

DECLARATION OF NOTIFICATION

Date: March 5, 2020

The undersigned, Sara Van Wouwe, Device Compliance Assistant of Qarad BVBA, hereby declares that:



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):

Name Device	Catalogue numbers
Wondfo SARS-CoV-2 Antibody Test (Lateral Flow Method)	W195
Finecare™ SARS-CoV-2 IgM Test	W277
Finecare™ SARS-CoV-2 Antibody Test	W276

The notification to the Belgian Competent Authorities has been carried out on March 5, 2020 by Qarad BVBA, 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.

Sara Van Wouwe

Device Compliance Assistant

Qarad BVBA

Authorized Representative