

## 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 21<sup>st</sup>, 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