



NEWS RELEASE

# BioVie to Host Conference Call and Webcast Wednesday November 1, 2023 at 8:00 AM ET

2023-10-30

CARSON CITY, Nev., Oct. 30, 2023 (GLOBE NEWSWIRE) -- BioVie Inc., (NASDAQ: BIVI) ("BioVie" or the "Company") a clinical-stage company developing innovative drug therapies for the treatment of advanced liver disease and neurological and neurodegenerative disorders, today announced that the Company's management will host a conference call and webcast on Wednesday, November 1 2023 at 8:00 AM Eastern Time. BioVie's President and CEO Cuong Do, Chief Medical Officer Dr. Joseph Palumbo, and Dr. Steven Arnold of the Harvard Medical School will review recent data on NE3017 presented at the 16th Clinical Trials on Alzheimer's Disease (CTAD) conference.

Registration details are as follows:

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|-------------------------|--|
| Date                    | November 1, 2023   |
| Time                    | 8:00 AM Eastern Time   |
| Conference Call Details | 1-877-407-3982 (U.S. Investors Dial-in)<br>1-201-493-6780 (International Investors Dial-in)<br>13742478 (Conference ID Code) |
| Webcast Link            | <a href="#">Click Here</a>   |
| Call-ME™ Feature        | <a href="#">Click Here</a>   |

## About BioVie

BioVie Inc. (NASDAQ: BIVI) is a clinical-stage company developing innovative drug therapies for the treatment of neurological and neurodegenerative disorders and advanced liver disease. In neurodegenerative disease, the Company's drug candidate NE3107 inhibits inflammatory activation of ERK and NFkB (e.g., TNF signaling) that leads to neuroinflammation and insulin resistance, but not their homeostatic functions (e.g., insulin signaling and neuron growth and survival). Both are drivers of Alzheimer's and Parkinson's diseases. The Company is conducting a potentially pivotal Phase 3 randomized, double-blind, placebo-controlled, parallel-group, multicenter study to evaluate NE3107 in patients who have mild to moderate Alzheimer's disease (NCT04669028). Results of a Phase 2

investigator initiated trial (NCT05227820) showing NE3107-treated patients experienced improved cognition and biomarker levels were presented at the Clinical Trial in Alzheimer's Disease (CTAD) annual conference in December 2022. An estimated six million Americans suffer from Alzheimer's. A Phase 2 study of NE3107 in Parkinson's disease (NCT05083260) has completed, and data presented at the International Conference on Alzheimer's and Parkinson's Disease and Related Neurological Disorders conference in Gothenburg, Sweden in March 2023 showed significant improvements in "morning on" symptoms and clinically meaningful improvement in motor control in patients treated with a combination of NE3107 and levodopa vs. patients treated with levodopa alone, and no drug-related adverse events. In liver disease, the Company's Orphan drug candidate BIV201 (continuous infusion terlipressin), with FDA Fast Track status, is being evaluated and discussed with guidance received from the FDA regarding the design of Phase 3 clinical testing of BIV201 for the treatment of ascites due to chronic liver cirrhosis. The active agent is approved in the U.S. and in about 40 countries for related complications of advanced liver cirrhosis. For more information, visit <http://www.bioviepharma.com/>.

#### Forward-Looking Statements

This press release contains forward-looking statements, including statements regarding the Company's strategy, plans and objectives, such as statements regarding the Company's anticipated timeline for announcing results from the NE3107 Phase 2 trial and details regarding the launch of its Phase 3 potential pivotal trials. Forward-looking statements may generally be identified by words such as "expect," "look forward to," "anticipate," "intend," "plan," "believe," "seek," "estimate," "will," "project" or words of similar meaning. Although BioVie Inc. believes such forward-looking statements are based on reasonable assumptions, it can give no assurance that its expectations will be attained. Actual results may vary materially from those expressed or implied by the statements herein due risks associated with conducting and completing clinical trials, including our reliance on third parties to conduct our clinical trials, to successfully defend potential future litigation, our ability to raise capital when needed on reasonable terms, changes in local or national economic conditions as well as various additional risks, many of which are now unknown and generally out of the Company's control, and which are detailed from time to time in reports filed by the Company with the SEC, including quarterly reports on Form 10-Q, reports on Form 8-K and annual reports on Form 10-K. BioVie Inc. does not undertake any duty to update any statements contained herein (including any forward-looking statements), except as required by law.

For Investor Relations Inquiries:

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Contact:  
Bruce Mackle  
Managing Director  
LifeSci Advisors, LLC  
[bmackle@lifesciadvisors.com](mailto:bmackle@lifesciadvisors.com)  
For Media Inquiries

Melyssa Weible  
Managing Partner  
Elixir Health Public Relations  
Ph: +1 201-723-5705  
[mweible@elixirhealthpr.com](mailto:mweible@elixirhealthpr.com)

