

EC - CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 04 12 19502 017



Product Service

Manufacturer: Haemonetics Corporation
400 Wood Road
Braintree, MA 02184
USA

EC-Representative: Haemonetics UK Ltd.
5 Ashley Dr.
Bothwell, Scotland
G71 8BS
UNITED KINGDOM

Product Category(ies): Plasma, Apheresis,
Cell Saving and Processing
Blood Infusion Products

The Certification Body of TÜV Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

Report No.: DM405872

Valid until: 2010-06-04

Date, 2005-06-05



TÜV Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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EC-Certificate

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Facility(ies):

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