



# Coronavirus (2019-nCoV)-Antigentest - Summary Data



Beijing Hotgen Biotech Co., Ltd.

# 目录

(Contents)

1、产品彩页（Product Brochure） .....	1-2
2、公司介绍（Company Profile） .....	3
3、产品照片（Product Photos） .....	4
4、包装信息（Packing Information） . .....	5
5、热景CE自测证（HOTGEN CE self test certificate） .....	6
6、ISO13485 认证（ISO13485 Certificate） .....	8

# Company Profile

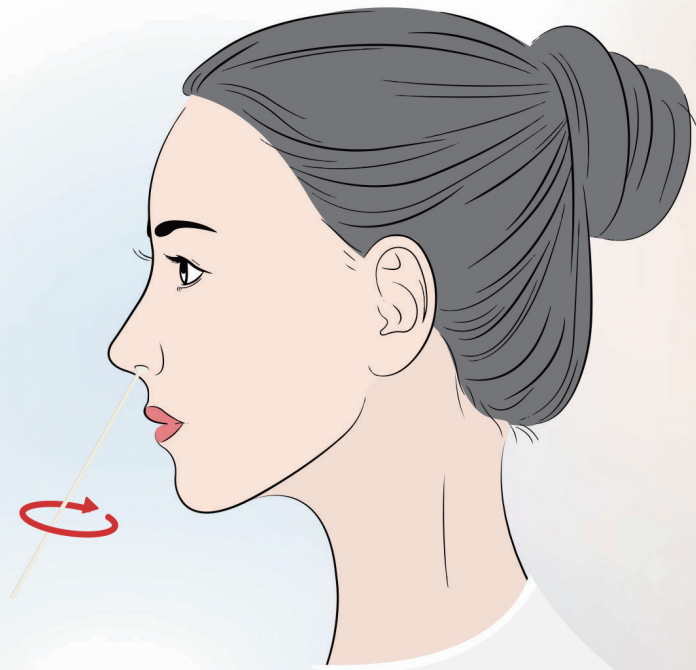
Beijing Hotgen Biotech Co., Ltd. (abbreviated as Hotgen Biotech, stock code: 688068) was established in June 2005, which is a high-tech enterprise focusing on the research& development, manufacture and sales of medical and public safety inspection products of in vitro diagnostics (IVD) in the field of biomedicine, as well as landed on the China Sci-Tech innovation board (STAR Market) in September 2019.

After several years of Research& development, Hotgen Biotech has developed an in vitro diagnostic reagent bioactive raw material development platform, a sugar chain abnormal protein detection (sugar capture) R&D technology platform, a Magnetic particles chemiluminescence R&D technology platform, a Up-converting Phosphor R&D technology platform, and a colloidal gold immune layer. The eight major technology platforms, such as the precipitation R&D platform, enzyme-linked immunoassay R&D technology platform, molecular diagnostics R&D platform, and instrument R&D technology platform, form a closed-loop system for in vitro diagnostic R&D and production. Hotgen Biotech has established a complete full level immunodiagnostic technology platform, from high-precision Up-converting Phosphor POCT (UPT series) to small, medium and large single- cartridge chemiluminescence platforms (MQ60 series), and then to large-scale full-automatic chemiluminescence Platform (C2000), which realizes the application of the immune diagnostic platform in the field of full diagnostic scenarios. Supporting products are widely used in the clinical and public safety fields. Specific users include hospitals at all levels, township health centers, third-party testing centers, and medical institutions, as well as medical and health institutions, as well as disease control centers, public security, fire protection, military, ports, food and medicine. Supervision, food and feed enterprises and other public safety fields.

Hotgen Biotech has won the second prize of the National Technology Invention Award, the Gold Medal of Independent Innovation, and the second prize of the Chinese Medical Science and Technology Award; In 2018, Hotgen Biotech was awarded the second prize of the "Technical Invention Category of China Rare Earth Science and Technology Award" by the China Rare Earth Society; Top 100 Private Scientific and Technological Innovations "and" Top 100 Medical Enterprises of the Future "; and" Postdoctoral Scientific Research Workstation "; major science and technology projects in the 12th and 13th five years, 863 plan, science and technology projects of the Beijing Science and Technology Commission, and Zhongguancun High Precision The project's major cutting-edge original technological achievements transformation and industrialization projects.

In the face of the COVID-19 epidemic situation, Beijing Hotgen Biotech Co.,Ltd has organized R&D developed a variety of Covid-19 detection reagents, including Novel Coronavirus 2019-nCoV Antigen Test (Colloidal Gold), Novel Coronavirus 2019-nCoV Antigen Test (Up-converting Phosphor Immunochromatographic Technology), Coronavirus disease(COVID-19) Antibody Test (Colloidal Gold), Coronavirus disease(COVID-19) IgM/IgG Antibody Rapid Test (Colloidal Gold), Novel Coronavirus 2019-nCoV Antibody Test (Up-converting Phosphor Immunochromatographic Technology), Novel Coronavirus (SARS-CoV-2) Neutralizing Antibodies Test (Colloidal Gold), Coronavirus disease( COVID-19) Nucleic Acid Test Kit (PCR-Fluorescent Probe Method), Disposable virus sampling tube, Nucleic acid Automatic Purification System, Nucleic acid extraction reagent, Biological Sample Releaser kit, etc.It is imperative to fight the epidemic Helping the global fight against epidemics!

