Certificate Of FDA Registration

2016-2020

This is certified that:
At The Address Stated Below Has Completed U.S. FOOD And DRUG ADMINISTRATION Medical Device Registration Through MANTONG.

VIVO OPTICS CO., LTD

ROOM 502~504, BLOCK B CENTRAL TOWERS,
NO. 567 LANGAO ROAD, PUTUO DISTRICT, SHANGHAI, CHINA

Owner/Operator Number: 10050758 **Registration Number** 3013072273

1. Proprietary Device Name	Optical Glass; Plastic Lens; Resin Lens
Device Name(s)	Lens, spectacle, non-custom (prescription)
Device Listing Number	D256900
2. Proprietary Device Name	Frame Spectacle
Device Name(s)	Frame, Spectacle
Device Listing Number	D353971
3. Proprietary Device Name	Magnifying Spectacles
Device Name(s)	Spectacle, Magnifying
Device Listing Number	D354482

The annual establishment registration fee must be paid between Oct. 1, and Dec. 31,

MTG FDA REGISTER CENTER CORPORATE SEAL CALIFORNIA

Jacky M. Chuang

Executive Director
Date: 12-10-2019

MTG STANDARDS TESTING & CERTIFICATION CENTER www.fdacn.org

This certification affirm that the above device and company was registered with U.S. Food and Drug Administration pursuant to section 305 of the United States Public Health and Bioterrorism Preparedness and response Act of 2002, P.L. 107-188, on the date stated above, and 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 whole sole benefit it is issued. MTG CO., Inc. assumes no liability to any person or entity in connection with the foregoing. MTG is a private registration agent not affiliated with the U.S. Food and Drug Administration