EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Deutschland GmbH
Health Care Business
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 is intended to be used on intact skin as a surgical tape and/or a dressing cover.
Reference	2991/1, 2991/2, 2991/3, 2991/4, 2991NP-3, 2991P-1, 2991P-2
Basic UDI-DI	06082232761050000000008GP

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.

3rd March 2020

Harald Ceschinski
Manager Regulatory Affairs and
Quality Management System
Health Care Business EMEA
3M Deutschland GmbH

Date