Cyclo Therapeutics' Clinical Data Webinar Recording Available on Company Website

GAINESVILLE, FL – (BusinessWire) – May 26, 2020 – Cyclo Therapeutics, Inc. (OTCQB: CTDH), a clinical stage biotechnology company that develops cyclodextrinbased products for the treatment of Niemann-Pick Disease Type C (NPC) and Alzheimer's Disease, issued a press release titled "Cyclo Therapeutics Inc. Announces Positive Top Line Results from Phase I Trial and Interim Analysis of Phase I/II Trial using Trappsol[®] Cyclo[™] Intravenously to Treat Patients with Niemann-Pick Disease Type C1 (NPC1)" and held a webinar to discuss the data reported on Wednesday, May 20, 2020.

The presentation materials and a recording of the webinar are now available on the company's website: www.cyclotherapeutics.com

About Cyclo Therapeutics:

Cyclo Therapeutics, Inc. is a clinical-stage biotechnology company that develops cyclodextrin-based products for the treatment of Niemann-Pick Disease Type C and Alzheimer's Disease. The company's Trappsol[®] Cyclo[™], an orphan drug designated product in the United States and Europe, is the subject of three formal clinical trials for Niemann-Pick Disease Type C, a rare and often fatal genetic disease, (ClinicalTrials.gov NCT02939547, NCT02912793 and NCT03893071). The company is planning an early phase clinical trial using Trappsol[®] Cyclo[™] intravenously in Alzheimer's Disease based on encouraging data from an Expanded Access program for late-onset Alzheimer's Disease (NCT03624842). Additional indications for the active ingredient in Trappsol[®] Cyclo[™] are in development. For additional information, visit the company's website: 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. 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. Also, the uncertainty around the severity and duration of the impact of COVID-19 on our business and operations. 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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