

Fiscal Year 2020
CERTIFICATION OF REGISTRATION

This certifies that:

Minghe Lixin Medical Device Co., Ltd
Daxipen, Yingping Road, 315880 Yuyao City Ningbo Province
China Yuyao Zhejiang, China 315488

has completed the FDA Establishment Registration (as manufacturer, design expert, contract manufacturer) and Device Listing with the U.S. Food & Drug Administration, being:

U.S. Agency (FDA)
Commissioner

MINHE TECHNICAL SERVICE INC.
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Registration Number: 1412900228

Device Listing: See annex

Minghe Technical Service Inc. will confirm that each registration/extension effective upon completing presentation of this certificate until the end of the calendar year listed above, unless said registration is terminated after expiry of the complete MINHE Technical Service Inc. model for other representations or activities, nor does the certificate state any representations or activities in any period 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. MINHE Technical Service Inc. assumes no liability in any period or entity in connection with the drug(s).

Annex to 21 CFR 812.18: "Registration of a device establishment or assignment of a registration number does not in any way denote approval of the establishment or its products, any representations that create an impression of official approval, because a registration or assignment of a registration number is exclusively administrative workmanship." The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. MINHE Technical Service Inc. is not affiliated with the U.S. Food and Drug Administration.



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