LEPU MEDICAL

SARS-CoV-2

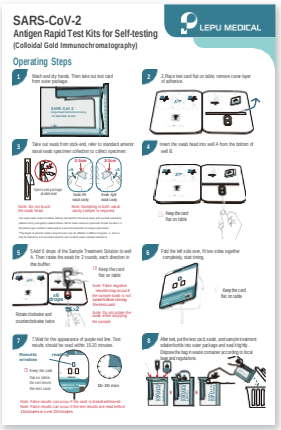
Antigen Rapid Test Kits for Self-testing

(Colloidal Gold Immunochromatography)

info@lepu-medical.it

Swab

buffer



Operation Card

Questo prodotto è un test immunologico rapido a flusso laterale destinato alla rilevazione qualitativa di antigeni nucleocapsidici SARS-CoV-2 da tamponi nasali anteriori auto-raccolti da un individuo di età pari o superiore a 18 anni o raccolti da un adulto da un individuo di età inferiore ai 18 anni. Questo test è destinato all'uso in soggetti con sintomi o altri motivi epidemiologici ove si sospetta un'infezione da COVID-19. Questo prodotto è destinato ad essere utilizzato come ausilio nella diagnosi dell'infezione da SARS-CoV-2.

Product Feature

- Non-invasivo
- Facilità Uso
- SELF- TEST
- Risultato in 15 minuti
- Alta attendibilità
- Economico

Performance Clinica

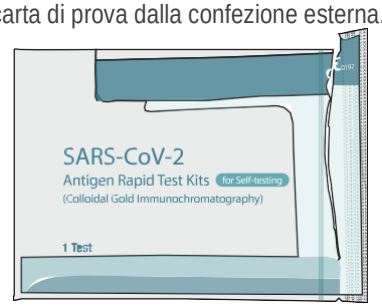
Lo studio sulle prestazioni cliniche del kit per il test rapido dell'antigene SARS-CoV-2 è stato condotto in Germania. Per eseguire il test sono stati utilizzati un totale di 222 campioni clinici. I campioni positivi e negativi sono stati tutti confermati mediante PCR. La sensibilità diagnostica e la specificità diagnostica del prodotto erano rispettivamente del 95,9% (90,8-98,2%) e del 100% (96,3-100,0%).

I risultati con correlazione al valore Ct dei campioni positivi sono stati mostrati nella tabella seguente

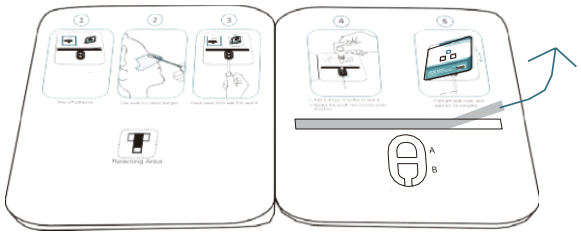
Ct Value	Diagnostic sensitivity	95%CI
≤ 30	96.2 %	88.3-98.7%
≤ 32	96.0 %	90.0-98.4%
≤ 34	95.5%	90.0-98.1%
≤ 36	95.9 %	90.8- 98.2%

