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	I Rule Number: 1
Intended Purpose	The product is intended to fill in cavities/folds in the intact skin around the stoma. This provides a seal between the stoma and the ostomy baseplate and protects the skin against the stoma output.
Basic UDI-DI	57089322976739D
Conformity to Common Specification(s)	No relevant Common Specification to list
Conformity to other Union Legislation(s)	No relevant Union Legislation to list

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

Catalogue Number	Product Name	Original CE Marking Date yyyy-mm-dd
120307 / 12030	Brava Mouldable Ring	2010-01-12
120425 / 12042	Brava Mouldable Ring	2010-01-12
120305	Brava Mouldable Ring	2010-01-12
120427 / 12042	Brava Mouldable Ring	2010-01-12

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.

Date of signature:	2023-09-11 yyyy-mm-dd
Place of signature:	Humlebaek, Denmark Place, Country
Signed on behalf of Coloplast A/S:	DKBENB, Benedikte Blom, Head of Regulatory Affairs  Name, Title