EUROPEAN MEDICAL DEVICE REGULATION

Declaration of Conformity

As Legal Manufacturer, we

3M Company
Single Registration Number US-MF-000014086
2510 Conway Ave. St. Paul, MN 55144 USA

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

Trade Name*	Nasogastric Securement Device
Intended Purpose	The adhesive side of the device affixes medical devices (such as tubing) to the skin. The product is non-invasive, non-sterile, and is intended for prolonged use (>24 hours to 30 days) on intact skin.
Reference	1500NG, 1501NG
Basic UDI-DI	06082238401010000000206AG

are classified per rule 1 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 DE-AR-000011642
Carl-Schurz-Str. 1
41453 Neuss, Germany

DocuSigned by:
Dianne Gibbs
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10/14/2022

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

Date