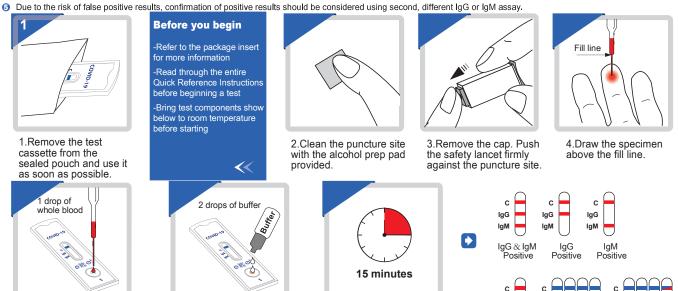
Quick Reference Instructions of Assure COVID-19 IgG/IgM Rapid Test Device

- 1 This test has not been FDA cleared or approved. This test has been authorized by FDA under an EUA for use by authorized laboratories. It is a point of care test for fingerstick whole blood specimens only. The user should be trained in the procedure. Wear appropriate protective attire for your safety when handling patient samples.
- This test has been authorized only for the presence of IgM and IgG antibodies against SARS-CoV-2, not for any other viruses or pathogens. This test is only authorized for ค 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 authorization is terminated or revoked sooner. 3 Read the complete Quick Reference Instructions before performing the test. For technical assistances, please call (800) 618-5829.
- O There should be a blue line in the control region (next to "C") before testing, discard the device if there is no blue line.



5.Transfer 1 drop of specimen into the specimen well. Adding more or less than 1 drop of specimen may lead to erroneous results.

6. Add 2 drops of buffer and start the timer. Adding 1 drop of buffer or more than 4 drops of buffer may lead to erroneous results. Wait for the blue line change to red line

15 at

Read result 7 adding minutes after buffer, do not interpret the result after 30 minutes.

lgG lgG lgG laM laM laN

Negative

Invalid

Invalid

Quick Reference Instructions of COVID-19 IgG/IgM Control

