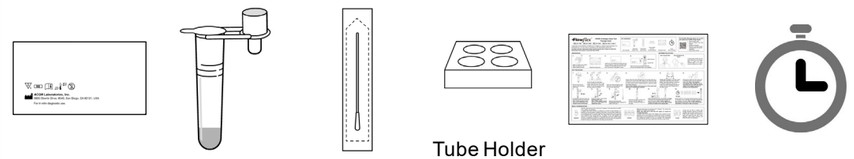
Flowflex™ COVID-19 Antigen Home Test Package Insert

REF L031-118B5	REF L031-125M5	REF L031-125N5	REF L031-125P5	English

A rapid test for the detection of SARS-CoV-2 antigens in anterior nasal specimens. For self-testing use. For use under an Emergency Use Authorization (EUA) only.

Carefully read the instructions before performing the test. Failure to follow the instructions may result in inaccurate test results.

KIT CONTENTS



Test Cassette | Extraction Buffer Tube | Disposable Nasal Swab | Tube Holder (only for 25 test quantity) | Package Insert | Timer (Not included)

Flowflex QR CODE

The Flowflex Web App allows you to track and report your COVID-19 test results

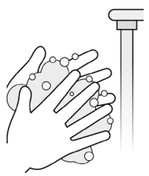
- The Web App is optional and not required to run a COVID-19 test. It will assist you in interpreting your visual test result and report your result to local health authorities.
- Ensure you have an internet connection and scan the Flowflex QR code or go to www.flowflexcovid.com prior to starting the test.
- Ensure you are using a compatible web browser (Chrome, Firefox, or Safari) and your electronic device has a camera.
- Click on "Report Your Test Result"
- Create an account

To perform a COVID-19 test

1. **Log in to the Flowflex Web App** - Ensure you are connected to the internet during your test.
2. **Answer a few questions on the Web App.**
3. **Follow step-by-step instructions for your test**
4. **Read result**

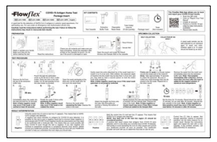
PREPARATION

1.



Wash or sanitize your hands. Make sure they are dry before starting the test.

2.

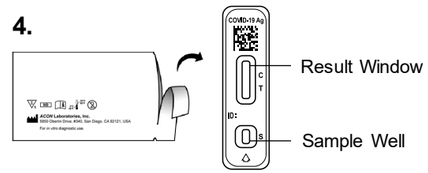


Read the instructions.

3.

Check your kit contents and make sure you have everything. Check the expiration date printed on the cassette foil pouch. Do not use if the pouch is damaged or open.

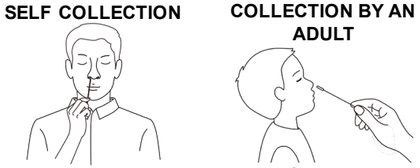
4.



Open the pouch and lay the cassette on a clean, flat surface. Locate the Result Window and Sample Well on the cassette.

SPECIMEN COLLECTION

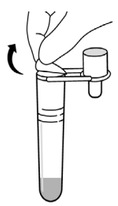
SELF COLLECTION | **COLLECTION BY AN ADULT**



A 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.

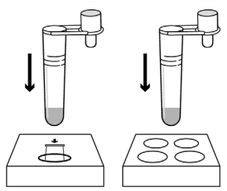
TEST PROCEDURE

1.



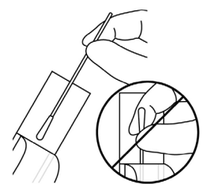
Remove the foil from the top of the extraction buffer tube.

2.



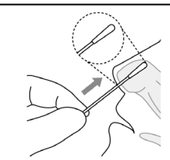
Punch through the perforated circle on the kit box to form a tube holder. Place the tube in the tube holder. For 25 test quantity kit box the tube holder is provided.

3.



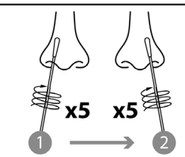
Open the swab packaging at the stick end, not the swab tip. Do not touch the swab tip.

4.



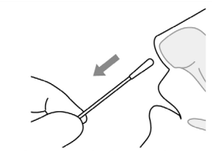
Gently insert the entire absorbent tip of the swab into 1 nostril (½ to ¾ of an inch). With children, the maximum depth of insertion into the nostril may be less than ¾ of an inch, and you may need to have a second person to hold the child's head while swabbing.
Note: A false negative result may occur if the nasal swab specimen is not properly collected.

5.



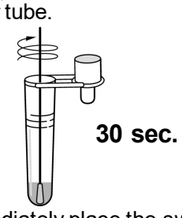
Firmly rub the **swab** in a circular motion around the inside wall of the nostril **5 times**. Take approximately 15 seconds to collect the specimen. Be sure to collect any nasal drainage that may be present onto the swab. Repeat this in the other nostril.

6.



