

May 20, 2021



# Cyclo Therapeutics to Participate in the M-Vest Virtual Conference Series: Alzheimer's Disease Panel

*- Panel discussion around innovating Alzheimer's Disease ("AD") drug development on Wednesday, May 26<sup>th</sup> 11:00 AM ET*

GAINESVILLE, Fla.--(BUSINESS WIRE)-- [Cyclo Therapeutics, Inc.](#) (Nasdaq: CYTH) ("Cyclo Therapeutics" or the "Company"), a clinical stage biotechnology company dedicated to developing life-changing medicines through science and innovation for patients and families suffering from diseases, today announced it will participate in the [M-Vest Virtual Conference Series: Alzheimer's Disease Panel](#) on Wednesday, May 26, 2021 at 11:00 AM ET.

Cyclo Therapeutics is currently advancing its lead product candidate [Trappsol<sup>®</sup> Cyclo<sup>™</sup>](#) in an AD program, the second pipeline candidate program generated from the Company's platform technology. The Company recently announced that it received feedback from the U.S. Food and Drug Administration ("FDA") supporting the Company's development strategy to submit an investigational new drug ("IND") application for a Phase 2 study of Trappsol<sup>®</sup> Cyclo<sup>™</sup> in the treatment of early AD, following a positive Type B interaction. Cyclo Therapeutics remains on track to file its IND for a Phase 2 study of AD in the second half of this year.

The panel discussion will be moderated by Jason McCarthy, PhD, Head of Biotechnology Research at Maxim Group. Sharon H. Hrynkow, PhD, Chief Scientific Officer, Senior Vice President for Medical Affairs will participate on behalf of Cyclo Therapeutics. Additional participating companies in the panel discussion will be Cassava Sciences, Vivoryon Therapeutics, Annovis Bio, and INmune Bio.

To RSVP for the M-Vest Virtual Alzheimer's Disease Panel please visit the event website, [here](#).

## **About Maxim Group LLC**

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## **About Cyclo Therapeutics**

Cyclo Therapeutics, Inc. is a clinical-stage biotechnology company dedicated to developing life-changing medicines through science and innovation for patients and families suffering from disease. The Company's Trappsol<sup>®</sup> Cyclo<sup>™</sup>, an orphan drug designated product in the United States and Europe, is the subject of three ongoing formal clinical trials for Niemann-Pick Disease Type C, a rare and fatal genetic disease, ([www.ClinicalTrials.gov](http://www.ClinicalTrials.gov) [NCT02939547](https://clinicaltrials.gov/ct2/show/study/NCT02939547), [NCT02912793](https://clinicaltrials.gov/ct2/show/study/NCT02912793), [NCT03893071](https://clinicaltrials.gov/ct2/show/study/NCT03893071) and [NCT04860960](https://clinicaltrials.gov/ct2/show/study/NCT04860960)). The Company is planning an early phase clinical trial using Trappsol<sup>®</sup> Cyclo<sup>™</sup> intravenously in Alzheimer's Disease based on encouraging data from an Expanded Access program for late-onset Alzheimer's Disease ([NCT03624842](https://clinicaltrials.gov/ct2/show/study/NCT03624842)). Additional indications for the active ingredient in Trappsol<sup>®</sup> Cyclo<sup>™</sup> are in development. For additional information, visit the Company's website: [www.cyclotherapeutics.com](http://www.cyclotherapeutics.com).

## **Safe Harbor Statement**

This press release contains "forward-looking statements" about the company's current expectations about future results, performance, prospects and opportunities, including, without limitation, statements regarding the satisfaction of closing conditions relating to the offering and the anticipated use of proceeds from the offering. Statements that are not historical facts, such as "anticipates," "believes" and "expects" or similar expressions, are forward-looking statements. These statements are subject to a number of risks, uncertainties and other factors that could cause actual results in future periods to differ materially from what is expressed in, or implied by, these statements. The factors which may influence the company's future performance include the company's ability to obtain additional capital to expand operations as planned, success in achieving regulatory approval for clinical protocols, enrollment of adequate numbers of patients in clinical trials, unforeseen difficulties in showing efficacy of the company's biopharmaceutical products, success in attracting additional customers and profitable contracts, and regulatory risks associated with producing pharmaceutical grade and food products. These and other risk factors are described from time to time in the company's filings with the Securities and Exchange Commission, including, but not limited to, the company's reports on Forms 10-K and 10-Q. Unless required by law, the company assumes no obligation to update or revise any forward-looking statements as a result of new information or future events.

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