

QUALITY SYSTEM

EC-CERTIFICATE

Directive 93/42/EEC

Manufacturer: GE Healthcare Finland Oy
Kuortaneenkatu 2
FI-00510 Helsinki
FINLAND

Coverage of Certificate: Design, manufacture and final inspection

Product category: Patient monitoring systems and
related accessories, anaesthesia
software

Valid until: 27th May 2024

The manufacturer's quality system for the design, manufacture and final inspection of the aforesaid product category has been evaluated and meets the provisions of Council Directive 93/42/EEC as set out in Annex II Section 3. This approval is valid until the expiry date provided that the manufacturer fulfils the obligations imposed by Annex II in Directive 93/42/EEC. Products covered by the certificate are specified in the attachment(s).

Valid from: 6th September 2019


Tuomas Toivonen


Anniina Mäkelä

Certificate no.
C-01-1004-698-19

Notified Body no. 0537:
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