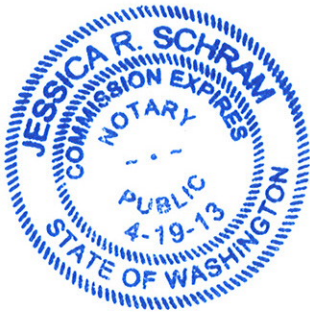


STATE OF WASHINGTON)
)
COUNTY OF SNOHOMISH) ss.

I certify that the attached certificate is a true and correct copy of a document in the possession of SonoSite, Inc. as of this date.

Dated this 29 day of June, 2009.



Jessica R Schram

Jessica R Schram
Printed Name

Notary Public in and for the State of Washington
My appointment expires 4-19-13

Manufacturer's Declaration of Conformity
Full Quality Assurance Procedure
 According To Australian Therapeutic Goods (Medical Devices) Regulations 2002



This is a declaration made in accordance with the requirements of Clause 1.8 of Schedule 3 of the Australian Therapeutic Goods (Medical Devices) Regulations 2002 relating to the devices stated in following Schedule.

Manufacturer's Name & Address:

SonoSite, Inc.
 21919 - 30th Drive SE
 Bothell, Washington 98021-3904 USA

Australasia Sponsor's Name & Address:

SonoSite Australasia, Ltd.
 Australasia Headquarters
 Suite 9, 13A Narabang Way, Belmore NSW 2085, Australia

Declares that the product(s):

Item	Part Number(s)	GMDN Code	Manufacturer
SonoSite NanoMaxx™ series ultrasound system	P11111, V11111	40761, 40762, 40763, 40764	Manufactured by SonoSite, Inc.

Complies with:

Each medical device part number listed above complies with European Community Council Directive (Medical Device Directive) 93/42/EEC as amended by 2007/47/EC, Annex II, and are Class IIa devices in accordance with Annex IX, Rule 10. EC Certificate CE 02429 has been issued by the British Standards Institution (BSI) for compliance with Annex II, section 3.2 of the European Council Directive 93/42/EEC. In addition, each kind of medical device to which the Full Quality Assurance Procedures have been applied complies with the applicable provisions of the essential principles, the classification rules, at each stage, from the design of the device until its final inspection before being supplied. In addition, said devices comply with the applicable requirements of:

Safety:

- EN60601-1: 1990, 2nd Edition including Amendment 1 and 2
- EN60601-1-1: 2001
- EN60601-2-37: 2001 + A1 (2004)

EMI / EMC:

- EN60601-1-2:2007

Supplementary Information:

Included are the following Accessories and OEM Products:

Accessories:

Low Noise Power Supply
 Battery Pack, Panasonic cells
 Battery Pack, Sanyo cells
 Biopsy/Needle Guides
 NanoMaxx™ Dock

Y-Universal Stand
 SonoCalc® IMT
 C11m8-5 Transducer
 C60m5-2 Transducer
 L25m13-6 Transducer
 L38m10-5 Transducer
 P21m5-1 Transducer

OEM Products:

Aquasonic 100 Acoustic Gel
 CIVCO, Transducer covers
 Sony, Black and White Video Printer, UP897MD

Quality Management and Quality Assurance:

- ISO 9001:2000 Quality management systems -- Requirements International Organization for Standardization (2000)
- ISO 13485:2003, Medical devices -- Quality management systems -- Requirements for regulatory purposes (2003)

Daina L. Graham

June 22, 2009
 Bothell, Washington USA

Daina L. Graham
 Vice-President, Regulatory Affairs and Quality Assurance