



**Declaration of Conformity**

As Legal Manufacturer  
We, 3M Company,  
2510 Conway Ave  
St. Paul, MN 55144 USA

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

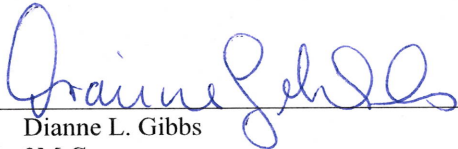
1818	Tie-On Surgical Mask
1818FS	Tie-On Surgical Mask with Face Shield
1835	High Fluid Resistant Surgical Mask
1835FS	High Fluid Resistant Surgical Mask with Face Shield
1838R	Filtron High Performance Surgical Mask
1826	Standard Earloop Face Mask

are classified,  
per Rule 1 of Annex IX of the Medical Device Directive 93/42/EEC, as amended per 2007/47/EC,  
as Class I devices  
and

are in accordance with Annex VII and all other applicable provisions of the 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.

EU Representative Address  
3M Deutschland GmbH  
Health Care Business  
Carl-Schurz-Str. 1  
41453 Neuss, Germany

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

  
Dianne L. Gibbs  
3M Company  
Division Regulatory Affairs Manager  
Medical Solutions Division

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



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