

EC DECLARATION OF CONFORMITY

The following EC declaration of conformity exemplifies the required content according to Directive 93/42/EEC.

EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: SONOSCAPE MEDICAL CORP.
4/F, 5/F, 8/F, 9/F & 10/F, Yizhe Building, Yuquan Road, Nanshan, Shenzhen, 518051, Guangdong, China

Name and address of the European Representative: SonoScape Europe S.r.l.
Via Luigino Tandura, 74-00128 Rome, Italy

We declare under our sole responsibility that

the medical device: Digital Color Doppler Ultrasound System
Model: P20, P20Pro, P15, P22, P25, P10 Plus
(Supported Probes: 3C-A, C1-6, 13L-A, L742, L752, ML3-18, 3P-A, 10L-I, 10I2, 8P1, 6V3, 6V3A, EC9-5, 6V1, VC6-2, VE9-5, C322, C613, PW2.0, CWD5.0, LAP7, 12LT-A, 12LI-A, 6CT-A, 6CI-A, BCC9-5, BCL10-5, 6V7, S1-5, L741, 7P-B, 12C-ER, MPTEE, MPTEE mini, 12L-A, 12L-B, 9L-A, 18L-A, VC2-9)

of class: / Ila

according to annex IX of directive 93/42/EEC

meets the provisions of the directive 93/42/EEC and its transpositions in national laws which apply to it. The declaration is valid in connection with the "final inspection report" of the device.

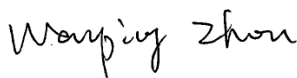
Conformity assessment procedure: / **Directive 93/42/EEC Annex II, excluding Section 4**

Registration No.: **HD 60128046 0001**

Notified Body: **TÜV Rheinland LGA Products GmbH
Tillystraße 2
90431 Nürnberg
Deutschland
CE 0197**

Shenzhen, May 31, 2018

Place, date /



Vice President

Name and function