



Product Service - iCe

EC Certificate

Production Quality Assurance System
Directive 93/42/EEG on Medical Devices (bgDD), Annex V
(Devices in Class Ia, Ib or III)

No. G2 14 09 52227 014

Manufacturer: Shanghai Carelife International
Trading Co., Ltd.
1707 Yinqiao Bldg., 58 Jinxin Rd.
201206 Shanghai
PEOPLE'S REPUBLIC OF CHINA

EC-Representative: Shanghai International Holding
Corp. GmbH (Europe)
EiffestraCe 80
20537 Hamburg
GERMANY

Product Category(ies): Disposable Syringe, Disposable Infusion Sets,
Disposable Blood Transfusion Sets,
Saafp Vein SetG, Blood Lancets for Single Use,
Disposable Surgical Blades,
Sterile f2calpels with Plastic Handle,
Sterile Dental Injection Needle for Single Use,
Sterile Insulin Needles for Single Use

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

Report No.: SH14289EXT01

Valid from: 2016-11-18

Valid until: 2021-11-19

Hans-Heiner Junker



Date, 2014-10-28

TÜV SÜD Product Service GmbH is Notified Body with identification no. 012Z

Page 1 of 2