# CERTIFICATE OF U.S. FDA REGISTRATION

This certifies that

## AP MDS CO.,LTD.

*M-1804/3203, 32, SONGDOGWAHAK-RO, YEONSU-GU INCHEON, 21984, KOREA, SOUTH*

Owner/Operator Number: 10083007

Registration Number: Not assigned yet

is registered and has listed the following medical device with the U.S. Food and Drug Administration for FY 2024 pursuant to Title 21, CFR Part 807 of the United States Code of Federal Regulations

### Device Listing:

| Listing Number | Proprietary Name | FDA Product Code |
|---|---|---|
| D454027 | Two-way and Three-way Disposable Silicone Foley catheters | EZL |
| D533266 | Silicone Bandage | KGX |

Expiration Date: April 11, 2024 ~ December 31, 2024

Above registration remains effective upon request and presentation of this certificate until the end of the year stated above, unless said registration is terminated after issuance of this certificate.
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. WISE COMPANY Inc. is not affiliated with the U.S. Food and Drug Administration.

Date: April 11, 2024
Place of Issue: Seoul, Korea

CERTIFIED

Sanglok, LEE
Director of WISE COMPANY Inc.