Transcept Pharmaceuticals to Host Conference Call to Discuss FDA Approval of Intermezzo®

Call Scheduled for Wednesday, November 23, 2011 at 2:30 p.m. ET

POINT RICHMOND, Calif., Nov. 23, 2011 /PRNewswire/ -- Transcept Pharmaceuticals, Inc. (Nasdaq: TSPT), a specialty pharmaceutical company focused on the development and commercialization of proprietary products that address important therapeutic needs in the field of neuroscience, announced that it will host a conference call to discuss the approval of Intermezzo® (zolpidem tartrate sublingual tablet) by the U.S. Food and Drug Administration.

Conference Call Information
Date: Wednesday, November 23, 2011
Time: 2:30 p.m. ET
Dial-in (U.S.): 877-638-4558
Dial-in (International): 914-495-8537

A simultaneous web cast of the conference call will be available on the Investors section of the Transcept website at www.transcept.com. A replay of the call will be available on the website shortly after the conclusion of the call until December 31, 2011.

A telephone replay of the conference call will be available shortly after the conclusion of the call until end of business November 30, 2011. The replay telephone numbers are 855-859-2056 (U.S.) and 404-537-3406 (International), replay passcode: 30305793.

About Transcept

Transcept Pharmaceuticals, Inc. is a specialty pharmaceutical company focused on the development and commercialization of proprietary products that address important therapeutic needs in the field of neuroscience. Intermezzo (zolpidem tartrate sublingual tablet) is the first FDA approved Transcept product and is indicated for use as needed for the treatment of insomnia when a middle-of-the-night awakening is followed by difficulty returning to sleep. Transcept and Purdue have entered into a collaboration agreement for the development and commercialization of Intermezzo in the United States.

Transcept is currently conducting a Phase 2 study of TO-2061, a low dose ondansetron augmentation therapy for patients with obsessive compulsive disorder (OCD) who have not adequately responded to treatment with approved first-line pharmacotherapy. For further information, please visit the Transcept website at: www.transcept.com.

Contact:
Transcept Pharmaceuticals, Inc.
Greg Mann
Director, Corporate Communications
(510) 215-3567
gmann@transcept.com

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