



February 22, 2017

Nektar to Announce Financial Results for the Fourth Quarter and Year-Ended 2016 on Wednesday, March 1, 2017, After Close of U.S.-Based Financial Markets

SAN FRANCISCO, Feb. 22, 2017 /PRNewswire/ -- Nektar Therapeutics (Nasdaq: NKTR) will announce its financial results for the fourth quarter and year-ended December 31, 2016, on Wednesday, March 1, 2017, after the close of U.S.-based financial markets. Howard Robin, president and chief executive officer, will host a conference call to review the results beginning at 5:00 p.m. Eastern Time (ET)/2:00 p.m. Pacific Time (PT).

The press release and a live audio-only Webcast of the conference call can be accessed through a link that is posted on the home page and Investors section of the Nektar website: <http://www.nektar.com>. The web broadcast of the conference call will be available for replay through Monday, April 3, 2017.

To access the conference call, follow these instructions:

Dial: (877) 881-2183 (U.S.); (970) 315-0453 (international)
Passcode: 75461730 (Nektar Therapeutics is the host)

About Nektar

Nektar Therapeutics has a robust R&D pipeline and portfolio of approved partnered medicines in oncology, pain, immunology and other therapeutic areas. In the area of oncology, Nektar is developing NKTR-214, an immuno-stimulatory CD122-biased agonist, that is in Phase 1/2 clinical development for patients with solid tumors. ONZEALD™ (etirinotecan pegol), a long-acting topoisomerase I inhibitor, is being developed for patients with advanced breast cancer and brain metastases and is partnered with Daiichi Sankyo in Europe. In the area of pain, Nektar has an exclusive worldwide license agreement with AstraZeneca for MOVANTIK™ (naloxegol), the first FDA-approved once-daily oral peripherally-acting mu-opioid receptor antagonist (PAMORA) medication for the treatment of opioid-induced constipation (OIC), in adult patients with chronic, non-cancer pain. The product is also approved in the European Union as MOVENTIG® (naloxegol) and is indicated for adult patients with OIC who have had an inadequate response to laxatives. The AstraZeneca agreement also includes NKTR-119, an earlier stage development program that is a co-formulation of MOVANTIK and an opioid. NKTR-181, a wholly owned mu-opioid analgesic molecule for chronic pain conditions, is in Phase 3 development. In hemophilia, Nektar has a collaboration agreement with Baxalta for ADYNOVATE® [Antihemophilic Factor (Recombinant)], a longer-acting PEGylated Factor VIII therapeutic approved in the U.S. and Japan for patients over 12 with hemophilia A. In anti-infectives, the company has two collaborations with Bayer Healthcare, Cipro Inhale in Phase 3 for non-cystic fibrosis bronchiectasis and Amikacin Inhale in Phase 3 for patients with Gram-negative pneumonia.

Nektar's technology has enabled nine approved products in the U.S. or Europe through partnerships with leading biopharmaceutical companies, including AstraZeneca's MOVANTIK™, Baxalta's ADYNOVATE®, UCB's Cimzia® for Crohn's disease and rheumatoid arthritis, and Amgen's Neulasta® for neutropenia.

Nektar is headquartered in San Francisco, California, with additional operations in Huntsville, Alabama and Hyderabad, India. Further information about the company and its drug development programs and capabilities may be found online at <http://www.nektar.com>.

MOVANTIK™ is a trademark and MOVENTIG® is a registered trademark of the AstraZeneca group of companies.

ADYNOVATE® is a registered trademark of Baxalta Incorporated, a wholly owned, indirect subsidiary of Shire plc.

Cimzia® is a registered trademark of UCB.

ONZEALD™ is a trademark of Nektar Therapeutics.

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