

CERTIFICATE OF REGISTRATION

FISCAL YEAR 2021



This certifies that: **Zhejiang Ailebao Medical Technology CO., LTD**

No.192 Xingxian Rd,
Longgang, Wenzhou,
Zhejiang, CHINA, 325802

is registered with the U.S. Food and Drug Administration for FY 2021 pursuant to Title 21, 807 et seq. of the United States Code of Federal Regulations:

Establishment Registration:
Device Listing:
Owner / Operator Number:
U.S. Agent for FDA Communications:

3017448136
Scan FDA Device Listing QR Code
10076832
PureVision Ai, Inc.
111 Town Square Place, Suite 1203, Jersey City, NJ 07310
Telephone: +1-201-503-5758 | E-mail: us-agent@purefda.com



FDA REGISTERED
ESTABLISHMENT

PureVision Ai, Inc. 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. PureVision Ai, Inc. 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. PureVision Ai, Inc. 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. PureVision Ai, Inc. is not affiliated with the U.S. Food and Drug Administration.



DJ Fang
Executive Director
Issued: January 11, 2021
PureFDA Certificate No.: 2021USTTAR613
Expiration Date: December 31, 2021



FDA DEVICE
LISTING