Istruzioni di Utilizzo

1 Lavare e asciugare le mani. Quindi estrarre la carta di prova dalla confezione esterna.

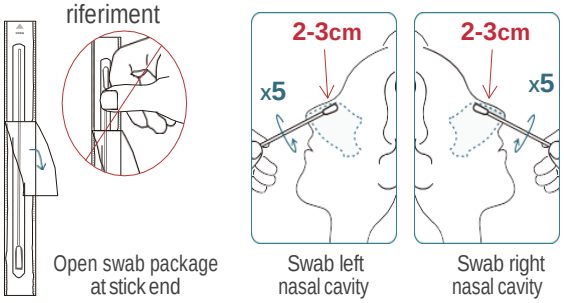


2 Posizionare la test card in piano sul tavolo, rimuovere lo strato di copertura dell'adesivo.



3 Estrarre il tampone dall'estremità del bastoncino, fare riferimento alla raccolta di campioni di

riferiment



Open swab package at stick end

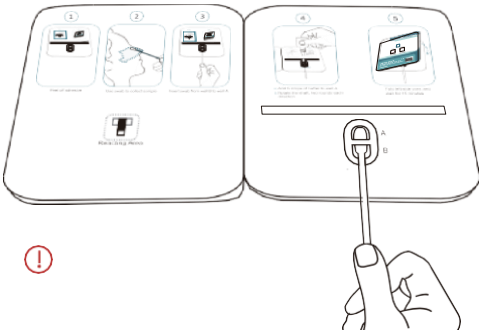
Swab left nasal cavity

Swab right nasal cavity

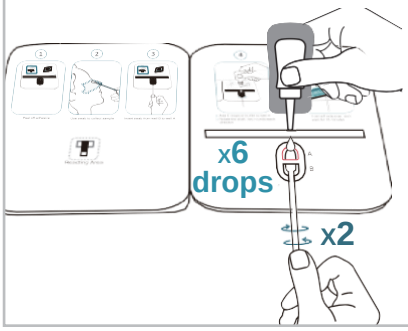
Nota: non toccare la testina del tampone. Nota: è richiesto il campionamento in entrambe le cavità nasali.

La testina del tampone nasale deve essere inserita completamente nella cavità nasale fino a quando non si avverte resistenza (circa 2-3 cm) e ruotata delicatamente 5 volte. Quando è stato rimosso, il campione deve essere prelevato allo stesso modo in un'altra cavità nasale per garantire la raccolta di campioni sufficienti. La lunghezza della cavità nasale anteriore degli utenti può essere diversa nelle diverse regioni, 2~3 cm è solo per riferimento. Si consiglia

4 Inserire la testina del tampone nel pozzetto A dal fondo del pozzetto B.

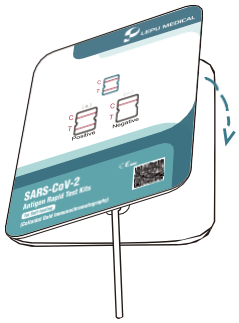


5 Aggiungere 6 gocce della soluzione per il trattamento del campione nel pozzetto A. Quindi ruotare il tampone per 2 giri, in ciascuna direzione

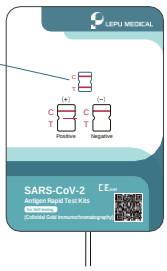


Nota: possono verificarsi risultati falsi negativi se il tampone del campione non viene girato prima di chiudere la test card.

6 Piegare il lato sinistro, unire completamente due lati, iniziare a cronometrare.

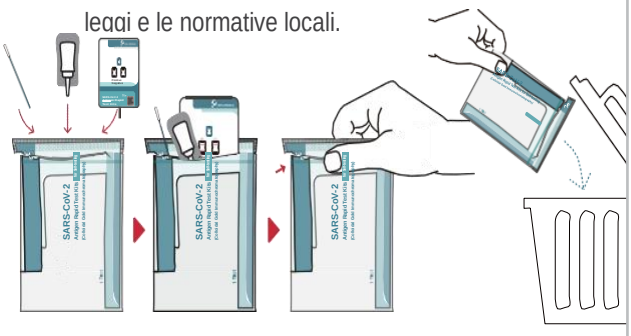


7 Attendi la comparsa della linea rosso porpora. I risultati del test devono essere letti entro 15-20 minuti



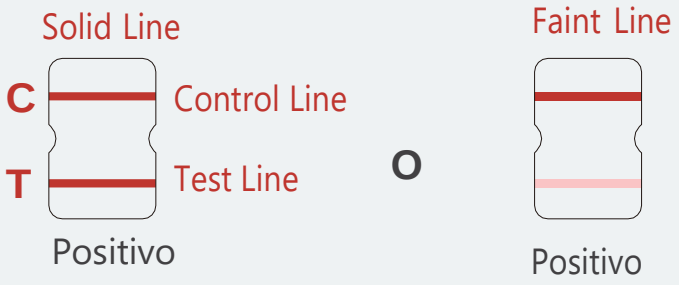
15-20 min

8 Dopo il test, inserire la scheda di prova, il tampone e il flacone della soluzione di trattamento del campione nella confezione esterna e sigillarlo ermeticamente. Smaltire il sacco nel contenitore dei rifiuti secondo le leggi e le normative locali.



