

## DECLARATION OF NOTIFICATION

Date: November 6, 2020

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

Guangzhou Wondfo Biotech Co. Ltd.  
No. 8 Lizhishan Road, Science City Luogang District,  
Guangzhou 510663  
PR China

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 product (for professional use only):

Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) (REF: W196)

The notification to the Belgian Competent Authorities has been carried out on August 11, 2020 by Qarad BV, the appointed Authorized Representative of Guangzhou Wondfo Biotech Co. Ltd. On November 6<sup>th</sup>, a notification of change was carried out during which the new product name Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) was notified.

Sara Van Wouwe  
Device Compliance Assistant  
Qarad BV  
Authorized Representative