March 19, 2014

CytoDyn Announces Investor Call to Provide Update on Clinical Development Strategy

-Company to host live conference call and webcast with investment community on Tuesday, March 25 at 1:30 p.m. PT-

VANCOUVER, Wash.-- **CytoDyn Inc. (OTCQB: CYDY)**, a biotechnology company focused on the development of new therapies for combating infection with immune deficiency viruses, announced today that the Company will host a live conference call and webcast with the investment community on Tuesday, March 25, 2014 at 1:30 p.m. PT to provide an update on the Company's clinical development strategy for lead product candidate, PRO 140, including progress with its preparations for its treatment substitution Phase 2b study in patients with Human Immunodeficiency Virus (HIV). PRO 140 is a humanized monoclonal antibody directed against CCR5, a molecular portal that HIV uses to enter cells, and belongs to a class of HIV therapies known as entry inhibitors that block HIV from entering into and infecting certain cells.

Conference Call and Webcast Instructions

CytoDyn's management team will host a conference call and live audio webcast on Tuesday, March 25, 2014 at 1:30 p.m. PT.

Interested participants and investors may access this conference call by dialing 877-407-2986 (U.S./Canada) or 201-378-4916 (international).

A live audio webcast may also be accessed via the Investors section of CytoDyn's corporate web site at www.cytodyn.com, and will be archived for 30 days. Web participants are encouraged to go to the web site 15 minutes prior to the start of the call to register, download and install any necessary software.

A replay of the conference call will be available until April 1, 2014. To access the replay, interested parties may dial 877-660-6853 (U.S./Canada) or 201-612-7415 (International); Conference ID: 13578723.

About PRO 140

PRO 140 belongs to a new class of HIV/AIDS therapeutics -- viral-entry inhibitors -- that are intended to protect healthy cells from viral infection. PRO 140 is a humanized monoclonal antibody directed against CCR5, a molecular portal that HIV uses to enter cells.

PRO 140 has been the subject of four Phase 1/1b and two Phase 2a clinical trials, each of which demonstrated PRO 140's ability to significantly reduce HIV viral load in human test

subjects, and has also been designated a “fast track” product candidate by the FDA. The PRO 140 antibody appears to be a powerful antiviral agent while not being a drug, potentially leading to fewer side effects and less frequent dosing requirements as compared to daily drug therapies currently in use.

About CytoDyn

CytoDyn is a biotechnology company focused on developing subcutaneously delivered humanized cell-specific monoclonal antibodies (mAbs) as entry inhibitors for the treatment and prevention of Human Immunodeficiency Virus (HIV). The Company has one of the leading mAbs under development for HIV infection, PRO 140, which is a Late Stage 2 humanized mAb with demonstrated antiviral activity in man. PRO 140 blocks the HIV co-receptor CCR5 and clinical trial results thus far indicate that it does not affect the normal function of the cell. Results from Phase 1/1b and Phase 2a human clinical trials have shown that PRO 140 can significantly reduce viral burden in people infected with HIV. CytoDyn intends to continue to develop PRO 140 as a therapeutic anti-viral agent in persons infected with HIV. For more information on the Company please visit www.cytodyn.com.

Forward-Looking Statements

This press release includes forward-looking statements and forward-looking information within the meaning of United States securities laws. These statements and information represent CytoDyn’s intentions, plans, expectations, and beliefs and are subject to risks, uncertainties and other factors, many beyond CytoDyn’s control. These factors could cause actual results to differ materially from such forward-looking statements or information. The words “believe,” “estimate,” “expect,” “intend,” “attempt,” “anticipate,” “foresee,” “plan,” and similar expressions and variations thereof identify certain of such forward-looking statements or forward-looking information, which speak only as of the date on which they are made.

CytoDyn disclaims any intention or obligation to publicly update or revise any forward-looking statements or forward-looking information, whether as a result of new information, future events or otherwise, except as required by applicable law. Readers are cautioned not to place undue reliance on these forward-looking statements or forward-looking information. While it is impossible to identify or predict all such matters, these differences may result from, among other things, the inherent uncertainty of the timing and success of and expense associated with research, development, regulatory approval, and commercialization of CytoDyn’s products and product candidates, including the risks that clinical trials will not commence or proceed as planned; products appearing promising in early trials will not demonstrate efficacy or safety in larger-scale trials; future clinical trial data on CytoDyn’s products and product candidates will be unfavorable; funding for additional clinical trials may not be available; CytoDyn’s products may not receive marketing approval from regulators or, if approved, may fail to gain sufficient market acceptance to justify development and commercialization costs; competing products currently on the market or in development may reduce the commercial potential of CytoDyn’s products; CytoDyn, its collaborators or others may identify side effects after the product is on the market; or efficacy or safety concerns regarding marketed products, whether or not scientifically justified, may lead to product recalls, withdrawals of marketing approval, reformulation of the product, additional pre-clinical testing or clinical trials, changes in labeling of the product, the need for additional marketing applications, or other adverse events.

CytoDyn is also subject to additional risks and uncertainties, including risks associated with the actions of its corporate, academic, and other collaborators and government regulatory agencies; risks from market forces and trends; potential product liability; intellectual property litigation; environmental and other risks; and risks that current and pending patent protection for its products may be invalid, unenforceable, or challenged or fail to provide adequate market exclusivity. There are also substantial risks arising out of CytoDyn's need to raise additional capital to develop its products and satisfy its financial obligations; the highly regulated nature of its business, including government cost-containment initiatives and restrictions on third-party payments for its products; the highly competitive nature of its industry; and other factors set forth in CytoDyn's Quarterly Report on Form 10-Q for the quarter ended November 30, 2013 and other reports filed with the U.S. Securities and Exchange Commission.

Investors and Media
Jenene Thomas Communications, LLC
Jenene Thomas, 908-938-1475
Investor Relations and Corporate Communications Advisor
jthomas@cytodyn.com

Source: CytoDyn Inc.