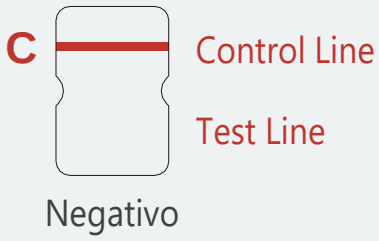
Interpretazione dei risultati

·Positivo (+): appare una banda rosso porpora nella linea di controllo (C) e nella linea di test (T).

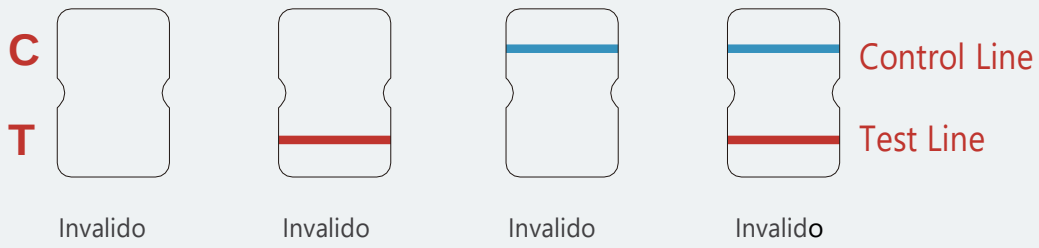


La linea di fondo può essere molto debole Qualsiasi linea rosa/viola visibile qui è positiva.

· Negativo (-): solo la linea di controllo (C) mostra una banda rosso porpora. Non appare alcuna banda rosso porpora nella linea di prova (T).



· Non valido: se "non appare alcuna banda rosso porpora nella linea di controllo (C)" e "appare una banda blu nella linea di controllo (C)", indica che il processo operativo non è corretto o che la carta di prova è stata danneggiata. In questo caso, leggere di nuovo attentamente il manuale di istruzioni e riprovare con una nuova carta reattiva. Se il problema persiste, interrompere immediatamente l'utilizzo di questo lotto di prodotti e contatta il tuo fornitore locale.



9Specifiche di Prodotto

1 Test

5 Tests

10 Tests

25 Tests

Self-testing | Home | Test Site | Airport | Hotel | Corporation | Mass Screening

LEPU MEDICAL

CE 0197 EN

SARS-CoV-2 Antigen Rapid Test Kits for Self-testing (Colloidal Gold Immunochromatography)

For In Vitro Diagnostic Use Only

[Packing Specifications]

1 test/kit, 5 tests/kit, 10 tests/kit, 25 tests/kit, 50 tests/kit

No.	Catalogue number	Spec.
1	CG3601	1 test/kit
2	CG3605	5 tests/kit
3	CG3610	10 tests/kit
4	CG3625	25 tests/kit
5	CG3650	50 tests/kit

【Produktname】

SARS-CoV-2 Antigen Rapid Test Kits for Self-testing (Colloidal Gold Immunochromatography)

【Intended Use】

This product is a rapid, lateral flow immunoassay intended for the qualitative detection of SARS-CoV-2 nucleocapsid antigens from anterior nasal swabs that are self-collected by an individual aged 18 years or older or are collected by an adult from an individual younger than 18 years old. This test is intended for use in individuals with symptoms or other epidemiological reasons to suspect a COVID-19 infection.
This product is intended to be used as an aid in the diagnosis of SARS-CoV-2 infection.
Results are for the identification of the SARS-CoV-2 nucleocapsid protein antigen. The antigen is generally detectable in anterior nasal swab specimens during the acute phase of infection. Positive results indicate the presence of viral antigens, but the clinical correlation with past medical history and other diagnostic information is necessary to determine infection status. Positive results do not rule out bacterial infection or co-infection with other viruses. The agent detected may not be the definite cause of disease.
Negative results should be treated as presumptive and confirmation with a molecular assay, if necessary, for patient management, may be performed. Negative results do not rule out SARS-CoV-2 infection and should not be used as the sole basis for treatment or patient management decisions including infection control decisions. Negative results should be considered in the context of a patient's recent exposures, history and the presence of clinical signs and symptoms consistent with COVID-19.
Individuals who test negative and continue to experience COVID-like symptoms should seek follow up care from their healthcare provider.

