

EU DECLARATION OF CONFORMITY

Manufacturer / Fabrikant:
(name + address)

STR Teknikk, Einar Stranden PhD
Skåtalia 15
N-6010 Ålesund
Norway

– herewith declare that the produkt:

– bekrefter med dette at produkt:

Model/type:

VenoPulse

Technical data / Tekniske data:

1A, 230VAC, 50Hz, Class 1 equipment.

Standards used / Standarder benyttet:

MDD-12
Medical Device Regulation (MDR)

– is in conformity with the provisions of mentioned Directives:

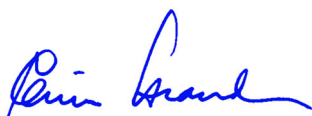
– er produsert i henhold til krav i følgende direktiver:

- Medical Device Directive 93/42/EEC
- REGULATION (EU) 2017/745, Annex I
- REGULATION (EU) 2017/745, Annex II
- REGULATION (EU) 2017/745, Annex III

CE marking is affixed from year:
CE merked er påført fra år:

April 2004

Place: Ålesund, Norway
Date : 25 February 2025



Einar Stranden PhD
Technical, and Managing Director