Declaration of Conformity

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

Product
G-PGD

Intended Use
Medium for embryo biopsy

Product category
Medical Device in Risk Class III

Attestation
We hereby declare that the product G-PGD 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 10 April 2015

Signature
On behalf of Vitrolife Sweden AB

Hans Lehmann
Director QA/QC

Vitrolife Sweden AB
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