Our Ref: SH-FDA-2024102401
November 05, 2024

FDA REGISTRATION CONFIRMATION

Thank you for appointing Frosa Technology LLC as your U.S. Agent, we hereby state that we accepts the appointment to be the U.S. Agent for your medical device facility, and we have completed the registration activation confirmation for the FDA Medical Device Registration and Listing Module with the US Food & Drug Administration for the Year 2025 of

The Registration information is as below:

Owner/Operator Number	**10090804**
Establishment Registration Number	**3031589938**

listing No	Product Name(s)	Product code	Activites
D548713	Tape And Bandage, Adhesive - (Foot Pad; Slimming Patch; Navel Patch; Detox Foot Patch; Foot Patch; Warm Foot Patch; Rheumatism Patch)	KGX	Foreign Exporter; Manufacturer
D536080	Pack, Hot Or Cold, Disposable - Ankle Patch; Body Warmer; Cold Compress Eye Mask; Cool Eye Mask; Foot Warmer; Hand Warmer; Heat Patch; Hot Or Cold Disposable Pack; Hot Or Cold Disposable Pad; Hot Patch; Knee Warmer; Menstrual Warmer Patch; Neck Warmer; Steam Eye Mask; Steam Eye Patch; Warm Neck Patch; Warm Patch; Warm Waist Patch; Warm Waistband; Warming Patch	IMD	Foreign Exporter; Manufacturer
D548712	Shield, Eye, Ophthalmic (Including Sunlamp Protective Eyewear And Post-Mydriatic Eyewear) - STEAM EYE MASK	HOY	Foreign Exporter; Manufacturer

NOTE: This file is the rendering of the Registration of Medical devices Product Facility page, not the Registration certificate. This file makes no other presentations or warranties, nor does this certificate make any representation or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. This file 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.