



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

MANUFACTURER: Edan Instruments, Inc.  
3/F - B, Nanshan Medical Equipments Park, Nanhai Rd  
1019#, shekou, Nanshan Shenzhen, 518067 P.R. China

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

PRODUCT/MODEL: **Ultrasonic Pocket Doppler/ SONOTRAX Lite,  
SONOTRAX Basic,  
SONOTRAX Basic A,  
SONOTRAX Pro,  
SONOTRAX II,  
SONOTRAX II Pro,  
SONOTRAX Vascular**

*The accessories are used together with the product*

UMDNS [NAME/CODE]: Fetal heart detectors, Ultrasonic /11696  
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, EN 60601-2-37:2008, EN ISO14971:2012, EN ISO10993-1:2009, EN ISO 10993-5:2009, EN ISO 10993-10:2010, EN 62304:2006, EN 62366:2008, EN 980: 2008, EN 1041: 2008**

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

IDENTIFICATION NUMBER **CE**<sub>0123</sub>

(EC) CERTIFICATE(S): G1 15 03 44180 040 VALID UNTIL: 2017-09-17

START OF CE-MARKING: 2002-09-25

PLACE, DATE OF ISSUE: SHENZHEN, 2016.5.26

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MANAGEMENT REPRESENTATIVE