Declaration of Conformity

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

Product
G-IVF/G-IVF PLUS

Intended Use
Medium for preparation and handling of gametes and for in vitro fertilisation

Product category
Medical Device in Risk Class III

Attestation
We hereby declare that the product G-IVF/G-IVF PLUS fulfil 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 07 April 2015

Signature
On behalf of Vitrolife Sweden AB

Hans Lehmann
Director QA/QC

Vitrolife Sweden AB
Reg. No. 556546-6298

Box 9080
SE-400 92 Göteborg
Sweden
Visting adress: Gustaf Werners gata 2
SE-421 32 V Frölunda
Sweden

Tel +46-31-721 80 00
Fax +46-31-721 80 90
www.vitrolife.com