DECLARATION OF NOTIFICATION

Date: January 21, 2021

The undersigned, Teresa Batet i Solà, Senior Consultant of Qarad BV, hereby declares that:

Guangzhou Wondfo Biotech Co. Ltd.
No. 8 LizhÍshan Road, Science City Luogang District,
Guangzhou 51 0663, PR China

has signed the EC Declaration of Conformity in agreement with the Annex III excluding 6 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):

Name Device	Catalogue number Device
Influenza/2019-nCoV Antigen Combo Test	W630P0002, W630P0003, W630P0004, W630P0005, W630P0006, W630P0007, W630P0008, W630P0009, W630P0010, W630P0011
2019-nCoV Antigen Saliva/Sputum Test	W633P0001, W633P0002, W633P0003, W633P0004, W633P0005, W633P0006
Wondfo 2019-nCoV Antigen Test (Lateral Flow Method)	W196P0005, W196P0006, W196P0007, W196P0008, W196P0009, W196P0010, W196P0011, W196P0012
Finecare™ 2019-nCoV Antigen Test	W286P0002, W286P0003

The notification to the Belgian Competent Authorities has been carried out on January 21st, 2021 by Qarad BV, the appointed Authorized Representative of Guangzhou Wondfo Biotech Co. Ltd

Teresa Batet i Solà
Senior Consultant

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Qarad BV
Authorized Representative