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

MANUFACTURER:	TANITA CORPORATION 1-14-2, MAENO-CHO, ITABASHI-KU, TOKYO, JAPAN SRN: JP-MF-000022755
EUROPEAN AUTHORIZED REPRESENTATIVE:	TANITA EUROPE B.V. HOOGOORDDREEF 56-E 1101 BE AMSTERDAM, THE NETHERLANDS SRN: NL-AR-000003401
PRODUCT NAME:	Body Composition Analyzer
DEVICE CODE:	DC-240MA
BASIC UDI-DI:	4904785BCA01RC
CLASSIFICATION:	CLASS IIa (RULE10, ACCORDING TO MDR ANNEX VIII)
CONFORMITY ASSESSMENT:	ANNEX IX (Excluding Chapter II)

WE UNDER OUR SOLE RESPONSIBILITY, HEREWITH DECLARE THAT ABOVE MENTIONED PRODUCTS MEET PROVISIONS OF THE REGULATION (EU) 2017/745 CONCERNING MEDICAL DEVICES.
ALL SUPPORT DOCUMENTATION IS RETAINED AT THE PREMINESS OF THE MANUFACTURER.

STANDARD APPLIED:	SEE ATTACHED LIST OF "HARMONIZED STANDARD LIST"
NOTIFIED BODY:	Regulation(EU) 2017/745: TÜV SÜD PRODUCT SERVICE GMBH RIDLERSTR 65, D-80339 MÜNCHEN, GERMANY CE 0123
ADDITIONAL INFORMATION:	
Place, Date:	JAPAN, 25-NOV, 2024
SIGNATURE:	K. Nishibayashi Kenji Nishibayashi General Manager Production Engineering Department TANITA CORPORATION