



Shenzhen Tian Hai Test Technology Co.,Ltd.

CE Certificate of Conformity

Certification number

Report number:

Shenzhen Tian Hai Test Technology Co.,Ltd. hereby declares that testing has been completed and reports have been generated for:

Applicant:

Address: 118-3 leye street, shenbei new district, shenyang city, liaoning province, China

Manufacturer:

Address: 118-3 leye street, shenbei new district, shenyang city, liaoning province, China

Product: Medical Protective Mask

Trade mark:



Model: Folding type

And, in accordance with the following applicable directives:

(EU) 2016/425 Personal Protective Equipment(PPE)

This product has been assessed against the following applicable standards;

Standard: EN 149:2001+A1:2009

Therefore, Shenzhen Tian Hai Test Technology Co.,Ltd. hereby acknowledges that the applicant may issue a DECLARATION of CONFORMITY and apply the CE marking in accordance with European Union Rules.

Attestation by:



Thomas Wong

Date of Issue: March 10, 2020



4F,A3 BLDG,The Silicon Valley Power intelligent terminal industrial park,Guan lan street,Longhua district,Shenzhen
Tel:+86-755-86615100 Fax:+86-755-86615105 <http://www.tianhaitest.com>



Fiscal Year 2020 CERTIFICATION OF FDA REGISTRATION

Cert. No.:

This certifies that:

Establishment:

Registered Address: 118-3 Leye Street, Shenbei New District Shenyang City,
Liaoning, 110141, CHINA

Registration Number: Not Yet Assigned

has completed the FDA Establishment Registration (as manufacturer) and Device Listing with the US Food & Drug Administration, such registration having been verified as currently effective on the date hereof by SPICA MEDTECH CORP.

Owner/Operator Number: (Device Listing#: See annex)

U.S Agent for FDA: SPICA MEDTECH CORP

Communications: 2255 EMERSON ST Denver,CO,80205,USA

Phone: 720 6176666 Ext. Email: spica_us@yahoo.com

SPICA MEDTECH CORP will confirm that such registration remains effective upon request and presentation of this certificate until the end of the calendar year stated above, unless said registration is terminated after issuance of this certificate.

SPICA MEDTECH CORP makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. This certificate does not denote endorsement or approval of the certificate-holder's device or establishment by the U.S. Food and Drug Administration.

Pursuant to 21 CFR 807.58, "Registration of a device establishment or assignment of a registration number does not in any way denote approval of the establishment or its products."

SPICA MEDTECH CORP (in USA)
SHANGHAI SPICA MANAGEMENT CONSULTING CO.,LTD (in China)
SPICA MEDTECH CERTIFICATION COMPANY LIMITED (in UK)
spica_us@yahoo.com, www.spicaglobal.com



Spica Consulting
Regulation Services

For and on behalf of
SPICA MEDTECH CORP

Authorized Signatory

SPICA CERT. NO. 218 Rd/Reg.No. 593 Eshai Road, Shanghai 200120, P.R. China

