Declaration of Conformity

Manufacturer
Vitrolife Sweden AB
Gustaf Werners gata 2
SE-421 32 Västra Frölunda
Sweden

Product
Rapid-i Kit

Intended Use
Rapid-i Kit is intended to be used to contain, vitrify and maintain embryos and/or oocytes.

Product Category
Medical Device in Risk Class IIa

Attestation
We hereby declare that the product Rapid-i Kit fulfills the Essential Requirements as stated in Annex I to the Council Medical Devices Directive 93/42/EEC as amended by 2007/47/EC.

Place and Date
Göteborg 05 April 2016

Signature
On behalf of Vitrolife Sweden AB

Hans Lehmann
Director QA/RA

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