【Introduction】

Coronavirus, as the broad family of viruses, is a single strand plus RNA virus with an envelope. The virus is known to cause major diseases such as cold, Middle East Respiratory Syndrome (MERS) and Severe Acute Respiratory Syndrome (SARS). The core protein of SARS-CoV-2 is N protein (Nucleocapsid), which is a protein component inside the virus. It is relatively conservative among β-coronaviruses and is commonly used as a diagnostic tool for coronavirus. As the key receptor for SARS-CoV-2 to enter cells, ACE2 is significant for the study of viral infection mechanisms.

【Principle】

The current test kit is based on specific antibody-antigen reaction and immunoassay technique. The test strip consists of gold marked pad (coated with gold-marked SARS-CoV-2 N protein mouse anti human monoclonal antibody), sample pad, NC membrane (paired SARS-CoV-2 N protein mouse anti human monoclonal antibody coated on the test line (T) and goat anti-mouse IgG polyclonal antibody coated on the quality control line (C)) and absorbing paper.
During the test, the N protein in the specimen binds to the gold-marked SARS-CoV-2 N protein mouse anti human monoclonal antibody pre-coated on the gold marked pad, and the conjugate moves upward under the capillary effect, and then is trapped by N protein mouse anti human monoclonal antibody conjugate fixed in the Test Line (T). The higher the N protein content in the specimen, the more conjugates are trapped, and the darker the color of the Test Line (T). If there is no SARS-CoV-2 in the specimen or the virus content is below the detection limit, no color appears in the Test Line (T). A purple-red band will appear in the Control Line (C) regardless of whether there is a virus in the specimen. The purple-red band that appears in the Control Line (C) is the criteria for determining whether there is enough specimen and whether the chromatography process is normal.

【Main Components】

The product includes test cards, instruction for use, operation card, disposable sterile swabs and sample treatment solution. Each reagent kit contains 1 novel coronavirus (SARS-CoV-2) antigen test card and 1 bag of desiccant.

Disposable sterile swab information:
Nasal Swab could be provided based on customer's requirement.

Name	Application
Disposable sterile swab information	Nasal swab

CE 0123 MDD 93/42/EEC
Manufacturer 1: Zhejiang Gongdong Medical Technology Co., Ltd.Beicheng Industrial Area 318020 Huangyan China

CE 0197 MDD 93/42/EEC
Manufacturer 2: Jiangsu Changfeng Medical Industry Co., Ltd. Touqiao Town,Guangling District Yangzhou 225109 Jiangsu China

CE 0197 MDD 93/42/EEC
Manufacturer 3: Shenzhen KangDaAn Biological Technology Co., Ltd.Liuxiandong industrial zone,Xili street Nanshan district,Shenzhen 518055 Guangdong China

CE 0413 MDD 93/42/EEC
Manufacturer 4: Medico Technology Co., Ltd.Zhangbei Industrial Park, Longcheng Street, Longgang district, Shenzhen, 518100 Guangdong, China

CE 0197 MDD 93/42/EEC
Manufacturer 5: Goodwood Medical Care Ltd. 1-2 Floor, 3-919, Yongzheng Street, Jinzhou District, Dalian 116100 Liaoning, China

Spec.	Test card	Instruction manual	Operation card	Sample treatment solution	Swabs
1 test / kit	1 test	1	1	300μl×1	1 piece
5 tests / kit	5 tests	1	1	300 μl×5	5 pieces
10 tests / kit	10 tests	1	1	300 μl×10	10 pieces
25 tests / kit	25 tests	1	1	300 μl×25	25 pieces
50 tests / kit	50 tests	1	1	300 μl×50	50 pieces