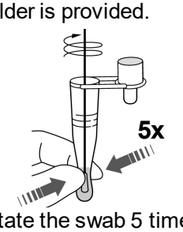
Remove the swab from the nostril and immediately place into the extraction buffer tube. ***Note: Test samples immediately after collection, and no more than one hour after the swab is added to the reagent solution, if stored at room temperature.***

7.



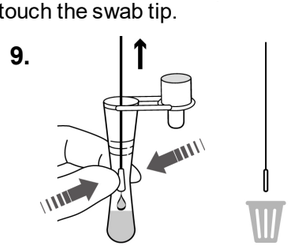
Immediately place the swab into the tube and swirl for 30 seconds. ***Note: A false negative result may occur if the swab is not swirled at least 30 seconds.***

8.



Rotate the swab 5 times while **squeezing the tube**.
Note: A false negative result may occur if the swab is not rotated five times.

9.

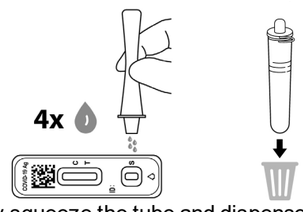


Remove the swab while **squeezing the tube**. Dispose the swab in the trash.

10.

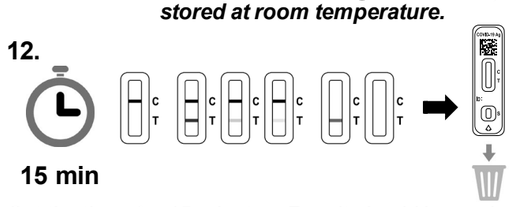
Attach the dropper tip firmly onto the tube. Mix thoroughly by swirling or flicking the bottom of the tube.

11.



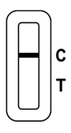
Gently squeeze the tube and dispense **4 drops** of solution into the Sample Well. Dispose the tube in the trash. ***Note: A false negative or invalid result may occur if less than 4 drops of fluid are added to the Sample Well.***

12.



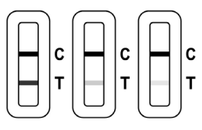
Set the timer for 15 minutes. Result should be read at 15 minutes. Do not read after 30 minutes. Dispose the test cassette in the trash. ***Note: A false negative or false positive result may occur if the test result is read before 15 minutes or after 30 minutes.***

RESULT INTERPRETATION



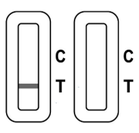
Negative

Only the control line (C) and no test line (T) appears. This means that no SARS-CoV-2 antigen was detected.
A negative test result indicates that antigens from the virus that causes COVID-19 were not detected from the specimen. A negative result does not rule out COVID-19. There is a higher chance of false negative results with antigen tests than with laboratory-based molecular tests. This means that there is a higher chance this test will give you a negative result when you have COVID-19. If you test negative and continue to experience COVID-19 like symptoms of fever, cough, and/or shortness of breath you should seek follow up care with your healthcare provider.



Positive

Both the control line (C) and test line (T) appear. This means that SARS-CoV-2 antigen was detected.
NOTE: Any faint red or pink line in the test line region (T) should be considered positive.
A positive test result means that the virus that causes COVID-19 was detected in your sample and it is very likely you have COVID-19 and are contagious. Please contact your doctor/primary care physician or your local health authority immediately and 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). Your healthcare provider will work with you to determine how best to care for you based on your test results along with medical history and your symptoms.



Invalid

Control line (C) fails to appear. If a control (C) line is not visible, the test is invalid. Re-test with a new swab and new test cassette. If the problem persists, call (800) 838-9502 for assistance.

FOR FDA EMERGENCY USE AUTHORIZATION (EUA) ONLY

- This product has not been FDA cleared or approved but has been authorized by FDA under an EUA.
- This product has been authorized only for the detection of proteins from SARS-CoV-2, 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 IVDs 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.
- For more information on EUAs please visit: https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization
- For the most up to date information on COVID-19, please visit: www.cdc.gov/COVID19
- For detailed instructions, please visit: www.flowflexcovid.com

INTENDED USE

The Flow*flex* COVID-19 Antigen Home Test is a lateral flow chromatographic immunoassay intended for the qualitative detection of the nucleocapsid protein antigen from SARS-CoV-2 in anterior nasal swab specimens directly from individuals within 7 days of symptom onset or without symptoms or other epidemiological reasons to suspect COVID-19. This test is authorized for non-prescription home use with self-collected anterior nasal swab specimens directly from individuals aged 14 years and older or with adult-collected anterior nasal samples directly from individuals aged 2 years or older. The Flow*flex* COVID-19 Antigen Home Test does not differentiate between SARS-CoV and SARS-CoV-2.

Results are for the identification of SARS-CoV-2 nucleocapsid protein antigen. This antigen is generally found in anterior 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 definite cause of disease.

Individuals who test positive should self-isolate and consult their healthcare provider as additional testing may be necessary and for public health reporting.

