



## NEWS RELEASE

### **For Further Information Contact:**

Roy Wolvin  
Nymox Pharmaceutical Corporation  
1-800-93NYMOX  
[www.nymox.com](http://www.nymox.com)

### **For Immediate Release:**

## **Nymox Announces New Positive Results from 3 ½ Year Study of NX-1207 for Benign Prostatic Hyperplasia**

HASBROUCK HEIGHTS, NJ (May 7, 2007) Nymox Pharmaceutical Corporation (NASDAQ: NYMX) today announced positive results from a new 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 Phase 1-2 studies initiated in 2003. 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 3 ½ years after NX-1207 treatment.

Overall, patients treated with NX-1207 showed a mean improvement of 8.6 points in the primary outcome endpoint of AUA Symptom Score value 42 months after NX-1207 treatment. 50% of these patients reported no additional treatment for the BPH during this period and had a mean improvement of 10.0 points in AUA Symptom Score. 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.

Paul Averback, CEO of Nymox, said, "This new data on NX-1207 showing sustained symptomatic improvement after 3 ½ years clearly confirms the enduring benefits of NX-1207 treatment for BPH."

The Company previously completed three U.S. trials for NX-1207, including most recently a Phase 2 double-blind, placebo controlled, randomized multi-site U.S. study, which showed positive efficacy and safety results for NX-1207 after 3 months in patients with BPH. Overall, patients treated with NX-1207 showed after 3 months a mean improvement of 9.35 points in AUA Symptom Score values, the standard scale used to evaluate BPH drugs and treatments. This improvement compares favorably to the 3.5 to 5 point reported in published studies of currently approved drugs for BPH and reached statistical significance when compared to placebo. Subjects treated with NX-1207 also showed an overall statistically significant reduction in mean prostate volume. The results of the trial demonstrated the excellent safety and side effect profile of NX-1207. Subjects treated with NX-1207 had no serious side effects. In particular, patients given NX-1207 had no (0%) significant sexual side effects.

The company also recently reported positive long-term outcome results from an 8-19 month study of 116 unselected subjects from 26 U.S. clinical sites in a blinded placebo-controlled study, which reached statistical significance ( $p=.028$ ). In that study, overall without further NX-1207 treatment, patients initially treated with NX-1207 showed a total pooled mean improvement of 7.4 points in the primary outcome endpoint of AUA Symptom Score values.

The AUA Symptom Score is a standardized measurement of BPH symptoms and includes data on 1) sensations of incomplete emptying of the bladder; 2) need to urinate frequently; 3) stopping and starting during urination; 4) urgent need to urinate; 5) weakness of urinary stream; 6) need to push or strain during urination; and 7) urination during sleep (nocturia). Published studies of currently approved drugs for BPH show AUA Symptom Score improvement in the 3.5 to 5 point range.

BPH afflicts approximately half of men over age 50 and close to 90% of men by age 80. The disorder 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](http://www.nymox.com), email: [info@nymox.com](mailto: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. The conduct of clinical trials and the development of drug products involve substantial risks and uncertainties and actual results may differ materially from expectations. Promising early results do not ensure that later stage or larger scale clinical trials will be successful or will proceed as expected. Such factors are detailed from time to time in Nymox's filings with the United States Securities and Exchange Commission and other regulatory authorities.*