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

MANUFACTURER:	TANITA CORPORATION 1-14-2, MAENO-CHO, ITABASHI-KU, TOKYO, JAPAN
EUROPEAN REPRESENTATIVE:	EC REP TANITA EUROPE B.V. HOOGOORDDREEF 56-E 1101BE AMSTERDAM, THE NETHERLANDS
PRODUCT:	MC-780MA
SIMILAR MODEL:	NONE
UMDNS CODE:	17417
CLASSIFICATION:	*CLASS IIA, (RULE 10)* *RULE ACCORDING TO ANNEX IX OF THE MDD*
CONFORMITY ASSESSMENT ROUTE:	*ANNEX II*

WE HEREWITH DECLARE THAT THE ABOVE MENTIONED PRODUCT(S) MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE 93/42/EEC CONCERNING MEDICAL DEVICES.
ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER.

STANDARDS APPLIED:	*SEE ATTACHED LIST OF "HARMONIZED STANDARD LIST"*
NOTIFIED BODY:	TÜV SÜD PRODUCT SERVICE GMBH RIDLERSTR 65, D-80339 MÜNCHEN, GERMANY
IDENTIFICATION NUMBER	CE 0123
START OF CE-MARKING:	*THE APPLIANCE IS CE-MARKED 2013* SERIAL NO. 13020001 – (E.G. 13020001 = THE 1ST UNIT THAT HAS BEEN MANUFACTURED FEB, 2013
PLACE, DATE OF ISSUE:	*JAPAN, 15 - FEB , 2013 .*
SIGNATURE:	*Kenji Teranishi* *EXECUTIVE MANAGER* *MEDICAL DEPARTMENT*