimicrobial preservative. Any unused solution must be discarded. See package leaflet for further instructions and shelf life of reconstituted product. Batch No.: see "lot" printed on the back of the bag. ยาลิ้นอายุ: see "exp" printed on the back of the bag. Manufacturing date: see "manuf." printed on the back of the bag. Store below 30°C. Do not refrigerate. Do not freeze.

5000 mL

Product No.: 955679



TH Reg. no: 2C 15083/61 (N)

Imported by: Baxter Healthcare (Thailand) Co., Ltd., Bangkok

Manufactured by: Bieffe Medital S.p.A., Via Stelvio, 94-23035 Sondalo (So), Italy