



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 099730 0003 Rev. 02

Manufacturer

**Lyncmed Medical
Technology (Beijing) Co., Ltd.**

Room 1601, Building No. 2

Zhubang 2000 Business Building, Balizhuang Xili 99

Chaoyang District

100022 Beijing

PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Sterile Surgical Gowns, Sterile Surgical Packs.

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.:

BJ19205031

Valid from:

2020-03-19

Valid until:

2022-12-19

Date, 2020-03-19

C.D.H

Christoph Dicks
Head of Certification/Notified Body