qarad
European Regulatory Services

DECLARATION

The undersigned, Mrs. Joke Doutreloigne, 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:

IVD name	Catalogue nr.
Finecare™ T3 (Triidothyronine) Rapid Quantitative Test	W231
Finecare™ Progesterone Rapid Quantitative Test	W233
Wondfo One Step Multi-drug Urine Test Panel for Analyzer	W501-P to W510-P, Panel
One Step Barbiturates Urine Test (BAR200)	W008-S (200ng/ml), Strip W008-C (200ng/ml), Cassette W008-P (200ng/ml), Panel
One Step Benzodiazepines Urine Test (BZO100)	W009-S (100ng/ml), Strip W009-C (100ng/ml), Cassette W009-P (100ng/ml), Panel
One Step Marijuana Urine Test (THC25)	W019-S (25ng/ml), Strip W019-C (25ng/ml), Cassette W019-P (25ng/ml), Panel
Multi-drug Tests	W2002-P to W2016-P, Panel W2002-CU to W2012-CU, Cup W502-CU2 to W518-CU2, T-Cup W502-CU3 to W518-CU3, Q-Cup

The notification to the Belgian Competent Authorities has been carried out on September 22, 2016 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.

Digitally signed by Varman Irina
Date: 2022.03.07 12:22:40 EET
Reason: MoldSign Signature
Location: Moldova

Joke Doutreloigne

Qarad b.v.b.a.
Authorized Representative