# Aytu BioScience Announces Results from a National Cancer Institute Evaluation of the Company's Licensed COVID-19 IgG/IgM Rapid Test

***National Cancer Institute's (NCI) analysis of COVID-19 IgG/IgM Rapid Test Cassette Reported 100% Combined Sensitivity, and 97.5% Combined Specificity for the Lateral Flow Immunoassay***

**ENGLEWOOD, CO / June 9, 2020 /** Aytu BioScience, Inc. (NASDAQ:AYTU), a specialty pharmaceutical company (the "Company") focused on commercializing novel products that address significant patient needs announced today that the U.S. Food and Drug Administration (the "FDA") has published data from the National Cancer Institute's (NCI) Frederick National Laboratory for Cancer Research, and its evaluation of the COVID-19 IgG/IgM Rapid Test distributed by the Company.

A well-characterized panel of 30 confirmed SARS-CoV-2 antibody positive, and 80 SARS-CoV-2 antibody-negative samples collected prior to 2020 were tested in an independent validation study performed by the NCI. 96.7% and 100% sensitivity were estimated for IgG and IgM, respectively. 97.5% and 100% specificity were estimated for IgG and IgM, respectively. Furthermore, combined sensitivity and specificity were 100% and 97.5%, respectively.