

Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test

Healthcare Provider Instructions for Use

For use under an Emergency Use Authorization (EUA) Only

For *in vitro* diagnostic use

For use with anterior nasal swab specimens

1. Intended Use

The Speedy Swab Rapid COVID-19 + Flu A&B Antigen Self-Test is a lateral flow immunoassay 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-collected 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 with signs and symptoms of respiratory infection consistent with COVID-19 within the first five (5) days of symptom onset when tested at least twice over three days with at least 48 hours between tests.

Clinical signs and symptoms of respiratory viral infection due to SARS-CoV-2 and influenza can be similar.

Results are for the simultaneous identification of SARS-CoV-2, influenza A virus, and influenza B virus protein antigens, but do not differentiate between SARS-CoV and SARS-CoV-2 viruses and are not intended to detect influenza C antigens.

The viral antigens targeted by 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 with patient 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 definitive cause of the disease. Individuals who test positive with the Speedy Swab Rapid COVID-19 + Flu A&B Antigen Self-Test should self-isolate and seek follow-up care with their physician or 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 performed. Negative results do not rule out SARS-CoV-2, influenza A, and influenza B infection and should not be used as the sole basis for treatment or patient management decisions, including infection control decisions such as isolating from others and wearing masks. 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 SARS-CoV-2, influenza A, and influenza B infection.

Individuals who test negative and continue to experience SARS-CoV-2 and/or influenza-like symptoms of fever, cough, and/or shortness of breath may still have SARS-CoV-2 and/or influenza infection and should seek follow up care with their physician or healthcare provider.

The Speedy Swab Rapid COVID-19 + Flu A&B Antigen Self-Test is only for use under the Food and Drug Administration's Emergency Use Authorization.

2. Summary and Explanation

Along with the common cold, influenza is one of the most common acute respiratory infections, producing symptoms such as headache, chills, dry cough, body aches, and fever. It affects 5%-20% of the United States population annually, resulting in more than 200,000 hospitalizations and 36,000 deaths.¹ The influenza A virus is typically more prevalent and is associated with the most serious influenza epidemics, while influenza B infections usually present with milder symptoms. Diagnosis is difficult because the initial symptoms can be similar to those caused by other infectious agents. Considering that the influenza virus is highly contagious, accurate diagnosis and prompt treatment of patients can have a positive effect on public health. Accurate diagnosis and the ability to distinguish between A or B antigens can also help reduce the inappropriate use of antibiotics and gives the physician the opportunity to prescribe an antiviral therapy. Initiation of antiviral therapy should begin as soon as possible after onset, ideally within 48 hours of the appearance of symptoms, as treatment may reduce the duration of symptoms.²

Coronaviruses are enveloped RNA viruses that are found broadly among humans, other mammals, and birds. The viruses are known to cause mild symptoms, but sometimes severe respiratory, enteric, hepatic, and neurological diseases can occur. Seven coronavirus species are known to cause human disease, four of which (229E, OC43, NL63 and HKU-1) are quite prevalent and can cause mild cold symptoms, especially in immunocompetent people.³ There are three other strains that are known to cause severe acute respiratory disease. These strains include severe acute respiratory syndrome coronavirus (SARS-CoV), Middle East respiratory syndrome coronavirus (MERS-CoV), and the 2019 Novel Coronavirus (COVID-19). These strains are all zoonotic in origin and have been linked to sometimes fatal respiratory illness. The prevalence of SARS and MERS has been quite low in recent years; the Novel Coronavirus (COVID-19) was recently identified in December 2019. The main manifestations of illness include fever, fatigue, and dry cough. Nasal congestion, runny nose, sore throat, myalgia, and diarrhea are found in a few cases. Most epidemiological studies suggest a 1–14-day incubation period. The median incubation period is estimated at 5.1 days, with most developing symptoms before 11.5 days.⁴ Infected but asymptomatic people can also be an infectious source. The Speedy Swab Rapid COVID-19 + Flu A&B Antigen Self-

¹ US Department of Health and Human Services. National Institutes of Health. Influenza [Fact Sheet]. January 2011.

² Montalto N, Byrd R. An Office-Based Approach to Influenza: Clinical Diagnosis and Laboratory Testing. *American Family Physician*. January 2003; 67:11-118.

³ Cui J, Li F, Shi ZL. Origin and evolution of pathogenic coronaviruses. *Nat Rev Microbiol* 2019; 17: 181-192.

⁴ Lauer S, et al. The Incubation Period of Coronavirus Disease 2019 (COVID-19) From Publicly Reported Confirmed Cases: Estimation and Application. *Ann Intern Med* 2020 May 5;172(9):577-582.

Test can provide rapid detection of influenza A, influenza B, and/or SARS-CoV-2 viral antigens from symptomatic patients.

3. Principle of the Procedure

The Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test is a rapid, qualitative lateral flow immunoassay for the determination of the presence of SARS-CoV-2 and influenza A&B antigens in anterior nasal swab specimens. The test strip in each device contains mouse monoclonal capture antibodies to the nucleocapsid protein (NP) of SARS-CoV-2, Influenza A and Influenza B and goat anti-Mouse IgG control antibody immobilized in the test and control regions on the nitrocellulose membrane, respectively. The conjugate pad is coated with colloidal gold conjugated SARS-CoV-2, Influenza A and Influenza B antibody and colloidal gold conjugated IgY. Once the extracted specimen is added in the sample well of the test device, it migrates chromatographically on the membrane by capillary action. The formation of the specific antibody-antigen conjugate complex Au-IgY is visualized by the presence of a colored band in the control region, which validates the test results. If SARS-CoV-2 nucleocapsid antigen and influenza A and influenza B nucleoprotein antigens are present in a specimen, the specific antibody antigen colored conjugate complex is formed and a distinct color band in the labeled region (S, B, A) is observed. Absence of this colored band in any of the test regions (S, B, A) indicates a negative result (when the control band is present).

4. Materials Provided

The components supplied with the different test configurations are as follows:

Component	1 Test/Kit	2 Tests/Kit	4 Tests/Kit
Test Device	1	2	4
Sterile Nasal Swab	1	2	4
Test Tube with Buffer Solution vials containing 500uL of buffer	1	2	4
Tube Holder in Box Top	1	1	1
Quick Reference Instruction (QRI)	1	1	1

5. Materials Required but not Provided

- A timer: required to determine the time to read the test results after addition of the extracted specimen to the test device

