Certificate Of FDA Registration

This is certified that:
At The Address Stated Below Has Completed U.S. FOOD And DRUG ADMINISTRATION Medical Device Registration Through MANTONG.

Xiamen Wenatone Medical Technology Co., Ltd.

A805, Jianye Building,Torch Hi-Tech (Xiang An) Industrial Park, Xiamen, Fujian ,China.

Registered Establishment Number:3013539171 Owner/Operator Number:10054367

1、Proprietary Device Name： Charming Sound Series;Fairy Series;IHear Series;Dragon Series;Teana Series;MDHearing Aid Air;Angel Series; Flame Series;Honor Series;King Series;Roewe Series;Apollo Series;Pando Series;Lumino Series;Link Series; Luma Series;Libra Series;Legend Series;Pure Series;Polestar Series;Xsport Series;Mini Series;Super power Series; Mini; VOLTMAX;UFO; N-I002;N-I003;N-I004;N-I005;N-B001;N-B002;N-B003;N-B004;N-B005; TWS Series;Hearbuds Series;Metone Series;Dolphin Series;F201;F202;F203;F401;F402;F403;F404;F601;F602;F801;F802; N-I001

Device Name(s) HEARING AID, AIR-CONDUCTION, PRESCRIPTION

Device Listing Number **D290771**

Product Code ESD

2、Proprietary Device Name： Apollo Series; Libra Series; Legend Series; IHear Series; Lumino Series; Deft Series; Flame Series; Angel Series; Explorer ,Zone; King Series; QUALISON TC;QUALISON BT;QUALISON HA;Clarus P;Clarus XP;Clarus T;Clarus XT;Sound Flo 217P; N-I001,N-I002,N-I003,N-I004,N-I005,N-B001,N-B002,N-B003,N-B004,N-B005,Listener ,Luma ,Lava,Link ,Era ,Pure, Forza; Sound Flo 217SP; Sound Flo 317P; Sound Flo 317UP; Sound Flo 318X; Sound Flo 318XP; Sound Flo 217UP; VoiceBud V50

Device Name(s) Face Plate Hearing Aid

Device Listing Number **D290772**

Product Code LRB

3、Proprietary Device Name： Angel CN 830S B+; Dragon CF 430 BLE; Melink BLE; R95; TEANA CF 130 B+; TEANA CF 430 B+; Vista BLE; WOLONG

Device Name(s) HEARING AID, AIR-CONDUCTION WITH WIRELESS TECHNOLOGY, PRESCRIPTION

Device Listing Number **D461342**

Product Code OSM

This certificate is valid from to Oct,2020 to Oct,2023

The annual establishment registration fee must be paid between Oct. 1, and Dec. 31,

MTG FDA REGISTER CENTER CORPORATE SEAL CALIFORNIA

Jacky M. Chuang

Executive Director

Date: 12-07-2022

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

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Registered Establishment Number:3013539171 Owner/Operator Number:10054367

1、**Proprietary Device Name：** In-The-Ear Hearing Aid; Teana series

Device Name(s) HEARING AID, AIR-CONDUCTION WITH WIRELESS TECHNOLOGY, OVER THE COUNTER

Device Listing Number **D488674**

Product Code QUG

2、**Proprietary Device Name：** Angel series; Behind-The-Ear Hearing Aid; Honor series; In-The-Ear Hearing Aid; mini series; UFO series; WoLong series

Device Name(s) HEARING AID, AIR-CONDUCTION, OVER THE COUNTER

Device Listing Number **D488675**

Product Code QUF

This certificate is valid from to Oct,2020 to Oct,2023

The annual establishment registration fee must be paid between Oct. 1, and Dec. 31,

MTG FDA REGISTER CENTER CORPORATE SEAL CALIFORNIA

Jacky M. Chuang

Executive Director

Date: 12-07-2022

MTG STANDARDS TESTING & CERTIFICATION CENTER www.fdacn.org