



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-245.10.07



Product Service

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 089675 0006 Rev. 00

Manufacturer:

Beijing Hotgen Biotech Co.,Ltd

9th Building, No. 9 Tianfu Street, Biomedical Base

Daxing District

102600 Beijing

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_089675_0006_Rev_00

Report No.:

BJ21071201

Valid from:

2021-08-04

Valid until:

2024-05-26

Date,

2021-08-04

Christoph Dicks

Head of Certification/Notified Body



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Model(s): **Coronavirus (2019-nCoV)-Antigentest-**

Facility(ies): Beijing Hotgen Biotech Co.,Ltd
 9th Building, No. 9 Tianfu Street, Biomedical Base, Daxing District,
 102600 Beijing, PEOPLE'S REPUBLIC OF CHINA

| Model Name: | REF number: |
|--------------------------------------|--------------|
| Coronavirus (2019-nCoV)-Antigentest- | HGCG134S0101 |
| Coronavirus (2019-nCoV)-Antigentest- | HGCG134S0105 |
| Coronavirus (2019-nCoV)-Antigentest- | HGCG134S0120 |
| Coronavirus (2019-nCoV)-Antigentest- | HGCG134S0140 |

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