

## Declaration of Conformity to EU Medical Device Regulation 2017/745

| Legal Manufacturer                                | Coloplast A/S<br>Holtedam 1, 3050 Humlebaek, DK<br>SRN: DK-MF-000025526   |  |
|---|---|--|
| EU Product Classification according to Annex VIII | l<br>Rule Number: 1   |  |
| Intended Purpose                                  | Brava Skin Barrier Spray is intended for protection of intact skin from the damaging effects of body waste and medical adhesives. |  |
| Basic UDI-DI                                      | 57089322976859L   |  |
| 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<br>yyyy-mm-dd |
|-------------------------|--------------------------|--|
| 12020 / 120205 / 120201 | Brava Skin Barrier Spray | 2010-01-15                             |

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:

2023-05-11 yyyy-mm-dd

Place of signature:

Humlebaek, Denmark

Place, Country

DKBENB, Benedikte Blom, Head of Regulatory Affairs

Signed on behalf of Coloplast A/S:

Beredikte Blom

Name, Title