



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