

Declaration of Conformity

Manufacturer

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

Product

G-ThawKit Blast

Intended Use

Media for thawing of blastocyst stage embryos

Product category

Medical Device in Risk Class III

Attestation

We hereby declare that the product G-ThawKit Blast 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 07 April 2015

Signature

On behalf of Vitrolife Sweden AB



Hans Lehmann
Director QA/QC