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 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 an individual's recent exposures, history and the presence of clinical signs and symptoms consistent with COVID-19.

Individuals should provide all results obtained with this product to their healthcare provider for public health reporting. Healthcare providers will report all test results they received from individuals who use the authorized product to relevant public health authorities in accordance with local, state, and federal requirements using appropriate LOINC and SNOMED codes, as defined by the Laboratory In Vitro Diagnostics (LIVD) Test Code Mapping for SARS-CoV-2 Tests provided by CDC.

The Flow*flex* COVID-19 Antigen Home Test is intended for self-use or lay user testing another in a non-laboratory setting. The Flow*flex* COVID-19 Antigen Home Test is only for use under the Food and Drug Administration's Emergency Use Authorization.

SUMMARY

The novel coronaviruses belong to the β genus. COVID-19 is an acute respiratory infectious disease. People are generally susceptible. Currently, the patients infected by the novel coronavirus are the main source of infection; asymptomatic infected people can also be an infectious source. Based on the current epidemiological investigation, the incubation period is 1 to 14 days, mostly 3 to 7 days. The main manifestations include fever, fatigue and dry cough. Nasal congestion, runny nose, sore throat, myalgia and diarrhea are found in a few cases.

WARNINGS, PRECAUTIONS, AND SAFETY INFORMATION

- Read the Flow*flex* COVID-19 Antigen Home Test Package Insert carefully before performing a test. Failure to follow directions may produce inaccurate test results.
- The Test is intended to aid in the diagnosis of active COVID-19. Please consult a healthcare professional to discuss your results and if any additional testing is required.
- Keep test kit and materials out of the reach of children and pets before and after use.
- Do not use on anyone under two years of age.
- Children aged 2 to 13 years of age should be tested by an adult.
- Wear a safety mask or other face-covering when collecting a specimen from a child or another individual.
- Leave the test cassette sealed in its pouch until just before use. Once opened, the test cassette should be used within 60 minutes.
- Do not use the test after the expiration date shown on the test cassette pouch.
- Do not use if any of the test kit contents or packaging is damaged or open.
- Test components are single use. Do not re-use. Do not use with multiple specimens.
- Make sure there is sufficient light when testing.
- Do not use nasal sprays for at least 30 minutes before collecting a nasal sample.
- Remove any piercings from the nose before starting the test.
- Do not use on anyone who is prone to nosebleeds or has had facial injuries or head injuries/surgery in the past six months.
- False negative test results may occur if a specimen is incorrectly collected or handled.
- Do not touch the swab tip when handling the swab.
- The test is intended to be read at 15 minutes. If the test is read before 15 minutes or after 30 minutes, false negative or false positive results may occur, and the test should be repeated with a new test cassette.
- Do not ingest any kit components.
- Avoid exposure of your skin, eyes, nose, or mouth to the solution in the extraction tube.
- The Reagent Solution contains a harmful chemical (see table below).

Hazard Category (mixture)	Hazard Statement for mixture	Labeling of Harm(s)
Not classified	Acute oral or dermal toxicity	None
Category 2	Eye irritation	May cause eye irritation
Category 3	Skin irritation	Causes mild skin irritation

- If the reagent solution contacts the skin or eye, flush with plenty of water. If irritation persists, seek medical advice. https://www.poisonhelp.org or 1-800-222-1222.

FREQUENTLY ASKED QUESTIONS

Q: WHAT IS COVID-19?

A: COVID-19 is caused by the SARS-CoV-2 virus which is a new virus in humans causing a contagious respiratory illness. COVID-19 can present with mild to severe illness, although some people infected with COVID-19 may have no symptoms at all. Older adults and people of any age who have underlying medical conditions have a higher risk of severe illness from COVID-19. Serious outcomes of COVID-19 include hospitalization and death. The SARS-CoV-2 virus can be spread to others even before a person shows signs or symptoms of being sick (e.g., fever, coughing, difficulty breathing, etc.). A full list of symptoms of COVID-19 can be found at the following link:
https://www.cdc.gov/coronavirus/2019-ncov/symptoms-testing/symptoms.html

Q: WILL THIS TEST HURT?

A: No, the nasal swab is not sharp, and it should not hurt. Sometimes the swab can feel slightly uncomfortable. If you feel pain, please stop the test and seek advice from your healthcare provider.

Q: WHAT ARE THE KNOWN POTENTIAL RISKS AND BENEFITS OF THIS TEST?

A: Potential **risks** include:

- Possible discomfort during sample collection.
- Possible incorrect test results (see Warnings and Result Interpretation section).

Potential **benefits** include:

- The results, along with other information, can help you and your healthcare provider make informed decisions about your care.
- The results of this test may help limit the spread of COVID-19 to your family and others in your community.

Q: WHAT IS THE DIFFERENCE BETWEEN AN ANTIGEN AND MOLECULAR TEST?

