HITACHI
Inspire the Next

EC-DECLARATION OF CONFORMITY

Manufacturer: Hitachi, Ltd.
Address: 2-16-1, Higashi-Ueno, Taito-ku, Tokyo, 110-0015, Japan

Selected conformity assessment procedure: Annex ll excluding (4) RoHS Article 7 (b), Module A

EU Authorized representative: Hitachi Medical Ultrasound Logistics,
Address: Zweigniederlassung der Hitachi Medical Systems Europe Holding AG
Carl-Zeiss-Strasse 5
D-72555 Metzingen
Germany

Product: ***S22 Probe***

Model Code:: ***S22***

Classification (MDD, Annex IX): ll a Categories (RoHS(Ⅱ), Annex I): No.8
Classification rule (MDD, Annex IX): rule 10

We herewith declare that the above mentioned products including all its accessories meet the provisions of the following EC Council Directives and Standards. All supporting documentations are retained under the premises of the manufacturer.

DIRECTIVES

General applicable directives:

Medical Device Directive : Council Directive 93/42/EEC of 14 June 1993
Medical Devices Directive (93/42/EEC as amended by 2007/47/EC;

Standards : MDD Harmonized Standards (published in the Office Journal of the European Communities) applicable to this product are :

EN 60601-1:2006/AC:2010, EN 60601-1-2:2007/AC:2010, EN 60601-2-37:2008
EN 62366:2008, EN 60601-1-6:2010, EN 1041:2008, EN ISO 14971:2012
EN ISO 10993-1:2009/AC:2010, EN ISO 14937:2009, EN ISO 17664:2004

Other Standards : IEC 62079:2001, ISO 7010:2003/A6:2010, IEC 60601-1-9:2007, ISO 3864-2:2004, ISO 15223-1:2012

Notified body : TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123
Address (for MDD): Ridlerstraße 65, 80339 München, Germany

RoHS Directive : Directive 2011/65/EU of 8 June 2011 concerning on the restriction of the use of certain hazardous substances in electrical and electronic equipment.

Standards : RoHS Directive Harmonized Standards (published in the Office Journal of the European Communities) applicable to this product is

EN 50581:2012

Production facility : ***Hitachi Healthcare Manufacturing, Ltd. Tokyo Works***
Address: ***3-7-19 Imai, Ome-shi, Tokyo, 198-8577, JAPAN***

Starting of CE Marking: ***204W3531***

Date: ***Jul. 11. 2016***

Signature: 

Name of issuer : Ryosuke Maeda Place: ***Tokyo, JAPAN***
Position : Department Manager
Quality Assurance Department 1