

## TO WHOM IT MAY CONCERN

Date: March 22, 2021

The undersigned, Kirsten Van Garsse, Director AR Operations of Qarad BV, hereby declares that Qarad BV with registered address at Ciplastraat 3, 2440 Geel, Belgium has been appointed as the European Authorized Representative in accordance with the European Directive 98/79/EC on In Vitro Diagnostic Medical Devices by:

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

for the following IVD products which have applied for special approval according to §11 paragraph 1 of the German Medical Devices Act (MPG) of antigen tests for self-administration by laypersons (self-tests) for the detection of SARS-CoV-2:

**Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) for nasal swab**

**2019-nCoV Antigen Saliva/Sputum Test**

Kirsten Van Garsse

Director AR Operations

Qarad BV

Authorized Representative