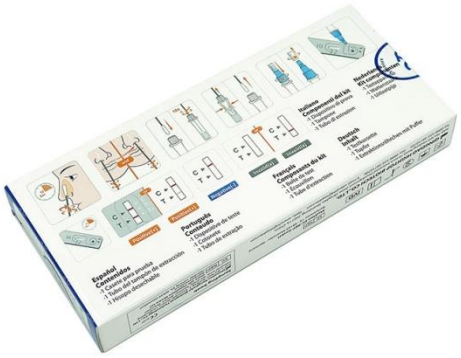
KINGSTAR



Safe Collection Covid-19 Self Test Rapid Antigen Test

KINGSTAR INC is Safe Collection Covid-19 Self Test Rapid Antigen Test manufacturers and suppliers in China who can wholesale Safe Collection Covid-19 Self Test Rapid Antigen Test. Safe collection covid-19 self test rapid antigen test is used for the detection of SARS-CoV-2 antigens in samples from the human anterior nasal cavityarea.

As a professional high quality Safe Collection Covid-19 Self Test Rapid Antigen Test manufacturers, you can rest assured to buy Safe Collection Covid-19 Self Test Rapid Antigen Test from KINGSTAR INC and we will offer you the best after-sale service and timely delivery.

Product Introduction

Safe collection covid-19 self test rapid antigen test is based on GICA principle, whereby the nitrocellulose membrane is coated with monoclonal coronavirus antibody 2 and goat-anti-mouse IgG ANTIBODY. No need to go for professional medical staffs, you can operate by yourselves.

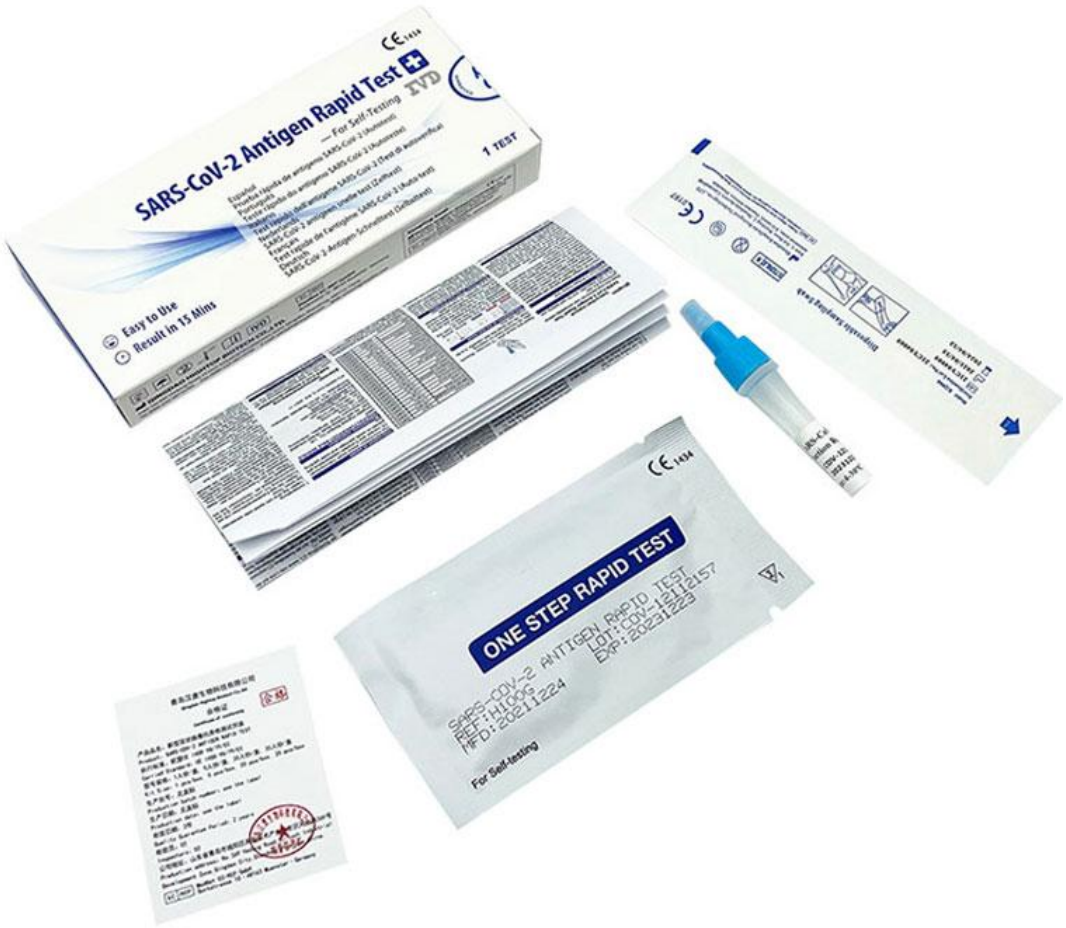
Product Parameter (Specification)

