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## Provisional Authorisation for COVID-19 Tests

<i>Applicant</i>	Siemens Healthcare Pte Ltd
<i>Name of test</i>	Siemens ADVIA Centaur® SARS-CoV-2 IgG (sCOVG)
<i>Intended purpose (As per manufacturer's information for use)</i>	<p><i>The ADVIA Centaur® SARS-CoV-2 IgG (sCOVG) assay is for in vitro diagnostic use in the qualitative and quantitative detection of IgG antibodies, including neutralizing antibodies, to SARS-CoV-2 in human serum and plasma (lithium heparin) obtained by venipuncture or capillary puncture using the ADVIA Centaur® CP, ADVIA Centaur® XP and ADVIA Centaur® XPT systems.</i></p> <p><i>This assay is intended as an aid in the diagnosis of patients with suspected SARS-CoV-2 infection and as an aid in identifying patients with an adaptive immune response to SARS-CoV-2, indicating recent or prior infection.</i></p>
<i>Date of Provisional Authorisation</i>	5 February 2021

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