EU 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:
Shanghai International Holding Corp. GmbH (Europe)
Eiffestrasse 80, 20537 Hamburg, Germany

We, the manufacturer, herewith declare that the product

Product name: *COOL PATCH*
Brand: *DermaPlast*
Model No.: *04.04.01*
CND Code: M9001
Basic UDI-DI: *6945296056499CM*

meets the General Safety and Performance requirements of EU Medical Device Regulation 2017/745.

The medical device, which was first placed on the market by us, has been assigned to class I (non-sterile) according to Rule 1 of Annex VIII. It bears the mark

CE

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 EU Medical Device Regulation 2017/745 (EU MDR) which apply to them. All supporting documentations are retained under the premises of the manufacturer.

Hubei, China, April 15,2020 Duanyun,chen , General Manager
Place : B-F11-3, Huazhong International Industrial Park, Yangluo Port, 430415, Wuhan, Hubei, China 430415

Date: May 26, 2020

武汉华卫科技有限公司
WUHAN HUAWEI TECHNOLOGY CO.,LTD