Declaration of Conformity

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

Product
IVF

Intended Use
Medium for in vitro fertilisation, culture and transfer of embryos

Product category
Medical Device in Risk Class III

Attestation
We hereby declare that the product IVF fulfils the Essential Requirements as stated in Annex 1 to the Council Medical Devices Directive 93/42/EEC as amended by 2007/47/EC.

Place and Date
Göteborg on 14 April 2015

Signature
On behalf of Vitrolife Sweden AB

Hans Lehmann
Director QA/QC

Vitrolife Sweden AB
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