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Zentralstelle der Länder  
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bei Arzneimitteln und  
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Product Service

# EC Certificate

Production Quality Assurance System  
Directive 93/42/EEC on Medical Devices (MDD), Annex V  
(Devices in Class IIa, IIb or III)

**No. G2 002537 0001 Rev. 00**

**Manufacturer:** **Jiangsu Yishun Medical  
Equipment Co., Ltd.**

No. 25, Haiyan Road  
High-Tech Industrial Zone  
226200 Qidong, Jiangsu Province  
PEOPLE'S REPUBLIC OF CHINA

**Facility(ies):** Jiangsu Yishun Medical Equipment Co., Ltd.  
No. 25, Haiyan Road, High-Tech Industrial Zone, 226200 Qidong,  
Jiangsu Province, PEOPLE'S REPUBLIC OF CHINA

**Product  
Category(ies):** **Instant Cold Pack,  
Instant Hot Pack, Hot Pack**

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

**Report No.:** SH1962710

**Valid from:** 2019-12-23

**Valid until:** 2024-05-26

**Date,** 2019-12-23

Christoph Dicks  
Head of Certification/Notified Body

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