Fiscal Year 2020
FDA REGISTRATION CERTIFICATE

Certificate No.: JF-FDA-0328-0116

Certificate Holder:

SHENZHEN CENTURION TECHNOLOGY CO.,LTD

301.BUILDING A,38 MINAN ROAD,PINGHU STREET,LONGGANG DISTRICT

Shenzhen Guangdong,cn 518011

has completed the FDA Establishment Registration (as manufacturer , foreign exporter, contract manufacturer) and Device Listing with the US Food & Drug Administration.

Registration Number: N

Owner/Operator Number: 10066564

Device Listing:

Device#	Product Codes	Device Name
D384209	QKR	Disposable Medical Mask,KN95 Disposable respirator,KN95 protective mask

Registration Expiration Date: 2020-12-31

J&F TECHNOLOGY SERVICES LLC has verified and declares that the above stated facility is registered with the US Food & Drug Administration, Center for Drug Evaluation and Research, Office of Drug Registration and Listing pursuit to the Code of Federal Regulation 21 CFR 207, on the data state above, and makes no other representations and warranties, nor does this certificate makes other representations and warranties to other person or entity other than the name certificate holder, for whose sole benefit it is issued. J&F TECHNOLOGY SERVICES LLC assumes no liability to any person or entity in connection with the foregoing. J&F TECHNOLOGY SERVICES LLC is a private registration agent and is not affiliated with the US Food and Drug Administration.

J&F TECHNOLOGY SERVICES LLC.

2424 Morris Ave 818 Union

NEW JERSEY 07083

United States

Review Report –审查报告–검토 보고서– Rapport d'Evaluation

CE Documentation Review

No. 4Q200407M.SCTUU98

Holder: Shenzhen Centurion Technology Co., Ltd.
301, Building A, 38minan Road, Pinghu Community, Pinghu Street, Longgang District, Shenzhen

Review goal: Verification of the presence of Technical Documentation compatible with the Medical Devices Directive 93/42/EEC Annex VII

Product: Single-Use Medical Face Mask (Not Sterile)
Model(s): YY01, YY02
Classification: Class I (Not Sterile)
(accordingly to the Manufacturer's declaration)

Review output: This document has been issued on a voluntary basis and upon request of the manufacturer. It is our opinion that the Technical Documentation shared with us by the manufacturer is compatible with the European Standard for Medical Devices.
The manufacturer is responsible for the CE Marking process, and not exempted to carry out all necessary compliance activities. This document has been issued on the basis of the regulation on ECM Voluntary Mark for the certification of products. RG01_ECM rev.3 available at: www.entecerma.it

Date of issue 07 April 2020
Approver
ECM Service Director
Luca Bedonni

Expiry date 06 April 2025

Technical Expert
Amanda Pavone

Ente Certificazione Macchine
Via Cà Bella, 243 - 40053 Valsamoggia Loc. Castello di Serravalle (Bo) Italy
☎ +39.0516705141 🖷 +39.0516705156 ✉ info@entecerma.it 🌐 www.entecerma.it