Declaration of Conformity

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

Product
G-MM

Intended Use
Medium for G-MM contains Recombinant Human Albumin solution (50 mg/mL) and is intended for use in assisted reproductive procedures which include gamete and embryo manipulation. These procedures include the use of G-MM as a supplement for culture medium.

Product category
Medical Device in Risk Class III

Attestation
We hereby declare that the product G-MM 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 20 March 2015

Signature
Hans Lehmann
Director QA/QC

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