EC REP CERTIFICATE

CMC MEDICAL DEVICES & DRUGS SL
NO. CMC/CE/2020/26082020.5

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

Shenzhen Shiqiao Science and Technology Co.LTD
B2904,Nanshan software park,No.10128,Shennan Avenue, Liancheng community, Nantou street, Nanshan district, Shenzhen city, Guangdong 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.

Complies with the applicable essential requirements of the council directive 93/42/EEC on medical devices as amended.

The products in Annex I was registered in Spanish MOH with number **RPS/2010/2020**

CE

Issued on: 26/08/2020

Valid until: 25/08/2021

Authorized Signatory
CMC Medical Devices & Drugs SL

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ANNEX I Medical Device Products

Medical Nitrile Inspection Gloves

CE