

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® Classic 5cm x 5m: 1000, 1010, 1020, 1030, 1060,
1062, 1063, 1064, 1065
CureTape® Classic 1cm x 5m: 1070-5, 1080-5
CureTape® Classic 2,5cm x 5m: 1040-2
CureTape® Classic 7,5cm x 5m: 1050
CureTape® Sports 5cm x 5m: 2101, 2102, 2103, 2104, 2105
CureTape® Art 5cm x 5m: 2201, 2202, 2203, 2204, 2205,
2206
CureTape® Punch 5cm x 5m: 1200, 1210, 1220, 1230
CureTape® Giant 5cm x 5m: 1001, 1011, 1021, 1031, 1061,
1066, 1067, 1068
- 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: July 2021

Place: Enschede, The Netherlands

Signature Manufacturer:

Name: Hyunsuk Lee

Function: President

Signature European Representative:

Name: G.J. Olthoff

Function: CEO