Test card consists of paper shell, test strip, sample well and adhesive tape. The test strip, sample well and adhesive tape are attached on the paper shell.
The test strip consists of gold marked pad (coated with gold-marked SARS-CoV-2 N protein mouse anti human monoclonal antibody), sample pad, NC membrane (paired SARS-CoV-2 N protein mouse anti human monoclonal antibody coated on the test line (T) and goat anti-mouse IgG polyclonal antibody coated on the quality control line (C)) and absorbing paper.
The main compositions of sample treatment solution include tris, tritonX-100, sodium caseinate.

【General description】

The SARS-CoV-2 Antigen Rapid Test Kits for Self-testing (Colloidal Gold Immunochromatography) contains 3 core elements for operation:

- Test card: Test card which is book-shaped hinged test cardboard containing the test strip (for single use)
- Sample Treatment Solution: Bottle containing sample treatment solution (for single use)
- Nasal Swabs: Sterile swab (for single use)

1 Test Card

1 Sample Treatment Solution(buffer)

1 Swab

【Material required but not provided】
Clock or timer or stopwatch, Waste container

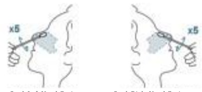
【Storage Conditions and Validity Period】

1. The test kit should be stored in a dry and dark place with temperature of 4-30℃, valid for 18 months.
2. The validity period of the test card is 1 hour after opening its inner package and it is suggested that the storage temperature should be 4 ~ 30 ℃ and the humidity should not exceed 70%.
3. The sample treatment solution should be used immediately after opening.

See package label for date of manufacture and expiration.

【Specimen Requirements】

This test kit is suitable for testing human anterior nasal swab specimens:
Specimen collection: During the collection process, relevant personnel should be well protected to avoid direct contact with the specimen. In case of accidental contact, timely disinfection should be carried out and necessary measures should be taken.
Anterior nasal swab specimen collection: During sampling, the nasal swab head should be entirely inserted into the nasal cavity until you feel resistance (about 2-3cm), and gently rotated 5 times. When it was removed, specimen should be taken in the same way in the other nasal cavity to ensure the collection of enough specimens.



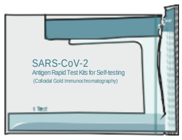
Swab Left Nasal Cavity   Swab Right Nasal Cavity

Specimen preservation: After the specimen are collected, please test immediately after sampling. Do not complete the test over 1 hour.

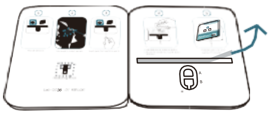
【Test Method】

Please read the instruction for use completely before performing any test, and use the reagents and specimens after returning to room temperature.

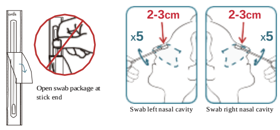
1. Wash and dry hands. Then take out test card from outer package.



2. Place test card flat on table, remove cover-layer of adhesive.



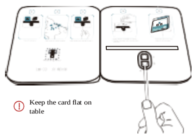
3. Take out swab from stick-end, refer to standard anterior nasal swab specimen collection to collect specimen: The nasal swab head should be entirely inserted into the nasal cavity until you feel resistance (about 2-3cm), and gently rotated 5 times. When it was removed, specimen should be taken in the same way in another nasal cavity to ensure the collection of enough specimens.
*The length of anterior nasal cavity of users may be different in different regions, 2~3cm is only for reference. It is recommended for user to insert swab until feel resistance.



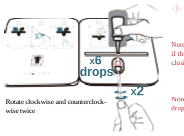
Note: Do not touch the swab head.

Note: Sampling in both nasal cavity sample is required.

4. Insert the swab head into well A from the bottom of well B.



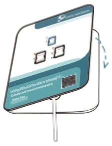
5. Add 6 drops of the Sample Treatment Solution to well A. Then rotate the swab for 2 rounds, each direction in the buffer.



