VERIFICATION OF FDA REGISTRATION

This certifies that:

Shenzhen Kingho Technology Co., Ltd
No.117, Zhangbei Road, Zhangbei Community, Longcheng Sub-district, Longgang District, Shenzhen, Guangdong, 518115, CHINA

has completed the FDA Establishment Registration and Device Listing with the US Food & Drug Administration, through UGK-LVM UNITED INC.

Owner/Operator Number: 10057591
Registration Number: 3014515315

Listing Number	Product Code(s)	Device Name(s)	Activities
D320801	HQZ	Frame, spectacle	Manufacturer;
D320800	HQY	Sunglasses (non-prescription including photosensitive)	

This is a formal notice upon your company that your product applied has been successfully registered by the U.S. Food and Drug Administration. The registration remains effective unless the said registration is terminated by the U.S. Food and Drug Administration. We make no other representations or warranties, nor does this certificate make sole benefit as it is issued. This notice does not denote endorsement or approval of the certificate-holder`s device or establishment by the U.S. Food and Drug Administration. We assume no liability to any person or entity in connection with the foregoing. 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. We are not affiliated with the U.S. Food and Drug Administration.

Cert. No.: E20835
Issued Date: 3 January 2022
Expiration Date: 31 December 2022