

Production Quality Assurance System

(Annex V of the Directive 93/42/EEC on Medical Devices)

No. G2S 08 09 64733 002

Manufacturer: **Shandong Zibo Guangda Medical Products Co., Ltd.**

No. 15, East 1 Rd. Liangxiang Industry Park,
Zhangdian, Zibo, China.

EC-Representative: **Shanghai International Holding Corp. GmbH (Europe)**

Eiffelstraße 80
20537 Hamburg
GERMANY

Product

Category(ies): Latex Household Gloves, Disposable Nitrile Gloves, Disposable Latex Gloves

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture according to Annex V, section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective product, product categories and conforms to the provisions of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.: SH0825001

Valid until: 2018-11-05



Hans-Heiner Jurker

Date: 2015-11-06

TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

Page 1 of 2

ZERTIFIKAT ◆ CERTIFICATE ◆ CERTIFICADO ◆ СЕРТИФИКАТ ◆ 認 証 証 書