3M Health Care Business

3M Center
2510 Conway Ave, Bldg. 275-5W-06
St. Paul, MN 55144 U.S.A.
651 733 1110

Declaration of Conformity

As Legal Manufacturer
We,

3M Company
3M Health Care
2510 Conway Ave
Saint Paul, MN 55144 USA

hereby declare under our sole responsibility
that the CE marked products to which this declaration relates,

3M™ Curos™ Disinfecting Cap for Needleless Connectors
3M™ Curos™ Disinfecting Cap Strip for Needleless Connectors
3M™ Curos™ Disinfecting Cap for Tego® Hemodialysis Connectors

Product Numbers
CFF1-270R, CFF10-250R, CTG1-270R

is classified,
per Rule 15 of Annex IX of the Medical Device Directive 93/42/EEC, as amended per 2007/47/EC
as a Class IIa device
and

is in accordance with Annex V of Directive 93/42/EEC, as amended per 2007/47/EC
on the approximation of the laws of the European Union Member States concerning medical devices.

In addition, we declare that the above mentioned devices fulfill the applicable provisions of the Directive 93/42/EEC, as amended per 2007/47/EC.

This declaration is made on the basis of the quality assurance certificate, CE 00493 delivered by BSI, 2797

EU Representative Address
3M Deutschland GmbH
Health Care Business
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41453 Neuss, Germany

Signature: Dianne Gibbs Date: 21 May 2020

Dianne Gibbs
3M Health Care
Division Regulatory Affairs Manager
Medical Solutions Division

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