

# EU Technical Documentation Assessment Certificate

Certificate no.:  
C677116

Initial certification date:  
21 May 2025

Valid Until:  
20 May 2030

This is to certify that:  
**InterDry**

Manufactured by:

**Coloplast A/S**

Holtedam 1, 3050 Humlebaek, Denmark

SRN: DK-MF-000025526

Has been assessed and found to comply with respect to:

**Technical Documentation Assessment as described in Annex IX  
(Chapter II) of Regulation 2017/745 on Medical Devices**

Place and date:  
Høvik, 21 May 2025

For the issuing office:  
DNV Product Assurance AS – Notified Body 2460  
Veritasveien 1, 1363 Høvik, Norway



*Mariann Jeremiassen*

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Management Representative

### Jurisdiction

Application of Regulation 2017/745 on medical devices, implemented in Norway by Act 7 May 2020 no. 37 on medical devices and Regulation 9 May 2021 no.1476 on medical devices by the Norwegian Ministry of Health and Care Services.

### Certificate history:

Revision	Description	Report No.	Issue Date
0.0	Original Certificate	2499116	21 May 2025

### Products covered by this Certificate:

Type of product and identification no., Basic UDI-DI		Class	EMDN code
<b>Basic UDI-DI</b>	57089322853167R	III	M04040801
<b>InterDry</b> 679190 / 67919 Pouch 25 cm x 91 cm			
<b>InterDry</b> 679180 / 67918 Roll 25 cm x 366 cm			
<b>Intended purpose of the Product</b>			
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.			

Conformity Assessment for devices listed is covered by separate EU Quality Management System Certificate No.: 10000376655-PA-NoMA-DNK

EU Representative
N/A

## Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall inform the Notified Body of any intended change of the products detailed above and the Notified Body will assess the changes and decide if the certificate remains valid.

The following may render this Certificate invalid:

- Changes in the design of the products to which this Certificate refers.
- Changes in requirements of the scheme to which this Certificate refers.

## Conformity declaration and marking of product

This Certificate must be accompanied with a valid EU Quality Management System Certificate.

When meeting with the terms and conditions above, the producer may draw up an EU declaration of conformity and legally affix the CE mark followed by the Notified Body identification number.

