Questcor Pharmaceuticals to Present at the Oppenheimer Annual Healthcare Conference

December 5, 2013

ANAHEIM, Calif., Dec. 5, 2013 /PRNewswire/ -- Questcor Pharmaceuticals, Inc. (NASDAQ: QCOR) announced today that executive management will present at the Oppenheimer 24th Annual Healthcare Conference in New York City on December 11, 2013 at 8:20 AM ET/5:20 AM PT.

A live webcast and subsequent archived replay of the presentation will be accessible at http://ir.questcor.com/events.cfm. The replay will be available for approximately 90 days after the event.

About Questcor

Questcor Pharmaceuticals, Inc. is a biopharmaceutical company focused on the treatment of patients with serious, difficult-to-treat autoimmune and inflammatory disorders. Questcor also provides specialty contract manufacturing services to the global pharmaceutical industry through its wholly-owned subsidiary BioVectra Inc. Questcor's primary product is H.P. Acthar® Gel (repository corticotropin injection), an injectable drug that is approved by the FDA for the treatment of 19 indications. Of these 19 indications, Questcor currently generates substantially all of its net sales from the following approved indications: the treatment of proteinuria in the nephrotic syndrome (NS) of the idiopathic type, the treatment of acute exacerbations of multiple sclerosis (MS) in adults, the treatment of certain rheumatology related conditions, and the treatment of infantile spasms (IS) in infants and children under two years of age. With respect to NS, the FDA has approved Acthar to "induce a diuresis or a remission of proteinuria in the nephrotic syndrome without uremia of the idiopathic type or that due to lupus erythematosus." Questcor has announced its intent to initiate a pilot commercialization effort for Acthar for the treatment of respiratory manifestations of symptomatic sarcoidosis. The FDA approved package insert for Acthar includes "symptomatic sarcoidosis" under the heading "Respiratory Diseases." Questcor is also exploring the possibility of developing markets for other on-label indications and the possibility of pursuing FDA approval of additional indications not currently on the Acthar label where there is high unmet medical need. Questcor also has agreed to acquire certain international rights for Synacthen® (tetracosactide) and Synacthen Depot®, and has licensed the right to develop and seek FDA approval for these products in the United States. For more information about Questcor, please visit www.questcor.com.

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