Quality System Approval Certificate

Medical Devices Directive 93/42/EEC

*The National Standards Authority of Ireland as a duly designated Notified Body, (identification number **0050**), for the purposes of the European Communities (Medical Devices) Regulations (S.I. No. 252 of 1994)*

APPROVES THE QUALITY SYSTEM APPLIED BY

Delfi Medical Innovations, Inc.

106-1099 West 8th Avenue
Vancouver
BC V6H IC3
Canada

to the Product Family

Tourniquet cuffs, Reusable and Sterile Single Use

on the basis of examination under the requirements of Annex II, Section 3 of Directive 93/42/EEC.
*The use of the NSAI Notified Body identification number **0050** in conjunction with CE Marking of Conformance for this product family is hereby authorised.*

Registration Number:	**252.889**
Original Approval:	**20 December 2013**
Last Amended on:	**07 August 2014**
Remains valid until:	**09 January 2017**

Signed:

Approved by:
Kevin D. Mullaney
Chief Executive Officer - NSAI Inc.

Approved by:
Susan Murphy
Risk Management Officer

This certificate remains valid on condition that the Approved Quality System is maintained in an adequate and efficacious manner .
Details of the current product range and operational locations included within the scope of this approval can be obtained from NSAI

National Standards Authority of Ireland, 1 Swift Square, Northwood, Santry, Dublin 9, Ireland.