QuickFinder™ COVID-19/Flu Antigen Self Test
QUICK REFERENCE INSTRUCTIONS

PERFORMING THE TEST

9. Open swab package from the stick end, and remove the swab by the stick side.

DO NOT touch the swab head. DO NOT contaminate the swab head with any liquid gel soap as this can lead to false results.

10. Gently insert the swab head 1/2 to 3/4 inch into a nostril. For young children, swab should not be inserted more than 1/2 inch. DO NOT insert the swab any farther if you feel any resistance.

Using medium pressure, rub and rotate the swab against the inside of the nostril, making at least 5 circles. REPEAT IN THE OTHER NOSTRIL USING THE SAME SWAB.

NOTE: If you are swabbing others, please wear a face mask. With children, you may not need to insert the swab as far into the nostril. For very young children, you may need another person to steady the child’s head while swabbing.

STOP: Did you swab BOTH nostrils? Inaccurate test results may occur if the nasal sample is not properly collected.

11. Place the swab into the extraction buffer tube and completely immerse the swab head in the solution. Vigorously mix the solution by rotating the swab forcefully against the side of the tube at least 10 times, keeping the swab tip submerged in the buffer solution the entire time.

12. Remove the swab while squeezing the tube with your fingers to ensure that the sample on the swab is fully mixed into the buffer solution.

13. Attach the filter cap onto the test tube. Mix 10x each nostril.

14. Squeeze only 4 DROPS of the buffer solution into the sample well. DO NOT squeeze more than 4 drops from the tube into the sample well.

15. Set a timer and read the test result after 15 minutes. DO NOT disturb the cassette during this time. Inaccurate results can occur if the cassette is disturbed. DO NOT interpret test result before 15 minutes or after 30 minutes.

INTERPRETATION OF RESULTS

Test results are read and interpreted visually. Read result at 15–30 minutes with good lighting. Test results should not be read until after the sample has been added and the test has been allowed to run for 15 minutes, and only test lines that appear at the correct position and in the correct color should be read and interpreted.

POSITIVE RESULTS

If the control line at C is visible, and any other line or multiple lines on S, B and/or A appear, the test is positive.

FOR EASE OF USE, HOLD TEST CASSETTE NEXT TO THE EXAMPLE RESULT IMAGES ON THIS SHEET

INVALID RESULTS

If a control line is not visible at “C” after 15 minutes, even if any other line is visible in the results window, THE TEST HAS FAILED and is considered invalid.

NOTE: The images displayed above are examples only; additional invalid outcomes are possible. For a complete set of invalid results, see https://www.osanghc.com/covid-19-flu-combo-self-testing

NEGATIVE RESULTS

If the control line at “C” is visible and you do not see a line at ‘A’, ‘B’ or ‘S’, the test is negative.

If you still have COVID-19, Flu A, or Flu B symptoms, you should seek follow-up care with your healthcare provider. To increase the chance that the negative result for COVID-19, Flu A and Flu B is accurate, you should test again in 48 hours if this is your first test and you have symptoms on the first day of testing.

INTERESTING YOUR RESULTS

INVALID RESULT: This test did not work. The result should not be used. The test cannot determine if you have COVID-19, influenza A (Flu A), or influenza B (Flu B). The test needs to be repeated with a new kit and sample.

NEGATIVE RESULT: The virus from COVID-19, Flu A, and/or Flu B were not detected in the sample. A negative result does not mean that you do not have COVID-19, Flu A and/or Flu B infection. There is a higher chance of false negative results with antigen tests compared to laboratory-based molecular tests. If you tested negative and continue to experience COVID-19, Flu A and/or Flu B-like symptoms, you should seek follow-up care with your healthcare provider.

POSITIVE RESULT: The COVID-19, Flu A and/or Flu B virus(es) were detected in your sample. It is very likely that you have the respective infection(s) and are contagious. Please contact your healthcare provider or your local health authorities and follow local guidelines for self-isolation. There is a small chance that this test can give you a positive result that is incorrect (a false positive). Report your test result(s) at MakeMyTestCount.Org – this voluntary and anonymous reporting helps public health teams understand COVID-19 spread in your area and across the country and informs public health decisions.

AFTER TEST IS COMPLETED, DISPOSE OF USED MATERIALS IN HOUSEHOLD TRASH.
**INTENDED USE**

The QuickFinder™ COVID-19/Flu Antigen Self Test is a lateral flow immunassay intended for the qualitative detection and differentiation of SARS-CoV-2, influenza A, and influenza B protein antigens. This test is authorized for non-prescription home use with self-collection and self-testing of anterior nasal swab specimens from individuals aged 14 years or older, or with adult-collected anterior nasal swab specimens from individuals two (2) years or older. This test is only authorized for individuals who are suspected of having respiratory infection consistent with COVID-19 within the first four (4) days of symptom onset when tested at least twice over three days with at least 48 hours between tests. Internal controls will be provided to assist in performing serial testing should be performed in all individuals with negative results at least twice over three days (with 48 hours between tests) for a total of at least two more times with 48 hours in between tests for a total of three tests. If you have a negative result, it does not rule out COVID-19 and Flu A and/or Flu B. The clinical performance has not been established for all circulating variants but is anticipated to be reflective of the prevalent variants in circulation at the time and location of sample testing. Performance at the time of testing may vary depending on the variants circulating, including newly emerging strains of COVID-19 and influenza and their prevalence in your community. For more information on EUAs go here: https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization

**HOW TO USE THIS TEST**

Serial testing should be performed in all individuals with negative results, individuals who are suspected of having respiratory infection consistent with SARS-CoV-2, influenza A, and influenza B and should seek follow up care with their healthcare provider or physician. The QuickFinder™ COVID-19/Flu Antigen Self Test is only for use under the Food and Drug Administration’s Emergency Use Authorization.

