SLENDERTONE®

DECLARATION OF CONFORMITY

We **Bio-Medical Research Ltd.**
Parkmore Business Park West
Galway, H91 NHT7, Ireland.

Declare under our sole responsibility that the product:

Slendertone® Connect Abs Type: **570**

to which this declaration relates, is in conformity with Directive 2014/53/EU of the European Parliament and of the Council of 16 April 2014 (Radio Equipment Directive).

	Standards:
Article 3.1(a):	IEC 60601-1:2005 + CORR. 1:2006 + CORR. 2:2007 +AM1 2012
	IEC 60601-2-10:2015
	IEC 60601-1-11:2010
Article 3.1 (b)	EN 55022:2009 +A1:2010
	EN 60601-1-2:2007
	IEC 60601-2-10:2012
	EN 301 489-1 v2.1.1
	EN 301 489-17 v3.1.1
Article 3.2	EN 300 328 v2.1.1

The object of the declaration described above is in conformity with Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment. The devices are in conformity with the standard: EN 50581:2012.

Signed for and behalf of Bio-Medical Research Ltd.

Signature: Mike Kilkelly Date: 15 Jun 2017

Michael Kilkelly

Title: Quality Manager

Location: Bio-Medical Research Ltd. BMR House, Parkmore Business Park West, Galway, H91 NHT7, Ireland