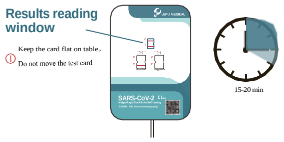
Note: False negative results may occur if the sample swab is not turned before closing the test card.

Note: Do not rotate the swab while dropping the sample

6. Fold the left side over, fit two sides together completely, start timing.

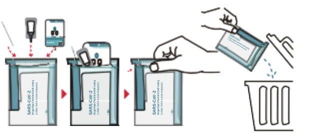


7. Wait for the appearance of purple-red line. Test results should be read within 15-20 minutes.

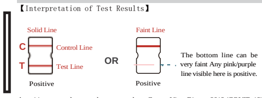


Note: False results can occur if the card is disturbed/moved.
Note: False results can occur if the test results are read before 15minutes or over 20 minutes.

8. After test, put the test card, swab, and sample treatment solution bottle into outer package and seal it tightly. Dispose the bag in medical waste container according to local laws and regulations.



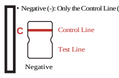
【Interpretation of Test Results】



The bottom line can be very faint Any pink/purple line visible here is positive.

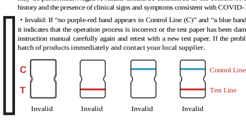
A positive test result means that you may have Corona Virus Disease 2019 (COVID-19). It is important to inform your health care provider of the results. Your healthcare provider will work with you to further confirm the diagnosis of COVID-19 and determine how best to care for you based on your test result(s) along with your medical history, symptoms and other related medical tests.

- Negative (-): Only the Control Line (C) shows a purple-red band. No purple-red band appears in the Test Line (T).



A negative test result means that proteins from SARS-CoV-2 which causes COVID-19 were not found in your sample.
Negative results are presumptive and confirmation with a molecular assay, if necessary, for patient management may be performed. Negative results should be considered in the context of an individual's recent exposures, history and the presence of clinical signs and symptoms consistent with COVID-19.

- Invalid: If "no purple-red band appears in Control Line (C)" and "a blue band appears in the Control Line (C)", it indicates that the operation process is incorrect or the test paper has been damaged. In this case, please read the instruction manual carefully again and retest with a new test paper. If the problem persists, please stop using this batch of products immediately and contact your local supplier.



【Limitations of the testing method】

1. The test results of this product should be combined with other clinical information and comprehensively judged by physicians, and should not be used as the only criterion.
2. This product is only used to determine the novel coronavirus (SARS-CoV-2) antigen in the specimen.
3. A negative test result may occur if the level of antigen in a sample is below the detection limit of the test.
4. False negative results may occur if a specimen is improperly collected or handled.
5. Invalid results may occur if inadequate sample treatment buffer is used (e.g., <6 drops). False negative results may occur if excessive sample treatment buffer is used (e.g., > 6 drops).
6. False negative results may occur if specimen swabs are not twirled within the test card.
7. False negative results may occur if swabs are stored in their paper sheath after specimen collection.
8. False negative results are more likely after seven days or more of symptoms.
9. This test detects both viable (live) and non-viable SARS-CoV-2. Test performance depends on the amount of virus (antigen) in the sample and may or may not correlate with viral culture results performed on the same sample.
10. The performance of the SARS-CoV-2 Antigen Rapid Test Kits for Self-testing (Colloidal Gold Immunochromatography) was evaluated using the procedures provided in this product insert only. Modifications to these procedures may alter the performance of the test.
11. The presence of high concentration mupirocin may interfere with the product and may cause false positive
12. Positive test results do not rule out co-infections with other pathogens.
13. Negative test results are not intended to rule in other non-SARS viral or bacterial infections.
14. Negative results do not rule out COVID-19 infection and it may be necessary to obtain additional testing with a molecular assay, if needed for patient management.
15. Positive test results do not differentiate between SARS-COV and SARS-CoV-2.

