

## EU DECLARATION OF CONFORMITY

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Manufacturer: Medico Electrodes International Ltd.  
Plot 142A/11, 12, 27, 28 & 29, Noida Special Economic Zone, Noida,  
Uttar Pradesh, 201305, India.

SRN: IN-MF-000021002

Product Name: Abrasive pad

Name/Type: 2121MED

Intended Purpose: Abrasive pad for skin preparation

UDI-DI: 8906035282121MEDNY

Device Classification: Class I (in accordance with Annex VIII, Rule 1- Non-invasive devices of EU MDR 2017/745)

Packaging: Various

GMDN Code: 47102 (Electrode Skin Abrasive Pad)

We Herewith Declare In Our Own Responsibility that The Above-Mentioned Products Meet The General Safety and Performance Requirements, Annex I, of Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on Medical Devices. All supporting documentation and Technical file is retained under the premises of the manufacturer (Head of Quality Department) & Authorized European Representative and will be kept at disposal of the Notified Body and Competent Authorities for a period of ten years from the production date of the device.

We further declare that a suitable procedure is maintained which grants the application of the post-marketing surveillance procedure required by EU MDR 2017/745.

General applicable regulation(s): Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on Medical Devices

Authorized European Representative: Medico Electrodes Ltd.  
Blvd. Saint-Michel 65/6  
1040 Brussels, Belgium  
SRN: BE-AR-000020093  
[www.medicoelectrodes.com](http://www.medicoelectrodes.com)

Person keeping the technical documentation: Amit Seth, Vice-President.



Signed  
For Medico Electrodes International Ltd.,  
Authorized Signatory.

Date: **08-03-2022**  
Place: NOIDA, U.P., INDIA

On the basis of this declaration, the product packaging will bear the following marking:

