qarad
European Regulatory Services

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 6th, 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

Digitally signed by Sara Van Wouwe (Signature)
Date: 2020.11.06 15:33:05 +01'00'