FDA & CE CERTIFICATE

Our products have been FDA Registered and CE Certificate.

Medi-19 brand has also been FDA registered, and is in the process of receiving a CE Certificate as well.

CERTIFICATE OF MEDICAL DEVICE REGISTRATION
2020

This Certifies that:

NAM ANH EQUIPMENT TRADING PRODUCTION CO., LTD

Add: No. 8-8A Le Lang Street
Phu Tho Hoa Ward, Tan Phu District, Ho Chi Minh City, Viet Nam

Owner/Operator Number: 10072150

Device Listing:

D397234	Face mask (except N95 respirator) for general public/healthcare personnel per IIE guidance
D397235	Gown, isolation, surgical
D397236	Shield, protective, personnel
D397237	Patient examination glove
D397238	Bariatric bed
D39723	Cabinet, table and tray, anesthesia
D397241	Disinfectant, medical devices

Proprietary Name: FAMAPRO, MEDINATION
Certificate Issue Date: 04/26/2020

Disclaimer:
This Certificate is only notification of Medical Device Registration currently in active and effective.
This Certificate makes no other representative or warranties nor representative or warranties any proven of entry.
This Certificate will be expired one year from the Certificate Issue Date.
This Certificate validity can be checked by owner/operation number and below link:
https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfRLM.cfm

FDA

BRANDS IN USA OFFICIAL SEAL LOS ANGELES CA

Certificate

CERTIFICATE of CONFORMITY

APPLICANT : NAM ANH EQUIPMENT TRADING PRODUCTION COMPANY LIMITED
No 8-8A Le Lang Street, Phu Tho Hoa Ward, Tan Phu District, Ho Chi Minh City, Vietnam

MANUFACTURER : NAM ANH EQUIPMENT TRADING PRODUCTION COMPANY LIMITED
No 8-8A Le Lang Street, Phu Tho Hoa Ward, Tan Phu District, Ho Chi Minh City, Vietnam

Product Name : Non sterile medical mask
Trade mark : Famapro
Product Classification : Class I
Applied Directive(s) : 93/42/EEC Medical Device Directive

This certificate has been issued according to the voluntary application of the manufacturer. This certificate is valid only for the product(s) in above mentioned. This certificate confirms that the declaration of conformity which is prepared by manufacturer is available. It's manufacturer's sole responsibility to meet all the necessary conformity assessment activities according to 93/42/EEC Medical Devices Directive and related standards or the mentioned product(s) before placing them on the market. It is the manufacturer's responsibility to comply a full technical file according to Annex VII of 93/42/EEC and the manufacturer is the responsible to take all necessary actions before affixing CE mark on the product(s)

Date of issue of this certificate : 02.04.2020
Expiry date : 02.04.2021
Certificate Number : 26.02.530-02

CE

Certificate of Registration

This certifies that the above-named facility is registered with the U.S. Food and Drug Administration pursuant to the regulations required by the US Laws. The registration has been completed effective by Willow Glen Consultancy on the date below, and any registration has been issued after issuance of this certificate. Willow Glen Consultancy makes no additional representations or warranties, and neither this certificate nor any of its contents may be used by any person or entity other than the named facility without prior written approval of Willow Glen Consultancy.

MEDI 19 CORP

1961 Capital Ave, Garden Grove, CA 92843, USA

Owner Operator Number: 10033202

US Agent: Willow Glen Consultancy LLC
Willow Glen Number: 3002871209
Registration Date: December 31, 2020

Willow Glen Consultancy is a private agent not affiliated with the U.S. Food and Drug Administration.

Amanda Do

Willow Glen

Amanda Do, Operation Director
Willow Glen Consultancy LLC
+1-415-569-1389
Date: May 25, 2020

Willow Glen Consultancy
Email: Mail@willowglen.com
3755 Avocado Blvd, Suite 206
La Mesa, CA 91941, USA

QSA INTERNATIONAL

Certificate

CE

CGA