- Personal protective equipment: Disposable masks and gloves if swabbing others

6. Warnings and Precautions

- 1 Read all instructions carefully before performing the test. Failure to follow the instructions may result in inaccurate test results.
- 2 In the USA, this product has not been FDA cleared or approved, but has been authorized by FDA under an Emergency Use Authorization. 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. The emergency use of this product is 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.
- 3 **Serial testing should be performed in individuals with SARS-CoV-2 negative results at least twice over three days (with 48 hours between tests) for symptomatic individuals. You may need to purchase additional tests to perform this serial (repeat) testing.**
- 4 **Consistent with serial testing recommendations for SARS-CoV-2, for multi-analyte tests, symptomatic individuals who test positive for influenza A or B on the initial test but test negative for SARS-CoV-2 should be tested again in 48 hours to evaluate for co-infection with SARS-CoV-2 infection.**
- 5 This test may only be used in symptomatic individuals.
- 6 An anterior nasal swab sample can be self-collected by an individual aged 14 years and older. Children aged 2 to 13 years should be tested by an adult.
- 7 Do not use on anyone under 2 years of age.
- 8 Wear a safety mask or other face-covering when collecting a specimen from a child or another individual.
- 9 Do not use if any of the test kit contents or packaging is opened or damaged.
- 10 Test components are for single use. Do not re-use the test strip, buffer liquid, or swab.
- 11 If any liquid spills from the buffer tube, discard test components and re-start test using new test components.
- 12 Do not use kit past its expiration date.
- 13 Only use the nasal swabs provided in the kit. Do not touch the swab head prior to testing.
- 14 Exposure to hand sanitizer may cause false positive results with this test.
- 15 The Buffer vial contains only enough liquid for one test. Do not use the same Buffer vial with an additional test as invalid or incorrect results may occur.
- 16 Do not interchange or mix components from different kit lots.
- 17 Inadequate or inappropriate sample collection, storage, and transport may yield false test results.

- 18 Once opened, the test device should be used within 60 minutes.
- 19 **Do not read test results before 15 minutes or after 20 minutes. Results read before 15 minutes or after 20 minutes may lead to a false positive, false negative, or invalid result.**
- 20 Ensure that there is sufficient lighting for testing and interpretation.
- 21 **Keep testing kit and kit components away from children and pets before and after use. Avoid contact with your skin, eyes, nose, or mouth. Do not ingest any kit components. The reagent solution contains harmful chemicals (see table below). If the solution contacts your skin, eyes, nose, or mouth, flush with large amounts of water. If irritation persists, seek medical advice: <https://www.poisonhelp.org> or 1-800-222-1222.**

Chemical Name	GHS Code for each Ingredient	Concentrations
Proclin 300	H317, allergic skin reaction	0.1%
Trimethylsilyl acetamide	H316, mild skin irritation	0.03%

- 22 For more information on EUAs please visit: <https://www.fda.gov/emergency-preparedness-and-response/mcmlegal-regulatory-and-policy-framework/emergency-use-authorization>
- 23 For the most up to date information on COVID-19, please visit: www.cdc.gov/COVID19

7. Kit Storage and Stability

Store the Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test at room temperature (15 to 30°C / 60 to 86°F) in the original packaging, away from direct sunlight. Kit contents are stable in the unopened box until the expiration date printed on the kit box.

- Do not freeze any of the test kit components.
- Ensure all test components are at room temperature before use (15 – 30 °C / 60 – 86 °F).
- Do not use test kit after expiration date.
- The Test Device must remain in the sealed foil pouch until use. Once the pouch has been opened, the test device should be used within 60 minutes. Use beyond one hour may not produce accurate results.

8. Sample Handling, and Storage

- Negative results can occur from inadequate sample collection and/or handling.
- To obtain accurate results, do not use visually bloody or overly viscous samples.
- Test components should not be used for any other purpose other than performing this test.
- Test samples immediately after collection. Swabs should be placed into buffer within 60 minutes of collection. Inoculated buffer should be added to the device within 30 minutes of swab addition and mixing.
- Only nasal swabs can be used with this test. Use of nasal washes, nasal aspirates, or nasopharyngeal swabs has not been validated for use with this test.
- When collecting anterior nasal swab specimens, make sure to use only the swab included in the test kit.


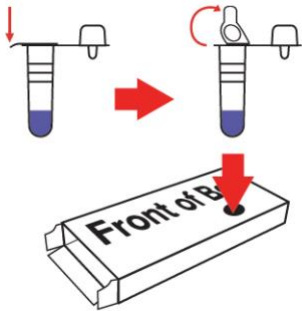
9. Quality Control

Each Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test has a built-in internal procedural control. The reddish/pink line appearing at the “C” position is an internal procedural control. This procedural control line indicates that sufficient flow has occurred. A distinct reddish/pink Control line should always appear if the test has been performed correctly. If the Control line does not appear, the test result is invalid, and a new test should be performed using a new swab and new test kit.

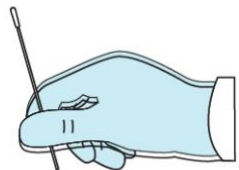
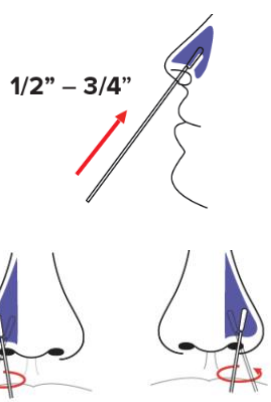
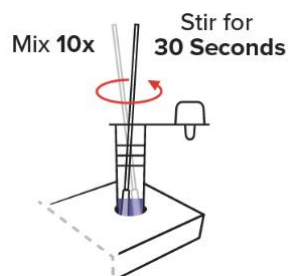
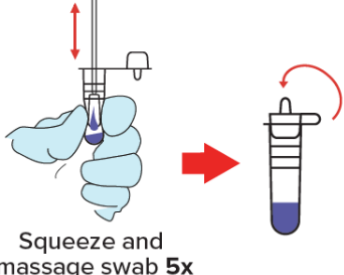
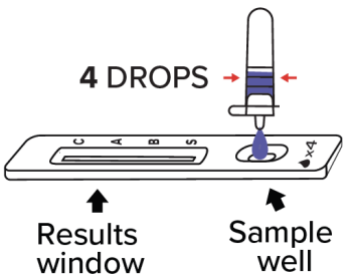
Contact Watmind USA at +1 866-928-6463 or technicalsupport@watmindusa.com if you experience a problem.


10. Test Procedure

Steps outlining the test are as follows:

Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test Steps		
Before Getting Started		Step 1: Check expiration date on the outside of the box. Do not use beyond the expiration date.
		Step 2: Wash hands thoroughly for at least 20 seconds before and after handling nasal swab samples.
		Step 3: Clean the tabletop on which the test will be performed. Before testing, read the User Instructions carefully.
Prepare the Materials		NOTE: It is recommended that gloves are used during testing. A face mask should be worn if swabbing others.
		Step 4: Arrange the materials on a clean, dry, flat surface. Use only one of each of the materials provided for each test.
		Step 5: Pick up the Test Tube and remove the sealing foil of the tube.
		Step 6: Locate the tube holder on the front of the box labeled "Push Tube Here" and insert the buffer tube into the tube holder.
		Step 7: Remove the Test Device from its foil pouch.
		NOTE: Use the Test Device within one hour of opening the test pouch.


Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test Steps

<p>Performing the Test</p>		<p>Step 8: Open swab package from its stick end and remove the swab from this end. NOTE: DO NOT touch the swab head or lay the swab on any surface.</p>
	 <p align="center">5x for 15 seconds, each nostril</p>	<p>Step 9: Gently insert the swab $\frac{1}{2}$ to $\frac{3}{4}$ inch into a nostril. NOTE: DO NOT insert the swab any farther if you feel any resistance. Using medium pressure, rub and rotate the swab against the inside walls of the nostril, making at least 5 circles for at least 15 seconds. REPEAT IN THE OTHER NOSTRIL USING THE SAME SWAB. NOTE: 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.</p>
		<p>Step 10: Place the swab into the buffer solution as soon as possible after collection and completely immerse the swab head in the sample. Swirl the swab in the solution by rotating the swab forcefully against the side of the tube at least 10 times for 30 seconds, keeping the swab tip submerged in the buffer solution the entire time.</p>
		<p>Step 11: Squeeze the tube 5 times with your fingers to ensure that the sample on the swab is fully mixed into the buffer solution. Attach the dropper cap to the test tube.</p>
		<p>Step 12: Squeeze only 4 DROPS of the Buffer Solution into the sample well. NOTE: DO NOT squeeze more than 4 drops from the tube. Additional sample volume may yield inaccurate results.</p>

Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test Steps		
Performing the Test		<p>Step 13: Set a timer and read the test result at 15 minutes.</p> <p>NOTE: DO NOT disturb the card during this time. Inaccurate results can occur if the card is disturbed.</p>

11. Interpretation of Results

Test results are read and interpreted visually. Read results at 15 minutes with good lighting.
WARNING: Do not read the result before 15 minutes or after 20 minutes. Inaccurate test interpretations may occur.



C = Control Line
A = Influenza A Line
B = Influenza B Line
S = COVID-19 (SARS-CoV 2)

Look for lines next to 'C' (Control), 'A', 'B' and 'S'.

INVALID RESULTS

If a pink or red 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**.

STOP: If the control (C) line is not visible, the test is invalid. Do not continue reading the results. Re-test with a new swab and new test device.

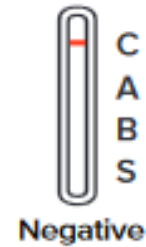


NEGATIVE RESULTS

At 15 minutes, the appearance of **ONLY** the Control Line indicates that Influenza A, Influenza B, or SARS-CoV-2 has **NOT** been detected.

To increase the chance that the negative result for COVID-19 is accurate, you should test again in 48 hours after the first day of testing.

If respiratory symptoms persist, you should seek follow up care with your healthcare provider.



POSITIVE RESULTS

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

NOTE: Any pink to red line, no matter how faint, should be considered an indication of a positive result when the control line is also present.



COVID-19
Positive



Flu A
Positive



Flu B
Positive



Flu A+
COVID-19
Positive



Flu B+
COVID-19
Positive



Flu A+
Flu B
Positive



Flu A+B+
COVID-19
Positive

Consult your healthcare provider to discuss your positive test result. Self-isolate at home per CDC recommendations to stop spreading virus to others.

Understanding Your Results

Invalid Result

If the control (C) line is not visible, the test is invalid. Re-test with a new swab and new test device.

COVID-19 Positive (+) Result

Repeat testing does not need to be performed if patients have a positive result at any time.

A positive test result means that the virus that causes COVID-19 was detected in the sample, and it is very likely the individual has COVID-19 and is contagious. Please contact the patient's doctor/primary care physician (if applicable) and the local health authority immediately and instruct your patient to adhere to the local guidelines regarding self-isolation. There is a very small chance that this test can give a positive result that is incorrect (a false positive).

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. Individuals who test positive with the Speedy Swab Rapid COVID-19 + Flu A&B Antigen Self-Test should self-isolate and seek follow up care with their physician or healthcare provider as additional confirmatory testing with a molecular test for positive results may also be necessary, if there is a low likelihood of COVID-19, such as in individuals without known exposures to COVID-19 or residing in communities with low prevalence of infection.

COVID-19 Negative (-) Result

To increase the chance that the negative result for COVID-19 is accurate, you should test again in 48 hours if the individual has symptoms on the first day of testing.

A negative test result indicates that the virus that causes COVID-19 was not detected in the sample. A negative result does not rule out COVID-19. There is a higher chance of false negative results with antigen tests compared to laboratory-based tests such as PCR tests. If the test is negative but COVID-19-like symptoms, e.g., fever, cough, and/or shortness of breath continue, follow up testing for SARS-CoV-2 with a molecular test or testing for other respiratory disease should be considered. If applicable, seek follow up care with the primary health care provider.

All negative results should be treated as presumptive and confirmation with a molecular assay may be necessary if there is a high likelihood of SARS-CoV-2 infection, such as in an individual with a close contact with COVID-19 or with suspected exposure to COVID-19 or in communities with high prevalence of infection. 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.

Repeat Testing is needed to improve test accuracy for negative SARS-CoV-2 results. Please follow the table below when interpreting test results with symptoms. Serial (repeat) SARS-CoV-2 testing does not need to be performed if patients have a positive SARS-CoV-2 result.

Status on First Day of Testing	Day 0 (First Test)	Serial Testing?	Day 2 (Second Test)	Final Interpretation
With Symptoms	SARS-CoV-2 (+) Influenza A and B (-)	NO	Not needed	Positive for COVID-19 Presumptive negative for Influenza
	SARS-CoV-2 (+) Influenza A and/or B (+)	NO	Not needed	Positive for COVID-19 Positive for Influenza A and/or B
	SARS-CoV-2 (-) Influenza A and/or B (-)	YES	SARS-CoV-2 (+) Influenza A and/or B (-)	Positive for COVID-19 Presumptive Negative for Influenza
	SARS-CoV-2 (-) Influenza A and/or B (+)	YES	SARS-CoV-2 (+) Influenza A and/or B (+)	Positive for COVID-19 Positive for Influenza A and/or B
	SARS-CoV-2 (-) Influenza A and/or B (-)	YES	SARS-CoV-2 (-) Influenza A and/or B (+)	Presumptive Negative for COVID-19 Positive for Influenza A and/or B
	SARS-CoV-2 (-) Influenza A and/or B (-)	YES	SARS-CoV-2 (-) Influenza A and/or B (-)	Presumptive Negative for COVID-19 Presumptive Negative for Influenza
	SARS-CoV-2 (-) Influenza A and/or B (-)	YES	SARS-CoV-2 (+) Influenza A and/or B (+)	Positive for COVID-19 Positive for Influenza A and/or B
	SARS-CoV-2 (-) Influenza A and/or B (+)	YES	SARS-CoV-2 (-) Influenza A and/or B (-)	Presumptive Negative for COVID-19 Positive for Influenza A and/or B
	SARS-CoV-2 (-) Influenza A and/or B (+)	YES	SARS-CoV-2 (-) Influenza A and/or B (+)	Presumptive Negative for COVID-19 Positive for Influenza A and/or B
	SARS-CoV-2 (-) Influenza A and/or B (+)	YES	SARS-CoV-2 (+) Influenza A and/or B (+)	Positive for COVID-19 Positive for Influenza A and/or B

12. Limitations

- The performance of this test was established based on the evaluation of a limited number of clinical specimens collected between June 2023 and March 2024. 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 the clinical evaluation. Performance at the time of testing may vary depending on the variants circulating, including newly emerging strains of SARS-CoV-2 and their prevalence, which change over time.
- 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 this test will give a false negative result in an individual with COVID-19 as compared to a molecular test, especially in samples with low viral load.
- All COVID-19 and influenza A&B antigen test negative results are presumptive and confirmation with a molecular assay may be necessary.
- If you continue to have symptoms of COVID-19 or influenza, and both your first and second tests are negative, you may not have COVID-19 or influenza.
- If the test is positive, then proteins from the virus that causes COVID-19 or influenza infection have been found in the sample and the individual likely has a respiratory infection with SARS-CoV-2 or influenza.
- This test is read visually and has not been validated for use by those with impaired vision or color-impaired vision.
- Incorrect test results may occur if a specimen is incorrectly collected or handled.
- Based on sequence analysis, a potential for cross-reactivity between the SARS-CoV-2 test and HKU1 exists. Wet testing for HKU1 coronavirus was not conducted and therefore, cross-reactivity between SARS-CoV-2 and HKU1 coronavirus cannot be ruled out.
- Individuals who recently received nasally administered influenza A or influenza B vaccine may have false positive test results after vaccination.