A:.There are different kinds of tests for the virus that causes COVID-19. Molecular tests detect genetic material from the virus. Antigen tests, such as the Flow*flex* COVID-19 Antigen Home Test, detect proteins from the virus. Antigen tests are very specific for the SARS-CoV-2 virus but are not as sensitive as molecular tests. This means that a positive result is highly accurate, but a negative result does not rule out infection. If your test result is negative, you should discuss with your healthcare provider whether an additional test is necessary and if you should continue isolating at home. There is a higher chance of false negative results with antigen tests than with laboratory-based molecular tests. This means that there is a higher chance this test will give you a negative result when you have COVID-19 than a molecular test would.

Q: HOW ACCURATE IS THIS TEST?

A: The performance of Flow*flex* COVID-19 Antigen Home Test was established in an all-comers clinical study conducted between March 2021 and May 2021 with 172 nasal swabs self-collected or pair-collected by another study participant from 108 individual symptomatic patients (within 7 days of onset) suspected of COVID-19 and 64 asymptomatic patients. All subjects were screened for the presence or absence of COVID-19 symptoms within two weeks of study enrollment. The Flow*flex* COVID-19 Antigen Home Test was compared to an FDA authorized molecular SARS-CoV-2 test. The Flow*flex* COVID-19 Antigen Home Test correctly identified 93% of positive specimens and 100% of negative specimens.

Q: WHAT IF YOU TEST POSITIVE?

A: A positive result means that it is very likely you have COVID-19 because proteins from the virus that causes COVID-19 were found in your sample. You should self-isolate from others and contact a healthcare provider for medical advice about your positive result. Your healthcare provider will work with you to determine how best to care for you based on your test result, medical history, and symptoms.

Q: WHAT IF YOU TEST NEGATIVE?

A: A negative test result indicates that antigens from the virus that causes COVID-19 were not found in your sample. If you have symptoms, you likely do not have COVID-19. However, negative results do not rule out SARS-CoV-2 infection.

It is possible for this test to give a negative result that is incorrect (false negative) in some people with COVID-19. This means that you could possibly still have COVID-19 even though the test is negative. For example, you may get a false negative result if you did not perform the test correctly or if the level of antigen from the virus causing COVID-19 was below the test limits. The amount of antigen in a sample may decrease the longer you have symptoms of infection. If you test negative and continue to experience symptoms of fever, cough and/or shortness of breath you should seek follow up care with your healthcare provider. Your healthcare provider will consider the test result together with all other aspects of your medical history (such as symptoms, possible exposures, and geographical location of places you have recently traveled) in deciding how to care for you. Your healthcare provider may suggest you need another test to determine if you have contracted the virus causing COVID-19. It is important that you work with your healthcare provider to help you understand the next steps you should take.

Q: WHAT DOES AN INVALID TEST RESULT MEAN?

A: An invalid result means the test was not able to tell if you have COVID-19 or not. If the test is invalid, a new swab should be used to collect a new nasal specimen and the test should be run again, using a new test cassette and extraction buffer tube.

IMPORTANT

This test is intended to be used as an aid in the clinical diagnosis of active COVID-19. Do not use this test as the only guide to manage your illness. Please consult your healthcare provider if your symptoms persist or become more severe, or if you are concerned at any time.

Individuals should report the test result through the Flow*flex* Web App or provide all results obtained with this product to their healthcare provider for public health reporting.