**FREQUENTLY ASKED QUESTIONS**

Q: WHAT OCCURS IF BOTH COVID-19 AND INFuenza A OR B ARE DETECTED?

If both COVID-19 and influenza A or B are detected, you should self-isolate from others and contact your healthcare provider for medical advice about your positive result.

Q: WHAT IF I HAVE A POSITIVE TEST RESULT?

A: A positive result means that it is very likely you have COVID-19 or influenza because proteins from the virus that causes COVID-19 or influenza have been found in your sample. 844.760.0556

Email: covidhomeuri@osanglc.com

**LIMITATIONS**

- There is a higher chance of false negative results with antigen tests than with laboratory-based molecular tests due to the sensitivity of the test technology. This means that there is a higher chance of having COVID-19 or Flu and testing negative.
- You should follow up with your healthcare provider if you have symptoms of COVID-19 or Flu and test negative.
- You should follow up with your healthcare provider if you test negative but continue to have symptoms of COVID-19 or Flu.
- If you test negative but continue to have symptoms of COVID-19 or Flu, you should test again in 48 hours since you are instructed to test negative in 48 hours between tests. Performing serial testing for COVID-19 and influenza can be similar. Results are for the simultaneous identification of SARS-CoV-2, influenza A, and influenza B protein antigens but do not differentiate between SARS-CoV-2 and SARS-CoV-2 viruses and are not intended to detect influenza C virus. The targets of this test are generally detectable from specimens collected using nasal swabs during the acute phase of infection. Positive results indicate the presence of viral antigens, but clinical correlation and diagnostic information is necessary to determine infection status. Positive results do not rule out bacterial infection or co-infection with other viruses. The antigen detected may not be the definitive cause of disease. Individuals who test positive with the QuickFinder™ COVID-19/Flu Antigen Self Test should self-isolate and seek follow-up care with your healthcare provider as additional testing may be necessary. All negative results are presumptive and confirmation with a molecular assay, if necessary for patient management, may be required. Incorrect test results may occur if a specimen is incorrectly collected or handled.

**INDEX OF SYMBOLS**

- Manufacturer:
- Date of manufacture:
- Contains allergens:
- Catalogue number:
- Temperature limit:
- Use-by date:
- Batch code:
- Contraindication:
- Intra vitreous:
- Consult instructions for use:

If you have any questions about using the test or reading the results, please call our customer care hotline. Telephone: 844.760.0556

Email: covidhomeuri@osanglc.com

**MANUFACTURING SITE**

- OSANG Healthcare, Ltd.
- 132, Anyangcheon-eudong-ro, Dongan-gu, Anyang-si, Gyeonggi-do, Korea (14040) www.osanghc.com

**SARISSA COVID-19/Flu Test**

- 100 Prospect St., 3rd Fl.
- Pasadena, CA 91101

**REFUND POLICY**

- 100% money-back guarantee if not satisfied as molecular tests. To be eligible for refund, the test must have been performed properly, and you should follow-up with your healthcare provider if you have symptoms of COVID-19 or influenza.
QuickFinder™ COVID-19 / Flu Antigen Self Test

HOW TO USE

1. SWAB
2. MIX
3. READ

The Kit Contains:
- 1 Test Cassette
- 1 Sterile Swab
- 1 Extraction Buffer Tubes & Filter Caps
- 1 QR (Quick Reference Instructions)

NEXUS DX, INC.
6759 Mesa Ridge Road San Diego, CA 92121
Manufactured for OSANG LLC
215 N Marengo Ave. 3rd Floor. Pasadena, CA 91101
Tel: 1-844-760-0556
Technical Support: hometest@osangllc.com

Manufacturing Site:
OSANG Healthcare Co., Ltd.
132, Anyangcheondong-ro, Dongan-gu, Anyang-si, Gyeonggi-do, 14040, S.Korea

Scan this QR Code for more information.

