Fiscal Year 2021

CERTIFICATION OF REGISTRATION

This certifies that:

Dongguan Quanding Medical Supplies Co., Ltd
3 Yongfa East Road, Qishi Town, Dongguan City, Guangdong , China

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

GUANGZHOU LETA TESTING TECHNOLOGY CO.,LTD

Owner/Operator Number: 10075777
Device Listing#:

Listing Number	Listing Status	Product Code(s)	Device Name	Product Number	Reference
D439743	Active	GXY	ELECTRODE, CUTANEOUS/ Adhesive Electrode pads	TE-A20, TE-A21, TE-A22, TE-A23, TE-A24; TE-B20,TE-B21, TE-B22, TE-B23,TE-B24;; TE-C20, TE-C21, TE-C22, TE-C23, TE-C24;TE-D20, TE-D21, TE-D22,TE-D23, TE-D24; TE-E20,TE-E21, TE-E22, TE-E23, TE-E24.	K171381

LETA Testing Technology will confirm that such registration remains effective upon request and presentation of this certificate until thr end of calendar year stated abouve,unless said registion is terminated after issuance of this certificate.LETA Testing Technology makes no other represe ntations 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 isissued. This certificate does not denote endorsement or approval of the certificate-holder,advice or establish ment by the U.S.Food and Drug Administration.LETA Testing Technology assumes no liability to any person or entity in connection with the foregoing.

FDA

GUANGZHOU LETA TESTING TECHNOLOGY CO.,LTD
6F, No.1 TianTai Road, Science City, LuoGang District
Guangzhou,CN

Executive Director
Jan.26,2021
CERT2021022601