ACADIA Pharmaceuticals to Announce Second Quarter Financial Results on August 6, 2015

July 30, 2015

ACADIA to Host Conference Call and Webcast on Thursday, August 6, 2015, at 5:00 p.m. Eastern Time

SAN DIEGO--(BUSINESS WIRE)--Jul. 30, 2015-- ACADIA Pharmaceuticals Inc. (NASDAQ: ACAD), a biopharmaceutical company focused on the development and commercialization of innovative medicines that address unmet medical needs in neurological and related central nervous system disorders, today announced that it will report its unaudited financial results for the second quarter ended June 30, 2015 on Thursday, August 6, 2015, after the U.S. financial markets close. ACADIA's management will host a conference call and webcast on Thursday, August 6, 2015, at 5:00 p.m. Eastern Time to discuss ACADIA's financial results and development programs.

The conference call may be accessed by dialing 855-638-4820 for participants in the United States or Canada and 443-877-4067 for international callers (reference passcode 92110444). A telephone replay of the conference call may be accessed through August 20, 2015 by dialing 855-859-2056 for callers in the United States or Canada and 404-537-3406 for international callers (reference passcode 92110444). The conference call also will be webcast live on ACADIA's website, www.acadia-pharm.com under the investors section and will be archived there until August 20, 2015.

About ACADIA Pharmaceuticals

ACADIA is a biopharmaceutical company focused on the development and commercialization of innovative medicines to address unmet medical needs in neurological and related central nervous system disorders. ACADIA has a pipeline of product candidates led by NUPLAZID™ (pimavanserin), for which we have reported positive Phase III trial results in Parkinson's disease psychosis and which has the potential to be the first drug approved in the United States for this disorder. Pimavanserin is also in Phase II development for Alzheimer's disease psychosis and has successfully completed a Phase II trial in schizophrenia. ACADIA also has clinical-stage programs for chronic pain and glaucoma in collaboration with Allergan, Inc. All product candidates are small molecules that emanate from internal discoveries. ACADIA maintains a website at www.acadia-pharm.com to which we regularly post copies of our press releases as well as additional information and through which interested parties can subscribe to receive e-mail alerts.

Forward-Looking Statements

Statements in this press release that are not strictly historical in nature are forward-looking statements. These statements include but are not limited to statements related to the progress and timing of ACADIA's drug discovery and development programs, either alone or with a partner, including clinical trials, the benefits to be derived from ACADIA's product candidates, in each case including NUPLAZID (pimavanserin), and the potential for NUPLAZID to be the first drug approved in the United States for Parkinson's disease psychosis. These statements are only predictions based on current information and expectations and involve a number of risks and uncertainties. Actual events or results may differ materially from those projected in any of such statements due to various factors, including the risks and uncertainties inherent in drug discovery, development, approval and commercialization, and collaborations with others, and the fact that past results of clinical trials may not be indicative of future trial results. For a discussion of these and other factors, please refer to ACADIA's annual report on Form 10-K for the year ended December 31, 2014 as well as ACADIA's subsequent filings with the Securities and Exchange Commission. You are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date hereof. This caution is made under the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. All forward-looking statements are qualified in their entirety by this cautionary statement and ACADIA undertakes no obligation to revise or update this press release to reflect events or circumstances after the date hereof, except as required by law.

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