1. Test Cassette.
2. Extraction Tube (with Extraction Solution).
3. Swab.
4. Instructions for Use.

Safe collection covid-19 self test rapid antigen test

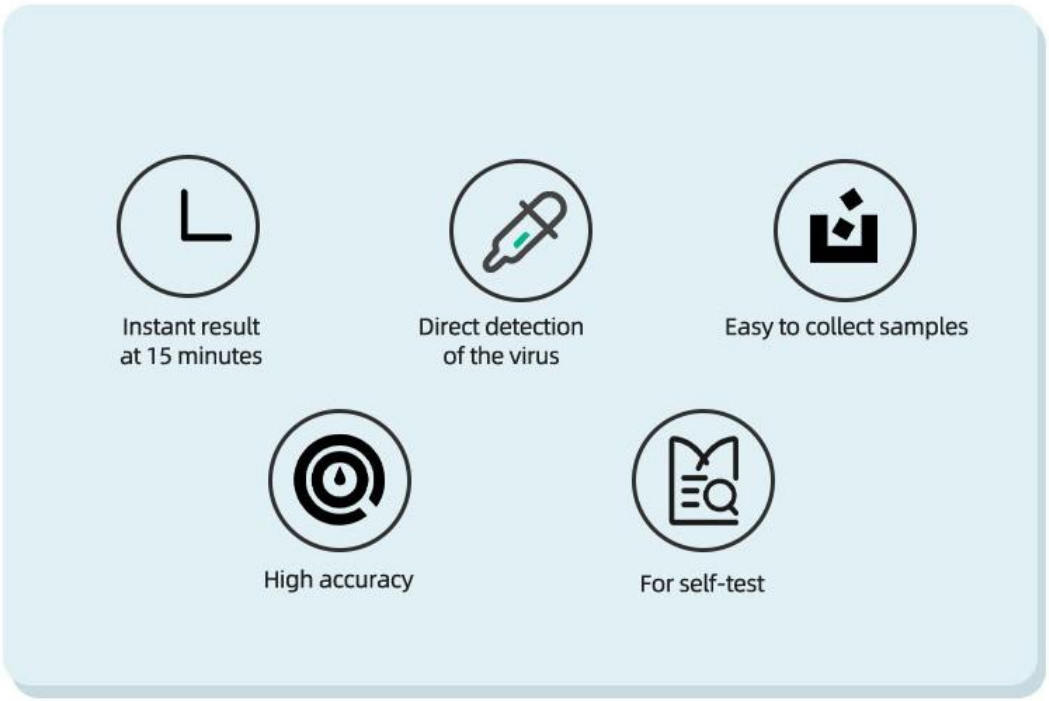
Tel: +86-574-62338878 **Phone:** +86-13858300666 **E-Mail:** info@nbkingstar.com

