Certificate

EC-Certificate

(full quality assurance system)
according to annex II (excluding section 4) of
Medical Devices Directive 93/42/EEC

It is herewith confirmed by

BSI Group Deutschland GmbH
Eastgate, Hanauer Landstrasse 115
60314 Frankfurt am Main
Germany

in its function as Notified Body (0535), that the manufacturer:

NSK WT GmbH
Portitzer Straße 69d
04425 Taucha
Germany

concerning the medical devices

Disinfector
UMDNS: 11-278
(products/variants specified in appendix)

fulfils the requirements according to Annex II (excluding section 4) Medical Devices Directive 93/42/EEC. The manufacturer has established a quality assurance system for the design, production and final inspection of the specified devices.
For the placing on the market of class III products an additional Annex II section 4 certificate is required.

The appendix is part of this certificate and contains 1 page.

Report No.: SMO7957250/SMO7957255
Certificate No.: CE 595189

Current Issue Date: October 21, 2013

Certification Body

Notified by
Zentralstelle der Länder
für Sicherheitstechnik
ZLS-NB-67/12

First Issue Date:
October 21, 2013.

Based on periodical surveillance
this certificate is valid until
July 16, 2018.

Translation

Appendix of EC-Certificate

(full quality assurance system)
according to annex II (excluding section 4) of Medical Devices Directive 93/42/EEC
Certificate No.: CE 595189

Medical devices of the manufacturer:

NSK WT GmbH
Portitzer Straße 69d
04425 Taucha
Germany

Name of product	Variant	Item	UMDNS/ GMDN	Class
iCare+	C3	S103001	11-278	IIb

Notified by
Zentralstelle der Länder für Sicherheitstechnik
ZLS-NB-67/12

Certification and Testing · BSI Group Deutschland GmbH · bsi.

Frankfurt am Main, October 21, 2013

Certification Body