

The Commonwealth of Massachusetts

William Francis Galvin
Secretary of the Commonwealth

APOSTILLE

(Convention de La Haye du 5 octobre 1961)

1. *Country:* United States of America

This public document

2. *has been signed by:* Rebecca M. Davis

3. *acting in the capacity of:* Notary Public

4. *bears the seal/stamp of:* Rebecca M. Davis

whose commission expires on: June 5, 2026

Certified

5. *at:* Boston, Massachusetts

6. *the:* 26 February, 2020

7. *by:* the Secretary of the Commonwealth

8. *No.:* 2194808

9. *Seal/stamp:*

Great Seal of the Commonwealth

10. *Signature:*



William Francis Galvin
Secretary of the Commonwealth





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Certify a Copy:

Commonwealth of Massachusetts

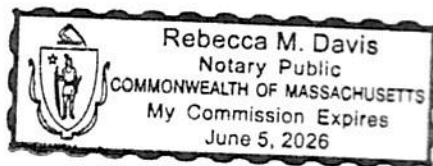
County of Middlesex

On this 21 day of February, 2020, I certify that the preceding or attached document, is a true, exact, complete, and unaltered copy made by me Rebecca Davis, presented to me by the document's custodian, Joline Palasek, and that, to the best of my knowledge, the photocopied document is neither a public record nor a publicly recordable document, certified copies of which are available from an official source other than a notary.

- Declaration of Conformity- ZOLL M2

Rebecca Davis
Notary Public Signature

Notary Stamp



Notary Seal



Declaration of Conformity



Suzhou ZOLL Medical Technology Co., Ltd.
Room102-2, Block 19, No.8 Jinfeng Road
Suzhou New District, 215163 Suzhou
Jiangsu, P.R. China

EC REP

ZOLL International Holding B.V.
Newtonweg 18
6662 PV ELST
The Netherlands

Product: ZOLL M2 Monitor/Defibrillator
UMDNS/GMDN Code: 37806
Classification (MDD, Annex IX): IIb, rule 9
Conformity Assessment Route: Annex II.3

We herewith declare that the above mentioned products meet the transposition into national law, the provisions of the following EC Council Directives and Standards. All supporting documentations are retained under the premises of the manufacturer. We are exclusively responsible for the Declaration of Conformity.

DIRECTIVES

General applicable directives:

Medical Device Directive: COUNCIL DIRECTIVE 93/42/EEC of 14 June 1993 concerning medical devices (MDD 93/42/EEC), amended by DIRECTIVE 2007/47/EC.

Notified Body: TÜV SÜD Product Service GmbH
Ridlerstr. 65, 80339 MÜNchen, Germany
Identification number: CE0123
(EC) Certificate(s): G1 103242 0002
Expire date of the Certificate: 2024-05-26
Start of CE Marking: 2019-07-25
Place, Date of Issue: Suzhou, 2019-07-25

Signature:

Name:

Anna Zhu

Title : Sr. QA/RA Manager