

## DECLARATION

The undersigned, Dr. Dirk Stynen, President of 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 III 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 (for professional use only):

Finecare™ FIA Meter	cat. nr. FS-131
Finecare™ hsCRP Rapid Test	cat. nr. W201
Finecare™ NT-proBNP Rapid Test	cat. nr. W202
Finecare™ Troponin I Rapid Test	cat. nr. W203
Finecare™ Myoglobin Rapid Test	cat. nr. W204
Finecare™ CK-MB Rapid Test	cat. nr. W205
Finecare™ PCT Rapid Test	cat. nr. W210

The notification to the Belgian Competent Authorities has been carried out on May the 17<sup>th</sup>, 2013 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.



Dirk Stynen, Ph. D.

President Qarad b.v.b.a.  
Authorized Representative