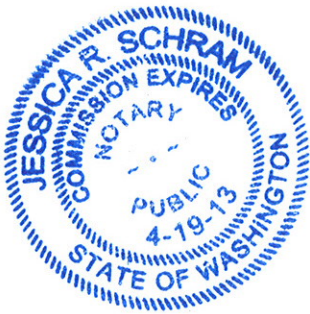


STATE OF WASHINGTON        )  
  )  
COUNTY OF SNOHOMISH    )        ss.

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:**

SonoSite, Inc.

**Australasia Sponsor's Name:**

SonoSite Australasia, Ltd.

**Manufacturer's Address:**

21919 - 30<sup>th</sup> Drive SE  
Bothell, Washington 98021-3904 USA

**Australasia Sponsor's Address:**

Australasia Headquarters  
Suite 9, 13A Narabang Way, Belrose NSW 2085, Australia

**Declares that the product(s):**

Item	Part Number(s)	GMDN Code	Manufacturer
Transducer, C11n/8-5	P11880, V11880	40767	SonoSite, Inc.
Transducer, C60n/5-2	P11878	40767	SonoSite, Inc.
Transducer, L25n/13-6	P12092, V12092	40767	SonoSite, Inc.
Transducer, L38n/10-5	P11877	40767	SonoSite, Inc.
Transducer, P21n/5-1, Phased Array	P11879	40767	SonoSite, Inc.

**Scope and Compliance:**

Each medical device part number listed above complies with European Community Council Directive (Medical Device Directive) 93/42/EEC, 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, 2<sup>nd</sup> Edition including Amendment 1 and 2
- EN60601-1-1: 2001
- EN60601-2-37: 2001 + A1 (2004)

**EMI / EMC:**

- EN60601-1-2:2007

**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)

June 22, 2009  
Bothell, Washington USA

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