

EC Declaration of Conformity

Manufacturer:
Wuhan Huawei Technology Co., Ltd.
B-F11-3, Huazhong International Industrial
Park, Yangluo Port, 430415, Wuhan, Hubei,
China 430415

whose single Authorized Representative:
Humiss International B.V.
Address: Joop Geesinkweg 701, 1114
AB Amsterdam-Duivendrecht, the
Netherlands

We, the manufacturer, herewith declare that the product

Product name: *Cool Patch*
Brand name: *DermaPlast*
Model No.: *04.04.01*
CND Code: M9001
Basic UDI-DI: 69452960Coglpatch001V4



meets the provisions of EU Medical Device Regulation 2017/745 (EU MDR) which
apply to them.

The medical device has been assigned to class I (non-sterile) according to Rule 1 of Annex VIII. It bears the mark



Conformity assessment procedure: The Conformity Assessment Procedure according to Article 52 (7) has been performed and the Technical Documentation is kept available.

We, the manufacturer, herewith declare under our sole responsibility that the above-mentioned products meet the provisions of the Regulation (EU) 2017/745 on Medical Devices (MDR). All supporting documentations are retained under the premises of the manufacturer.

Hubei, China, Dec 10, 2021 Duanyun.chen , General Manager
Place : B-F11-3, Huazhong International Industrial Park, Yangluo Port, 430415,
Wuhan, Hubei, China 430415



Date: Dec 10, 2021

