



3M Deutschland GmbH

Carl-Schurz-Straße 1  
41453 Neuss  
GERMANY

+49 (0)2131/140  
+49 (0)2131/142649  
Internet: [www.3M.com/de](http://www.3M.com/de)  
E-Mail: [innovation.de@mmm.com](mailto:innovation.de@mmm.com)  
WEEE-Reg.-Nr. DE 36963167  
VAT-ID: DE 120679179

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## ***Declaration of Conformity***

We

**3M Deutschland GmbH  
Health Care Business  
Carl-Schurz-Str. 1  
41453 Neuss  
Germany**

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

<b>3M™ Coban™ Self-Adherent Wrap Hospital Packs</b>	<b>1581, 1581B, 1581K, 1582, 1582B, 1582K, 1583, 1583B, 1583G, 1583K, 1583R, 1583W, 1584, 1584L, 1584B, 1584K, 1584W, 1586</b>
<b>3M™ Coban™ Self-Adherent Wrap EU Pharmacy Packs</b>	<b>1582/S, 1582/B, 1582/W 1583/S, 1583/B, 1583/W 1584/S, 1584L/S,</b>
<b>3M™ Coheban™ Self-Adherent Wrap</b>	<b>1581, 1582, 1583, 1583/2, 1583/B, 1584, 1584/2</b>
<b>3M™ Nexcare™ Athletic Wrap</b>	<b>N1675W, N1650T</b>

are classified per rule 1 of Annex IX of the Medical Device Directive 93/42/EEC,  
as **Class I** devices  
and

are in accordance with  
***Annex VII and all other applicable provisions of the Directive 93/42/EEC***  
on the approximation of the laws of the European Union Member States concerning medical devices.

Signature:

*Margaret Bessenbach*  
Margaret Bessenbach  
Manager Regulatory and Quality  
Health Care Business West Europe & CEE  
3M Deutschland GmbH, Health Care Business

*November 10, 2015*  
Neuss / Date

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