EC Declaration of Conformity according to Medical Device Regulation MDR 2017/745

This EU Declaration of Conformity is issued under the sole responsibility of the manufacturer in accordance with Annex IV of Regulation (EU) 2017/745.

MANUFACTURER
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CH AUTHORISED REPRESENTATIVE (CH-REP) – Switzerland
CONFINIS AG
CH-3186 Düdingen
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PRODUCT NAME
Kinesiology Tape (Elastic Bandage Tape)

INTENDED PURPOSE
Demaplast is designed to be applied to the skin.
It supports the body's natural recovery process. It is also suitable for fixation and targeted support.

MODEL NAME
DermaPlast
DermaPlast Active
DermaPlast Sensor patches
DermaPlast Xtreme

CLASSIFICATION
Medical Device Class I, according to Rule 1 of Annex VIII, MDR 2017/745

BASIC UDI-DI

871917226elastic-tapeY8

EMDN CODE

M030402030102

Declaration

We hereby declare that the above-mentioned products meet all applicable requirements of Regulation (EU) 2017/745.

Date of issue: 2026-02-18

Expiry date: 2028-02-17

Place of issue: Enschede

Signature:

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