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: /	IIa 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

Wanping Zhou

Vice President

Place, date / Name and function