HertART at the beginning of life

Declaration of Conformity

Manufacturer
HertART ApS
Korskildelund 6
2670 Greve
Denmark

Product
'Vitrolife' Micro well group culture dish, 9 well

Intended Use
For In Vitro Fertilisation, handling and embryo culture

Product Category
Medical Device in Risk Class IIa

Notified Body
BSI UK, Notified Body Number 0086

Attestation
I hereby declare that the product 'Vitrolife' Micro well group culture dish, 9 well, fulfills the Essential Requirements as stated in Annex I to the Council Medical Device Directive 93/42/EEC as amended by 2007/47/EC.

Place and Date
Copenhagen, Denmark, 14. August 2014.

Signature
Tony Winslöf, CEO