EC Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 103703 0001 Rev. 00

Manufacturer:

Shenzhen AOJ Medical Technology Co., Ltd.
601, 6th floor, B2 Building
An'le Industrial Park
#172 Hangcheng Avenue, Sanwei Community, Hangcheng Street
Bao'an
518126 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Infrared Thermometer and Fetal Doppler

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

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Valid from: 2020-01-15
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Date, 2020-01-15

Christoph Dicks
Head of Certification/Notified Body