



Declaration of Conformity

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

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

3M Red Dot Monitoring Electrodes

Product numbers:

2230, 2231, 2235, 2237, 2237-3, 2237-5, 2238, 2238-3, 2238-5, 2239, 2245-50, 2248-50, 2249, 2249-50, 2255-50,
2256, 2258-3, 2259-3, 2259-50, 2268-3, 2268-5, 2270-3, 2270-5, 2270-50, 2271-3, 2271-5, and 2271-50

are classified,

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

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

3M Health Care Business self-declares conformity with Directive 2011/65/EU of the European Parliament
and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances
in electrical and electronic equipment and compliance to the requirements of EN 50581:2012.

EU Representative Address
3M Deutschland GmbH
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Carl-Schurz-Str. 1
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Signature: _____

Dianne L. Gibbs
3M Health Care
Division Regulatory Affairs Manager
Infection Prevention Division

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