

# 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 6.6 of Schedule 3 of the Australian *Therapeutic Goods (Medical Devices) Regulations 2002* relating to the devices stated in following Schedule.

### Manufacturer's Name & Address:

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

### Australasia Sponsor's Name & Address:

FUJIFILM SonoSite Australasia Pty, Ltd.  
Suite 9, 13A Narabang Way  
Belrose, NSW 2085, Australia

### Declares that the product(s):

| Item                               | Classification    | Part Number(s) | GMDN Code    |
|------------------------------------|-------------------|----------------|--------------|
| Battery Pack                       | Class I, Rule 2.1 | P15051         | 36534        |
| ECG Cable Assembly                 | Class I, Rule 2.1 | P03241, P01592 | 31684        |
| Edge MiniDock                      | Class I, Rule 2.1 | P15078         | 40811        |
| Edge Stand                         | Class I, Rule 2.1 | P15800, P16000 | 37341        |
| Low Noise Power Supply             | Class I, Rule 2.1 | P09823         | 36529        |
| PowerPack                          | Class I, Rule 2.1 | P13559         | 17115, 36534 |
| PowerPark Dock Module              | Class I, Rule 2.1 | P12834         | 37341        |
| PowerPark Stand Module             | Class I, Rule 2.1 | P12822         | 37341        |
| Triple Transducer Connect Module   | Class I, Rule 2.1 | P15922         | 39128        |
| USB Bar Code Scanner Kit           | Class I, Rule 2.1 | P14166         | 17220        |
| Wireless Kit                       | Class I, Rule 2.1 | P12240         | 16902        |
| SiteLink™ Image Management System  | Class I, Rule 4.1 | P10117         | 40224        |
| SonoCalc® IMT Measurement Software | Class I, Rule 4.1 | P14699         | 40224        |
| SonoSite® Workflow Solutions       | Class I, Rule 4.1 | P13321         | 40224        |

### Scope and Compliance:

Each medical device part number listed above complies with the Therapeutic Goods (Medical Devices) Regulations 2002 and are Class I medical devices in accordance with Schedule 2, Part 1, Rule 2.1 and Rule 4.1, as demonstrated in accordance Schedule 3, Part 6, Clause 6.6 and Schedule 3, Part 1. 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.

### Conformity Assessment Certificate:

CE 02429 Full Quality Assurance Certificate first issued 13 September 1999 by BSI in respect of the design, development and manufacture of portable diagnostic ultrasound equipment, transducers and other associated accessories.

### Standards Applied:

- EN60601-1: 1990, 2<sup>nd</sup> Edition including Amendment 1 and 2
- EN60601-1-1: 2001
- EN60601-1: 2006, 3<sup>rd</sup> Edition
- EN60601-2-37: 2001 + A1 (2004)
- EN60601-2-37: 2008
- EN60601-1-2: 2007

### 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
- EN ISO 13485:2012, Medical devices – Quality management systems – Requirements for regulatory purposes

October 24, 2013  
Bothell, Washington USA

Ken Hansen  
Director, Regulatory Affairs and Quality Systems