Since its establishment, the company has continuously grown its business and has now achieved group development. At present, Hotgen (Langfang), Hotgen (Jilin), Weikekang Technology, Shunjing Biological and many other subsidiaries have been established. Hotgen Biotech marketing and service network has covered all regions of the country. Each province is equipped with professional technical service engineers, academic engineers, etc. who are responsible for pre-sales and after-sales work to meet customer needs. The company takes "developing biotechnology and benefiting human health" as its mission, "quality determines the company's life and death, customers determine the company's success or failure, talents determine the company's rise and fall, innovation determines the company's future" as its core values, and "tests because of me advanced" as its philosophy , High ambitions, technological entrepreneurship, and industrial prosperity!



## Coronavirus (2019-nCoV)-Antigentest-



### Product Features

- High Accuracy, Specificity and Sensitivity
- No need instrument, get results in 15 minutes
- Room temperature storage
- Sample : Human Anterior Nares Swab
- Detect the presence of viral proteins
- Identify acute or early infection

# Specimen Requirements and Test Procedure

## Specimen Requirements

**1**

Please wash and dry your hands thoroughly before the test.



**2**

Read the Instructions for Use carefully.



**3**

Please scan the QR code to watch the operation video.



**4**

Take the swab out of the package and do not touch the sampling end.




**5**

Carefully insert the swab 1.5 cm into the nostril until resistance is felt. Under moderate pressure, swab the surface of the nostril 4 to 6 times for at least 15 seconds.




**6**

Repeat sampling with the same swab in the other nostril.



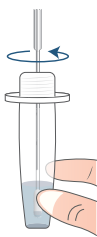
**7**

Open the larger end of the sampling tube, and insert the swab after collecting the sample into the sampling tube.




**8**

Soak the swab in the buffer for at least 15 seconds, stir the swab several times, and squeeze the swab head 3 times.




**9**

When remove the swab, squeeze the sides of the sampling tube.




**10**

Close the sampling tube with the tube lid




**11**

Open the foil pouch and place the test cassette on a flat surface.




**12**

Open the tube lid at the front end of the sampling tube, and drop 4 drops of the treated sample into the sample well (S) of the test cassette.




**13**

After reacting at room temperature for 15 minutes, observe the results. A result after 30 minutes is invalid.




**14**

After the test, pack all the components of this test in the Biohazard specimen bag for contaminated waste and dispose of this bag closed with the residual waste. Not reusable.

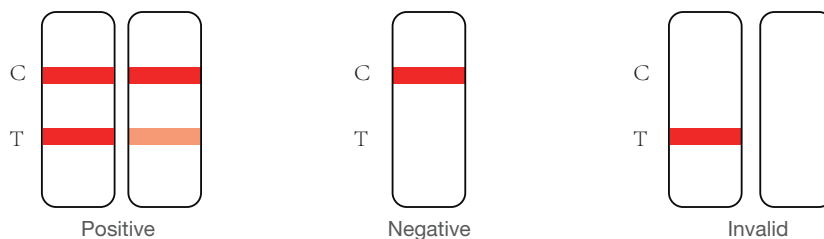


**15**

Wash or disinfect hands again.



