

Valued customer,

We are very pleased to announce that the following products are CE marked in conformity with the new European Medical Device Regulation (MDR, 2017/745).

- FertiCult™ IVF medium

As summarized in our previous newsletter, CE marking according to the MDR does result in the following changes:

- Use of 3-layered labels (For more information: [see previous newsletter](#))
- New version of instruction for use
- No specific changes for FertiCult™ IVF medium

If additional information is required or additional registration actions in your jurisdiction(s) are required, please send your question to ra@fertipro.com



COMPANY ADDRESS

FertiPro nv
Industriepark Noord 32
B-8730 Beernem / Belgium

tel. +32 (0)50 79 18 05
fax +32 (0)50 79 17 99
info@fertipro.com
www.fertipro.com

RPR Brugge
VAT BE 0448 767 035
IBAN BE 98 7374 4024 9293
BIC KREDBEBB