13. Analytical Performance

13.1 Analytical Sensitivity: Limit of Detection (LoD) - Analytical Sensitivity

The limit of detection (LoD) for the Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test was established using dilutions of one SARS Coronavirus 2 (SARS-CoV-2) Isolate USA-WA1/2020 strain (BEI Resources Catalog number NR-52287), two influenza A strains (Influenza A H1N1: Influenza A/Indiana/02/2020, Influenza A H3N2: Influenza A/Alaska/232/2015) and two influenza B strains (Influenza B/Washington/02/2019, Influenza B/Florida/4/2006) in negative clinical matrix. The isolate dilutions were tested by adding fifty (50) μ L to the head of the nasal swab and extracting the swab per the Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test Instructions for Use.

In this study, range finding testing was followed by final dilution testing to determine the LoD of the assay. Range finding involved testing a series of 10-fold dilutions in replicates of three (3) to determine the starting point for the dilution series to determine LoD. The dilution of each virus which resulted in the lowest concentration that generated 100% positive detection rate was set as the target for the next dilution series, which involved testing three (3) replicates of two (2)-fold dilutions. In the final dilution testing, the lowest concentration that generated $\geq 95\%$ positive detection rate was set as the LoD concentration. Confirmatory testing was performed by testing twenty (20) replicates at the preliminary (1X) LoD concentrations.

Virus Strains	Stock Concentration	LoD Concentration	LOD per Swab	# Positive / # Total Tested	Percent Detected
SARS-CoV-2 USA-WA1/2020	7.9×10^5 TCID ₅₀ /mL	7.9×10^2 TCID ₅₀ /mL	4.0×10^1 TCID ₅₀	20/20	100%
Influenza A A/Indiana/02/2020 (H1N1)	9.7×10^8 CEID ₅₀ /mL	9.7×10^6 CEID ₅₀ /mL	4.9×10^5 CEID ₅₀	20/20	100%
Influenza A A/Alaska/232/2015 (H3N2)	2.6×10^8 CEID ₅₀ /mL	2.6×10^4 CEID ₅₀ /mL	1.3×10^3 CEID ₅₀	20/20	100%
Influenza B B/Washington/02/2019 (Victoria)	2.1×10^9 CEID ₅₀ /mL	2.1×10^6 CEID ₅₀ /mL	1.1×10^5 CEID ₅₀	19/20	95%
Influenza B B/Florida/4/2006 (Yamagata)	7.3×10^7 CEID ₅₀ /mL	1.8×10^4 CEID ₅₀ /mL	9.2×10^2 CEID ₅₀	20/20	100%

13.2 Inclusivity (Analytical Reactivity)

The analytical reactivity of the Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test was evaluated with currently available influenza strains and SARS-CoV-2 strains using a dilution series. Concentrations listed in the table below indicate the lowest detectable concentrations in pooled nasal wash (PNW) and/or pooled nasal swab matrix (PNSM) for which all replicates were positive.

Virus	Strain	Concentration in PNW	Concentration in PNSM	Concentration Units
Influenza A, H1N1	A/Michigan/45/2015	1.0×10^7	3.9×10^2	CEID ₅₀ /mL
	A/St. Petersburg/61/2015	1.2×10^6	4.7×10^5	CEID ₅₀ /mL
	A/Hawaii/66/2019	1.9×10^7	1.9×10^7	CEID ₅₀ /mL
	A/Wisconsin/588/2019	2.8×10^3	2.8×10^3	FFU/mL
	A/Idaho/07/2018	8.0×10^1	1.6×10^2	TCID ₅₀ /mL
	A/Brownsville/39H/2009	N/A	8.0×10^1	TCID ₅₀ /mL
	A/Massachusetts/15/2013	N/A	1.6×10^6	CEID ₅₀ /mL
	A/Bangladesh/3002/2015	N/A	6.5×10^4	CEID ₅₀ /mL
	A/Dominican Republic/7293/2013	N/A	1.3×10^3	TCID ₅₀ /mL
	A/Iowa/53/2015	N/A	2.9×10^6	CEID ₅₀ /mL
Influenza A, H3N2	A/New York/21/2020	6.5×10^4	6.5×10^4	FFU/mL
	A/Tasmania/503/2020	3.3×10^4	3.3×10^4	FFU/mL
	A/Hong Kong/2671/2019	1.6×10^6	6.2×10^5	CEID ₅₀ /mL
	A/Hong Kong/45/2019	3.8×10^3	3.8×10^3	FFU/mL
	A/Singapore/INFIMH-16-0019/2016	1.1×10^5	5.5×10^4	CEID ₅₀ /mL
Influenza A, H1N1v	A/Ohio/09/2015	N/A	7.0×10^5	CEID ₅₀ /mL
Influenza A, H1N2	A/Minnesota/19/2011	N/A	4.00×10^6	CEID ₅₀ /mL
Influenza A, H3N2v	A/Indiana/08/2011	N/A	2.0×10^2	TCID ₅₀ /mL
Influenza A, H7N3	A/northern pintail/Illinois/100S3959/2010	N/A	2.8×10^5	CEID ₅₀ /mL
Influenza A, H5N1	A/mallard/Wisconsin/2576/2009	N/A	1.6×10^5	CEID ₅₀ /mL
Influenza B, Non-Victoria, Non-Yamagata	B/Maryland/1/1959	4.5×10^2	4.5×10^2	CEID ₅₀ /mL
Influenza B, Victoria	B/Colorado/6/2017	1.6×10^5	1.6×10^5	CEID ₅₀ /mL
	B/Florida/78/2015	1.7×10^6	1.7×10^6	CEID ₅₀ /mL

Virus	Strain	Concentration in PNW	Concentration in PNSM	Concentration Units
Influenza B, Yamagata	B/Texas/06/2011	8.0×10^6	4.0×10^6	CEID ₅₀ /mL
	B/Wisconsin/1/10	7.05×10^1	3.53×10^1	TCID ₅₀ /mL
SARS-CoV-2	hCoV-19/USA/MD-HP40900/2022 (Lineage XBB.1.5; Omicron Variant)	N/A	8.0×10^1	TCID ₅₀ /mL

13.3 Analytical Specificity: Cross-reactivity and Microbial interference

Cross-Reactivity

Cross-reactivity and microbial interference with related pathogens, high prevalence disease agents, and normal or pathogenic flora that are reasonably likely to be encountered in the clinical specimen were evaluated with the Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test. Each organism was tested in replicates of three (3) at the concentration listed in the following table of test results.

