





EC Certificate

EC Design-Examination Certificate
Directive 98/79/EC on In Vitro Diagnostic Medical Devices (IVDD), Annex III (6)
(Devices for self-testing)

No. V9 058008 0037 Rev. 00

Manufacturer: GUANGZHOU WONDFO BIOTECH CO.,

LTD.

No. 8 Lizhishan Road, Science City Luogang District

510663 Guangzhou

PEOPLE'S REPUBLIC OF CHINA

Product: In Vitro diagnostic devices for self testing

The Certification Body of TÜV SÜD Product Service GmbH declares that a design examination has been carried out on the respective devices in accordance with IVDD Annex III (6). The design of the devices conforms to the requirements of this Directive. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:V9 058008 0037 Rev. 00

Report No.: SH2114101

Valid from: 2021-06-02

Valid until: 2024-05-26

Date, 2021-06-02

Christoph Dicks

Head of Certification/Notified Body



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Model(s): Wondfo 2019-nCoV Antigen Test

(Lateral Flow Method)

Facility(ies): GUANGZHOU WONDFO BIOTECH CO., LTD.

No. 8 Lizhishan Road, Science City, Luogang District, 510663

REF W634P0027

Guangzhou, PEOPLE'S REPUBLIC OF CHINA

Model Name: Model No.:

Wondfo 2019-nCoV Antigen Test (Lateral Flow Method)

Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) REF W634P0024
Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) REF W634P0028
Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) REF W634P0029
Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) REF W634P0025
Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) REF W634P0026

