

EC Declaration of Conformity

Certificate No.: EU202110219

Manufacturer: **ARI Medical Technology Co., Ltd.**

Address: **Block C, CC Park, No. 728 Lanzhou Road, Baohe Industrial Zone, Hefei City, Anhui, China**

Products: **Sterilizer**

Model: **LX-B Series, YX-280 Series, YXQ.DY.250A Series, YXQ.DY.250B Series, AAL-L-B Series, AAL-L-N Series, GRF Series, WS-YDA Series, WS-YDB Series, WS-YDC Series, WS Series, WS-YV Series. APS Series, N Series, GRF Series, ZXC-II Series**

According to EC Declaration of Conformity set out in Annex VII of the Council Directive 93/42/EEC (MDD93/42/EEC) concerning medical devices.

We herewith declare that the above mentioned products meet the transposition into national law, the essential requirements set out in the Council Directive 93/42/EEC (MDD93/42/EEC) Annex VII.

The manufacturing site for the above products comply with EN ISO13485:2016 Quality Management System Requirements for Medical Devices.

UMDNS No: **13746**

Risk Class: **II**

Applied harmonized standards, national standards or other normative documents

Medical Devices - Quality Management Systems ISO 13485:2016

Identification: **CE**

Place of Issue: Hefei, Anhui.

Date of Issue: 2021-06-01

Valid Until: 2023-06-01

Signature and Stamp:

