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中国国际贸易促进委员会

China Council for the Promotion of International Trade
China Chamber of International Commerce

证明书
CERTIFICATE

号码 No. 211100B0/035082

兹证明： 所附文件的影印件与原件相符。

THIS IS TO CERTIFY THAT: the annexed photostated copy of DOCUMENT is in conformity with the original.

China Council for the Promotion of International Trade

授权签字：

Authorized Signature: Lyu Cuilian

日期：2021年06月02日

(Date: Jun. 02, 2021)

证书查询网址 Website for verifying the certificate: http://www.rzccpit.com/validate.html

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

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 057996 0014 Rev. 01

Manufacturer:

Weifang KAWA Medical Products Co., Ltd.
No.2 Xingan Road
Shouguang Beiluo Industrial Park
262700 Weifang, Shandong
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Sterile syringe for single use, Sterile Infusion sets for single use, Sterile Transfusion set for single use, Sterile Volumetric Burette infusion set, Sterile scalp vein set for single use, Sterile IV catheter for Single Use, Sterile stomach catheter for single use, Sterile urine catheter for single use, Sterile Phlegm Suction Tube for single use, Sterile feeding tube for single use, Sterile redon wound drainage system for single use, Disposable Air Filter, Disposable Chest drainage system, Sterile Manifold, Sterile and Non-sterile nebulizer masks, Sterile and Non-sterile oxygen masks, Sterile and Non-sterile venturi masks.

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.: BJ19832071

Valid from: 2020-02-24
Valid until: 2024-05-26

Date, 2020-02-24

Christoph Dicks
Head of Certification/Notified Body

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

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Facility(ies):

Weifang KAWA Medical Products Co., Ltd.
No.2 Xingan Road, Shouguang Beiluo Industrial Park, 262700
Weifang, Shandong, PEOPLE'S REPUBLIC OF CHINA

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERT

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