

Declaration of Conformity



Manufacturer: Shenzhen Mindray Bio-Medical Electronics Co., Ltd.
Mindray Building, Keji 12th Road South, Hi-tech Industrial
Park, Nanshan, Shenzhen, 518057, P. R. China

EC-Representative: Shanghai International Holding Corp. GmbH (Europe)
Eiffestraße 80
20537 Hamburg, Germany

Product: Ultrasonic Diagnostic System

Model: TE7 /TE7T /TE7S /TE7 Pro /TE7 Super /TE5 /TE5T /TE5S
/TE5 Pro /TE5 Super

Supplementary information: Included are following transducers: C5-2s, C11-3s, L12-4s,
L14-6Ns, L14-6s, L7-3s, P4-2s, P7-3Ts, V11-3Ws, 7LT4s,
L16-4Hs, P10-4s, L20-5s, SC6-1s, SP5-1s, 6CV1s, 7L4s, P7-3s,
L12-3RCs, L14-5Ws, C4-1s, C5-1s, L11-3VNs, L9-3s
and following needle -guided brackets: NGB-004, NGB-007,
NGB-010, NGB-011, NGB-015, NGB-016, NGB-018,
NGB-022, NGB-034, NGB-035, NGB-036, NGB-043.

We herewith declare under our sole responsibility that the above mentioned products meet the provisions of the Council Directive 2011/65/EU. All supporting documentations are retained under the premises of the manufacturer.

Standards Applied:
EN 50581:2012.

Start of CE-Marking: 2014.12.2

Place, Date of Issue: Shenzhen, 2018.9.20

Signature: _____

Name of Authorized Signatory: Mr. Wang Xinbing

Position Held in Company: Manager, Technical Regulation Department