

# Adial Pharmaceuticals Announces Closing of \$5 Million Private Placement Priced at a Premium to Market

Financing fully funds ONWARD™ Phase 3 trial until data read-out

CHARLOTTESVILLE, Va., Aug. 05, 2021 — **Adial Pharmaceuticals, Inc. (NASDAQ: ADIL; ADILW)** (“Adial” or the “Company”), a clinical-stage biopharmaceutical company focused on developing therapies for the treatment and prevention of addiction and related disorders, today announced that it has completed the \$5,000,002 private placement of 1,666,668 shares of common stock at a price of \$3.00 per share (the “Shares”) on August 4, 2021. The private placement was led by Bespoke Growth Partners, Inc., which invested \$2,500,000, and is a company controlled by Mark Peikin, Adial’s Chief Strategy Officer, and also included Richard Gilliam, founder of Cumberland Resources, and Keystone Capital Partners LLC, all of whom were previous investors in the Company. No warrants were issued in this financing, and Brookline Capital Markets, a division of Arcadia Securities, LLC, acted as an advisor on the transaction.

As previously disclosed, the Company received \$500,002 upon the parties’ execution of their respective Securities Purchase Agreements and has now received the balance of \$4,500,000 following the U.S. Securities and Exchange Commission declaring the registration statement on Form S-3, registering the resale of the private placement shares, effective on July 29, 2021.

In combination with the private placement previously announced on June 3, 2021, this transaction marks the completion of a set of financings totaling \$7.1 million.

William Stilley, Adial’s Chief Executive Officer, commented, “We are pleased to close this last piece of our \$7.1 million in private placements, and we appreciate the support of the participating investors. We believe this latest financing illustrates the tremendous confidence and support of our existing shareholders, as well as our conviction in the positive outlook for the business. This funding is anticipated to provide us more than sufficient capital to complete the ONWARD™ Phase 3 trial of AD04 as a genetically targeted treatment for Alcohol Use Disorder through data read-out. Additionally, it is expected to allow us to achieve key milestones for our pre-clinical adenosine program for non-opiate pain relief.”

## About Adial Pharmaceuticals, Inc.

Adial Pharmaceuticals is a clinical-stage biopharmaceutical company focused on the development of treatments for addictions. The Company’s lead investigational new drug product, AD04, is a genetically targeted, serotonin-3 receptor antagonist, therapeutic agent for the treatment of Alcohol Use Disorder (AUD) and is currently being investigated in the Company’s landmark ONWARD™ pivotal Phase 3 clinical trial for the potential treatment of AUD in subjects with certain target genotypes, which are to be identified using the Company’s proprietary companion diagnostic genetic test. A Phase 2b clinical trial of AD04 for the treatment of AUD showed promising results in reducing frequency of drinking, quantity of drinking and heavy drinking (all with statistical significance), and no overt safety concerns (there were no statistically significant serious adverse events reported). AD04 is also believed to have the potential to treat other addictive disorders such as Opioid Use Disorder, gambling, and obesity. The Company is also developing adenosine analogs for the treatment of pain and other disorders. Additional information is available at [www.adialpharma.com](http://www.adialpharma.com).

## About the Landmark ONWARD™ Pivotal Phase 3 Clinical Trial

The ONWARD trial is a 24-week, multicenter, randomized, double-blind, placebo-controlled, parallel group, Phase 3 clinical study to evaluate the efficacy, safety and tolerability of AD04 in patients with Alcohol Use Disorder (AUD) and selected polymorphisms in the serotonin transporter and receptor genes. Patients are genetically screened prior to enrollment in the ONWARD trial so that only genetically positive patients are enrolled. The primary endpoint for analysis of efficacy is the change from baseline in the monthly number of heavy drinking days during the last 8 weeks of the 24-week treatment period. ONWARD is currently being conducted in 25 clinical sites in seven countries in Scandinavia and Central

and Eastern Europe (Sweden, Finland, Poland, Latvia, Estonia, Bulgaria and Croatia). The principal investigator is Professor Hannu E.R. Alho, Emeritus Professor of Addiction Medicine at the University of Helsinki.

## Forward Looking Statements

*This communication contains certain “forward-looking statements” within the meaning of the U.S. federal securities laws. Such statements are based upon various facts and derived utilizing numerous important assumptions and are subject to known and unknown risks, uncertainties and other factors that may cause actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by such forward-looking statements. Statements preceded by, followed by or that otherwise include the words “believes,” “expects,” “anticipates,” “intends,” “projects,” “estimates,” “plans” and similar expressions or future or conditional verbs such as “will,” “should,” “would,” “may” and “could” are generally forward-looking in nature and not historical facts, although not all forward-looking statements include the foregoing. The forward-looking statements include statements regarding the financing illustrating the tremendous confidence and support of our existing shareholders, as well as our conviction in the positive outlook for the business, the funding providing us more than sufficient capital to complete the ONWARD™ Phase 3 trial of AD04 as a genetically targeted treatment for Alcohol Use Disorder through data read-out, this funding allowing us to achieve key milestones for our pre-clinical adenosine program for non-opiate pain relief and the potential of AD04 to treat other addictive disorders such as opioid use disorder, gambling, and obesity. Any forward-looking statements included herein reflect our current views, and they involve certain risks and uncertainties, including, among others, our ability to maintain the confidence and support of our existing shareholders, as well as our conviction in the positive outlook for the business, our ability to complete the ONWARD™ Phase 3 trial of AD04 as a genetically targeted treatment for Alcohol Use Disorder through data read-out as planned, our ability to achieve key milestones for our pre-clinical adenosine program for non-opiate pain relief, our ability to enroll patients within the timelines anticipated and complete clinical trials on time and achieve desired results and benefits as expected, our ability to obtain regulatory approvals for commercialization of product candidates or to comply with ongoing regulatory requirements, regulatory limitations relating to our ability to promote or commercialize our product candidates for specific indications, acceptance of its product candidates in the marketplace and the successful development, marketing or sale of products, our ability to maintain our license agreements, the continued maintenance and growth of our patent estate, our ability to establish and maintain collaborations, our ability to obtain or maintain the capital or grants necessary to fund its research and development activities, and our ability to retain our key employees or maintain our Nasdaq listing. These risks should not be construed as exhaustive and should be read together with the other cautionary statement included in our Annual Report on Form 10-K for the year ended December 31, 2020, subsequent Quarterly Reports on Form 10-Q and current reports on Form 8-K filed with the Securities and Exchange Commission. Any forward-looking statement speaks only as of the date on which it was initially made. We undertake no obligation to publicly update or revise any forward-looking statement, whether as a result of new information, future events, changed circumstances or otherwise, unless required by law.*

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