Declaration of Conformity

Manufacturer

Vitrolife Sweden AB
Box 9080
SE-400 92 Göteborg
Sweden

Product: Disposable Devices for IVF

- Follicle Aspiration Set, Single Lumen
- Follicle Aspiration Set, Double Lumen
- Follicle Aspiration Set, Single Lumen, Luer
- Follicle Aspiration Set, Single Lumen, Luer with Tubing
- Follicle Aspiration Set, Reduced Single Lumen
- Amniocentesis Needle
- Chorion Biopsy Needle
- Cyst Puncture Needle with Stylet
- Cyst Puncture Needle without Stylet
- Pre-ovarian Block (POB) Needle
- Rapid-i Kit
- VitroLoop

Product category

Medical Device in Risk Class IIa

Attestation

I hereby declare that the products mentioned above fulfil the Essential Requirements as stated in Annex 1 to the Council Directive 93/42/EEC as amended by 2007/47/EC

Place and Date

Gothenburg on 15 August 2012

Signature

Nina Arvidsson
Regulatory Affairs Manager
Vitrolife Sweden AB