

STATE OF WASHINGTON        )  
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COUNTY OF SNOHOMISH    )

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 1 day of September, 2010.

Jessica R Schram

Jessica R Schram

Printed Name  
Notary Public in and for the State of Washington  
My appointment expires 4-19-2013





*Declaration of Conformity*  
According to ISO/IEC 17050

**Manufacturer's Name & Address:**

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

**European Authorized Representative's Name & Address:**

SonoSite, Ltd.  
European Headquarters  
Alexander House, 40A Wilbury Way, Hitchin, Herts, SG4 0AP UK

**Declares that the CE-marked product(s):**

Item	Part Number(s)	GMDN Code	Manufacturer
Low Noise Power Supply	P09823	36529	International Components Corporation
Battery Pack, Panasonic Cells	P12889	36534	MicroPower Electronics, Inc.
Battery Pack, Sanyo Cells	P12890	36534	MicroPower Electronics, Inc
NanoMaxx™ Dock	P09723	40811	SonoSite, Inc.
PowerPack, Kit	P13559	17115, 36534	SonoSite, Inc
PowerPark Dock Module	P12834	37341	SonoSite, Inc
PowerPark Stand Module	P12822	37341	SonoSite, Inc.
V-Universal Stand	P12738	37341	SonoSite, Inc.
SonoCalc® IMT Measurement Software	P12807	40224	SonoSite, Inc.
SonoSite® Workflow Solutions	P13321, P14047	40224	SonoSite, Inc.
Wireless Kit	P12240	16902	SonoSite, Inc.

**Complies with:**

European Community Council Directive (Medical Device Directive) 93/42/EEC as amended by 2007/47/EC, VII, and are Class I medical device accessories in accordance with Annex IX, Rule 12. In addition, that said products comply with the applicable requirements of:

**Safety:**

- EN60601-1: 1990, 2<sup>nd</sup> Edition including Amendment 1 and 2
- EN60601-1-1: 2001

**EMI / EMC:**

- EN60601-1-2:2007 (when tested as part of the ultrasound system)

**Quality Management and Quality Assurance:**

- ISO 9001:2008 Quality management systems -- Requirements International Organization for Standardization
- ISO 13485:2003 Medical devices – Quality management systems – Requirements for regulatory purposes

*Mary K. Moore*

September 1, 2010  
Bothell, Washington USA

Mary K. Moore  
Vice-President of Regulatory Affairs