

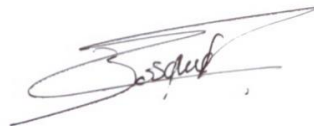
EC DECLARATION OF CONFORMITY - DIRECTIVE 93/42/EEC

We, MANATEC 21 RUE DU STADE PETIT-EBERSVILLER 57 730 FOLSCHVILLER FRANCE, hereby certify that the PhysioFlow devices (devices of class IIa according to the rule 10 of the appendix IX of the EC Directive 93/42/EEC, devices whose list can be found in the attached appendix), non-invasive medical devices for hemodynamic measurements by transthoracic impedance signal analysis, are in conformity with the book Vbis L665-4 of the French Public Health Code and the decree 95-292 of March 16th 1995 which concerns those products (decree transposing the EC Directive 93/42/EEC into the French national legislation).

This declaration is based on the following elements:

- Technical files, established in accordance with the appendix II section 3.2(c) and demonstrating the conformity of the product to the standards of the Directive 93/42/EEC and in particular to the Essential Requirements (see reference for each product in attached appendix)
- EC certificate 11826 rev. 5 of approval of the full Quality Assurance System in accordance with the appendix II section 3 of the Directive 93/42/EEC and granted on December 23rd 2016 by the notified body LNE/G-MED, identification number N°0459 (1 rue Gaston Boissier – 75 724 PARIS cedex 15, FRANCE)

In Petit-Ebersviller, December 23rd 2016



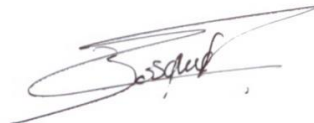
Benoît Bosquet – Co-Manager

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APPENDIX : LIST OF THE PHYSIOFLOW PRODUCTS concerned by the EC Declaration of Conformity

Device designation	Device commercial name or article code	Technical file reference
PhysioFlow PF07 Enduro	PhysioFlow Enduro	PF07 Enduro
PhysioFlow PF07 Q-Link	PhysioFlow Q-Link	PF07 Q-Link

In Petit-Ebersviller, December 23rd 2016



Benoît Bosquet – Co-Manager