

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

: TD-4302 Product model

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

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

: V1 052126 0069 Rev.03 EC Certificate Number

: MedNet EC-REP GmbH European Representative

Borkstraße 10, 48163 Münster, Germany

: TÜV SÜD Product Service GmbH Notified Body (CE0123)

Ridlerstraße 65, 80339 München, Germany

GMDN code : 53307

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

EN 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 ISO 15223-1:2021	Medical devices. Symbols to be used with information to be supplied by the manufacturer. 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 18113-2:2011	In vitro diagnostic medical Devices. Information supplied by the manufacturer (labelling). In vitro diagnostic reagents for professional use
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 23640:2015	In vitro diagnostic medical devices. Evaluation of stability of in vitro diagnostic reagents
EN ISO 20417:2021	Medical devices - Information to be supplied by the manufacturer
EN 13532:2002	General requirements for in vitro diagnostic medical devices for self-testing
EN 13612:2002 /AC:2002	Performance evaluation of in vitro diagnostic medical devices

2022, 5, 20

Date of Issue

Management Representative