

泰博科技股份有限公司 | 新北市24888五級區五工二路127號 1-7樓 TaiDoc Technology Corp. | New Taipei City 24888, Taiwan

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## **EC Declaration of Conformity**

We, TaiDoc Technology Corporation

B1-7F, No.127, Wugong 2nd Road, Wugu Dist, 24888 New Taipei City, TAIWAN declare under our sole responsibility that the product

**Product Name** 

: Blood Glucose Test Strip

Product model

: TD-4302

Classification

: 98/79/EC (IVDD), Annex II, List B

Conformity Assessment Route

: 98/79/EC (IVD), Annex IV excluding section 4 & 6

EC Certificate Number

: V1 052126 0042 Rev.01

European Representative

: MedNet GmbH

Borkstraße 10, 48163 Münster, Germany

Notified Body (CE0123)

: TÜV SÜD Product Service GmbH

Ridlerstraße 65, 80339 München, Germany

GMDN code

: 62537

to which this declaration relates is in conformity with the following standard(s) or other normative document(s):

ISO 13485:2016	Medical devices. Quality management systems. Requirements for regulatory purposes
EN ISO 14971:2012	Medical devices. Application of risk management to medical devices
EN ISO 15197:2015	In vitro diagnostic test systems. Requirements for blood-glucose monitoring systems for self-testing in managing diabetes mellitus
EN 15223-1:2016	Medical devices. Symbols to be used with medical device labels, labelling and information to be supplied. General requirements
EN ISO 18113-1:2011	In vitro diagnostic medical devices. Information supplied by the manufacturer (labelling). Terms, definitions and general requirements
EN ISO 18113-4:2011	In vitro diagnostic medical Devices. Information supplied by the manufacturer (labelling). In vitro diagnostic reagents for self-testing
EN ISO 18113-5:2011	In vitro diagnostic medical Devices. Information supplied by the manufacturer (labelling). In vitro diagnostic instruments for self-testing
EN ISO 23640:2015	In vitro diagnostic medical devices. Evaluation of stability of in vitro diagnostic reagents



Let #880-2-5615-8888 as: 1886-2-6675-028E

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EN 1041:2008+A1:2013	Information supplied by the manufacturer of medical devices
EN 13532:2002	General requirements for in vitro diagnostic medical devices for self-testing
	Performance evaluation of in vitro diagnostic medical devices

2019.3.11-

Date of Issue

Um Jan

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Management Representative