

Date 2021-05-25

EC Declaration of Conformity for Medical Devices Class IIa

We herewith declare that the medical devices of the following product groups

| Group no. | Product group | Class acc. to 93/42/EEC | Rule | UMDNS | GMDN |
|-----------|----------------------|-------------------------|--------|--------|-------|
| 3.01 | Hydrocolloids | IIa | 4 (3.) | 10-288 | 43186 |
| 3.02 | Hydrogels | IIa | 4 (3.) | 10-288 | 47764 |

which are first placed on the market by CMC Consumer Medical Care GmbH, meet the applicable provisions, especially the essential requirements of the Council Directive 93/42/EEC of 14th June 1993.

The required conformity assessment procedure acc. to Annex II excluding (4) has been performed and the technical documentation is kept available.

The EC Declaration of Conformity is issued under the sole responsibility of the CMC Consumer Medical Care GmbH.

The conformity assessment procedure is under the supervision of the Notified Body:
MTIC InterCert S.r.l., Via G.Leopardi, 14, 20123 - Milano (MI), Italy, No. 0068.

ppa.

Dr. Benjamin Wenzel
Director R&D RA

i.V.

Susanne Wolpert
Head of R&D & RA
Person Responsible for Regulatory Compliance acc.
to Art. 15 MDR

This document is valid until: 2024-05-25

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