

EC Declaration of Conformity

According to Medical Device Regulation MDR 2017/745

MANUFACTURER

: THYSOL HI-TECH LTD. (Ecoworkskorea)

17-40, Hwaseong-ro 1424, Gyeonggi-do, Seoul, South-Korea.

SRN-number: KR-MF-000008975

EUROPEAN REPRESENTATIVE

: FysioTape BV, Josink Kolkweg 18, 7545 PR, Enschede, The Netherlands

SRN-number: NL-AR-000002061

PRODUCT NAME

: Elastic balance tape

MODEL NAME + CODE

: CureTape ® Patches: 8717624161773, 8717624161780, 8717624161797,

8717624161926, 8717624161933, 8717624161940, 8717624161957,

8717624161964, 8717624161971

CLASSIFICATION

: Medical Device Class I, by rule 1 of annex 8, section III

Basic UDI:

: 87176241elastic-tapeWX

This declaration of conformity is issued under the sole responsibility of FysioTape BV. We hereby declare that the above mentioned products meet the provisions of the Medical Device Regulation MDR 2017/745. All supporting documentation is retained under the premises of the manufacturer.

Date of issue: March 2023 Place: Enschede, The Netherlands

Signature Manufacturer:

0/000

Signature European Representative:

Name: Hyunsuk Lee

Nama.

Function: President

Function:

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