HEALTHCARE PROVIDERS

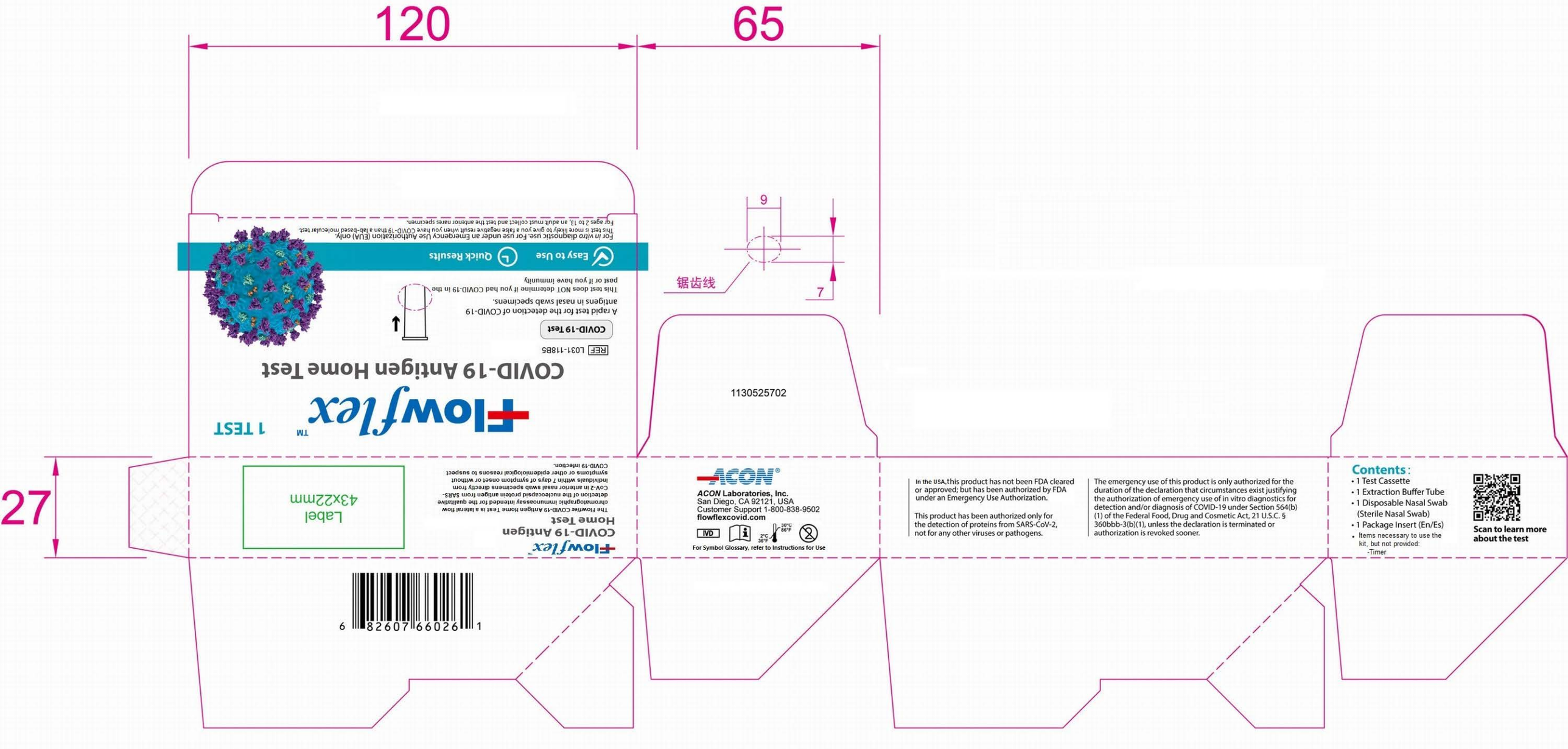
Please visit www.flowflexcovid.com to obtain the complete instructions for use and fact sheet for healthcare providers.

