EC Certificate Production Quality Assurance System: Certificate CN18/41007

The management system of

SHENZHEN SHENCHAO TRANSDUCER CO., LTD.

5th Floor, Building A, Jingchengda Industrial Park, Keji 4th Road, Langxin Community, Shiyan Street, Bao'an District, Shenzhen City, 518108, P.R. China

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

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 IIb or class III medical device(s), a valid EC Type Examination Certificate according to Annex III is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 28 April 2018 until 15 April 2023 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 30 December 2020
Issue 2. Certified since 16 April 2015

Certification is based on reports numbered CN/SZX 49947

Authorised by

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