qarad
European Regulatory Services

# DECLARATION

The undersigned, Mrs. Agnes Goris, Consultant at Qarad b.v.b.a.,

hereby declares that Guangzhou Wondfo Biotech Co. Ltd. has signed the EC Declaration of Conformity in agreement with the Annex IV of the European Directive 98/79/EC on In Vitro Diagnostic Medical Devices and has submitted the required technical documentation, for the following IVD products:

| | |
|---|---|
| Finecare™ CRP PCT Rapid Test | W218 |
| Finecare™ Cystatin C Rapid Test | W219 |
| Finecare™ βhCG Rapid Test | W225 |
| Finecare™ CRP Control | W806-L, W806-M, W806-H |
| Finecare™ Cystatin C Control | W816-L, W816-M, W816-H |
| Finecare™ Alpha-Fetoprotein Control | W815-L, W815-M, W815-H |

The notification to the Belgian Competent Authorities has been carried out on March 2nd, 2015 by Qarad b.v.b.a., the appointed Authorized Representative of Guangzhou Wondfo Biotech Co. Ltd.

Information on the notification to the competent Authorities of other European countries is available upon request.

Agnes Goris.

Qarad b.v.b.a.
Authorized Representative

Digitally signed by Varman Irina
Date: 2022.04.21 23:34:16 EEST
Reason: MoldSign Signature
Location: Moldova