Fiscal Year 2020

CERTIFICATION OF REGISTRATION

This certifies that:

Name: ANHUI [redacted] TECHNOLOGY CO., LTD.

Add: 106 Innovation Avenue,High-Tech Development Zone

Hefei Anhui, CN 230088

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

The Owner/ Operator Number for this Registration is :3013482554

Listing No	Code	Premarket Submission NO.	Device Name
D374223	QJR		COVID-19(SARS-CoV-2)IgM/IgG Antibody Test Kit(Colloidal Gold); COVID-19(SARS-CoV-2)IgM Antibody Test Kit(Colloidal Gold); COVID-19(SARS-CoV-2)IgG Antibody Test Kit(Colloidal Gold)

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

ABmed assumes no liability to any person or entity in connection with foregoing.

Date of verification:Feb 12, 2020

Date of expiration: Dec 31, 2020

SH OFFICE

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