CRYO IQ

CE

# EC – Declaration of Conformity

**Manufacturer**

CryoIQ global
Apelrödsvägen 1
439 32 Onsala
Sweden

**Products**

CryoIQ DERM Liquid
CryoIQ DERM Contact

CryoIQ DERM plus Liquid
CryoIQ DERM plus Contact

CryoIQ PRO Liquid
CryoIQ PRO Contact

CryoIQ PRO Aesthetics Liquid
CryoIQ PRO Aesthetics Contact

**Classification**

Class IIa

**Declaration**

We here with declare that the above mentioned products meets the provisions of the Council Directive 92/42/ECC for medical devices. All supporting documentation is retained under the premises of the manufacturer.

**Standard applied**

DIN EN ISO 13485:2016 (ISO 13485:2016 + AC:2017)

**Notified body**

Eurofins Expert Services Oy
Hermiankatu 6-8H
33720 Tampere
Finland
Identification number: 0537

**EC Certificate(s):**

EN ISO 13485:2016, cert.no. EUFI29-21000195-S
92/42/ECC, Annex V, section 4, cert.no. C-01-1187-768-21

Onsala 24 May 2021

Stefan Skafte
president

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