# CE Declaration of conformity

The Manufacturer:

**CenterVue SpA**
Company

**Via San Marco 9H, 35129 Padova – ITALY**
Address

**+39 049 501 8399 / +39 049 501 8398**
Phone/fax

Declares under its responsibility that the medical device

# EIDON AF

class: IIa, rule 10
Conformity assessment procedure: Annex II.3 93/42/EC Directive (including 2007/47/EC)
Notified Body: TÜV SÜD Product Service GmbH, Ridlerstrasse 65, D-80339 München (N. 0123)

***Meets the essential requirements of Annex I***
***of Medical Devices Directive 93/42/EC (including 2007/47/EC).***

This medical device satisfies the applicable harmonized standards.
Technical documents are maintained by the Manufacturer for all the life time of the product and they could be inspected by the Competent Authority.

A post market surveillance system with recall provision is in place.
CE mark starting date: **June 28th 2016**.

**Management representative** — **Giuliano Barbaro**

centervue
CenterVue S.p.A.
Via S. Marco, 9/H
35129 Padova, Italy
C.F./P.IVA 04296580287

Name Surname

Signature

**Padova, 2018-06-27**
Place and date