Certificate Of FDA Registration

2018

This is certified that:

At The Address Stated Below Has Completed U.S. FOOD And DRUG ADMINISTRATION Medical Device Registration Through MANTONG.

WUHAN HUAWEI TECHNOLOGY CO., LTD

B-F11-3, Yangluo Port,Huazhong International Industrial Park, Wuhan, Hubei, China 430415

FEI Number: 3010370474 Owner/Operator Number: 10043381

1、Device Listing Number D183576

Device Name: Tape And Bandage, Adhesive

2、Device Listing Number D183584

Device Name: Pack, Hot Or Cold, Disposable

3、Device Listing Number D183585

Device Name: Bandage, Elastic

———The FDA annual establishment registration fee must be paid between Oct. 1 and Dec. 31 every year.

MTG FDA REGISTER CENTER CORPORATE SEAL CALIFORNIA

Jacky M. Chuang

Executive Director

Date: 01.11.2018

MTG STANDARDS TESTING & CERTIFICATION CENTER www.fdacn.org

This certification affirm that the above device and company was registered with U.S. Food and Drug Administration pursuant to section 305 of the United States Public Health and Bioterrorism Preparedness and response Act of 2002, P.L. 107-188,on the date stated above, and makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whole sole benefit it is issued. MTG CO., Inc. assumes no liability to any person or entity in connection with the foregoing. MTG is a private registration agent not affiliated with the U.S. Food and Drug Administration