

**EC Certificate**  
**Directive 93/42/EEC Annex II, excluding Section 4**  
**Full Quality Assurance System**  
**Medical Devices**

**Registration No.:** HD 60128046 0001

**Report No.:** 17032653 014

**Manufacturer:** SONOSCAPE MEDICAL CORP.  
4/F, 5/F, 8/F, 9/F & 10/F  
Yizhe Building  
Yuquan Road, Nanshan  
Shenzhen  
518051 Guangdong  
China

**Products:** Ultrasonic Diagnostic Systems, Medical Endoscope Systems  
(see attachment for additional site included)  
Replaces Approval, Registration No.: HD 60106282 0001

**Expiry Date:** 2023-06-18

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

**Effective Date:** 2018-06-19

**Date:** 2018-04-16



**TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg**

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.



**TÜV Rheinland**  
**LGA Products GmbH**  
**Tillystraße 2, 90431 Nürnberg**

Doc. 1/1, Rev. 0

**Attachment to  
Certificate**

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Shenzhen  
518051 Guangdong  
China

**Site included:**

4/F(B), 1/F(S), 5/F, Nanfeng Building, Nanshan Yungu  
Innovation Industrial Park, 4093 Liuxian Blvd., Taoyuan  
Subdistrict, Nanshan, Shenzhen, 518055, Guangdong, China

**Date:** 2018-04-16

**Notified Body**



**S. Liu**