ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-MDR-099

Product Service

EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III (Class IIa and Class IIb Devices)

No. G10 036336 0058 Rev. 01

Manufacturer:

Zhejiang Kindly Medical Devices Co., Ltd.
No.758, 5th Binhai Road
Binhai Industrial Park, Longwan District
325025 Wenzhou, Zhejiang Province
PEOPLE'S REPUBLIC OF CHINA

SRN Manufacturer: CN-MF-000007594

Authorized Representative:

Shanghai International Holding Corp. GmbH (Europe)
Eiffestraße 80, 20537 Hamburg, GERMANY

The Certification Body of TÜV SÜD Product Service GmbH certifies that the manufacturer has established, documented and implemented a quality management system as described in Article 10 (9) of the Regulation (EU) 2017/745 on medical devices. Details on device categories covered by the quality management system are described on the following page(s).
The Report referenced below summarises the result of the assessment and includes reference to relevant CS, harmonized standards and test reports. The conformity assessment has been carried out according to Annex IX Chapter I and III of this regulation with a positive result.
The quality management system assessment was accompanied by the assessment of technical documentation for devices selected on a representative basis.
The certified quality management system is subject to periodical surveillance by TÜV SÜD Product Service GmbH. The surveillance assessment shall also include an assessment of the technical documentation for the device or devices concerned on the basis of further representative samples.
All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with.
For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G10 036336 0058 Rev. 01

Report No.: BJ21081202

Preceding Certificate No.: G10 036336 0058 Rev. 00

Valid from: 2022-07-01
Valid until: 2026-04-14

Date of Initial Issuance: 2021-04-15

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2022-07-01

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123
TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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Classification:	IIa
Device Group:	A010105 - NEEDLES FOR COLLECTION UNDER VACUUM
Intended Purpose:	-

Classification:	IIa
Device Group:	A020106 - INSULIN SYRINGES, SINGLE-USE
Intended Purpose:	-

Classification:	IIb
Device Group:	C010101 - PERIPHERAL I.V. CATHETERS
Intended Purpose:	Sterile I.V. Catheter for Single Use is intended for medication infusion when assembled with appropriate matching medical devices such as disposable syringe, infusion set or pressure infusion device.

Classification:	IIb
Device Group:	A010102 - BUTTERFLY NEEDLES
Intended Purpose:	Scalp vein sets is intended to be used with disposable syringe, infusion set for intravenous medication infusion.

Classification:	IIa
Device Group:	A010101 - HYPODERMIC NEEDLES
Intended Purpose:	-

Classification:	IIa
Device Group:	A010401 - ARTERIOVENOUS FISTULA NEEDLES
Intended Purpose:	-

Classification:	IIa
Device Group:	A010601 - CARPULE NEEDLES
Intended Purpose:	-

The validity of this certificate depends on conditions and/or is limited to the following: - none -

Revision History:

Rev.	Dated	Report
00	2021-04-15	BJ20081202

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