

**DECLARATION OF CONFORMITY TO  
Regulation(EU) 2017/745  
CONCERNING MEDICAL DEVICES**

**MANUFACTURER:** Edan Instruments, Inc.  
#15 Jinhui Road, Jinsha Community, Kengzi Sub-District,  
Pingshan District, 518122 Shenzhen, P.R.China

**SRN:** CN-MF-000009957

**EUROPEAN REPRESENTATIVE:** Shanghai International Holding Corp. GmbH  
Eiffestrasse 80 20537 Hamburg Germany

**PRODUCT/MODEL:** **PC ECG / PADECG**

**EMDN [NAME/CODE]:** ELECTROCARDIOGRAPHS / Z120503

**Basic UDI-DI:** 69444138PADECGPK

**CLASSIFICATION:** Class II a, Rule 10 According To Annex VIII OF the MDR

**CONFORMITY ASSESSMENT ROUTE:** ANNEX IX EXCLUDING CHAPTER II.

WE, EDAN INSTRUMENTS, INC., HERE WITH DECLARE THAT THE ABOVE MENTIONED PRODUCT(S) MEET THE PROVISIONS OF REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL ON MEDICAL DEVICE.

ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER.

EU DECLARATION OF CONFORMITY IS ISSUED UNDER THE SOLE RESPONSIBILITY OF THE MANUFACTURER

**STANDARDS APPLIED: EN 60601-1:2006+A2:2021, EN 60601-1-2:2015+A1:2021, EN 60601-1-6:2010+A2:2021, EN 60601-2-25:2015, EN ISO 10993-1:2020, EN ISO 10993-5:2009, EN ISO 10993-10: 2023, EN ISO 10993-23:2021, EN ISO 14971:2019, EN 62304:2006+A1:2015, EN 62366-1:2015+A1:2020, EN ISO 15223-1:2021, EN 1041:2008+A1:2013, EN ISO 780:2015**

**NOTIFIED BODY:** TÜV SÜD PRODUCT SERVICE GMBH  
RIDLERSTR 65, D-80339 MÜNCHEN, GERMANY

**IDENTIFICATION NUMBER** 0123

**(EC) CERTIFICATE(S):** G10 091264 0025      **VALID UNTIL: 2026-02-17**

**START OF CE-MARKING:** 2013-07-31

**PLACE, DATE OF ISSUE:** SHENZHEN, 2024.3.26

**SIGNATURE:**   
NAME LIU YONGYING  
MANAGEMENT REPRESENTATIVE