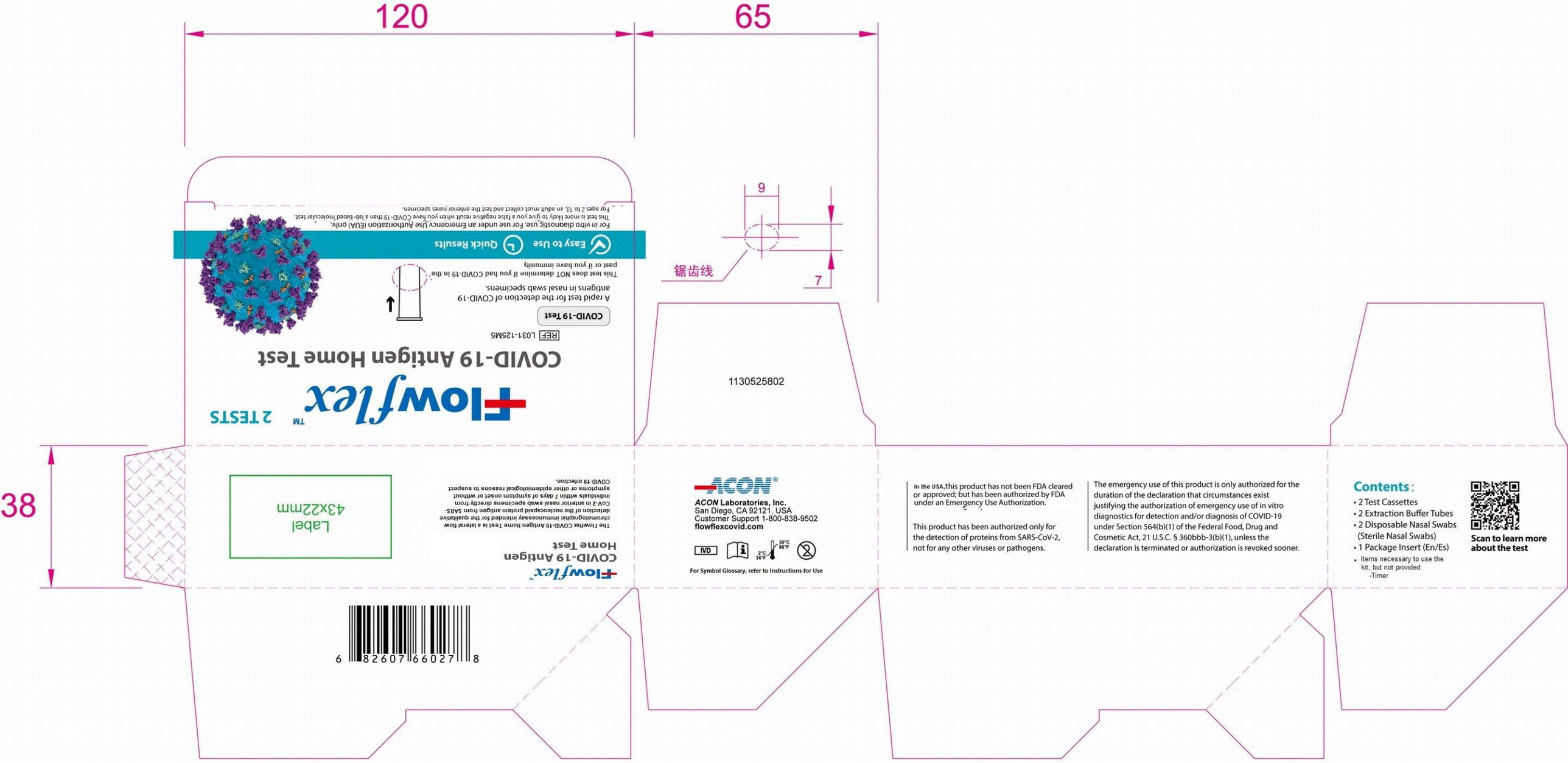
Index of Symbols

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

***ACON* Laboratories, Inc.**
San Diego, CA 92121, USA
flowflexcovid.com
Customer Support: 1-800-838-9502

Number: 1151297703
Effective Date: 2022-xx-xx

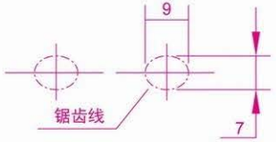

120
65
27
9
7
锯齿线
1130525702
Flowflex™
1 TEST
COVID-19 Antigen Home Test
REF L031-118B5
COVID-19 Test
A rapid test for the detection of COVID-19 antigens in nasal swab specimens.
This test does NOT determine if you had COVID-19 in the past or if you have immunity
Easy to Use
Quick Results
For in vitro diagnostic use. For use under an Emergency Use Authorization (EUA) only.
This test is more likely to give you a false negative result when you have COVID-19 than a lab-based molecular test.
For ages 2 to 13, an adult must collect and test the anterior nares specimen.
Flowflex™
COVID-19 Antigen Home Test
The Flowflex COVID-19 Antigen Home Test is a lateral flow chromatographic immunoassay intended for the qualitative detection of the nucleocapsid protein antigen from SARS-CoV-2 in anterior nasal swab specimens directly from individuals within 7 days of symptom onset or without symptoms or other epidemiological reasons to suspect COVID-19 infection.
Label
43x22mm
6 82607 66026 1
ACON®
ACON Laboratories, Inc.
San Diego, CA 92121, USA
Customer Support 1-800-838-9502
flowflexcovid.com
IVD
2°C 36°F
30°C 86°F
For Symbol Glossary, refer to Instructions for Use
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, 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.
Contents:
• 1 Test Cassette
• 1 Extraction Buffer Tube
• 1 Disposable Nasal Swab (Sterile Nasal Swab)
• 1 Package Insert (En/Es)
• Items necessary to use the kit, but not provided:
-Timer
Scan to learn more about the test


120
65
38
9
7
锯齿线
1130525802
Flowflex™ 2 TESTS
COVID-19 Antigen Home Test
REF L031-125M5
COVID-19 Test
A rapid test for the detection of COVID-19 antigens in nasal swab specimens.
This test does NOT determine if you had COVID-19 in the past or if you have immunity
Easy to Use
Quick Results
For in vitro diagnostic use. For use under an Emergency Use Authorization (EUA) only.
This test is more likely to give you a false negative result when you have COVID-19 than a lab-based molecular test.
For ages 2 to 13, an adult must collect and test the anterior nares specimen.
Flowflex™
COVID-19 Antigen Home Test
The Flowflex COVID-19 Antigen Home Test is a lateral flow chromatographic immunoassay intended for the qualitative detection of the nucleocapsid protein antigen from SARS-CoV-2 in anterior nasal swab specimens directly from individuals within 7 days of symptom onset or without symptoms or other epidemiological reasons to suspect COVID-19 infection.
Label
43x22mm
6 82607 66027 8
ACON®
ACON Laboratories, Inc.
San Diego, CA 92121, USA
Customer Support 1-800-838-9502
flowflexcovid.com
IVD
2°C 36°F
30°C 86°F
For Symbol Glossary, refer to Instructions for Use
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, 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.
Contents:
• 2 Test Cassettes
• 2 Extraction Buffer Tubes
• 2 Disposable Nasal Swabs (Sterile Nasal Swabs)
• 1 Package Insert (En/Es)
• Items necessary to use the kit, but not provided:
-Timer
Scan to learn more about the test

165

80

45



1130525902

Flowflex™ 5 TESTS

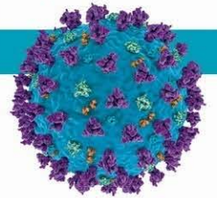
COVID-19 Antigen Home Test

REF L031-125N5

COVID-19 Test

A rapid test for the detection of COVID-19 antigens in nasal swab specimens.

