QS
QUALITY SERVICE

EC-CERTIFICATE

For Own Brand Labelling
(Directive 93/42/EEC on Medical Devices)
Registration No. 9322

TRIONARA TECHNOLOGIES AB

1st Fl., Lilla Badhusgatan 2, 41121 Gothenburg - Sweden

The Notified Body of QS Zürich AG hereby declare that an examination of the under mentioned quality assurance system documentation has been carried out following the requirements of the national legislation to which the undersigned is subjected, transposing of the Directive 93/42/EEC on medical devices. We certify that the quality assurance system conforms to the relevant provisions of the aforementioned directive for Conformity Assessment under Own Brand Labelling procedure.

This EC-Certificate for Own Brand Labelling is only valid for the following products:
(The products are specified in annex „Product range". This Certificate is only valid together with this annex)

Patient Care Monitors

During the validity of this Certificate, the requirements of the above mentioned regulations must be fulfilled permanently. This is supervised by QS ZÜRICH AG

For precise and updated information concerning possible changes occurred in the certification object of the present certificate, please contact info@QS-ME.com

CE 1254

Audit date: 26.02.2013

Date of issue: 05.03.2013

Expiration date: 25.02.2018

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