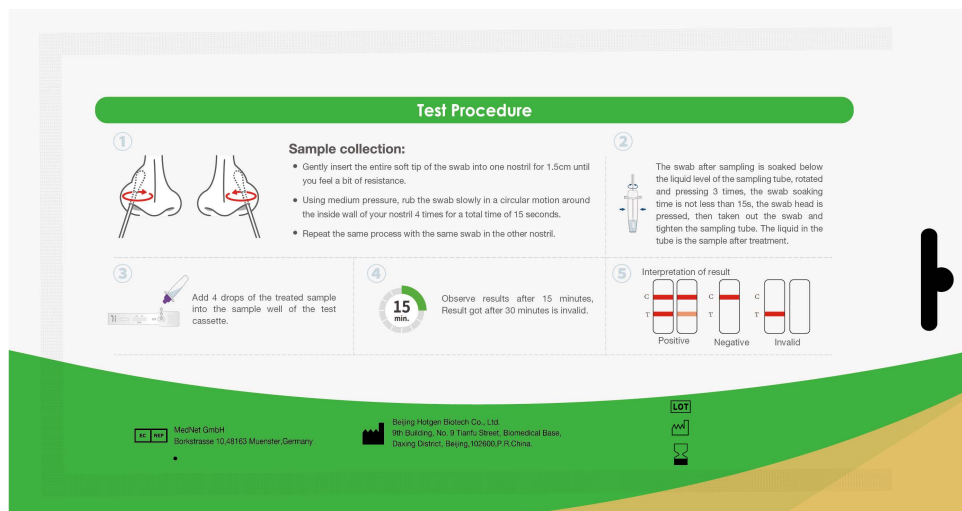
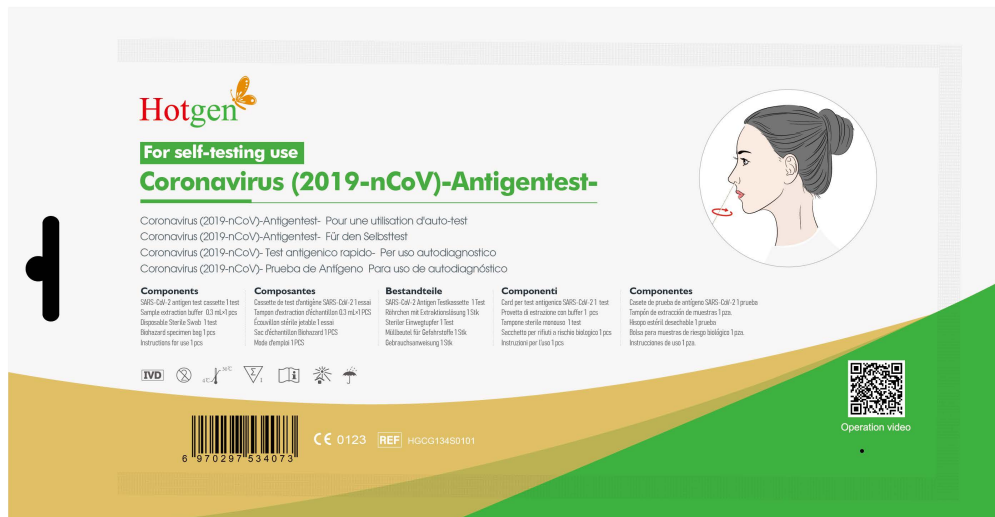
## Interpretation of result



## Product information

Product name	Test samples	Specification	Storage conditions
Coronavirus (2019-nCoV)-Antigen test-	Human Anterior Nares Swab	1T/kit	4-30 °C

# Coronavirus (2019-nCoV)-Antigentest - Product Photos



## 前鼻腔抗原胶体金检测试剂包装信息

### Coronavirus (2019-nCoV)-Antigentest - Packing Information

产品名称 Product name	规格/盒 Specifications	单位 Unit	单位包装毛重 Gross weight per unit package
Coronavirus (2019-nCoV)-Antigentest -	1T	袋/kit	0.037 kg/袋 0.037 kg / kit

前鼻腔抗原胶体金试剂盒出口包装箱 Coronavirus (2019-nCoV)-Antigentest - Export Packing Cartons							
包装箱 / 盒 Packing Carton/ box	长 length cm	宽 Width cm	高 height cm	每箱装袋 数 量 Kit quantity per carton	单盒试剂 净重 Net weight of single kit	整箱净重 Net weight of the whole carton	抛重 Throwing weight
纸箱 carton	72.5	38.5	25.8	400袋 400T	0.037 公斤 0.037 kg	11.5公斤 11.5kg	12公斤 12kg

For reference only, shipping cost according to the actual weight of the airline department





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Zentralstelle der Länder  
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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 II (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](http://www.tuvsud.com/ps-cert?q=cert:V9_089675_0006_Rev_00)

Report No.:

BJ21071201

Valid from:

2021-03-04

Valid until:

2024-05-26

Date,

2021-03-04

Christoph Dicks  
Head of Certification/Notified Body





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Medizinprodukten  
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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**

**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



Product Service

# Certificate

No. Q5 089675 0005 Rev. 01

**Holder of Certificate:** **Beijing Hotgen Biotech Co.,Ltd**  
9th Building, No. 9 Tianfu Street, Biomedical Base  
Daxing District  
102600 Beijing  
PEOPLE'S REPUBLIC OF CHINA

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

**Certification Mark:**



**Scope of Certificate:** Design and Development, Production, Distribution and Service of Automated Immunoassay Analyzer, Up-converting Phosphor Immunoassay Analyzer, Up-converting Phosphor Technology Test Kits, Colloidal Gold Test Kits, Chemiluminescence Immunoassay Test Kits, Enzyme-Linked Immunoassay Test Kits.

**Applied Standard(s):** EN ISO 13485:2016  
Medical devices - Quality management systems - Requirements for regulatory purposes (ISO 13485:2016)  
DIN EN ISO 13485:2016

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). 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:Q5 089675 0005 Rev. 01](http://www.tuvsud.com/ps-cert?q=cert:Q5 089675 0005 Rev. 01)

**Report No.:** BJ20071201

**Valid from:** 2020-12-05

**Valid until:** 2023-12-04

**Date,** 2020-09-01

Christoph Dicks  
Head of Certification/Notified Body