



Mölnlycke[®]

EU Declaration of Conformity according to MDR

Document ID: Created by: Approved by: Approval date:

Clumserry

PD-595102 Rev: 00 Åsa Isaksson Christina Lewing 2020-11-04

Title: Mepore Roll Page 1(2)

We, Mölnlycke Health Care AB, Gamlestadsvägen 3C, Box 13080, SE-402 52 Göteborg, Sweden 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 Medical Device Regulation 2017/745, concerning medical devices.

Other Union Legislation

Not applicable

applicable:

Trade name/
Product name:

Mepore Roll

Product classification: I

MDR Classification Rule: 4

Sterility: Non-sterile

Measuring function: No

This declaration is supported by a conformity assessment procedure in accordance with

Annex/es: IV

Common Specification: No CS is applicable

Certificate number: Not Applicable

Issued by: Not Applicable

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

Mölnlycke Health Care issues this declaration in recognition of all applicable harmonised standards.

Signed for and on behalf of Mölnlycke Health Care

Date: 2020-11-04 Function: Regulatory Affairs Director

Name: Christina Lewing Signature:



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Intended Purpose: Mepore Roll is a non-sterile, self-adhesive, absorbent

dressing for superficial skin damage such as injection sites.

Product(s) covered by this declaration:

Product Reference:	Product Descriptor:	Basic UDI-DI:	GMDN Code:
331900	Self-adhevise dressing roll	7332430000000000002JC	34864 Adhesive bandage
331980	Self-adhevise dressing roll	733243000000000002JC	34864 Adhesive bandage
332000	Self-adhevise dressing roll	733243000000000002JC	34864 Adhesive bandage
332080	Self-adhevise dressing roll	733243000000000002JC	34864 Adhesive bandage