无锡裕清医疗器械厂
WUXI YUQING MEDICAL INSTRUMENT FACTORY
NO.38,South Jingda Road, Zhangjing ,Xibei Town, Xishan District,Wuxi, China

EC Declaration of Conformity

Report Reference No. 23-10-20

We, the manufacturer, herewith declare that the products
STETHOSCOPE / ANEROID SPHYGMOMANOMETER / LATEX BULB / LATEX BLADDER / CUFF WITH(WITHOUT) BLADDER / GAUGE METER / EARPLUG / DIAPHRAGM
Model: *SINGLE HEAD / DUAL HEAD / 1 TUBE / 2 TUBE*
meet the provisions of MDR EU_2017/745 which apply to them.

The medical device has been assigned to *Class I with measuring function,* according to to the classification principle of Rule 1 in Chapter III of Annex VIII of the MDR EU_2017/745.
The product concerned has been manufactured under a quality management system according to Annex IV of the MDR EU_2017/745.
Following the procedure relating to the EC Declaration of Conformity set out in Annex VI of the MDR EU 2017/745.
This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices.
This Declaration of Conformity covers all medical devices as specified in the product list belonging to this declaration and is only valid in connection with a batch specific Certificate of Compliance for all products concerned bearing the CE mark.

The above mentioned declaration of conformity is exclusively under the responsibility of Company: *WUXI YUQING MEDICAL INSTRUMENT FACTORY*
Address: *No.38, South Jingda Road, ZhangJing, Xibei Town, Xishan District, 214194 Wuxi, China*
UDI-DI NO.:*697638463*

Place,*WUXI*
Date,*202310/20*

无锡裕清医疗器械厂 WUXI YUQING MEDICAL INSTRUMENT FACTORY

Legally Binding Signature,Management Representative
Chen Zhiqing
陈志清