FDA REGISTRATION NUMBER: 3013513780
LAST VALIDATED: December 13, 2023
STATUS: Valid

VALID **2024 U.S. FDA Medical Device Establishment Registration**

JIANGMEN K.K.PLASTIC FACTORY LTD.

ESTABLISHMENT INFORMATION

NO.193, 3rd Street Jian She RD
Jiangmen Guangdong, CN 529000

86-750-3412154
joyce@jkkp.com.cn
http://www.jkkp.com.cn/zh-CN/introduce.html

U.S. AGENT AND OFFICIAL CORRESPONDENT

Registrar Corp
144 Research Drive
Hampton VA, USA 23666

+1-757-224-0177
info@registrarcorp.com
www.registrarcorp.com

Registration of a device establishment, assignment of a registration number, or listing of a medical device does not in any way denote approval of the establishment or its products by U.S. FDA.

LISTINGS ON FILE WITH FDA

Device Name	Product Code	Status	Last Validated
Dispenser, Liquid Medication	KYX	Valid	December 12, 2023