



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.

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Date,

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Stefan Preiß

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