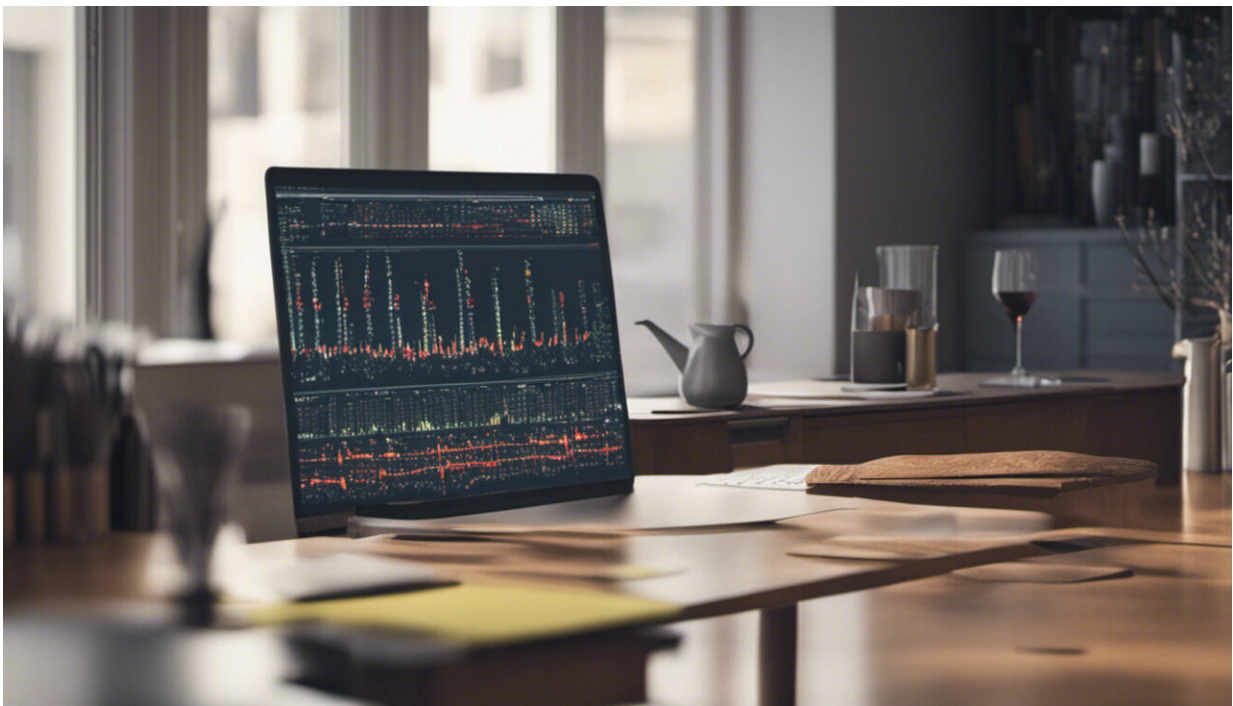


The new 15-minute COVID test has potential, but standard tests are still the best way to track COVID-19

September 25 2020, by Katherine Gibney, Deborah Williamson and Jodie McVernon



Credit: AI-generated image ([disclaimer](#))

Testing remains a vital component of Australia's success in managing COVID-19.

We need to diagnose people infected with SARS-CoV-2, the virus that causes COVID-19, as early as possible so they can be isolated from others and their contacts quarantined. Testing also helps us understand to what degree the virus is present in the population, so we can tailor public health measures accordingly.

If you've had a COVID-19 test, in all likelihood you received a PCR test. That's the one with the throat and nose swabs, and is regarded as the "gold standard" in COVID-19 testing.

But now the Therapeutic Goods Administration (TGA) has approved [a new kind of COVID-19 test](#), which can produce results in as little as 15 minutes, as opposed to a day or more for standard tests.

So is this new rapid test set to revolutionize COVID-19 testing in Australia? Not quite yet.

The traditional tests

Nucleic acid tests, or PCR tests, can detect ribonucleic acid (RNA) of SARS-CoV-2 from a day or two before symptoms start, and for a week or more afterwards, as symptoms resolve. Of course, some people will test positive without ever having symptoms.

PCR tests have been the backbone of SARS-CoV-2 testing worldwide. Because of the vast global experience with PCR tests and their high performance, they're considered [the most reliable](#) COVID-19 test.

PCR tests require specialized laboratory equipment and trained scientists and technicians to test the specimens; processing and testing take several hours.

Since January, we've performed an astonishing [7.4 million](#) SARS-CoV-2

PCR tests in Australia, which has needed a massive upscaling of capacity in laboratories nationally.

At times, demand for PCR testing has exceeded capacity, occasionally resulting in delays of up to several days in getting results back to patients. Meanwhile, laboratories swamped with COVID-19 tests may be limited in their capacity to perform their routine business, including diagnostic testing for other infectious diseases.

As people are required to isolate until they [receive a negative test result](#) and their symptoms resolve, these delays may come at a cost to the person waiting, their family, and the economy.

Recognizing these costs may lead some people to choose not to be tested, Victoria has offered [financial compensation](#) for people without leave entitlements awaiting test results.

But delayed case confirmation also increases the time to identification and quarantine of contacts, undermining public health efforts.

What can we expect from the antigen test?

Rapid antigen tests can diagnose COVID-19 [in 15 minutes](#). They're relatively inexpensive and require a swab from the nose.

These tests detect viral antigens, proteins on the surface of SARS-CoV-2. The [immune system](#) recognizes these proteins as foreign, and responds by making antibodies to SARS-CoV-2 ("anti-gen" means antibody generator).

Antigen tests perform best early in the infection when the amount of virus in a person's system is highest. For a person with symptomatic COVID-19, this would be in the first week of symptoms. So they only

pick up current infections – [unlike antibody tests](#), which can detect if a person was previously infected with SARS-CoV-2.

Four SARS-CoV-2 rapid antigen tests have been [licensed for use](#) in Australia in the past two months.

Unfortunately, rapid antigen tests for COVID-19 appear to be less sensitive than PCR tests, meaning they may give a negative result in someone who does actually have COVID-19. One of the recently licensed rapid antigen tests may give a false negative result in [up to 18.3%](#) of people with COVID-19 diagnosed by PCR.

While a positive rapid antigen test result is more reliable, widespread use of these tests in asymptomatic people will result in some false positive results—that is, a positive test result in someone who doesn't have COVID-19.

At this stage, [national COVID-19 guidelines](#) don't include information on antigen tests. So a person with a positive antigen test would need to undergo a PCR test to be counted in Australia's official COVID-19 case numbers.

Considering the pros and cons

We're faced with a trade-off between the potential benefits of the rapid antigen tests—the ability to test larger numbers of people, consuming fewer laboratory resources, and quicker results—and the potential to miss a few cases because of the lower test sensitivity.

Despite the lower sensitivity, increasing testing rates might result in an overall net increase in the proportion of COVID-19 cases diagnosed, and therefore a [public health benefit](#) by preventing onward transmission from these cases.

One possible strategic use of these tests may be in screening people without symptoms to detect asymptomatic and pre-symptomatic infection that might otherwise go undetected. This [could include](#) people in workplaces where ongoing exposure to colleagues and the public is unavoidable, including sectors of the food supply chain or other essential services.

Because of the lower test sensitivity for the rapid [antigen test](#), a PCR [test](#) remains most appropriate for people with symptoms, those at greater risk of poor outcomes from COVID-19, and people working in high-risk settings like aged care and health care.

While rapid antigen tests show promise, we'll need to evaluate their efficacy in Australia before we can determine their role in our fight against COVID-19.

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