Fiscal Year 2024

CERTIFICATION OF REGISTRATION

This certifies that:

DONG GUAN TOP YAO INDUSTRY CO.,LTD

Address: Rm,401,Building 1, NO.56, Dongcheng Duan, GuangzhangRoad , Dongcheng Street, Dongguan, Guangdong, 523122, China

has completed the FDA Establishment Registration and Device Listing with the US Food & Drug Administration, with below listing information:

Owner/Operator Number: 10079187

Device Listing #: See annex

through the U.S Agent

KingDaBio International Corp

Address: 3206 Erie Blvd E Syracuse, NY, De Witt, New York, 13214, UNITED STATES

KINGDABIO INTERNATIONAL CORP 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. KINGDABIO INTERNATIONAL CORP 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. KINGDABIO INTERNATIONAL CORP 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, KINGDABIO INTERNATIONAL CORP is not affiliated with the U.S. Food and Drug Administration.

FDA

Fiscal Year 2024

CERTIFICATION OF REGISTRATION

Registration Number: 3017961176
Annex to Device Listing# for Owner/Operator Number: 10079187
Product Code: ILY

Listing No.	Device Names	Proprietary Names	Activity
D427135	Far Infrared Heating Pad	FHP-20-4030 / FHP-20-6040 / FHP-20-8060 / FHP-20-12060 /FHP-20-16080/FHP-21-4030 / FHP-21-6040 / FHP-21-12060 /FHP-21-16080 / FHP-21-8060	Manufacturer
D427135	Red & Infrared Light Therapy Bulb Black & Silver	RLT-1137-S / RLT-1135-R / RLT-1136-R / RLT-1135-RI / RLT-1136-RI； RLT-S22-WT/ RLT-S21-WTP / RLT-S22-WTV24 / RLT-S22-WTV30	Manufacturer
D427135	Red Light & Near Infrared Light Therapy Devices	RLT-HAND-PB / RLT-F / RLT-F02 / RLT-F-PB / RLT-KNEE / RLTKNEE-PB/ RLT-WB01 / RLT-WB02 / RLT-WB-PB / RLT-HAND / RLT-HAND02/ RLT-D20-CF / RLT-S20-CF / RLT-FB01 / RLT-S1135-S /RLT-S1135-B/ RLT-S20-WT / RLT-S20-WT02 / RLT-S20-WT-PB /RLT-S20-HD / RLT-S20-HD02 / RLT-S20-FT / RLT-S20-FT02/RLTS20-FB / RLT-D21-SHD / RLT-S21-SHD/ RLT-8000B/RLTHELMET/RLT-S20-HELMET； Model: RLT-S20-FB/ RLT-S22-SHDNKP;RLT-S22-CFP02; RLT-HELMET / SCTM23-WB / HPT-V22-WT / RLT-FT001 / UV23-311-WB；RLT-S20-HDP / RLT-S20-HDP02；RLT-S21-FTPS / RLT-S21-FTP；RLT-S23-USBWT； RLT-MVS23-WTP；RLT-MS23-SHDP SCTM23-WB02；RLT-Helmet；UV2212	Manufacturer
FDA registration website: https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfRL/rl.cfm?lid=710151&lpcd=ILY			

Attestation by:

Execute Manager

FDA

Issued: Oct 19, 2024
Expiration Date: December 31, 2024