Declaration of Conformity

Manufacturer
Vitrolife Sweden AB
Box 9080 (Gustaf Werners gata 2)
SE-400 92 Göteborg
Sweden

Product
RapidVit™ Oocyte/RapidWarm™ Oocyte

Intended Use
Media for vitrification of oocytes/Media for warming of vitrified oocytes

Product category
Medical Device in Risk Class III

Attestation
We hereby declare that the product RapidVit™ Oocyte/RapidWarm™ Oocyte fulfils 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 on 4 November 2013

Signature
On behalf of Vitrolife Sweden AB,

Hans Lehmann
Director QA/QC

Vitrolife Sweden AB
Reg No 556546-6298

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