中国国际贸易促进委员会

中国国际商会

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

证明书

CERTIFICATE

243600B0/H00090

兹证明: 在所附文件上的江西蓝博医疗制品有限公司的印章属实。

THIS IS TO CERTIFY THAT: the seal of JIANGXI RAINBOW MEDICAL PRODUCTS CO., LTD on the annexed DOCUMENT is genuine.

China Council for the Propotion of International Trade

号码 No.

签证员:

Authorized

Official:

日期: 2024年06月20日

(Date: Jun. 20, 2024)

书查询网址 Website for venifying the certificates http://www.rzccpit.com/validate.html

Zhang Xin

Enterprise/Company Statement

The enterprise/company applied the certificate issued by CMC MEDICAL DEVICES & DRUGS S.L. on April 23, 2021 EC REP CERTIFICATION NO. CMC/CE/2021/23042021.11 , proving the conformity declaration of our company's CE.

We hereby declare that :our company guarantees that the certificate content attached to the statement is true, legal, and valid, and there is no forgery or tampering with any information. If there is a violation of relevant regulations, such as forgery, tampering, etc. that does not match the facts, our company shall bear the responsibility arising from this.

B

Attached : Copy of the certificate

Jiangxi Rainbow Medical Products CO., LTD

2024/...6/...19

13/61

C REP CERTIFICATE



CMC MEDICAL DEVICES & DRUGS SL NO. CMC/CE/2021/23042021.11

CONFIRMED THAT CMC MEDICAL DEVICES & DRUGS S.L. Is the European Authorized Representative of

Jiangxi Rainbow Medical Products Co.,Ltd
East side of Jianshe Road, Penghuwan Area, Jiujiang City,
Jiangxi Province, China

The certificate remains valid until the expiration agreement of EC REP, manufacturing conditions, the quality system or relevant legislation are changed. The validity is conditioned by positive results of periodic surveillance audits.

The product liability rests with the manufacturer in accordance with applicable directive and standard, after fulfilling of the relevant EU legislation requirements, the manufacturer shall affix relevant CE marking to all above mentioned models of the medical device.

Manufacturer declares that complies with the applicable Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April on Medical Devices

The products in Annex I was registered in Spanish MOH with number RPS/794/2021

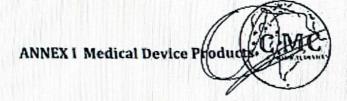
CE

Issued on: 23/04/2021

Valid until: 24/04/2026

CMC Medical Devices & Drugs SL

EC REP CERTIFICATE



Disposable Vinyl/Nitrile Blend Examination Gloves for Single Use (MDR)

Disposable Vinyl Examination Gloves for Single Use (MDR)



