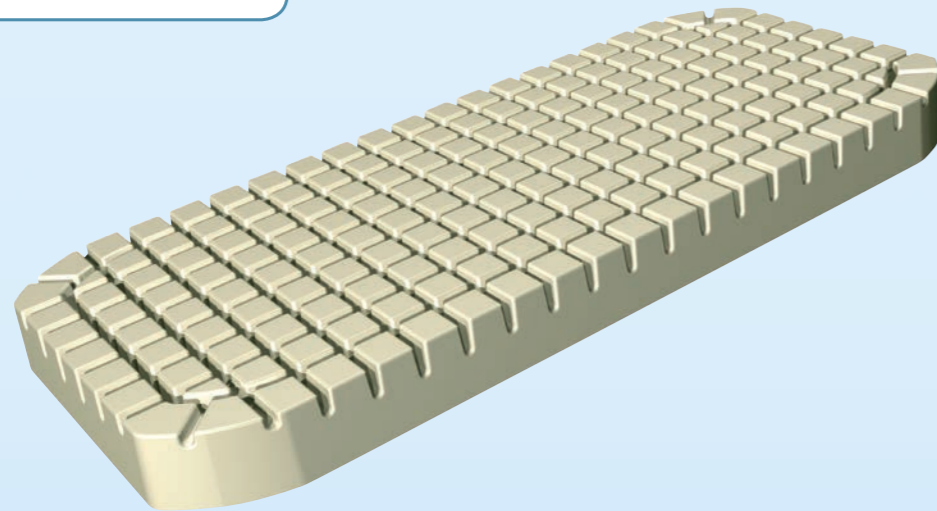
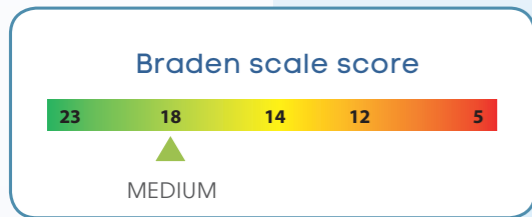
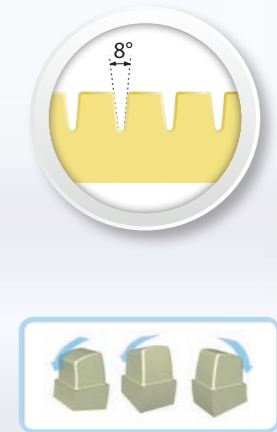


APLOT®



KEY FEATURES

- ▶ High Resilient foam (HR)
- ▶ Density: $\geq 34 \text{ Kg/m}^3$
- ▶ Hardness: 4.5 kPa
- ▶ Dimensions: 195 x 85 x 14 cm (76.7 x 33.4 x 5.5 inches)
- ▶ Mattress weight: 8.5 Kg
- ▶ Draw sheet and covers available: Dermalon draw sheet, Promust PU and Promust PU HD covers



KEY BENEFITS

- ▶ The castellated foam allows reducing shearing forces applied on the skin and underlying tissue.
- ▶ The mobility of castellated foam allows relieving pressure points to enhance blood circulation thus reducing pressure ulcers risk.

INDICATIONS

HELP IN PREVENTING PRESSURE ULCERS FOR PATIENT AT LOW TO MEDIUM RISK OF PRESSURE ULCER DEVELOPMENT, BEDRIDDEN BETWEEN 10 AND 15 HOURS A DAY.

HELP IN TREATING EXISTING PRESSURE ULCERS FROM STADE 1 TO 2 COMBINED WITH TECHNICAL AID DEVICES.

PATIENT WEIGHT: 40 TO 120 KG/ 6.2 TO 18.8 ST

- ▶ Single-piece molded High Resilient (HR) polyurethane foam mattress with 4 cut corners
- ▶ Castellated foam to prevent from shearing forces
- ▶ Removable and washable integral cover

DESCRIPTION	ITEM REFERENCES ACCORDING TO PACKAGING			FIRE STANDARDS
	CARDBOX	PALLET BOX	ROLLED AND COMPRESSED (RC)	
APLOT® Dermalon FR draw sheet	VA103MB14	VA103MB14-BOX (10 units)	VA103MB14RC1 (1 unit) VA103MB14RC (5 units)	EN 597-1 and 2 GPEM D1 90 classe D
APLOT® Promust PU cover (not installed)	VA103MB14 /HIP	VA103MB14 /HIP-BOX (10 units)	—	EN 597-1 and 2 GPEM D1 90 classe D
APLOT® Promust PU HD cover (not installed)	VA103MB14 /HIPH	VA103MB14 /HIPH-BOX (10 units)	—	

Cleaning instructions:
Dermalon FR draw sheet $\leq 1000\text{ppm}$

Cleaning instructions:
Promust PU (HD) cover $\leq 500\text{ppm}$

2 years warranty

Photos not contractually binding

FP000032/01.2018



WINCARE Group - 200 rue Charles Tellier - Actiparc de Grézan - 30034 Nîmes Cedex 1 France
Ph : +33 (0)4 66 02 15 15 - Fax : +33 (0)4 66 02 15 00 - Email : contact.askle@winnicare.fr - www.winnicare.fr

In the interest of constantly improving its products, the WINCARE Group reserves the rights to modify the technical specifications of any product without prior notice

Class I medical device in accordance with European Directive 93/42/EEC

