EU Certificate

Production Quality Assurance
REGULATION (EU) 2017/745 on Medical Devices
Annex XI Part A

Registration No.:	DZ 2028764-1
Manufacturer:	Jiangsu Intco Medical Products Co., Ltd. No. 298, Yandunshan Road, Zhenjiang New District 212132 Jiangsu P.R. China
EUDAMED Single Registration No.:	CN-MF-000016338
Products:	Products of class IIa: Z120602 - PHYSIOTHERAPY EQUIPMENT - Cold Packs - Hot Packs - Warmers - Heat Patches
Authorized representative(s):	MedNet EC-REP C IIb GmbH Borkstrasse 10, 48163 Münster, Germany

Certificate history		
Revision:	Description:	Issue date:
0	Initial certification	2025-10-17

The Notified Body hereby declares that the requirements of Annex XI, Part A of the REGULATION (EU) 2017/745 have been met for the listed products. The above named manufacturer has established and applies a production quality assurance, which is subject to periodic surveillance, defined by Annex XI, Part A, Section 7 of the aforementioned regulation.
If class III devices or class IIb devices are covered by this certificate, an EU type-examination certificate in accordance with Annex X of the aforementioned regulation is required before placing them on the market.

Report No.:	244498705-200
Effective date:	2025-10-17
Expiry date:	2030-10-16
Issue date:	2025-10-17

Jason Pan
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This certificate can be validated on https://www.certipedia.com

TÜV Rheinland LGA Products GmbH is a Notified Body according to REGULATION (EU) 2017/745 concerning medical devices with the identification number 0197.

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