For cross reactivity, the organisms listed below were tested in negative samples. No cross-reactivity was observed for any of the organisms tested, except for SARS-coronavirus which exhibited cross-reactivity when tested neat (7.9×10^3 TCID₅₀/mL). A titration of SARS-CoV was performed and it was shown that cross reactivity was no longer observed for SARS-CoV at 7.9×10^0 TCID₅₀/mL. These results are not unexpected in that the Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test targets the SARS-CoV-2 nucleocapsid protein which is present on both the SARS-CoV and SARS-CoV-2 viruses.

No cross reactivity was observed for any organism.

Microorganism Introduced	Concentration Tested for Cross Reactivity	Influenza A Test Results (Positive/Total)	Influenza B Test Results (Positive/Total)	SARS-CoV-2 Test Results (Positive/Total)
Human coronavirus 229E	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Human coronavirus OC43	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Human coronavirus NL63	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
SARS-coronavirus Urbani	7.9×10^3 TCID ₅₀ /mL	0/3	0/3	3/3
MERS-coronavirus	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Adenovirus Goman	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Adenovirus Adenoid 71	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Human metapneumovirus 4	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Parainfluenza virus 1	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Parainfluenza virus 2	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Parainfluenza virus 3	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Parainfluenza virus 4b	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Enterovirus	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3

Microorganism Introduced	Concentration Tested for Cross Reactivity	Influenza A Test Results (Positive/Total)	Influenza B Test Results (Positive/Total)	SARS-CoV-2 Test Results (Positive/Total)
Respiratory syncytial virus A	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Respiratory syncytial virus B	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Rhinovirus	1.0×10^5 pfu/mL	0/3	0/3	0/3
<i>Haemophilus influenzae</i>	1.1×10^6 cfu/mL	0/3	0/3	0/3
<i>Streptococcus pneumoniae</i>	1.1×10^6 cfu/mL	0/3	0/3	0/3
<i>Streptococcus pyogenes</i>	1.1×10^6 cfu/mL	0/3	0/3	0/3
<i>Candida albicans</i>	1.1×10^6 cfu/mL	0/3	0/3	0/3
<i>Bordetella pertussis</i>	$>1.0 \times 10^4$ cfu/mL	0/3	0/3	0/3
<i>Mycoplasma pneumoniae</i>	1.1×10^6 cfu/mL	0/3	0/3	0/3
<i>Chlamydia pneumoniae</i>	1.1×10^6 ifu/mL	0/3	0/3	0/3
<i>Legionella pneumophila</i>	1.1×10^6 cfu/mL	0/3	0/3	0/3
<i>Mycobacterium tuberculosis</i>	1.1×10^6 cfu/mL	0/3	0/3	0/3
<i>P. jiroveci</i> - <i>S. cerevisiae</i>	1.1×10^6 cfu/mL	0/3	0/3	0/3
<i>Staphylococcus aureus</i>	1.1×10^6 cfu/mL	0/3	0/3	0/3
<i>Staphylococcus epidermidis</i>	3.3×10^7 cfu/mL	0/3	0/3	0/3
<i>Corynebacterium</i> sp.	$>1.0 \times 10^4$ cfu/mL	0/3	0/3	0/3
<i>Escherichia coli</i>	$>2.5 \times 10^4$ cfu/mL	0/3	0/3	0/3
<i>Lactobacillus</i> sp.	$>1.0 \times 10^4$ cfu/mL	0/3	0/3	0/3
<i>Moraxella catarrhalis</i>	1.1×10^6 cfu/mL	0/3	0/3	0/3
<i>Neisseria meningitidis</i>	1.05×10^5 cfu/mL	0/3	0/3	0/3
<i>Neisseria</i> sp.	$>1.0 \times 10^4$ cfu/mL	0/3	0/3	0/3
<i>Pseudomonas aeruginosa</i>	1.1×10^6 cfu/mL	0/3	0/3	0/3
<i>Streptococcus salivarius</i>	$>1.0 \times 10^4$ cfu/mL	0/3	0/3	0/3
Measles Virus	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Mumps Virus	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3
Epstein Barr Virus	$>2.5 \times 10^3$ CEID ₅₀ /mL	0/3	0/3	0/3
Cytomegalovirus	1.5×10^5 TCID ₅₀ /mL	0/3	0/3	0/3

Microbial Interference

For evaluating microbial interference against the SARS-CoV-2, Influenza A (H1N1), Influenza B (Victoria) test lines, the organisms were tested with co-spiked SARS-CoV-2 Isolate USA-WA1/2020 strain (BEI Resources Catalog number NR-52287), Influenza A/Alaska/232/2015 (International Reagent Resource Catalog number FR-1540) or Influenza B/Florida/04/2006 (BEI Resources Catalog number NR-41795) diluted to 3x LoD concentration in negative clinical matrix. SARS-CoV-2, Influenza A and Influenza B were detected in all samples tested in the presence of potentially interfering organisms demonstrating no interference.

Microorganism Introduced	Microorganism Concentration	Influenza A Test Results (Positive/Total)	Influenza B Test Results (Positive/Total)	SARS-CoV-2 Test Results (Positive/Total)
Human coronavirus 229E	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Human coronavirus OC43	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Human coronavirus NL63	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
SARS-coronavirus Urbani	7.9 × 10 ⁰ TCID ₅₀ /mL	3/3	3/3	3/3
MERS-coronavirus	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Adenovirus Goman	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Adenovirus Adenoid 71	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Human metapneumovirus 4	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Parainfluenza virus 1	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Parainfluenza virus 2	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Parainfluenza virus 3	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Parainfluenza virus 4b	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Enterovirus	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Respiratory syncytial virus A	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Respiratory syncytial	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Rhinovirus	1.0 × 10 ⁵ pfu/mL	3/3	3/3	3/3
<i>Haemophilus influenzae</i>	1.1 × 10 ⁶ cfu/mL	3/3	3/3	3/3
<i>Streptococcus pneumoniae</i>	1.1 × 10 ⁶ cfu/mL	3/3	3/3	3/3
<i>Streptococcus pyogenes</i>	1.1 × 10 ⁶ cfu/mL	3/3	3/3	3/3
<i>Candida albicans</i>	1.1 × 10 ⁶ cfu/mL	3/3	3/3	3/3
<i>Bordetella pertussis</i>	>1.0 × 10 ⁴ cfu/mL	3/3	3/3	3/3
<i>Mycoplasma pneumoniae</i>	1.1 × 10 ⁶ cfu/mL	3/3	3/3	3/3
<i>Chlamydia pneumoniae</i>	1.1 × 10 ⁶ ifu/mL	3/3	3/3	3/3
<i>Legionella pneumophila</i>	1.1 × 10 ⁶ cfu/mL	3/3	3/3	3/3
<i>Mycobacterium tuberculosis</i>	1.1 × 10 ⁶ cfu/mL	3/3	3/3	3/3
<i>P. jiroveci-S. cerevisiae</i>	1.1 × 10 ⁶ cfu/mL	3/3	3/3	3/3
<i>Staphylococcus aureus</i>	1.1 × 10 ⁶ cfu/mL	3/3	3/3	3/3
<i>Staphylococcus epidermidis</i>	3.3 × 10 ⁷ cfu/mL	3/3	3/3	3/3

