



Certificate Of FDA Registration

Fiscal Year 2020

This is certified that:

At The Address Stated Below Has Completed U.S. FOOD AND DRUG ADMINISTRATION Medical Device Registration Through MANTONG.

GUILIN REFINE MEDICAL INSTRUMENT CO., LTD.

Address: NO.8-3, INFORMATION INDUSTRIAL PARK, HIGH-TECH ZONE, QIXING DISTRICT
GUILIN, GUANGXI 541004, CHINA

Owner/Operator Number 10064688

Device Listing Number See annex

- The FDA annual establishment registration fee must be paid between Oct. 1 and Dec. 31 of every year.
- You can search the FDA registration information on this website or scan QR code: <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfRL/rl.cfm>
- Cert No.: 2008290MTG



July M. Chuang

Executive Director

Date: 03-30-2020

MTG STANDARDS TESTING & CERTIFICATION CENTER www.fda.cn.org

This certification affirms that the above device and company was registered with the 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 certification make any representations or warranties to any person or entity other than the named certificate holder, for whose 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.



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Annex to Certificate No.: 209829MTG

Listing Number

D379683

Product Code

HOY

Device Name

Safety Goggle
Face Shield

END OF THE ANNEX



Jacky M. Chuang

Executive Director

Date: 03-30-2020

MTG STANDARDS TESTING & CERTIFICATION CENTER www.mtgcn.org

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