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Zentralstelle der Länder  
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bei Arzneimitteln und  
Medizinprodukten  
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ZLG-BS-244.10.08



Product Service

# 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 004536 0002 Rev. 00

**Manufacturer:**

**SHENZHEN CHINMED PLUS  
TECHNOLOGY CO., LTD.**

2#, 7th Floor, Building 7  
the 2nd Industrial Area of Xinghua  
Tang Wei, Gong Ming  
Guang Ming District  
518132 ShenZhen  
PEOPLE'S REPUBLIC OF CHINA

**Facility(ies):**

SHENZHEN CHINMED PLUS TECHNOLOGY CO., LTD.  
2#, 7th Floor, Building 7, the 2nd Industrial Area of Xinghua, Tang  
Wei, Gong Ming, Guang Ming District, 518132 ShenZhen,  
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): Spo2 sensor**

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.:** GZ1833601

**Valid from:** 2019-01-22

**Valid until:** 2024-01-21

**Date,** 2019-01-22

Stefan Preiß

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 ZERTIFIKAT ♦ CERTIFICATE ♦ 認證證書 ♦ CERTIFICADO ♦ CERTIFICAT