MANUFACTURER'S DECLARATION OF CONFORMITY TO COUNCIL DIRECTIVE 93/42/EEC

MANUFACTURER:	ImpediMed Limited Unit 1, 50 Parker Court Pinkenba Qld 4008 Australia
EUROPEAN REPRESENTATIVE:	Medimark Europe Sarl 11, Rue Emile Zola – BP 2332 38033, Grenoble Sedex 2 - France
PRODUCT:	Body Impedance Analyzer/Imp DF50
GMDNS CODE:	Analyzer, Fat/Lean [36022]
CLASSIFICATION:	Class IIa, Rule 10, according to Annex IX of the MDD
CONFORMITY ASSESSMENT ROUTE:	Annex II.3

We herewith declare that the above mentioned products meet the transposition into national law the provisions of Council Directive 93/42/EEC for medical devices - as amended by Directive 98/79/EC on in vitro diagnostic medical devices and Directive 2007/47 EC, and Directive 2007/47 EC.
All supporting documentation is retained at the premises of the manufacturer.

STANDARDS APPLIED:	IEC 60601-1:1988 +A1:1991 + A2:1995, IEC 60601-1-2: 2004, ISO 13485:2003
NOTIFIED BODY:	SGS United Kingdom Ltd 202B Worle Parkway, Weston-Super-Mare Somerset BS22 6WA UK
IDENTIFICATION NUMBER	CE 0120
(EC) CERTIFICATE(S):	AU05/64407
START OF CE-MARKING:	24 Oct 2005
PLACE:	Brisbane, Qld, Australia
DATE OF ISSUE:	15 APRIL 2011
SIGNATURE:	
NAME	Mr Alden Kay
POSITION	ImpediMed Limited – VP QA/RA