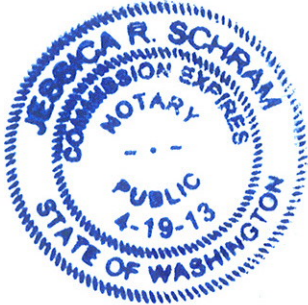


STATE OF WASHINGTON)
)
COUNTY OF SNOHOMISH) ss.

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 23rd day of August, 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 - 30th Drive SE
Bothell, Washington 98021-3904 USA

European Compliance 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 No.	GMDN Code	Manufacturer
SonoSite S Series™ Ultrasound System	P09041, P09417, P12057	40761, 40762, 40763, 40764	Manufactured by SonoSite, Inc.
SonoSite S-Cath™ Ultrasound System	P08778	40761, 40762, 40763, 40764	Manufactured by SonoSite, Inc.
SonoSite S-FAST™ Ultrasound System	P07578	40761, 40762, 40763, 40764	Manufactured by SonoSite, Inc.
SonoSite S-GYN™ Ultrasound System	P11991	40761, 40762, 40763, 40764	Manufactured by SonoSite, Inc.
SonoSite S-ICU™ Ultrasound System	P07577	40761, 40762, 40763, 40764	Manufactured by SonoSite, Inc.
SonoSite S-MSK™ Ultrasound System	P07579	40761, 40762, 40763, 40764	Manufactured by SonoSite, Inc.
SonoSite S-Nerve™ Ultrasound System	P07576	40761, 40762, 40763, 40764	Manufactured by SonoSite, Inc.
SonoSite S-VetMed™ Ultrasound System	P11995	40761, 40762, 40763, 40764	Manufactured by SonoSite, Inc.
SonoSite S-Women's Health™ Ultrasound System	P11993	40761, 40762, 40763, 40764	Manufactured by SonoSite, Inc.

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 02429 has been issued by the British Standards Institution for compliance with Annex II, section 3.2 of the European Council Directive 93/42/EEC. In addition, that said products comply with the applicable requirements of:

Safety:

- | | |
|--|--|
| <ul style="list-style-type: none"> EN60601-1: 1990, 2nd Edition including Amendment 1 and 2 EN60601-1-1: 2001 | <ul style="list-style-type: none"> EN60601-2-37: 2001 + A1 (2004) |
|--|--|

EMI / EMC:

- EN60601-1-2:2007

Supplementary Information:

Included are the following Accessories and OEM Products:

Accessories:

AC Power Supply
Bar Code Reader Kit
Basket
Battery Pack
Biopsy/Needle Guides
Bracket, Single Transducer
Keyboard Tray
PowerPack
PowerPark
SiteLink Image Management System
SonoCalc® IMTSonoSite® Workflow Solutions
Stand
Transducer Bracket Kit
Wireless Kit

Transducers:

C11x/8-5 Transducer
C60x/5-2 Transducer
HFL38x/13-6 Transducer
HFL50x/15-6 Transducer
ICTx/8-5 Transducer
L25x/13-6 Transducer
L38x/10-5 Transducer
L38xi/10-5 Transducer
L52x/10-5 Transducer
P10x/8-4 Transducer
P21x/5-1 Transducer
SLAx/13-6 Transducer
TEEx/8-3 Transducer

OEM Products:

Aquasonic 100 Acoustic Gel
ATP Electronics, Memory, USB, 2GB
CIVCO, Transducer covers
DSI, Keyboard, USB, KB-ASK-3100U-US
Sony, Black and White Video Printer, UP897MD
Sony, DVD Recorder, DVO-1000MD
Symbol, Bar Code Scanner, DS6707-SR20007ZZR

Declaration of Conformity
According to EN45014



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

August 23, 2010
Bothell, Washington USA

Mary K. Moore
Vice-President of Regulatory Affairs