

EU Declaration of Conformity according to MDR

Document ID: PD-563411 Rev:00

Created by: Kristin Ödling State: Released

Approved by: Karin Darle Olsson **Release date:** 2022-06-17 11:43:31

Dates and times in Greenwich Mean Time, 24 hours format

Title: MDR DoC Mepilex Border Ag

We, Mölnlycke Health Care AB, Gamlestadsvägen 3C, Box 13080, SE-402 52 Göteborg, Sweden, SRN SE-MF-000014042, being exclusively responsible manufacturer for conformity of the device/s; declare that the devices listed in the attached schedule are in conformity with the provisions of the Medical Device Regulation 2017/745, concerning medical devices.

Other Union Legislation applicable: Not applicable

Trade name / Product name:

Mepilex Border Ag

Product Classification: MDR Class III

MDR Classification Rule: 4, 14

Sterility: Sterile

Measuring Function: No

This declaration is supported by a conformity assessment procedure in accordance with Annex/es: IX

Common Specification: No CS is applicable

Certificate number: MDR 722028, MDR 758927

Issued by: BSi Id No: 2797

(Notified Body Name)

NB. For non-sterile, non-measuring Class I products, no certificate is issued by a Notified Body.

Mölnlycke Health Care AB issues this declaration in recognition of applied harmonized standards.

Signed for and on behalf of Mölnlycke Health Care AB

Place of Issue: Göteborg, Sweden

Date: 2022-06-17 Function: Regulatory Affairs Excellence

Manager

Name: Karin Darle Olsson Signature:



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Generic product information		
Intended Purpose:	Mepilex Border Ag is designed for the management of medium to high exuding leg and foot ulcers, pressure ulcers, malignant wounds, partial thickness burns, traumatic and surgical wounds where a moist environment, exudate handling, gentle fixation and an antimicrobial action is indicated. Mepilex Border Ag may be used on infected wounds as part of a treatment regimen under supervision of a qualified health care professional. Mepilex Border Ag can be used under compression bandaging.	
Basic UDI-DI:	73324300000000028JW	
GMDN Code:	47042 Wound - nonadherent dressing, absorbent, antimicrobial	

Product References Covered by this Declaration:	Product Descriptor:
382000	Self- adherent antimicrobial soft silicone foam dressing
382200	Self- adherent antimicrobial soft silicone foam dressing
382400	Self- adherent antimicrobial soft silicone foam dressing
395010	Self- adherent antimicrobial soft silicone foam dressing
395200	Self- adherent antimicrobial soft silicone foam dressing
395221	Self- adherent antimicrobial soft silicone foam dressing.
395260	Self- adherent antimicrobial soft silicone foam dressing
395300	Self- adherent antimicrobial soft silicone foam dressing
395360	Self- adherent antimicrobial soft silicone foam dressing
395400	Self- adherent antimicrobial soft silicone foam dressing
395410	Self- adherent antimicrobial soft silicone foam dressing
395460	Self- adherent antimicrobial soft silicone foam dressing
395600	Self- adherent antimicrobial soft silicone foam dressing
395660	Self- adherent antimicrobial soft silicone foam dressing



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395700	Self- adherent antimicrobial soft silicone foam dressing
395800	Self- adherent antimicrobial soft silicone foam dressing
395900	Self- adherent antimicrobial soft silicone foam dressing

Site(s): GLO