CE Certificate - 93/42/EEC

TUV

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 002317 0002 Rev. 00

Manufacturer: **Shen Zhen Moph Biotech CO. LTD**
No.63 Gulyue Road, Qiping Community
Guanlan Street, Longhua New Zone
518110 Shenzhen, Guangdong
PEOPLE'S REPUBLIC OF CHINA

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

Product Category(ies): Design and Development, Production and Distribution of Anaesthetic Mask, Protective Face Mask, Medical Face Mask and Disposable Surgical Mask

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: GZ1733001

Valid from: 2020-03-20
Valid until: 2021-06-19

Date, 2020-03-20

Stefan Preiß

Page 1 of 2
TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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Page 2 of 2
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