Microorganism Introduced	Microorganism Concentration	Influenza A Test Results (Positive/Total)	Influenza B Test Results (Positive/Total)	SARS-CoV-2 Test Results (Positive/Total)
<i>Corynebacterium</i> sp.	>1.0 × 10 ⁴ cfu/mL	3/3	3/3	3/3
<i>Escherichia coli</i>	>2.5 × 10 ⁴ cfu/mL	3/3	3/3	3/3
<i>Lactobacillus</i> sp.	>1.0 × 10 ⁴ cfu/mL	3/3	3/3	3/3
<i>Moraxella catarrhalis</i>	1.1 × 10 ⁶ cfu/mL	3/3	3/3	3/3
<i>Neisseria meningitidis</i>	1.05 × 10 ⁵ cfu/mL	3/3	3/3	3/3
<i>Neisseria</i> sp.	>1.0 × 10 ⁴ cfu/mL	3/3	3/3	3/3
<i>Pseudomonas aeruginosa</i>	1.1 × 10 ⁶ cfu/mL	3/3	3/3	3/3
<i>Streptococcus salivarius</i>	>1.0 × 10 ⁴ cfu/mL	3/3	3/3	3/3
Measles Virus	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Mumps Virus	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3
Epstein Barr Virus	>2.5 × 10 ³ CEID ₅₀ /mL	3/3	3/3	3/3
Cytomegalovirus	1.5 × 10 ⁵ TCID ₅₀ /mL	3/3	3/3	3/3

13.4 Endogenous Interfering Substances

A total of twenty nine (29) potentially interfering substances, either naturally present in respiratory specimens or artificially introduced into the nasal cavity or nasopharynx, were tested to evaluate the performance of the Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test when these substances were added to the nasal swab head in the absence (negative) and presence (positive) of co-spiked SARS-CoV-2 Isolate USA-WA1/2020 strain (BEI Resources Catalog number NR-52287), influenza A (Influenza A/Alaska/232/2015, International Reagent Resource Catalog number FR-1540), or two different strains of influenza B (Influenza B/Florida/04/2006, BEI Resources Catalog number NR-41795) diluted to 3x LoD concentration. Each substance was tested in replicates of three (3). For testing Zanamivir, pooled nasal swab matrix (PNSM) was used instead of pooled nasal wash (PNW).

At 15% v/v, hand sanitizer cream lotion produced a false positive and false negative result in three of three (3/3) replicates for all three target analytes. When diluted to 5% v/v, SARS-CoV-2 only false positive results were observed in all three (3) test replicates.

False positive results were also detected for Influenza A and Influenza B analytes in the presence of FluMist Quadrivalent Live Intranasal Influenza Virus Vaccine at 15% v/v and 1.5% v/v. One out of three (1/3) replicates tested positive for SARS-CoV-2 at 1.5% v/v.

Substance	Concentration Tested	Virus Unspiked Test Results (Positive/Total)		
		SARS-CoV-2	Flu A	Flu B
Biotin	3,500 ng/mL	0/3	0/3	0/3
Chloraseptic (Menthol/Benzocaine)	1.5 mg/mL	0/3	0/3	0/3

Substance	Concentration Tested	Virus Unspiked Test Results (Positive/Total)		
		SARS-CoV-2	Flu A	Flu B
Fluticasone Propionate	5% v/v	0/3	0/3	0/3
Homeopathic (Alkalol)	10% v/v	0/3	0/3	0/3
Human Whole Blood (EDTA tube)	4% v/v	0/3	0/3	0/3
Mucin (porcine stomach, type II)	0.5%	0/3	0/3	0/3
Mupirocin	10 mg/mL	0/3	0/3	0/3
Nasal Drops (Phenylephrine)	15% v/v	0/3	0/3	0/3
Nasal Spray (Cromolyn)	15% v/v	0/3	0/3	0/3
Nasal Spray (Oxymetazoline)	15% v/v	0/3	0/3	0/3
Naso GEL (NeilMed)	5% v/v	0/3	0/3	0/3
Sore Throat Phenol Spray	15% v/v	0/3	0/3	0/3
Tamiflu (Oseltamivir Phosphate)	5 mg/mL	0/3	0/3	0/3
Tobramycin	4 µg/mL	0/3	0/3	0/3
Zicam	5% v/v	0/3	0/3	0/3
FluMist Quadrivalent Live Intranasal Influenza Virus Vaccine	15% v/v	0/3	3/3	3/3
FluMist Quadrivalent Live Intranasal Influenza Virus Vaccine	1.5% v/v	1/3	3/3	3/3
FluMist Quadrivalent Live Intranasal Influenza Virus Vaccine	0.15% v/v	0/3	0/3	0/3
Body and Hand Lotion	0.5% w/v	0/3	0/3	0/3
Body Lotion with 1.2% Dimethicone	0.5% w/v	0/3	0/3	0/3
Hand Lotion	5% w/v	0/3	0/3	0/3
Hand Sanitizer with Aloe, 62% Ethyl Alcohol	5% v/v	0/3	0/3	0/3
Hand Sanitizer Cream Lotion	15% v/v	3/3	3/3	3/3
Hand Sanitizer Cream Lotion	5% v/v	3/3	0/3	0/3
Hand Sanitizer Cream Lotion	1.5% v/v	0/3	0/3	0/3
Hand Sanitizer, 80% Ethanol	15% v/v	0/3	0/3	0/3
Hand Soap Liquid Gel	10% w/v	0/3	0/3	0/3
Hand Soap Liquid Gel	5.0% w/v	0/3	0/3	0/3
Zanamivir	282 ng/mL	0/3	0/3	0/3

Substance	Concentration Tested	Virus Spiked Test Results (Positive/Total)		
		SARS-CoV-2	Flu A	Flu B
Biotin	3,500 ng/mL	3/3	3/3	3/3
Chloraseptic (Menthol/Benzocaine)	1.5 mg/mL	3/3	3/3	3/3
Fluticasone Propionate	5% v/v	3/3	3/3	3/3
Homeopathic (Alkalol)	10% v/v	3/3	3/3	3/3
Human Whole Blood (EDTA tube)	4% v/v	3/3	3/3	3/3
Mucin (porcine stomach, type II)	0.5%	3/3	3/3	3/3
Mupirocin	10 mg/mL	3/3	3/3	3/3
Nasal Drops (Phenylephrine)	15% v/v	3/3	3/3	3/3
Nasal Spray (Cromolyn)	15% v/v	3/3	3/3	3/3
Nasal Spray (Oxymetazoline)	15% v/v	3/3	3/3	3/3
Naso GEL (NeilMed)	5% v/v	3/3	3/3	3/3
Sore Throat Phenol Spray	15% v/v	3/3	3/3	3/3

