Item number 330225

ABRI•CELL

Underpad, ABENA Abri-Cell, 6 layers, 60x60cm, light blue

- ✓ Absorbent core with multilayer tissue
- ✓ Soft and smooth nonwoven surface
- ✓ Rapid liquid absorption
- ✓ Liquid safe back sheet and sealed edges for extra protection
- ✓ Available in different sizes
- ✓ Available with differing numbers of tissue layers

MD CE

Product Description

Abri-Cell is used as a protective disposable underpad for e.g. beds and chairs. It is suitable for non-sterile procedures, such as wound care and blood tests, but it can also be used for incontinent users. The underpad has sealed edges for optimum leakage protection, and the surface is covered with a soft nonwoven material for enhanced user comfort. The use of disposable underpads reduces the risk of contamination.

Abri-Cell is available in different sizes and with different layers of tissue.

Specifications

Base name	Underpad
Brand	ABENA
Sub-brand	Abri-Cell
Sub-color	Light blue
Features	Non-sterile
Single or multiple use	Single use
Material	Tissue, nonwoven, PE
Length/depth	60 cm
Width	60 cm
Weight, net	41 g
Tolerance	+/- 2 cm
Layer quantity	6 layers
Sterilization method	EO
Absorption Rothwell	550 ml
Certifications	CE, MD
CE Class (Medical Devices)	Class I
Product or test standards	ISO 11948-1
Directives, regulations and acts	MDR (EU) 2017/745
Because of the products composition it has an expiry date from the production date on:	5 years
Storage Instructions	Store dry, room temperature and no direct sunlight.

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Product Disposal Instructions	To be disposed of with ordinary household waste. If contaminated, dispose of as clinical waste.
Packaging Disposal Instructions	Can be recycled or incinerated.

Instructions for use/application

The underpad must be placed on the bed or chair with the blue side down. Ensure that the underpad is fully smoothed out before use.

Packaging data

Unit	Contains	Length	Width	Height	EAN
cll	300 pcs	67 cm	27 cm	42 cm	5703538453953
pck	25 pcs	27 cm	20 cm	11 cm	5703538453939

MD

Regulation (EU) 2017/745 is a regulation of the European Union on the clinical investigation and sale of medical devices for human use. It repeals Directive 93/42/EEC, which concerns medical devices, and Directive 90/385/EEC, which concerns active implantable medical devices, on 26 May 2021.

CE

The CE mark guarantees that a product is safe to use and complies with all safety precautions. CE stands for Conformité Européenne (European Conformity) and is mainly found on electronic equipment, safety equipment, construction products, and medical devices.

The product is only for single-use.