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	3M™ Curos™ Disinfecting Cap for Needleless Connectors 3M™ Curos™ Disinfecting Cap Strip for Needleless Connectors 3M™ Curos™ Disinfecting Cap for Tego® Hemodialysis Connectors 3M™ Curos Jet™ Disinfecting Cap Strip for Needleless Connectors 3M™ Curos Jet™ Disinfecting Cap for Needleless Connectors
Intended Purpose	Disinfecting cap for use on needleless connectors only prior to I.V. access and to act as a cover between line accesses. The cap will disinfect the needleless connector one (1) minute after application and protect from contamination between accesses for up to seven (7) days if not removed.
Reference	CFF1-270R: Individual caps, 270 caps per carton CFF10-250R: 10-cap strips, 250 caps per carton CTG1-270R: Individual caps, 270 caps per carton CFJ5-250R: 5-cap strips, 250 caps per carton CFJ1-270R: Individual caps, 270 caps per carton
Basic UDI-DI	06082238401010000000040A6

are classified per rule 16 of Annex VIII of the Medical Device Regulation (EU) 2017/745, as Class IIa devices in accordance with Annex XI Part A and all other applicable provisions of the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL concerning medical devices.

This declaration is made based on the quality assurance certificate
EC Certificate Number: MDR 725202
Issued by: BSI, 2797

EU Authorized Representative:
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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St. Paul 8/29/2022

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

Location/Date