



Aptinyx Raises \$70 Million in Series B Financing to Advance Growing Pipeline through Clinical Studies in Neurologic Disorders

Evanston, Ill., Dec 18, 2017 – Aptinyx Inc., a clinical-stage biopharmaceutical company developing transformative therapies for challenging neurologic disorders, today is announcing the completion of a \$70 million Series B financing to fund advancement of its expanding clinical-stage pipeline.

The financing was led by Bain Capital Life Sciences. Additional new investors include Adage Capital, Agent Capital, HBM Healthcare Investments, Nan Fung Life Sciences, Partner Fund Management, and Rock Springs Capital. Existing investors also participated in the Series B round, including New Leaf Venture Partners, Frazier Healthcare Partners, Longitude Capital, Osage University Partners, Adams Street Partners, LVP Life Science Ventures, PathoCapital, Goudy Park Capital, Beecken Petty O’Keefe & Company, and Northwestern University.

“This financing will provide fuel for our growing pipeline as we continue to develop our innovative and differentiated NMDA receptor modulators for the treatment of a variety of neurologic disorders with high unmet medical need,” said Norbert Riedel, Ph.D., president and chief executive officer of Aptinyx. “We are delighted by the support of this syndicate of highly respected global investors with extensive domain expertise, which will be valuable as we advance our pipeline toward late-stage development in multiple indications.”

Concurrent with the financing, Adam M. Koppel, M.D., Ph.D., a managing director of Bain Capital Life Sciences, has joined the Aptinyx board of directors.

“Aptinyx is leading a renaissance in neurotherapeutic drug development with its prolific drug discovery platform, unique mechanism of action relevant in a number of challenging neurologic disorders, and ability to quickly advance drug candidates,” said Dr. Koppel. “We are eager to support Aptinyx in the development of its portfolio of promising NMDA receptor modulators for the benefit of patients who suffer from these neurologic conditions.”

The funds will support clinical studies of Aptinyx’s drug candidates, including NYX-2925, in development for the treatment of neuropathic pain, and NYX-783, in development as a therapy for post-traumatic stress disorder (PTSD). The company also plans to advance a third proprietary compound into the clinic in 2018, initiate development programs in new indications, and continue discovery of additional novel, small-molecule modulators of N-methyl-D-aspartate (NMDA) receptors.

Aptinyx is currently conducting a Phase 2 [study](#) of NYX-2925 in patients with neuropathic pain associated with diabetic peripheral neuropathy (DPN), as well as an exploratory [study](#) in patients with fibromyalgia. The FDA has granted Fast Track designation to the development of NYX-2925 for neuropathic pain associated with DPN.

Aptinyx [recently](#) initiated a Phase 1 clinical study to evaluate the safety and tolerability of NYX-783. The company intends to develop NYX-783 for the treatment of PTSD, for which the FDA has granted Fast Track designation.

Aptinyx's chemistry and discovery platform has generated numerous small-molecule modulators of the NMDA receptor, including clinical drug candidates NYX-2925 and NYX-783. In studies to date, these molecules have demonstrated high oral bioavailability, diverse NMDA receptor subtype binding profiles, differentiated efficacy across preclinical models of various nervous system conditions, and very favorable safety.

About Aptinyx

Aptinyx Inc. is a clinical-stage biopharmaceutical company discovering and developing transformative therapies for challenging disorders of the brain and nervous system. Aptinyx has a proven platform for discovery of novel compounds that work through a unique mechanism to modulate – rather than block or over-activate – NMDA receptors and enhance synaptic plasticity, the foundation of neural cell communication. Drugs that modulate NMDA receptors in this distinct way have both robust efficacy and exceptionally favorable safety. The company's lead drug candidate, NYX-2925, is in Phase 2 clinical development as a therapy for neuropathic pain and its second drug candidate, NYX-783, is in Phase 1 clinical development for the treatment of post-traumatic stress disorder (PTSD). Both programs have received Fast Track designation by the FDA. Aptinyx is also advancing additional compounds from its proprietary chemistry platform, which continues to generate a rich and diverse pipeline of small-molecule NMDA receptor modulators with the potential to treat an array of neurologic disorders. For more information, visit www.aptinyx.com.

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