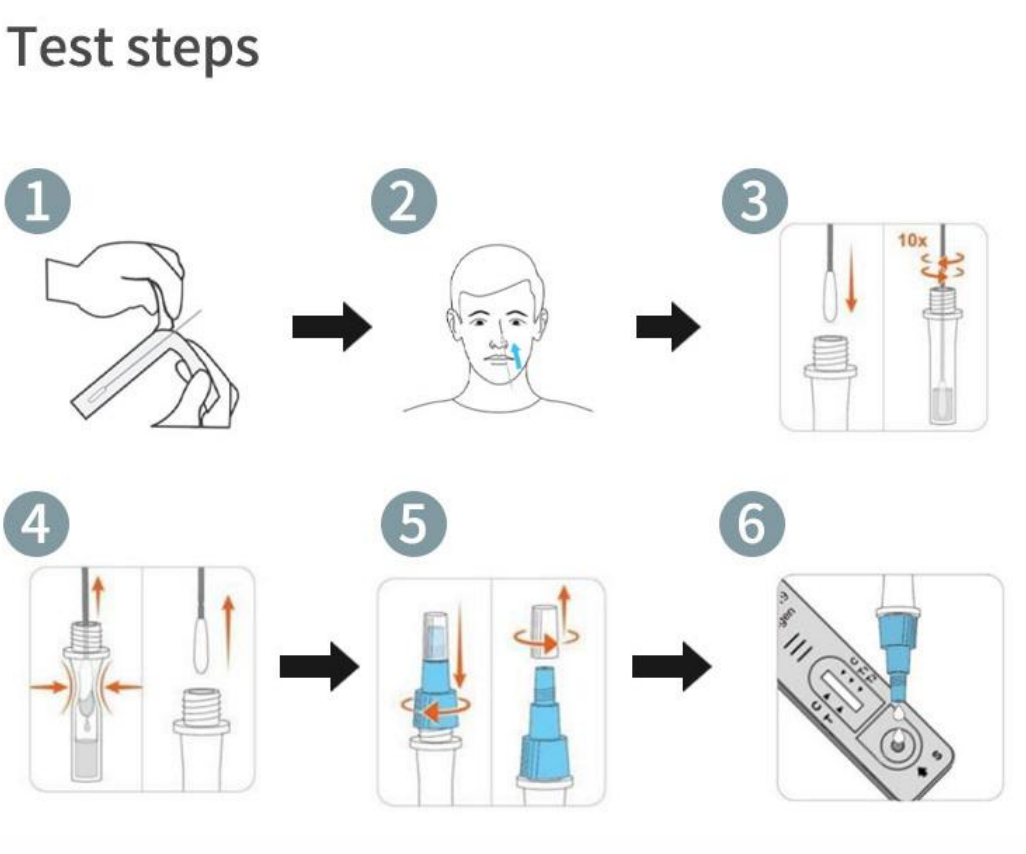

Español
Contenidos
Português
Conteúdo
Français
Composants du kit
Deutsch
Inhalt
Italiano
Componenti del kit
Positive(+)
Negative(-)
Invalid(-)


SARS-CoV-2 Antigen Rapid Test
For Self-Testing
1 TEST
Easy to Use
Result in 15 Mins
ONE STEP RAPID TEST
SARS-COV-2 ANTIGEN RAPID TEST
For Self-testing

