Coloplast

EC Declaration of Conformity to Council Directive 93/42/EEC and amendments concerning Medical Devices

Manufacturer	Coloplast A/S Holtedam 1 3050 Humlebaek Denmark
Product	Product family: Antimicrobial skin friction and moisture control fabric
	Product name: InterDry
Description	On primary packaging: See attachment
	Global Medical Device Nomenclature code/name: 62291 Antimicrobial skin friction/moisture control fabric
EC Product Class according to Annex IX	Class: Non-sterile class III Rule no.: 13
We hereby declare that the above-mentioned device complies with the relevant provisions of Annex 1 to Directive 2001/83/EC, as amended.	
Notified Body	Presafe Denmark A/S – (0543) Tuborg Parkvej 8, DK-2900 Hellerup, Denmark
ID No.	454

The undersigned, Vice President for Global RA Operations, declares that the following devices:
InterDry
Conform to the relevant provisions of the European Communities' Council Directive 93/42/EEC and amendments and are in accordance with Annex II Conformity Assessment Procedure (full quality assurance system) as verified by Presafe Denmark A/S – (0543).

This product was originally CE-marked: 2018-12-13

Date: 13/12/2018 By: _____

Caroline Jane Allen,
Vice President, Global RA Operations

Description on Primary Packaging for
Product Family:
Antimicrobial moisture wicking fabric

Product name:
InterDry

Item number	Product name	Product description	Date of original CE-marking
67918	InterDry	25 cm x 366 cm, roll	2018-12-13
67919	InterDry	25 cm x 91 cm, sheet	2018-12-13

Date: 13|12|2018 By: _____

Caroline Jane Allen,
Vice President, Global RA Operations