Substance	Concentration Tested	Virus Spiked Test Results (Positive/Total)		
		SARS-CoV-2	Flu A	Flu B
Tamiflu (Oseltamivir Phosphate)	5 mg/mL	3/3	3/3	3/3
Tobramycin	4 µg/mL	3/3	3/3	3/3
Zicam	5% v/v	3/3	3/3	3/3
FluMist Quadrivalent Live Intranasal Influenza Virus Vaccine	15% v/v	3/3	3/3	3/3
FluMist Quadrivalent Live Intranasal Influenza Virus Vaccine	1.5% v/v	3/3	3/3	3/3
FluMist Quadrivalent Live Intranasal Influenza Virus Vaccine	0.15% v/v	3/3	3/3	3/3
Body and Hand Lotion	0.5% w/v	3/3	3/3	3/3
Body Lotion with 1.2% Dimethicone	0.5% w/v	3/3	3/3	3/3
Hand Lotion	5% w/v	3/3	3/3	3/3
Hand Sanitizer with Aloe, 62% Ethyl Alcohol	5% v/v	3/3	3/3	3/3
Hand Sanitizer Cream Lotion	15% v/v	3/3	3/3	3/3
Hand Sanitizer Cream Lotion	5% v/v	3/3	3/3	3/3
Hand Sanitizer Cream Lotion	1.5% v/v	3/3	3/3	3/3
Hand Sanitizer, 80% Ethanol	15% v/v	3/3	3/3	3/3
Hand Soap Liquid Gel	10% w/v	3/3	3/3	1/3
Hand Soap Liquid Gel	5.0% w/v	3/3	3/3	3/3
Zanamivir	282 ng/mL	3/3	3/3	3/3

13.5 High Dose Hook Effect

No high-dose hook effect was observed with the Speedy Swab Rapid COVID-19 + FLU A&B Antigen Self-Test when testing high concentrations of SARS-CoV-2, Influenza A or Influenza B strains.

Virus Strain Tested	Concentration	Results (Positive/Total)
USA-WA1/2020	2.8×10^6 TCID ₅₀ /mL	3/3
Influenza A/Indiana/02/2020 (H1N1)	9.7×10^8 CEID ₅₀ /mL	3/3
Influenza A/Alaska/232/2015 (H3N2)	2.6×10^8 CEID ₅₀ /mL	3/3
Influenza B/Washington/02/2019 (Victoria)	2.1×10^9 CEID ₅₀ /mL	3/3
Influenza B/Texas/06/2011 (Yamagata)	1.6×10^9 CEID ₅₀ /mL	3/3

13.6 Competitive Interference

Competitive interference was evaluated to determine if high concentrations of one target virus interferes with the detection of low concentrations of another target virus. No competitive interference was observed between SARS-CoV-2 and influenza A and B as listed in the table below.

Combination Number	Concentration			Results (Positive/Total)		
	Influenza A	Influenza B	SARS-CoV-2	Influenza A	Influenza B	SARS-CoV-2
1	6.5×10^7 CEID ₅₀ /mL	2.2×10^5 CEID ₅₀ /mL	Negative	3/3	3/3	0/3

Combination Number	Concentration			Results (Positive/Total)		
	Influenza A	Influenza B	SARS-CoV-2	Influenza A	Influenza B	SARS-CoV-2
2	6.5 × 10 ⁷ CEID ₅₀ /mL	Negative	2.4 × 10 ³ TCID ₅₀ /mL	3/3	0/3	3/3
3	2.0 × 10 ⁵ CEID ₅₀ /mL	3.7 × 10 ⁷ CEID ₅₀ /mL	Negative	3/3	3/3	0/3
4	Negative	3.7 × 10 ⁷ CEID ₅₀ /mL	2.4 × 10 ³ TCID ₅₀ /mL	0/3	3/3	3/3
5	2.0 × 10 ⁵ CEID ₅₀ /mL	Negative	4.0 × 10 ⁵ TCID ₅₀ /mL	3/3	0/3	3/3
6	Negative	2.2 × 10 ⁵ CEID ₅₀ /mL	4.0 × 10 ⁵ TCID ₅₀ /mL	0/3	3/3	3/3

14. Clinical Evaluation

A prospective study was completed at seven sites in the United States between June 2023 and March 2024 for clinical validation of the detection of SARS-CoV-2, Influenza A, and Influenza B in subject-collected anterior nasal (AN) swab samples. The study evaluated the performance in symptomatic individuals (i.e., those with symptoms of upper respiratory infection). The testing of each enrolled subject included two AN swab samples: one was collected by a healthcare professional and sent for testing using an FDA-cleared molecular comparator method, and the other swab was either self-collected or collected from the study subject by another lay individual. Each subject then performed the test according to the Quick Reference Instructions (QRI) provided with the test kit in a simulated home-use environment. 1245 symptomatic subjects were enrolled with a total of 1171 subjects evaluable. Performance was determined by comparing Speedy Swab Rapid COVID-19 + Flu A&B Antigen Self-Test results to results from the highly sensitive RT-PCR molecular comparator. 1132 individuals met the eligibility criteria for SARS-CoV-2, in which 148 (13.1%) individuals tested positive for SARS-CoV-2 and 984 (86.9%) individuals tested negative for SARS-CoV-2. From the same study cohort, 1171 individuals met the eligibility criteria for Flu A and Flu B. 80 (6.8%) individuals tested positive and 1091 (93.2%) individuals tested negative for Flu A. The study also had 30 (2.6%) individuals who tested positive and 1141 (97.4%) individuals who tested negative for Flu B. At least 30 enrolled subjects were below the age of 14. The positive percent agreement (PPA) is 92.6% [87.2%, 95.8%] for SARS-CoV-2, 82.5% [72.7%, 89.3%] for Influenza A, and 90.0% [74.4%, 96.5%] for Influenza B. The negative percent agreement (NPA) is 99.4% [98.7%, 99.7%] for SARS-CoV-2, 99.7% [99.2%, 99.9%] for Influenza A, and 99.9% [99.7%, 100.0%] for Influenza B.