This test does NOT determine if you had COVID-19 in the past or if you have immunity



Easy to Use | Quick Results

For *in vitro* diagnostic use. For use under an Emergency Use Authorization (EUA) only.
This test is more likely to give you a false negative result when you have COVID-19 than a lab-based molecular test.
For ages 2 to 13, an adult must collect and test the anterior nares specimen.

Label
43x22mm

Flowflex™
COVID-19 Antigen
Home Test

The Flowflex COVID-19 Antigen Home Test is a lateral flow chromatographic immunoassay intended for the qualitative detection of the nucleocapsid protein antigen from SARS-CoV-2 in anterior nasal swab specimens directly from individuals within 7 days of symptom onset or without symptoms or other epidemiological reasons to suspect COVID-19 infection.

6 82607 66028 5

ACON®

***ACON* Laboratories, Inc.**
San Diego, CA 92121, USA
Customer Support 1-800-838-9502
flowflexcovid.com

IVD 2°C 36°F 30°C 86°F

For Symbol Glossary, refer to Instructions for Use

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, 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.

Contents:

- 5 Test Cassettes
- 5 Extraction Buffer Tubes
- 5 Disposable Nasal Swabs (Sterile Nasal Swabs)
- 1 Package Insert (En/Es)
- Items necessary to use the kit, but not provided:
  - Timer

Scan to learn more about the test

125

210

93

25 TESTS

Flowflex™

COVID-19 Antigen Home Test

REF L031-125P5

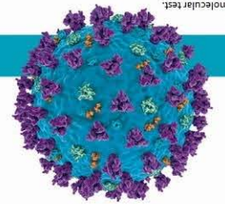
COVID-19 Test

A rapid test for the detection of COVID-19 antigens in nasal swab specimens.

This test does NOT determine if you had COVID-19 in the past or if you have immunity

Easy to Use | Quick Results

For in vitro diagnostic use. For use under an Emergency Use Authorization (EUA) only.
This test is more likely to give you a false negative result when you have COVID-19 than a lab-based molecular test.
For ages 2 to 13, an adult must collect and test the anterior nares specimen.



Flowflex™

COVID-19 Antigen Home Test

Label
43x22mm

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, 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.

1130553302

The Flow*flex* COVID-19 Antigen Home Test is a lateral flow chromatographic immunoassay intended for the qualitative detection of the nucleocapsid protein antigen from SARS-CoV-2 in anterior nasal swab specimens directly from individuals within 7 days of symptom onset or without symptoms or other epidemiological reasons to suspect COVID-19 infection.

