

DECLARATION OF CONFORMITY TO COUNCIL DIRECTIVE 93/42/EEC CONCERNING MEDICAL DEVICES

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

EUROPEAN REPRESENTATIVE: Shanghai International Holding Corp. GmbH (Europe)
Eiffestrasse 80 D-20537 Hamburg Germany

PRODUCT/MODEL: **Diagnostic Ultrasound System / U50**
The accessories are used together with the product

GMDN [NAME/CODE]: General ultrasound imaging system, line-powered / 40761

CLASSIFICATION: Class II a, Rule 10 According To Annex IX of the MDD

CONFORMITY ASSESSMENT ROUTE: Annex II excluding (4)

WE, EDAN INSTRUMENTS, INC., HEREWITH DECLARE THAT THE ABOVE MENTIONED PRODUCT(S) MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE 93/42/EEC OF 14 JUNE 1993 INCLUDING AMENDMENTS BY DERECTIVE 2007/47/EC.
ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER.

STANDARDS APPLIED: **EN 60601-1:2006/A1:2013, EN 60601-1-2: 2007/AC:2010, EN 60601-2-37:2008, EN62366:2008, EN 62304:2006, EN ISO14971:2012, EN ISO 10993-1:2009, EN ISO10993-5:2009, EN ISO10993-10:2010, EN 980:2008 , EN 1041: 2008**

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

IDENTIFICATION NUMBER

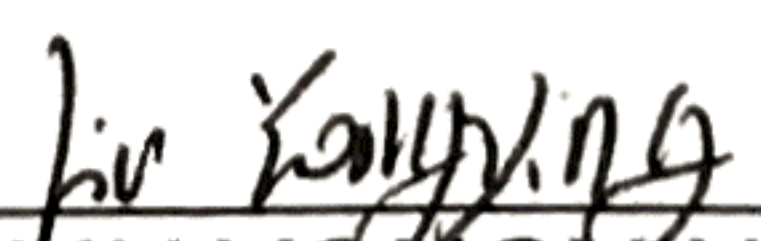


(EC) CERTIFICATE(S): G1 17 05 91264 006 VALID UNTIL: 2022-09-17

START OF CE-MARKING: 2012-04-12

PLACE, DATE OF ISSUE: SHENZHEN, 2017.9.5

SIGNATURE:


NAME **LIU YONGYING**
MANAGEMENT REPRESENTATIVE