EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Deutschland GmbH
Health Care Business
Single Registration Number DE-MF-000011641
Carl-Schurz-Str. 1
41453 Neuss
Germany

hereby declare under our sole responsibility that the following CE marked device(s)

Trade Name	Medipore™
Intended Purpose	Soft cloth surgical tape on liner
Reference	2991/1, 2991/2, 2991/3, 2991/4, 2991NP-3, 2991P-1, 2991P-2
Basic UDI-DI	06082327610100000000022CM

are classified per rule 1 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class I non-sterile devices in accordance with all applicable provisions of the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL concerning medical devices.

Margaret Bessenbach

May 31, 2022

Margaret Bessenbach
Manager Regulatory Affairs and Quality
Health Care Business EMEA
3M Deutschland GmbH

Date

Related to REG-STED-MDR-05-417495