VERIFICATION OF FDA REGISTRATION

This certifies that:

NANCHANG FIRSTOMATO MEDICAL DEVICES CO., LTD
No.555 Aixihu 2 road, NanChang, Jiangxi, 330000, CHINA

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

Owner/Operator Number: 10087316
Registration Number: /

Listing Number	Product Code(s)	Device Name(s)	Activities
D496114	JYS	PERFORATOR, EAR-LOBE - Firstomato Piercing Instruments	Manufacturer;

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.: E23025
Issued Date: 12 Feb. 2023
Expiration Date: 31 Dec. 2023