ACON®

***ACON* Laboratories, Inc.**
San Diego, CA 92121, USA
Customer Support 1-800-838-9502
flowflexcovid.com

IVD

For Symbol Glossary, refer to Instructions for Use

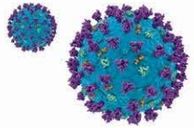
Flowflex™

COVID-19 Antigen Home Test

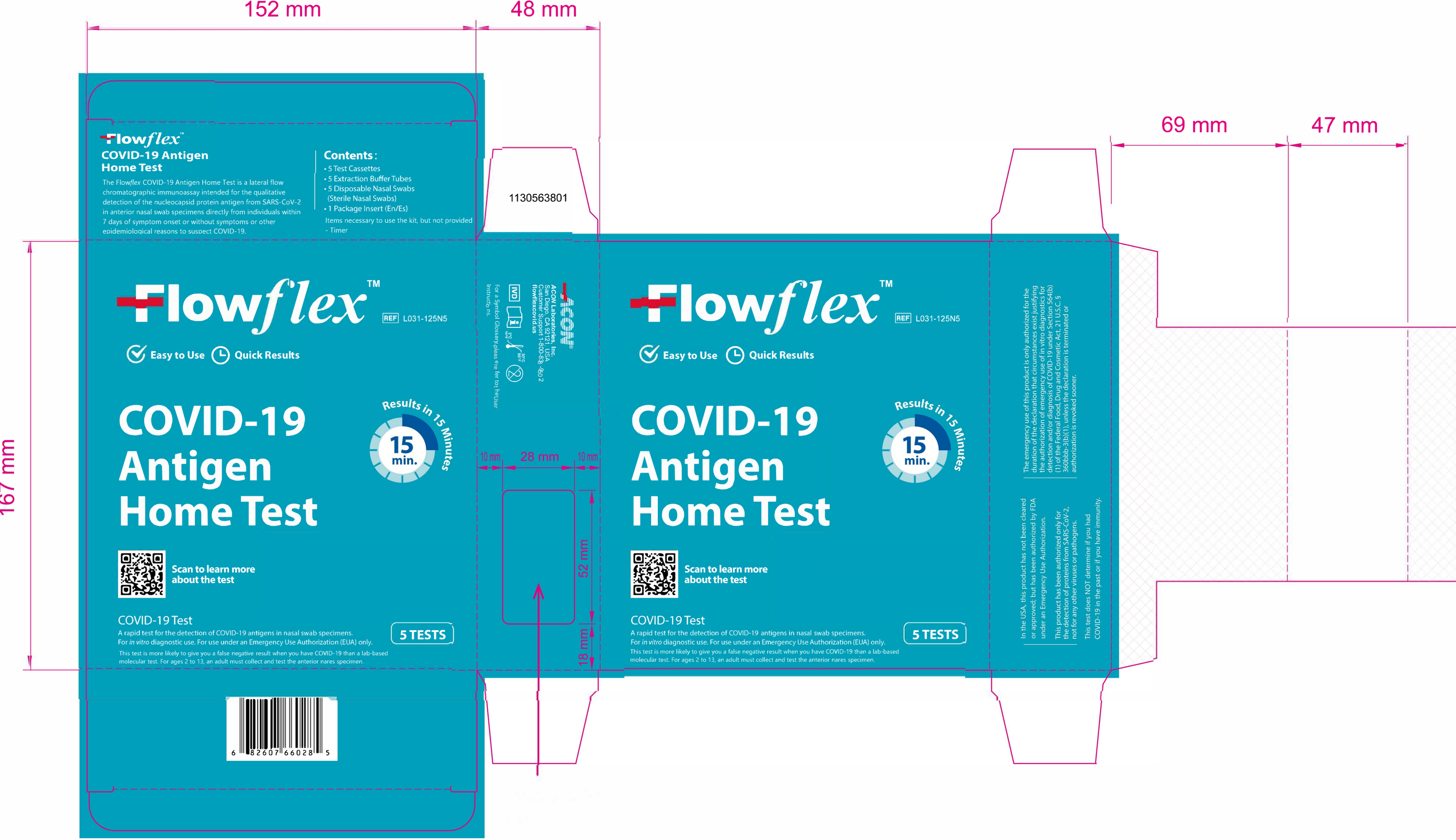
Scan to learn more about the test

Contents:

- 25 Test Cassettes
- 25 Extraction Buffer Tubes
- 25 Disposable Nasal Swabs (Sterile Nasal Swabs)
- 1 Tube Holder
- 1 Package Insert (En/Es)
- Items necessary to use the kit, but not provided:
  -Timer



6 82607 66047 6


152 mm
48 mm
69 mm
47 mm
167 mm
1130563801
10 mm
28 mm
10 mm
52 mm
18 mm
Flowflex™
COVID-19 Antigen Home Test
The Flowflex COVID-19 Antigen Home Test is a lateral flow chromatographic immunoassay intended for the qualitative detection of the nucleocapsid protein antigen from SARS-CoV-2 in anterior nasal swab specimens directly from individuals within 7 days of symptom onset or without symptoms or other epidemiological reasons to suspect COVID-19.
Contents:
• 5 Test Cassettes
• 5 Extraction Buffer Tubes
• 5 Disposable Nasal Swabs (Sterile Nasal Swabs)
• 1 Package Insert (En/Es)
Items necessary to use the kit, but not provided
- Timer
Flowflex™
REF L031-125N5
Easy to Use
Quick Results
COVID-19 Antigen Home Test
Results in 15 Minutes
15 min.
Scan to learn more about the test
COVID-19 Test
A rapid test for the detection of COVID-19 antigens in nasal swab specimens.
For in vitro diagnostic use. For use under an Emergency Use Authorization (EUA) only.
This test is more likely to give you a false negative result when you have COVID-19 than a lab-based molecular test. For ages 2 to 13, an adult must collect and test the anterior nares specimen.
5 TESTS
6 82607 66028 5
ACON Laboratories, Inc.
San Diego, CA 92121, USA
Customer Support 1-800-838-8502
flowflexcovid.us
IVD
For a Symbol Glossary please refer to the User Instructions.
Flowflex™
REF L031-125N5
Easy to Use
Quick Results
COVID-19 Antigen Home Test
Results in 15 Minutes
15 min.
Scan to learn more about the test
COVID-19 Test
A rapid test for the detection of COVID-19 antigens in nasal swab specimens.
For in vitro diagnostic use. For use under an Emergency Use Authorization (EUA) only.
This test is more likely to give you a false negative result when you have COVID-19 than a lab-based molecular test. For ages 2 to 13, an adult must collect and test the anterior nares specimen.
5 TESTS
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.
In the USA, this product has not been 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, 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.