

## **Aytu BioScience to Reschedule Fourth Quarter and FY 2020 Conference Call**

**ENGLEWOOD, CO / September 23, 2020** / Aytu BioScience, Inc. (NASDAQ:AYTU) (the “Company”), a specialty pharmaceutical company focused on commercializing novel products that address significant patient needs, announced today that it will reschedule its fourth quarter and FY 2020 conference call, previously scheduled for Thursday, September 24 at 4:30pm ET.

The adjustment was made to allow for additional time to finalize financial statements due to the additional workload created by the recently completed acquisition of Innovus Pharmaceuticals (the “Innovus Merger”) and the Cerecor, Inc. pediatric prescription portfolio (the “Pediatric Portfolio”) and to accommodate scheduling needs related to COVID-19. The company will make a further announcement in a subsequent press release to schedule the new date and time of the earnings conference call.

Aytu’s Chairman and Chief Executive Officer Josh Disbrow will host the call during which the Company expects to report record revenue for its fourth quarter and full fiscal year and progress on the Company’s business activities.

### **About Aytu BioScience, Inc.**

Aytu BioScience is a commercial-stage specialty pharmaceutical company focused on commercializing novel products that address significant patient needs. The company currently markets a portfolio of prescription products addressing large primary care and pediatric markets. The primary care portfolio includes (i) Natesto®, the only FDA-approved nasal formulation of testosterone for men with hypogonadism (low testosterone, or “Low T”), (ii) ZolpiMist®, the only FDA-approved oral spray prescription sleep aid, and (iii) Tuzistra® XR, the only FDA-approved 12-hour codeine-based antitussive syrup. The pediatric portfolio includes (i) AcipHex® Sprinkle™, a granule formulation of rabeprazole sodium, a commonly prescribed proton pump inhibitor; (ii) Cefaclor, a second-generation cephalosporin antibiotic suspension; (iii) Karbinal® ER, an extended-release carbinoxamine (antihistamine) suspension indicated to treat numerous allergic conditions; and (iv) Poly-Vi-Flor® and Tri-Vi-Flor®, two complementary prescription fluoride-based supplement product lines containing combinations of fluoride and vitamins in various for infants and children with fluoride deficiency. Aytu also distributes a COVID-19 IgG/IgM rapid antibody test and rapid antigen tests. These assays are used in the rapid, qualitative diagnostic assessment of the 2019 Novel Coronavirus. Additionally, Aytu recently licensed worldwide rights to develop the Healight™ technology platform from Cedars-Sinai Medical Center. Healight is a pre-clinical medical device being studied as a prospective treatment for COVID-19 and other respiratory infections.

Aytu also operates a consumer health subsidiary, Innovus Pharmaceuticals, Inc. (“Innovus”),

a specialty pharmaceutical company commercializing, licensing and developing safe and effective consumer healthcare products designed to improve men's and women's health and vitality. Innovus commercializes over thirty-five consumer health products competing in large healthcare categories including diabetes, men's health, sexual wellness and respiratory health. The Innovus product portfolio is commercialized through direct-to-consumer marketing channels utilizing the company's proprietary Beyond Human® marketing and sales platform.

Aytu's strategy is to continue building its portfolio of revenue-generating Rx and consumer health products, leveraging its focused commercial team and expertise to build leading brands within large therapeutic markets. For more information visit [aytubio.com](http://aytubio.com) and visit [innovuspharma.com](http://innovuspharma.com) to learn about the company's consumer healthcare products.

### **Forward-Looking Statement**

This press release includes forward-looking statements within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, or the Exchange Act. All statements other than statements of historical facts contained in this presentation, are forward-looking statements. Forward-looking statements are generally written in the future tense and/or are preceded by words such as 'may,' 'will,' 'should,' 'forecast,' 'could,' 'expect,' 'suggest,' 'believe,' 'estimate,' 'continue,' 'anticipate,' 'intend,' 'plan,' or similar words, or the negatives of such terms or other variations on such terms or comparable terminology. These statements are just predictions and are subject to risks and uncertainties that could cause the actual events or results to differ materially. These risks and uncertainties include, among others: the regulatory and commercial risks associated with introducing the COVID-19 rapid tests, the accuracy of the COVID-19 rapid tests as compared to other COVID-19 tests, market acceptance of the tests, the ability to obtain FDA approval or authorization for the tests, our ability to obtain sufficient tests to meet consumer demand, if any, the manufacturers' ability to scale up manufacturing to meet customer demand, if any, reputation risks if the tests are not as effective as anticipated, and that the current regulatory environment continues to permit the sale of the tests. We also refer you to the risks described in 'Risk Factors' in Part I, Item 1A of the company's Annual Report on Form 10-K and in the other reports and documents we file with the Securities and Exchange Commission from time to time.

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