【Internal Quality Control】

The product has a Test Line (T) and a Control Line (C) on the surface of the test card. Neither the Test Line (T) nor the Control Line (C) is visible in the result window before applying a specimen. The control line is used for procedural control and should always appear if the test procedure is performed properly and the test reagents of the control line are working.

【Product Performance Index】

1.Determination of the Limit of Detection
SARS-CoV-2 Antigen Rapid Test Kits limit of detection (LOD) was determined by evaluating different concentrations of inactivated new coronavirus culture medium. Negative natural nasal swab specimens were eluted in 6 drops of sample treatment solution. 20 Swab eluates were combined and mixed thoroughly to create a clinical matrix pool to be used as the diluent. Inactivated new coronavirus culture medium was diluted in this natural nasal swab matrix pool to generate virus dilutions for testing.
The testing was performed according to the test procedure, with the virus dilutions applied directly onto the Swab to prepare the contrived nasal swab samples.
The LOD was determined as the lowest virus concentration that was detected ≥ 95% of the time (i.e., concentration at which at least 19 out of 20 replicates tested positive). The SARS-CoV-2 Antigen Rapid Test Kits for Self-testing (Colloidal Gold Immunochromatography) LOD in natural nasal swab matrix was confirmed 200 TCID50/mL.

2. Analysis of specificity
2.1 Cross-reaction: No cross-reactivity was seen with the following microorganisms when tested at the concentration presented in the table below.

Potential Cross-Reactant	Test Concentration
Human coronavirus OC43	10[illegible] TCID50/mL
Human coronavirus 229E	10[illegible] TCID50/mL
Human coronavirus NL63	10[illegible] TCID50/mL
Human coronavirus HKU1 recombinant N protein	50μg/mL
adenovirus	10[illegible] TCID50/mL
Human metapneumovirus (hMPV)	10[illegible] TCID50/mL
MERS coronavirus recombinant N protein	50μg/mL
Parainfluenza virus 1	10[illegible] TCID50/mL
Parainfluenza virus 2	10[illegible] TCID50/mL
Parainfluenza virus 3	10[illegible] TCID50/mL
Parainfluenza virus 4	10[illegible] TCID50/mL
Influenza A	10[illegible] TCID50/mL
Influenza B	10[illegible] TCID50/mL
Enterovirus (EV68)	10[illegible] TCID50/mL
Respiratory syncytial virus	10[illegible] TCID50/mL
Rhinovirus	10[illegible] TCID50/mL
Measles virus	10[illegible] TCID50/mL
Varicella zoster virus	10[illegible] TCID50/mL
Haemophilus influenzae	10[illegible] CFU/mL
Chlamydia pneumoniae	10[illegible] CFU/mL
Legionella pneumophila	10[illegible] CFU/mL
Mycobacterium tuberculosis	10[illegible] CFU/mL
Streptococcus pneumoniae	10^7 CFU/mL
Streptococcus pyogenes	10^7 CFU/mL
Bordetella pertussis	10[illegible] CFU/mL
Mycoplasma pneumoniae	10[illegible] CFU/mL
Candida albicans	10[illegible] CFU/mL
Staphylococcus epidermidis	10^7 CFU/mL
Staphylococcus aureus	10^7 CFU/mL
Pneumocystis giraldi	10[illegible] CFU/mL
Staphylococcus salivarius	10[illegible] CFU/mL
Combined human nasal Lotion	/

2.2 Interfering substances: No interference was seen with the following substances when tested at the concentration presented in the table below.

Potential Interfering substances	Test Concentration
Mucin	0.5%
Human whole blood	4%
HAMA	60 ng/mL
Biotin	1.2μg/mL
Benzocaine	2 mg/mL
Zanamivir	18μg/mL
Ribavirin	25μg/mL
Lopinavir	20μg/L
Ritonavir	18μg/mL
Acetylsalicylic acid	2 mg/dL
Ibuprofen	25 mg/dL
Phenylephrine	15%
Oxazole (nasal spray)	15%
Fluticasone	5%
Sodium chloride (containing preservatives)	10 mg/mL
Beclomethasone	2μg/mL
Budesonide	4ng/mL
Mometasone	2ng/mL
Strepsils (flurbiprofen 8.75mg)	5%
Throat candy (Mint)	5%
Naso GEL (NeilMed)	5%

3. Clinical performance

The clinical performance study for SARS-CoV-2 Antigen Rapid Test Kit was conducted in Germany. A total of 222 clinical samples were used to perform the test. The positive and negative samples were all confirmed by PCR. The diagnostic sensitivity and diagnostic specificity of the product was 95.9% (90.8-98.2%) and 100% (96.3-100.0%) respectively.
Results with correlation to Ct value of the positive samples were shown in the table below.

