



**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™ Curoso™ Disinfecting Cap for Needleless Connectors  
3M™ Curoso™ Disinfecting Cap Strip for Needleless Connectors  
3M™ Curoso™ 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  
Carl-Schurz-Str. 1  
41453 Neuss, Germany

Signature: \_\_\_\_\_

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

Date: \_\_\_\_\_

21 May 2020