



EC Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 044992 0016 Rev. 01

Manufacturer

Beijing Reagent Latex

Products Co., Ltd.

Ciqu Industrial Zone, Tongzhou District 101111 Beijing

PEOPLE'S REPUBLIC OF CHINA

Facility(ies):

Beijing Reagent Latex Products Co., Ltd.

Ciqu Industrial Zone, Tongzhou District, 101111 Beijing,

PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Sterile Disposable Examination Gloves.

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.:

BJ1987907

Valid from: Valid until:

2019-07-03 2024-05-26

Date.

2019-07-03

Stefan Preiß

1. Purnil

Head of Certification/Notified Body