ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 15 09 74115 010

Manufacturer:
Christie Medical Holdings, Inc.
3175 Lenox Park Boulevard, Suite 200
Memphis TN 38115
USA

EC-Representative:
EMERGO EUROPE
Molenstraat 15
2513 BH The Hague
THE NETHERLANDS

Product Category(ies):
Noninvasive Vein Imaging Systems

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: 72108930

Valid from: 2015-12-10
Valid until: 2017-11-26

Date, 2015-12-18

Hans-Heiner Junker

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV®

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Facility(ies):

Christie Medical Holdings, Inc.
3175 Lenox Park Boulevard, Suite 200, Memphis TN 38115, USA

Christie Medical Holdings, Inc.
809 Wellington St North, Kitchener ON N2G-4Y7, CANADA

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