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:***	**VenoPulse**
Model/type:	.
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