HotSwap

EC DECLARATION OF CONFORMITY

According to the Medical Device Directive 93/42/EEC Annex II

Document No.:	1682-1RE004-03
Manufacturer:	Hammarplast Medical AB
Division:	
Address:	Kartåsgatan 8, SE-531 40 Lidköping, Sweden
Medical Device	Electronic tourniquet
Device identification (e.g. Part No.):	800-20, 800-21, 800-40, 800-41
Classification according to Annex IX (93/42/EEC):	Class IIa

We hereby declare that the devices mentioned above comply with the Swedish National Board of Health and Welfare Regulation and guidelines on medical devices LVFS 2003:11– transposing European Medical Devices Directive 93/42/EEC.

The Conformity according to Annex II is certified with the notified body of:	The identification number of the notified body for implementation of applicable parts of Annex IV of the Directive is:
Namn/Name Intertek SEMKO AB	Identification No: 0 4 1 3
Adress/Address Box 1103	
SE-164 22 Kista	Certificate No. / Date

Manager

Place and date Lidköping 11-12-30	
Signature Mikael D	Printed Name Mikael Dahlén