Fiscal Year 2020
CERTIFICATION OF REGISTRATION

This certifies that:

GUANGDONG AOZHEN PHARMACEUTICAL CO.LTD

28 South Yihe Boyi Road, BOLUO County Huizhou, Guangdong, 516100, CHINA

has completed the FDA Establishment Registration and Device Listing with the US Food & Drug Administration, through For any registration issues, please Contact SpecifyA@163.com

Owner/Operator Number: 10063671

Device Listing#: See annex

Shenzhen HRC Testing Technology Co., Ltd 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. Shenzhen HRC Testing Technology Co., Ltd 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.

Shenzhen HRC Testing Technology Co., Ltd assumes no liability to any person or entity in connection with the foregoing.

Pursuant to 21 CFR 807.39, "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 representation that creates an impression of official approval because of registration or possession of a registration number is misleading and constitutes misbranding." 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. Shenzhen HRC Testing Technology Co., Ltd is not affiliated with the U.S. Food and Drug Administration.

FDA

Chief engineer

Issued: March 22, 2020

Expiration Date: December 31, 2020