

EC Declaration of Conformity Issue 4

E & O Laboratories Ltd
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Scotland
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Declares that its diagnostic products have been classified and registered in accordance with Directive 98/79/EC of the European Parliament and the Council of 27 October 1998.

Each of the products meets the relevant provisions of the directive and have been classified as "all other IVD Medical devices" within Annex III of the directive.

E&O Laboratories Ltd has developed, implemented and maintains a documented post-production monitoring process, including the notification of reportable events in compliance with Article 11 Vigilance System Guidelines.

Medicines & Healthcare Products Regulatory Agency
Market Towers
1 Nine Elms Lane
London
SW8 5NQ
Registration Ref: IVD 000122, 28 Nov 2003

Virginia Lucey (Managing Director)

Date



25th August 2015
