EU Declaration of Conformity

Manufacturer:
Qualmedi Technology Co.,Ltd.
Address:
A302 Room, No.23, HangBu Road,
Feixi County Economic Development
Zone, Hefei City, Anhui Province，
China.

EC-Representative:
Kingsmead Service B.V.
Address:
Zonnehof, 36 2632 BE Nootdorp

We declare under our sole responsibility that

Products:	Vein Finder
Model:	QV-500，QV-600
UMDNS:	14346
Class:	I (According to Annex VIII Rule 13 of Regulation (EU) 2017/745）

Intended Use：Vein finder can help medical professionals to locate certain superficial veins. This equipment is intended to be used as a supplement to appropriate medical training and experience

. **BASIC UDI-DI code:** 16974011590007

Manufacturer SRN code: CN-MF-000028309

EC Representative SRN code: NL-AR-000002066

Conformity assessment procedure: Conformity assessment procedure According to Art. 52 section 1 and 7 Regulation EU 2017/745

meet the provisions of the Regulation (EU) 2017/745 in national laws which apply to it, and of RoHS Directive 2011/65/EU Annex II amending Annex (EU)2015/863 and amending Annex (EU)2017/2102

Applied standards:

EN ISO 15223-1:2021	EN ISO 14971:2019/A11:2021	ISO 13485:2016
EN ISO 20417:2021	IEC 60601-1-2:2014+A1:2020	IEC 60601-1:2005+AMD1:2012
MEDDEV 2.7.1: REV4.	EN ISO 10993-1:2020	

The CE Mark: CE

The above-mentioned declaration of conformity is exclusively under the responsibility of

Company: Qualmedi Technology Co.,Ltd.
Address: A302 Room, No.23, HangBu Road, Feixi County Economic Development Zone, Hefei City, Anhui Province，China.

Hefei，2022-9-14
Place，Date

General Manager, JASON ZHANG
Duly authorized to sign this Authorization on behalf of:
Qualmedi Technology Co., Ltd.