

Declaration of Conformity

Manufacturer

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

Product

OVOIL

Intended Use

For covering of medium during in vitro fertilisation and micro-manipulation procedures

Product category

Medical Device in Risk Class IIa

Attestation

We hereby declare that the product OVOIL 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 01 April 2015

Signature

On behalf of Vitrolife Sweden AB,



Hans Lehmann
Director QA/QC