Synaptogenix to Present Research Highlighting Potential Broad Applicability for Lead Compound Bryostatin at the International Conference on Alzheimer's and Parkinson's Diseases (AD/PD 2022)

PR Newswire

NEW YORK , March 15, 2022 /PRNewswire/ -- Synaptogenix, Inc. (Nasdaq: SNPX), an emerging biopharmaceutical company developing regenerative therapeutics for neurodegenerative disorders, announced today that Dr. Daniel Alkon , the Company's President and Chief Scientific Officer, will present recent clinical results today, Tuesday, March 15 th , at the International Conference on Alzheimer's (AD) and Parkinson's Diseases (AD/PD 2022). The titled abstract, "Neurotrophic Restoration for Neurodegenerative Disease" summarizes therapeutic improvement by Bryostatin for advanced AD patients in placebo-controlled trials and its potential benefits for other neurodegenerative disorders such as Parkinson's Disease.

Dr. Daniel Alkon , commented, "Synaptogenix's lead compound, Bryostatin-1, is currently in an ongoing National Institutes of Health sponsored six-month trial focused on Alzheimer's disease. Our work published today suggests we may be able to broaden the possible applicability of Bryostatin to more complicated dementias with more than one pathology beyond AD, such as Parkinson's, multi-infarct dementia, and Multiple Sclerosis."

Synaptogenix AD/PD 2022 presentation will go live on Tuesday, March 15th, 2022 , at 8:00 AM CET (3:00 AM EDT) and will be accessible to AD/PD 2022 registered meeting attendees in-person and on its virtual platform.

About Synaptogenix, Inc.

Synaptogenix is a clinical-stage biopharmaceutical company that has historically worked to develop novel therapies for neurodegenerative diseases. Synaptogenix has conducted clinical and preclinical studies of its lead therapeutic candidate, Bryostatin-1, in Alzheimer's disease. Preclinical studies have also demonstrated Bryostatin's regenerative mechanisms of action for the rare disease, Fragile X syndrome, and for other neurodegenerative disorders such as multiple sclerosis, stroke, and traumatic brain injury. The U.S. Food and Drug Administration has granted Orphan Drug Designation to Synaptogenix for Bryostatin-1 as a treatment for Fragile X syndrome. Bryostatin-1 has already undergone testing in more than 1,500 people in cancer studies, thus creating a large safety data base that will further inform clinical trial designs.

Additional information about Synaptogenix, Inc. may be found on its website: www.synaptogen.com .

Forward-Looking Statements

Any statements contained in this press release that do not describe historical facts may constitute forward-looking statements. These forward-looking statements include statements regarding the continued development of use of Bryostatin-1 for Alzheimer's disease, Parkinson's, multi-infarct dementia, Multiple Sclerosis and other cognitive diseases. Such forward-looking statements are subject to risks and uncertainties and other influences, many of which the Company has no control over. There can be no assurance that the clinical program for Bryostatin-1 will be successful in demonstrating safety and/or efficacy, that we will not encounter problems or delays in clinical development, or that Bryostatin-1 will ever receive regulatory approval or be successfully commercialized. Actual results and the timing of certain events and circumstances may differ materially from those described by the forward-looking statements as a result of these risks and uncertainties. Additional factors that may influence or cause actual results to differ materially from expected or desired results may include, without limitation, the Company's inability to obtain adequate financing, the significant length of time associated with drug development and related insufficient cash flows and resulting illiquidity, the Company's patent portfolio, the Company's inability to expand its business, significant government regulation of pharmaceuticals and the healthcare industry, lack of product diversification, availability of the Company's raw materials, existing or increased competition, stock volatility and illiquidity, and the Company's failure to implement its business plans or strategies. These and other factors are identified and described in more detail in the Company's filings with the Securities and Exchange Commission. The Company does not undertake to update these forward-looking statements.

Contact information:
Investors and Media

800-811-5591
ir@synaptogen.com

Robert Weinstein
Chief Financial Officer
Synaptogenix, Inc.
rweinstein@synaptogen.com

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