Ct Value	Diagnostic sensitivity	95%CI
≤ 30	96.2 %	88.3-98.7%
≤ 32	96.0 %	90.0-98.4%
≤ 34	95.5%	90.0-98.1%
≤ 36	95.9 %	90.8- 98.2%

【Warnings and Precautions】

1. For in vitro diagnostic use only. The product can be used for self-testing.
2. Do not eat or smoke while handling specimens.
3. The temperature and humidity of the experimental environment should be avoided to be too high, the reaction temperature should be 15-30°C and the humidity should be below 70%.
4. The packaging bag contains desiccant, do not eat.
5. It is recommended to test in a well-lit environment.
6. Before testing, please wash hands or wear clean gloves.
7. Please do not use the test card with damaged card bag packaging, unclear marking or beyond the expiration date.
8. A test card should be used within 1 hour after it is taken out from the aluminum foil bag.
9. Users shall take samples according to the instruction manual. Inadequate or inappropriate sample collection may yield error results and retesting with a new test may be required. Particular attention needs to be paid to appropriate sample collection technique.
10. Remove the covering layer of double-sided adhesive to prevent liquid splashing before testing.
11. Do not drop the dilution buffer into the wrong well.
12. In the process of testing, the test card should be placed on a horizontal table, and it should not be moved.
13. If the buffer solution makes contact with the skin or eye, wash/ flush with a large volume of water. If skin irritation, rash or other abnormal reaction occurs, please get medical advice/attention.
14. Avoid splashing or aerosol formation of specimen and buffer.
15. All users have to read the instruction prior to performing a test.
16. Do not mix or interchange different specimens.
17. Do not mix reagent of different lots or those for other products.
18. To avoid contamination, do not touch the head of provided swab when opening the swab pouch.
19. To avoid cross-contamination, do not reuse the sterilized swabs for specimen collection.
20. Do not dilute the collected swab with any solution except for the provided extraction buffer.
21. Keep foreign substances away from the test during the testing process. Contact with foreign substances, specifically bleach, may result in an incorrect test result.
22. Nasal swabs are not recommended for anyone who is prone to nosebleeds or has had facial or head injury/surgery in the last 6 months.
23. Patients with severe allergic rhinitis may have false positive results.
24. For patients with severe dry nasal mucosa, the sample volume may be insufficient due to the serious shortage of nasal secretions, resulting in inaccurate results.
25. Do not refrigerate or use after the expiration date (see packaging bag for expiration date).
26. Dispose of used specimens, test cards and other waste into waste container in accordance with relevant local laws and regulations.
27. It is suggested that the test should be performed in the company of people with normal vision for abnormal color vision users.

【Explanation of Symbols】

(symbol)	DO NOT USE IF PACKAGE IS DAMAGED	(symbol)	CONSULT INSTRUCTIONS FOR USE
(symbol)	DO NOT REUSE	(symbol)	USE-BY DATE
4℃ 30℃	TEMPERATURE LIMIT	(symbol)	DATE OF MANUFACTURER
(symbol)	MANUFACTURER	LOT	BATCH CODE
(symbol)	KEEP AWAY FROM SUNLIGHT	(symbol)	KEEP DRY
IVD	IN VITRO DIAGNOSTIC MEDICAL DEVICE	CE 0197	CE MARK
EC REP	AUTHORIZED REPRESENTATIVE IN THE EUROPEAN COMMUNITY	REF	CATALOGUE NUMBER

【Basic Information】

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