Declaration of Conformity to EU Medical Device Regulation 2017/745

Legal Manufacturer	Coloplast A/S Holtedam 1, 3050 Humlebaek, DK SRN: DK-MF-000025526
EU Product Classification according to Annex VIII	III Rule Number: 14
Intended Purpose	The product is intended to be used as a skin protectant in the management of superficial skin damage in skin-folds and/or other skin-to-skin contact areas.
Basic UDI-DI	57089322853167R
Conformity Assessment Procedure	Annex IX
Notified Body Name and Number	DNV Product Assurance AS - (2460)
Notified Body Certificate Type and Number	EU Quality Management System Certificate - 10000376655-PA-NoMA-DNK EU Technical Documentation Assessment Certificate - C677116
Conformity to Common Specification(s)	No relevant Common Specification to list
Conformity to other Union Legislation(s)	Community code relating to medicinal products for human use Directive 2001/83/EC, relevant provisions of Annex 1

This EU Declaration of Conformity is applicable for following catalogue numbers:

Catalogue Number	Product Name	Original CE Marking Date yyyy-mm-dd
67919 / 679190	InterDry	2018-12-13
67918 / 679180	InterDry	2018-12-13

This EU Declaration of Conformity is issued under the sole responsibility of Coloplast A/S. Coloplast A/S declares that the devices covered by the present declaration are in conformity with the Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices and the requirements specified herein are fulfilled.

The devices that are covered by the present declaration are in conformity with the relevant Union legislation(s) referenced in this declaration.

Date of signature: 2025-09-25

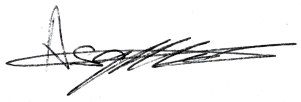
yyyy-mm-dd

Place of signature: Humlebaek, Denmark

Place, Country

DKADGR, Adam Gregory, Head of Regulatory Affairs, Wound & Skin Care

Signed on behalf of Coloplast A/S:



Name, Title