
Provisional Authorisation for COVID-19 Tests

<i>Applicant</i>	AMT Pte Ltd
<i>Name of test</i>	AMT Resolute 2.0 assay
<i>Intended purpose (As per manufacturer's information for use)</i>	<p><i>RESOLUTE 2.0 is a real-time RT-PCR test intended for the qualitative detection of nucleic acid from the SARS-CoV-2 virus in nasopharyngeal swabs specimen from individuals with signs and symptoms of infection who are suspected of COVID-19.</i></p> <p><i>RESOLUTE 2.0 is intended for the direct amplification of Coronavirus SARS-CoV-2 RNA from nasopharyngeal swabs (NPS) in Universal Transport Medium (UTM). RNA extraction is not required for the use of this kit.</i></p> <p><i>Results are for the detection of SARS-CoV-2 RNA. The SARS-CoV-2 RNA is generally detectable in respiratory specimens, including nasopharyngeal swabs during the acute phase of infection. A positive result is indicative of the presence of SARS-CoV-2 RNA. Correlation with clinical presentation and other clinical investigations is necessary to determine the patient's infection status. Positive results do not rule out bacterial infection or co-infection with other viruses.</i></p> <p><i>A negative result does not exclude SARS-CoV-2 infection. Clinical correlation is required and if indicated, repeat sampling and use of other clinical and laboratory investigations may be warranted.</i></p> <p><i>The test results from the RESOLUTE 2.0 SARS-CoV-2 detection assay should not be used as the sole basis for patient management decisions.</i></p>
<i>Date of Provisional Authorisation</i>	7 July 2020
