EU DECLARATION OF CONFORMITY

1. No. ECR-US-032-01-E

2. Name and address of the manufacturer and authorized representative:

Manufacturer
Hitachi Medical Corporation
4-14-1, Soto-Kanda, Chiyoda-ku
Tokyo, 101-0021, Japan

European Representative
Hitachi Medical Systems GmbH
Otto-von-Guericke-Ring 3
D-65205 Wiesbaden, Germany

3. This declaration of conformity is issued under the sole responsibility of the manufacture.

4. Object of the declaration

Product name in general: Ultrasound Transducer/Probe
Series name: EUP-F531
Category of RoHS Directive(ANNEX I): 8. Medical Devices

5. The object of the declaration described above is in 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 electric equipment.

6. Where applicable, references to the relevant harmonized standards used or references to the technical specifications in relation to which conformity is declared:

- EN 50581:2012
  technical documentation for the assessment of electrical and electronic products with respect to the restriction of hazardous substances

7. Additional information:

Signature N. Yoshida
N.Yoshida
Department Manager
Ultrasound Systems Design Office

Signature N. Kawabe
N.Kawabe
Division Manager
QA Division

Place and date of issue:

Place JAPAN Date 16/06/2014 (DD/MM/YYYY)

Table of product name in general:

1 Ultrasound Transducers/Probes
2 Ultrasound Diagnostic Systems
3 Sterile Puncture Adapters for Ultrasound Probes
4 MRI Systems
5 X-Ray CT Systems
6 Optical Encephalography Systems
7 Mobile X-ray Units
8 Radiographic Systems