Product Feature and Application

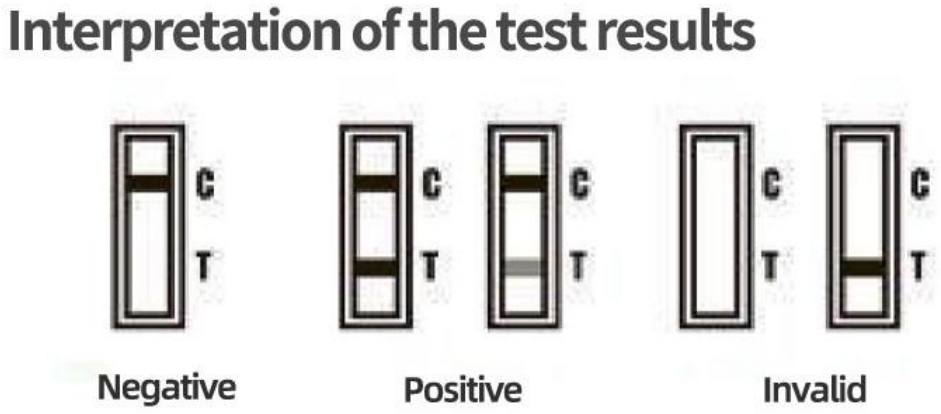
Safe collection covid-19 self test rapid antigen test is for own usage only. It is suitable for used by people who are suspected of having a COVID-19 disease over 18 years of age.





Product Details

Safe collection covid-19 self test rapid antigen test can help you get the result around 15 minutes. DO NOT read after 20 minutes.



Warning and precaution

1. Suitable for people aged 16 and over. Keep the test kits away from young children to reduce the risk of accidentally drinking the buffer liquid or swallowing small parts.
2. The test cassette should be used as soon as possible after removal from the foil bag to avoid prolonged exposure to moisture, as these could affect the test result.
3. Do not freeze the test kits.
4. The test set should be disposed after use in a lockable garbage bag in the household waste.
5. Incorrect operation may affect the accuracy of the results, such as e. g. too little effective time in the buffer solution, too little or too much buffer in the solution, insufficient sample addition, inaccurate detection time, etc.

6. False-negative results can occur when the swab is placed in a bag between sampling and evaluation.
7. Do not suck the sample with your mouth.
8. During the test, do not smoke, eat, drink alcohol, apply make-up or put in contact lenses, or take them out.
9. Disinfect spilled samples or reagents with disinfectant.
10. If the extraction reagent come into contact with the skin or eyes, wash / rinse the affected area with plenty of water. If irritation is found, contact your doctor.
11. After the test, stow all components in a sealable plastic bag and dispose of them in household or residual waste.
12. Wash hands thoroughly after test completion.

Storage and stability

1. The test kits should be stored at temperatures of 4-30°C and should not be exposed to direct sunlight or moisture. Before use, tests stored at low temperature should be brought to room temperature.
2. Do not use expired and damaged products. The expiry date is printed on the outer packaging.
3. Under room temperature (15-30°C) and humidity of less than 60%, the test kits must be used within half an hour after opening the packaging. If the humidity exceeds 60%, use immediately after opening the packaging.

We have single serving, 5 servings and 20 servings .

(If you have specific requirement, pls let us know.)

single serving

The following is the certificates of Safe collection covid-19 self test rapid antigen test.

CERTIFICATE

EC Certificate No. 1434-IVDD-450/2021

EC Design-examination
Directive 98/79/EC concerning
***in vitro* diagnostic medical devices**

Polish Centre for Testing and Certification certifies
that manufactured by:

[redacted]

in vitro diagnostic medical devices
for self-testing

SARS-CoV-2 Antigen Rapid Test

in terms of design documentation, comply with requirements
of Annex III (Section 6) to Directive 98/79/EC (as amended)
implemented into Polish law,
as evidenced by the audit conducted by the PCBC
Validity of the Certificate: from 13.08.2021 to 27.05.2024

The date of issue of the Certificate: 13.08.2021

The date of the first issue of the Certificate: 27.05.2021

CE 1434

Issued under the Contract No. MD-55/2021
Application No: 105/2021
Certificate bears the qualified signature.
Warsaw, 13/08/2021
Module A1

Vice-President
Mgr Anna Wyroba

POLISH CENTRE FOR TESTING AND CERTIFICATION 02-844 Warsaw, 469 Puławska Street, tel. +48 22 46 45 200, e-mail:pcbc@pcbc.gov.pl