

**EUROPEAN MEDICAL DEVICE REGULATION****Declaration of Conformity**

As Legal Manufacturer, we

3M Company
Single Registration Number (TBD)
2510 Conway Ave. St. Paul, MN 55144 USA

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

Trade Name*	3M Reston™ Self-Adhering Foam
Intended Purpose	3M™ Reston™ Self-Adhering Foam products are designed to protect against skin damage by adhering to splints, casts, and prostheses. The products may be used as padding on devices and surfaces such as beds, chairs, and operating room tables. This product is not designed, sold, or intended for use except as indicated.
Reference	1560M, 1561H, 1563L, 2851H
Basic UDI-DI	06082238401010000000053AF

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

The Authorized European Representative for the concerned device(s) is

3M Deutschland GmbH
Health Care Business
Single Registration Number (TBD)
Carl-Schurz-Str. 1
41453 Neuss, Germany



Dianne Gibbs, Division Regulatory Affairs Manager
3M Company
2510 Conway Ave. St. Paul, MN 55144 USA

12 August 2020
Date

3M and Reston are trademarks of 3M.