

# 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-4371  
 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 0069 Rev.03  
 European Representative : MedNet EC-REP GmbH  
 Borkstraße 10, 48163 Münster , Germany  
 Notified Body (CE0123) : TÜV SÜD Product Service GmbH  
 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 14971:2019	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 - Part 1: General requirements
ISO 18113-1:2022	In vitro diagnostic medical devices. Information supplied by the manufacturer (labelling). Terms, definitions and general requirements
ISO 18113-2:2022	In vitro diagnostic medical Devices. Information supplied by the manufacturer (labelling). In vitro diagnostic reagents for professional use
ISO 18113-4:2022	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

**2023.11.02.**

**Date of Issue**



**Jim Jan**

**Management Representative**