

## EC Declaration of Conformity

**Declaration Number:** DC - 1101

**Manufacturer:** MYOTON LIMITED  
20-22 Bedford Row  
London WC1R 4JS

**Product Family Name:** Myoton

**Type:** MyotonPRO – medical device for non-invasive *in vivo* assessment of soft biological tissues.

**Product Sizes/Codes:**

Name	Product Code Number
MyotonPRO	1308600502

**Classification (MDD, Annex IX)  
and Rule Number:**

**Class I , Annex IX, Rule Number 1**

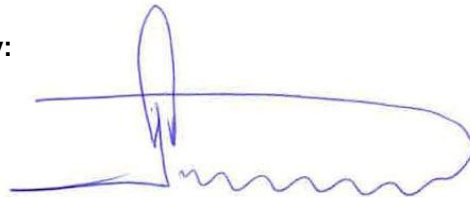
### Declaration

*Myoton AS hereby declares that the device specified above conforms with the Essential Requirements of the Medical Device Directive - 93/42/EEC of June 14, 1993 as amended by Directive 2007/47/EC of 21<sup>st</sup> September 2007.*

The stated products are designed and manufactured by Myoton AS, in accordance with the scope of a quality system which meets the requirements of the Medical Devices Directive - 93/42/EEC.

To ensure conformity with the provisions of the Directive applicable to Class I equipment, Myoton AS has designed and manufactured the device specified above in accordance with EN60601-1-2:2007 and EN60601-1:2006.

**Authorised by:**



**Aleko Peipsi**

**MYOTON Ltd – Director  
MYOTON AS – CEO**

**28<sup>th</sup> November 2011**

**UK Company:**

**MYOTON Ltd**  
20-22 Bedford Row,  
London WC1R 4JS  
Company no. 07816863

**Subsidiary of:**

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