Important safety information

🔔 Warning!

Follow all health and safety regulations.

- Use appropriate personal protective equipment.
- Handle all samples as if they contain infectious agents.
- Observe all precautions and warnings in the Instructions for Use.

Document information

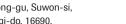
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For in vitro diagnostics use only Not for self testing

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SARS-CoV-2 Rapid Antibody Test

Quick Reference Guide

This guide is a reference for using the SARS-CoV-2 Rapid Antibody Test.

Read the Instructions for Use before using this test.

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1 Preparing for a test

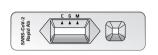
1. Carefully read the Instructions for Use for the SARS-CoV-2 Rapid Antibody Test.



2. Check the expiry date \square on the back of the foil pouch. Do not use the test, if the expiry date has passed.



- 3. Open the foil pouch and remove the test device and the desiccant package.
- 4. Ensure that the test device is undamaged and that the desiccant status indicator shows valid (yellow).

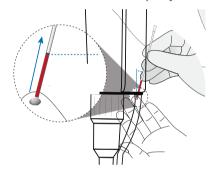




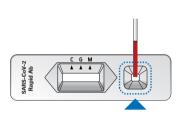
5. Perform a QC as required according to the Instructions for Use of the QC material and according to your local guidelines.

2 Performing a test with capillary whole blood

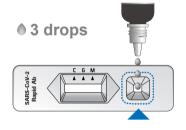
1. Using a 20 μL capillary tube, collect capillary whole blood to the black line on the capillary tube.



2. Apply the collected capillary whole blood to the specimen well of the test device.



3. Add 3 drops (90 μ L) of buffer vertically into the specimen well of the test device. Do not touch the surface of the specimen well with the tip of the



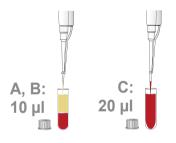
4. Read the test result at 10 to 15 min. Continue with 4 Interpreting results.

WARNING! Risk of incorrect results. Do not read the test result after 15 min.

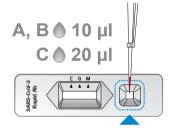


3 Performing a test with serum, plasma, or venous whole blood

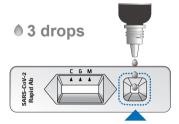
1. Using a micropipette, collect 10 µl of serum or plasma (A, B), or 20 µl of venous whole blood (C).



2. Apply the collected 10 µl of serum or plasma (A, B) or 20 µl of venous whole blood (C) to the specimen well of the test device.



3. Add 3 drops (90 μ L) of buffer vertically into the specimen well of the test device. Do not touch the surface of the specimen well with the tip of the



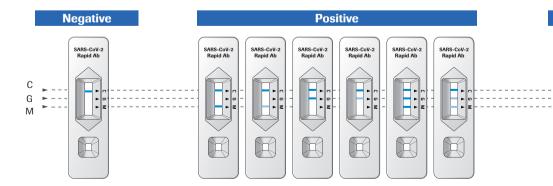
4. Read the test result at 10 to 15 min. Continue with 4 Interpreting results.

WARNING! Risk of incorrect results. Do not read the test result after 15 min.



4 Interpreting results

- 1. A colored band appears in the top section of the result window to show that the test is working properly. This is the control line (C). Even if the control line is faint, the test should be considered to have been performed properly. If no control line is visible the test is invalid.
- 2. Colored bands appear in the lower section of the result window. These are the test lines of IgM/ IgG (M, G). Even if the test lines are very faint or not uniform, the test result should be interpreted as a positive result.

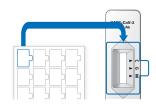


Perform a QC and repeat test with a new test device.

Ensuring test performance

Observe the following techniques to ensure best test performance:

1. If the test site is subject to strong ventilation, attach the provided film to the result window of the test device before applying the sample. This ensures sufficient migration of the sample.



2. Add the buffer immediately after applying the sample to the specimen well.

Note:

- SARS-CoV-2 Rapid Antibody Test can cross-react with antibody against SARS-CoV.
- For diagnostic purposes, the results should always be assessed in conjunction with the patient's medical history, clinical examination and other findings
- A negative test result does not completely rule out the possibility of an infection with SARS-CoV-2. This test cannot be used to diagnose an acute infection.