- This test can be used at home on people aged 2 years and older.
- You may need to purchase additional tests to perform serial (repeat) testing.
- This test is more likely to give you a false negative result when you have SARS-CoV-2, Influenza A, and Influenza B, than a lab-based molecular test.
- Do not use if you've had symptoms longer than 4 days or no symptoms at all.
- In the USA, this product has not been FDA cleared or approved; but has been authorized by FDA under an Emergency Use Authorization.
- This test is authorized for non-prescription home use with self-collected anterior nares nasal swab samples from individuals with symptoms of SARS-CoV-2, influenza A, and influenza B within the first 4 days of symptom onset.
- The emergency use of this test was only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostics for detection and/or diagnosis of COVID-19 under Section 564(b)(1) of the Federal Food, Drug, and Cosmetic Act, 21 U.S.C. § 360bbb-3(b)(1), unless the declaration is terminated or authorization is revoked sooner.
- This test has been authorized only for the detection of proteins from SARS-CoV-2, influenza A, and influenza B, not for any other viruses or pathogens.
- This test does NOT determine if you had COVID-19 in the past or if you have immunity.
- Determining a negative result requires multiple tests.

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QuickFinder™ COVID-19 / Flu Antigen Self Test

How to Use

The Same Test with 3 Results

1. Insert tube here
2. Wait 15 mins
3. Insert tube here

For Ages 2 and Up.

For Emergency Use Authorization (EUA) only.
For in vitro diagnostic use.

QuickFinder™ COVID-19 / Flu Antigen Self Test

Ⓒ OSANG Healthcare Co., Ltd.

Manufacturing Site:
OSANG Healthcare Co., Ltd.
132, Anyangcheondong-ro,Dongan-gu, Anyang-si, Gyeonggi-do, 14040, S.Korea

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HOW TO USE

SWAB

MX

TEST

READ

COVID-19

Flu

Flu A/Flu B

QuickFinder™ COVID-19 / Flu Antigen Self Test

The Same Test with 3 Results

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For in vitro diagnostic use.
For Ages 2 and Up.

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This product has been authorized only for the detection of proteins from SARS-CoV-2, influenza A and influenza B, not for any other viruses or pathogens.
This test does NOT determine if you had COVID-19 in the past or if you have immunity.
Determining a negative result requires multiple tests.

The Kit Contains
2 Test Cassettes 2 Sterile Swabs 2 Extraction Buffer Tubes & Filter caps 2 QRI (Quick Reference Instructions)  Needed but not provided: Timer

FOR SYMBOL GLOSSARY, REFER TO QUICK REFERENCE INSTRUCTIONS.
QuickFinder™ COVID-19 / Flu Antigen Self Test

- This test can be used at home on people aged 2 years and older.
- The emergency use of this product is only authorized for the duration
- You may need to purchase additional tests to perform serial (repeat)

- This test is more likely to give you a false negative result when you of COVID-19 under Section 564(b)(1) of the Federal Food, Drug, and
- This test does NOT determine if you had COVID-19 in the past or if
- This test is authorized for non-prescription home use with you have immunity.
- Determining a negative result requires multiple tests.

- Do not use if you've had symptoms longer than 4 days or no symptoms
- This product has been authorized only for the detection of proteins from
- In the USA, this product has not been FDA cleared or approved; but pathogens.
- This test has been authorized by FDA under an Emergency Use Authorization.
- This test does NOT determine if you had COVID-19 in the past or if
- A negative result may indicate the need for serial testing.
QuickFinder™ COVID-19 / Flu Antigen Self Test

The Box Contains
- 5 Test Cassettes
- 5 Sterile Swabs
- 5 Extraction Buffer Tubes & Filter caps
- 1 QRI (Quick Reference Instructions)

Needed but not provided: Timer

Manufacturing Site:
Manufactured for OSANG LLC
OSANG Healthcare Co., Ltd.
215 N Marengo Ave. 3rd Floor.
132, Anyangcheondong-ro, Pasadena, CA 91101
14040, S.Korea

Technical Support:
hometest@osangllc.com
NEXUS DX, INC.
6759 Mesa Ridge Road
San Diego, CA 92121

For Symbol Glossary, refer to Quick Reference Instructions.

Antigen Self Test

- This test can be used at home on people aged 2 years and older.

- The emergency use of this product is only authorized for the duration of emergency use of in vitro diagnostics for detection and/or diagnosis of COVID-19 under Section 564(b)(1) of the Federal Food, Drug, and Cosmetic Act, 21 U.S.C. § 360bbb-3(b)(1), unless the declaration is terminated or authorization is revoked sooner.

- Do not use if you've had symptoms longer than 4 days or no symptoms at all.

- This product has been authorized only for the detection of proteins from SARS-CoV-2, influenza A and influenza B, not for any other viruses or pathogens.

- In the USA, this product has not been FDA cleared or approved; but has been authorized by FDA under an Emergency Use Authorization.

- This test does NOT determine if you had COVID-19 in the past or if you have SARS-CoV-2, Influenza A, and Influenza B, than a lab-based molecular test.

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The test can be used at home on people aged 2 years and older. It is intended to be used in a single test only.

You may need to purchase additional tests to perform serial (repeat) testing.

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For Symbol Glossary, refer to Quick Reference Instructions.

For Emergency Use Authorization (EUA) only.
For in vitro diagnostic use.
For Ages 2 and Up.

QuickFinder™ COVID-19 / Flu Antigen Self Test

The same test with 3 results

For Counseling the Authorization (EUA) only.
For Quicker, simpler, and more accurate testing.

Scan this QR Code or more information

This test can be used at home on people aged 2 years and older. It is intended to be used in a single test only.

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