



Edgar Filing: VistaGen Therapeutics, Inc. - Form 8-K

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (17 CFR 230.405) or Rule 12b-2 of the Securities Exchange Act of 1934 (17 CFR 240.12b-2)

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act



Item 8.01 Other Events.

VistaGen Therapeutics, Inc. (the “Company”) today announced the initiation of ELEVATE, the Company’s double-blind, placebo-controlled Phase 2 clinical study to evaluate the efficacy and safety of AV-101 (L-4-chlorokynurenine) as an adjunctive treatment of Major Depressive Disorder (“MDD”) in patients with an inadequate response to current antidepressants approved by the U.S. Food and Drug Administration (“FDA”). AV-101, the Company’s oral N-methyl-D-aspartate (“NMDA”) receptor glycine B (“GlyB”) antagonist, belongs to a new generation of investigational medicines in neuropsychiatry known as glutamate receptor modulators that have the potential to treat MDD faster than current FDA-approved antidepressants. A copy of the Company’s press release is attached to this Current Report on Form 8-K as Exhibit 99.1 and is incorporated by reference herein.

Item 9.01 Financial Statements and Exhibits.

See Exhibit Index.



SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned thereunto duly authorized.

VistaGen Therapeutics, Inc.

Date: April 5, 2018 By: /s/ Shawn K. Singh  
Shawn K. Singh  
Chief Executive Officer



EXHIBIT INDEX

Exhibit No. Description

99.1 Press Release issued by VistaGen Therapeutics, Inc., dated April 5, 2018