Fiscal Year 2020
FDA REGISTRATION CERTIFICATE

Certificate No.: JF-FDA-0330-0101

Certificate Holder:

[redacted]

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

Registration Number: 3005070829

Owner/Operator Number: 10051617

Device Listing#: See annex

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

FDA

Fiscal Year 2020
FDA REGISTRATION CERTIFICATE

Annex to Cert. No.: JF-FDA-0323-0101

Device#	Product Codes	Device Name
D264142	FPP	STRETCHER, HAND-CARRIED (Hand-carried stretcher])
D264143	FPO	STRETCHER,WHEELED (Wheeled stretcher, power assisted stretche)
D377462	KHA	MASK, SCAVENGING (DisposableProtective mask, Disposable medical mask)
D381923	MSH	Respirator, surgical (N95)

END OF THE ANNEX

The 3rd Party Certificate of

FDA Facility Registration & Listing

Note: This document is Not issued by FDA. We,J*F, as a 3rd party, created it to make easier for customer to present information. The following contents, FDA registered establishment & product information, are excerpted from the database at www.fda.gov.

Establishment / Firm:

[redacted]

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

Registration Number: 3005070829

Owner/Operator Number: 10051617

Device Listing#: See annex

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J&F TECHNOLOGY SERVICES

19838 Moon Shadow Circle

Walnut, California, 91789

United States

The 3rd Party Certificate of

FDA Facility Registration & Listing

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Annex to Cert

Device#	Product Codes	Device Name
D393679	QKR	Face mask (except N95 respirator) for general public/healthcare personnel per IIE guidance (DisposableProtective mask, Disposable medical mask)
D381923	MSH	Respirator, surgical (N95, KN95 MASK)
D264142	FPP	STRETCHER, HAND-CARRIED (Hand-carried stretcher])
D264143	FPO	STRETCHER,WHEELED (Wheeled stretcher, power assisted stretche)
D403709	OEA	Non-surgical isolation gown （PROTECTIVE CLOTHING）
D403710	HOY	Shield, eye, ophthalmic (including sunlamp protective eyewear and post-mydriatic eyewear) （GOGGLE）
D381890	LYU	ACCESSORY, SURGICAL APPAREL

END OF THE ANNEX