



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 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™ H-FABP Control	W811-L, W811-N, W811-H
Finecare™ D-Dimer Control	W812-L, W812-N, W812-H
Finecare™ Microalbumin Control	W813-L, W813-N, W813-H

The notification to the Belgian Competent Authorities has been carried out on October 6th, 2014 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