

EC Certificate Full Quality Assurance System: Certificate CN15/30486

The management system of

SHENZHEN SHENCHAO TRANSDUCER CO.,LTD

West Section of 5F ,5th Building , Hongfa Jiateli Hi-tech Park, Jingtang Road, Shixin Community, Shiyan Street, Bao'an District Shenzhen, Guangdong Province, 518108, P.R.China

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Ultrasonic Transducer used for Ultrasonic Medical Diagnostic Systems (Convex Array Probe, Linear Array Probe, Cavity Probe, and Phase Array Probe)

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 13 May 2015 until 15 April 2020 and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 31 December 2017 Issue 2. Certified since 16 April 2015

Certification is based on reports numbered CNISZX 49947

Authorised by



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