precordior®

Declaration of Conformity

We, Precordior Ltd., Aurakatu 6, 20100 Turku, Finland ("the Manufacturer") hereby declare under our own responsibility that the following products:

- *CardioSignal Mobile application*
- *CardioSignal Cloud service*

meet the provisions of the Finnish Law (629/2010) and Council Directives 93/42/EEC and 2007/47/EC and

that it fulfils the requirements given by the European standards EN ISO 13485:2016, EN ISO 14971:2012, EN ISO 15223-1:2016, EN 62304:2006, EN 62366-1:2015.

The above-mentioned devices have been classified as Medical Device CE Class IIa according to rule 10.

This declaration is based on the conformity assessment of product to the requirements of Annex II of Council Directive 93/42/EEC. The notified body is Eurofins Expert Services Oy, number 0537.

This Declaration is valid for the products concerned listed above, bearing the CE mark, and manufactured by Precordior Ltd.

Turku, January 19th, 2022,

Juuso Blomster
CEO
Precordior Ltd.

PRECORDIOR OY FI28009277