



15 December 2021



TGA Registration Update

AnteoTech Ltd (ASX: ADO) ("AnteoTech" or "the Company") advises that it has received feedback from the Therapeutic Goods Administration ("TGA") on its EuGeni Reader and SARS CoV- 2 Ag Rapid Diagnostic Test (RDT)¹ submission.

The TGA have completed a preliminary review and have requested that AnteoTech supply additional information and clarification in support of its submission.

The information requested relates to the performance of AnteoTech's RDT in relation to the detection of SARS-CoV-2 variants. In addition, the TGA has requested that AnteoTech provide plans for the monitoring of the emergence of new strains of SARS-CoV-2 and performance testing of the RDT against these strains, as well as other technical and performance elements.

AnteoTech is working with the TGA to provide the information as quickly as possible.

The Company will provide further updates as our submission progresses.

This announcement has been authorised for release by the Board.

For more information, please contact:

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About AnteoTech - (ASX:ADO)

AnteoTech is a surface chemistry company with Intellectual Property ("IP") in its core technology product groups AnteoCoat™, AnteoBind™ and AnteoRelease™. The Company's purpose is to create shareholder value by identifying and solving important global industry problems by providing unique value-add solutions for its customers. Customers operate in the life sciences, diagnostics, energy and medical devices markets.

AnteoTech - Social Media Policy

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¹ The AnteoTech Antigen Rapid Diagnostic Test detects the SARS-CoV-2 active virus that causes the disease called COVID-19