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™ Alpha-fetoprotein Rapid Test	W208
Finecare™ PCT Control	W805-L, W805-N, W805-H
Finecare™ NT-proBNP Control	W807-L, W807-N, W807-H
Finecare™ troponin I Control	W808-L, W808-N, W808-H
Finecare™ Myoglobin Control	W809-L, W809-N, W809-H
Finecare™ CK-MB Control	W810-L, W810-N, W810-H

The notification to the Belgian Competent Authorities has been carried out on September 18th, 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

Digitally signed by Varman Irina
Date: 2021.06.24 11:37:07 EEST
Reason: MoldSign Signature
Location: Moldova