



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 18 04 82113 008

Manufacturer:**Suzhou Junwei Medical
Equipment Co., Ltd.**

Hall 91, No.116, Wutaishan Road
Suzhou Italy Industrial Park
Suzhou Science & Technology Town
Suzhou National Hi-Tech District
215153 Suzhou
PEOPLE'S REPUBLIC OF CHINA

EC-Representative:**Shanghai International Holding
Corp. GmbH (Europe)**

Eiffestraße 80
20537 Hamburg
GERMANY

**Product
Category(ies):****Dental Suction Systems**

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.:

SH23665EXT01

Valid from:

2023-05-25

Valid until:

2025-05-24

**Date,** 2023-04-25

S. Preiß
Stefan Preiß

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

Page 1 of 2



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