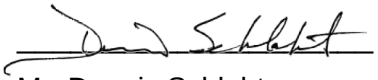
impedimed
Unit 1 50 Parker Court Pinkenba QLD 4008 Australia
www.impedimed.com

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 Cedex 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 under 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.
All supporting documentation is retained at the premises of the manufacturer.

STANDARDS APPLIED:	IEC60601-1 3rd Ed. Amendment 1, IEC60601-1-2:2004, ISO13485:2003
NOTIFIED BODY:	BSI Group, Kitemark Court, Davy Avenue, Knowlhill, Milton Keynes, MK5 8PP, United Kingdom
IDENTIFICATION NUMBER:	CE 0086
(EC) CERTIFICATE NUMBER:	654813
START OF CE-MARKING:	24 Oct 2005
PLACE:	Brisbane, Qld, Australia
DATE OF ISSUE:	14 September 2016
SIGNATURE:	
NAME:	Mr. Dennis Schlaht
POSITION:	ImpediMed Limited – SVP Quality