

## Declaration of Conformity

### Manufacturer

Vitrolife Sweden AB  
Box 9080 (Gustaf Werners gata 2)  
SE-400 92 Göteborg  
Sweden

### Product

RapidVit Blast/RapidWarm Blast

### Intended Use

Media for vitrification of blastocyst stage embryos/Media for warming of vitrified blastocyst stage embryos

### Product category

Medical Device in Risk Class III

### Attestation

I hereby declare that the products RapidVit Blast/RapidWarm Blast 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 28 June 2011

### Signature



Hans Lehmann  
Director QA/QC