Declaration of Conformity

Manufacturer:	Jousing Medical Co., Ltd.
Address:	301&401, Building 21, 200 Xingpu Road, Suzhou Industrial Park, 215000 Suzhou, Jiangsu Province, PEOPLE'S REPUBLIC OF CHINA
Authorised Representative:	Wellkang Ltd Enterprise Hub, NW Business Complex, 1 Beraghmore Road, Derry, BT48 8SE, Northern Ireland, UK.

We, the manufacturer, herewith declare that the product

Automated External Defibrillator

Model: iAED-S1

Basic UDI-DI: 697331555iAED1QC

Meet the transposition into national law, the provisions of council directive 93/42/EEC of 14 June 1993 concerning medical devices; including, at 21 March 2010, The amendments by council Directive 2007/47/EEC all supporting documentation is retained at the premises of the manufacture.

The medical device has been assigned to class IIb, rule 9 according to MDD, Annex IX. It bears the mark

CE 0123

Meet the Article 120.3c of MDR (as amended by (EU) 2023/607).

And the extension until 31 December 2027 has been completed by the EU competent authority.

Notified Body: TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339 MÜnchen, Germany

Ceitificate No: G1 096469 0002

Issued date: 2021-03-11

Expiry date: 2027-12-31

Following the procedure relating to the EC Declaration of Conformity set out in Annex IV of the Regulation(EU) 2017/745.

This declaration of conformity is valid in connection with the release document for the respective batch of produced devices.

The above mentioned declaration of conformity is exclusively under the responsibility of Company: Jousing Medical Co., Ltd.

Address: 301&401, Building 21, 200 Xingpu Road, Suzhou Industrial Park, 215000 Suzhou, Jiangsu Province, PEOPLE'S REPUBLIC OF CHINA

Yuxi Wang

QA Manager

2024-04-09

Signature: