



NEWS RELEASE

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For Immediate Release:

Nymox Announces New Positive Results at 5 Years After Single Treatment in Long Term Study of NX-1207 for Benign Prostatic Hyperplasia

HASBROUCK HEIGHTS, NJ (February 24, 2009) Nymox Pharmaceutical Corporation (NASDAQ: NYMX) today announced new positive results from a long term outcome study of NX-1207 for benign prostatic hyperplasia (BPH). The study evaluated symptomatic progress of U.S. patients involved in the Company's two initial 2003 Phase 1-2 studies of NX-1207. Patients treated with NX-1207 were followed-up on an unselected and as available basis and assessed for symptomatic improvement, treatment outcomes, and durability of efficacy 64 months after treatment.

Data was available for 75% of the subjects in the initial studies. Overall, 67% of the patients in the new outcome study treated with NX-1207 reported no current drug treatment for their BPH and had a mean improvement of 11 points in AUA Symptom Score. In addition, 46% of the patients reported no other approved treatments at any time for their BPH since their original treatment with NX-1207, with a mean improvement of 13 points. This sustained improvement in BPH symptom score after NX-1207 treatment compares favorably to the 3.5 to 5 points reported in published studies of currently approved BPH drugs, which, unlike NX-1207 treatment, require uninterrupted, daily administration to be effective.

NX-1207 has entered its Phase 3 development program, the last stage before filing with the FDA for approval. The drug involves a new targeted approach to the treatment of BPH. NX-1207 is injected by a urologist in an office setting directly into the zone of the prostate where the enlargement occurs. The entire procedure lasts on average 5-10 minutes, with the injection taking 1-2 minutes, does not require anesthesia or catheterization, and involves little or no pain or discomfort.

Reception to independent presentations of data from clinical trials of NX-1207 given at recent U.S. specialty urology (AUA) meetings has been very positive. In multicenter U.S. clinical trials to date NX-1207 has been found to produce significant improvements in BPH symptoms without the side effects associated with approved drugs, which can include sexual dysfunction, blood pressure changes and other adverse reactions.

BPH treatment represents a growing market with more than 100 million men worldwide being estimated to suffer from BPH symptoms. The disorder is a common affliction of older men, affecting approximately half of men over age 50 and close to 90% of men by age 80, and is associated with growth in prostate size as men age. BPH causes difficulties with urination associated with aging, such as urination at night, urge to void frequently, hesitancy, weak stream, and other problems.

More information about Nymox is available at www.nymox.com, email: info@nymox.com, or 800-936-9669.

This press release contains certain "forward-looking statements" as defined in the United States Private Securities Litigation Reform Act of 1995 that involve a number of risks and uncertainties. There can be no assurance that such statements will prove to be accurate and the actual results and future events could differ materially from management's current expectations. Such factors are detailed from time to time in Nymox's filings with the United States Securities and Exchange Commission and other regulatory authorities.

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