Subject Demographics

	Subjects (by lay-user collection and testing (N=140))	Self-collecting and testing (N=1032)	Overall (N=1172)
Age			
Mean (SD)	8.5 (5.9)	42.1 (16.0)	38.1 (18.6)
Median [Min, Max]	8 [2, 63]	40 [14, 89]	37 [2, 89]
Age Group			
≥ 2 - <14 years of age	129 (92.1%)	0 (0.0%)	129 (11.0%)
14 - 24 years of age	10 (7.1%)	156 (15.1%)	166 (14.2%)
> 24 - 64 years of age	1 (0.7%)	784 (76.0%)	785 (67.0%)
≥ 65 years of age	0 (0.0%)	92 (8.9%)	92 (7.8%)
Sex at Birth			
Female	71 (50.7%)	647 (62.7%)	718 (61.3%)
Male	69 (49.3%)	385 (37.3%)	454 (38.7%)
Ethnicity			
Hispanic/Latino	82 (58.6%)	619 (60.0%)	701 (59.8%)
Not Hispanic/Latino	58 (41.4%)	402 (39.0%)	460 (39.2%)
Unknown/Prefer not to answer	0 (0.0%)	11 (1.1%)	11 (0.9%)
Race			
American Indian or Alaskan Native	1 (0.7%)	2 (0.2%)	3 (0.3%)
Asian	0 (0.0%)	7 (0.7%)	7 (0.6%)
Black or African American	35 (25.0%)	305 (29.6%)	340 (29.0%)
Native Hawaiian/Pacific Islander	1 (0.7%)	0 (0.0%)	1 (0.1%)

	Subjects (by lay-user collection and testing (N=140))	Self-collecting and testing (N=1032)	Overall (N=1172)
White	91 (65.0%)	672 (65.1%)	763 (65.1%)
Unknown/Prefer not to answer	3 (2.1%)	2 (0.2%)	5 (0.4%)
Other (Mixed race/biracial)	9 (6.4%)	44 (4.3%)	53 (4.5%)

SARS-CoV-2 Performance

SARS-CoV-2	Comparator Positives	Comparator Negatives	Total
Investigational Positives	137	6	143
Investigational Negatives	11	978	989
Total	148	984	1132

Positive Percent Agreement = $(137/148) = 92.6\%$ (95% CI: 87.2% - 95.8%)

Negative Percent Agreement = $(978/984) = 99.4\%$ (95% CI: 98.7% - 99.7%)

Days Post Symptom Onset	Number of Subject Samples Tested	Investigational Positives	Comparator Positives	% Positive Rate (by Comparator)	PPA (95% CI)
0	15	0	0	0.0%	N/A
1	155	12	13	8.4%	92.3% (66.7% - 99.6%)
2	323	40	44	13.6%	90.9% (78.8% - 96.4%)
3	349	42	47	13.5%	89.4% (77.4% - 95.4%)
4	212	23	24	11.3%	95.8% (79.8% - 99.8%)
5	78	20	20	25.6%	100.0% (83.9% - 100.0%)
Total	1132	137*	148	13.1%	92.6% (87.2% - 95.8%)

* 6 subjects had a false positive result on the investigational test and were excluded.

Influenza A Performance

Influenza A	Comparator Positives	Comparator Negatives	Total
Investigational Positives	66	3	69
Investigational Negatives	14	1088	1102
Total	80	1091	1171

Positive Percent Agreement = $(66/80) = 82.5\%$ (95% CI: 72.7% - 89.3%)

Negative Percent Agreement = $(1088/1091) = 99.7\%$ (95% CI: 99.2% - 99.9%)

Influenza B Performance

Influenza B	Comparator Positives	Comparator Negatives	Total
Investigational Positives	27	0	27
Investigational Negatives	3	1141	1144
Total	30	1141	1171

Positive Percent Agreement = $(27/30) = 90.0\%$ (95% CI: 74.4% - 96.5%)

Negative Percent Agreement = $(1141/1141) = 100.0\%$ (95% CI: 99.7% - 100.0%)

15. Serial Testing

A prospective clinical study was conducted between January 2021 and May 2022 as a component of the Rapid Acceleration of Diagnostics (RADx) initiative from the National Institutes of Health (NIH). A total of 7,361 individuals were enrolled via a decentralized clinical study design, with a broad geographical representation of the United States. Per inclusion criteria, all individuals were asymptomatic upon enrollment in the study and at least 14 days prior to it and did not have a SARS-CoV-2 infection in the three months prior to enrollment. Participants were assigned to one of three EUA authorized SARS-CoV-2 OTC rapid antigen tests to conduct serial testing (every 48 hours) for 15 days. If an antigen test was positive, the serial-antigen testing result is considered positive.

At each rapid antigen testing time point, study subjects also collected a nasal swab for comparator testing using a home collection kit (using a 15-minute normalization window between swabs). SARS-CoV-2 infection status was determined by a composite comparator method on the day of the first antigen test, using at least two highly sensitive EUA RT-PCRs.

If results of the first two molecular tests were discordant, a third highly sensitive EUA RT-PCR test was performed, and the final test result was based upon the majority rule.

Study participants reported symptom status throughout the study using the MyDataHelps app. Two-day serial antigen testing is defined as performing two antigen tests 36 – 48 hours apart. Three-day serial antigen testing is defined as performing three antigen tests over five days with at least 48 hours between each test.

Out of the 7,361 participants enrolled in the study, 5,609 were eligible for analysis. Among eligible participants, 154 tested positive for SARS-CoV-2 infection based on RTPCR, of which 97 (62%) were asymptomatic on the first day of their infection, whereas 57 (39%) reported symptoms on the first day of infection.

Performance of the antigen test with serial testing in symptomatic individuals is described in the table below. Data establishing PPA of COVID-19 antigen serial testing compared to the molecular comparator single day testing throughout the course of infection with serial testing. Data is from all antigen tests in study combined.

DAYS AFTER FIRST PCR POSITIVE TEST RESULT	SYMPTOMATIC ON FIRST DAY OF TESTING		
	Ag Positive/PCR Positive (Antigen Test Performance % PPA)		
	1 Test	2 Tests	3 Tests
0	34/57 (59.6%)	47/51 (92.2%)	44/47 (93.6%)
2	58/62 (93.5%)	59/60 (98.3%)	43/43 (100%)
4	55/58 (94.8%)	53/54 (98.1%)	39/40 (97.5%)
6	27/34 (79.4%)	26/33 (78.8%)	22/27 (81.5%)
8	12/17 (70.6%)	12/17 (70.6%)	7/11 (63.6%)
10	4/9 (44.4%)	3/7 (42.9%)	
<p>1 Test= one (1) test performed on the noted days after first PCR positive test result. Day 0 is the first day of documented infection with SARS-CoV-2.</p> <p>2 Tests= two (2) tests performed an average of 48 hours apart. The first test performed on the indicated day and the second test performed 48 hours later.</p> <p>3 Tests= three (3) tests performed an average of 48 hours apart. The first test performed on the indicated day, the second test performed 48 hours later, and a final test performed 48 hours after the second test.</p>			

16. Technical Support

If you have questions regarding the use of this product, or if you want to report a problem with the Speedy Swab Rapid COVID-19 + Flu A&B Antigen Self-Test, please contact Watmind USA Technical Support at +1 866-928-6463

(Available Hours: Mon. to Fri. 8 am to 8 pm EST) or technicalsupport@watmindusa.com

17. Reorder











Speedy Swab Rapid COVID-19 + Flu A&B Antigen Self-Test – 1-pack (LFA0462-1N)

Speedy Swab Rapid COVID-19 + Flu A&B Antigen Self-Test – 2-pack (LFA0462-2N)

Speedy Swab Rapid COVID-19 + Flu A&B Antigen Self-Test – 4-pack (LFA0462-4N)

18. Symbols

The following symbols may be included on the labeling and packaging of the product:

	Manufacturer		Date of manufacture
	Contains sufficient for <n> tests		Catalogue number
	<i>In vitro</i> diagnostic medical device		Use-by date
	Consult instructions for use		Batch code
	Temperature limit		Do not reuse

LBL-020 Ver. 5, 06/04/24