

CytoDyn Inc.  
Form 8-K  
December 12, 2016

**UNITED STATES**  
**SECURITIES AND EXCHANGE COMMISSION**  
**WASHINGTON, DC 20549**

**FORM 8-K**

**CURRENT REPORT**  
**PURSUANT TO SECTION 13 OR 15(d)**  
**OF THE SECURITIES EXCHANGE ACT OF 1934**  
**Date of Report (Date of earliest event reported): December 12, 2016**

**CytoDyn Inc.**

**(Exact name of registrant as specified in its charter)**

**Delaware**  
**(State or other jurisdiction**

**of incorporation)**

**1111 Main Street, Suite 660**

**000-49908**  
**(SEC**

**File Number)**

**75-3056237**  
**(I.R.S. Employer**

**Identification No.)**

**98660**

**Vancouver, Washington**  
**(Address of principal executive offices)** **(Zip Code)**  
**Registrant's telephone number, including area code: (360) 980-8524**

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions (see General Instruction A.2. below):

Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

**Item 8.01. Other Events.**

On December 12, 2016, the Company announced the first several patients have been treated in the Company's Phase 3 clinical trial with PRO 140 as a single-agent maintenance therapy in virally suppressed subjects with HIV.

This multicenter, open-label trial will enroll 300 patients prequalified with CCR5-tropic HIV-1 infection who are clinically stable on standard-of-care highly active antiretroviral therapy (HAART). The objective of the monotherapy trial is to assess the efficacy, safety and tolerability of PRO 140 as a long-acting, single-agent maintenance therapy for the chronic suppression of HIV. Patients enrolled in the trial will be shifted from daily HAART regimens to weekly PRO 140 subcutaneous injections for 48 weeks.

**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 hereunto duly authorized.

CytoDyn Inc.

December 12, 2016

By: */s/ Michael D. Mulholland*  
Name: Michael D. Mulholland  
Title: Chief Financial Officer