



NEWS RELEASE

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For Immediate Release:

Nymox Announces Clinical Study Presentation on BPH Drug at American Urological Association Meeting

HASBROUCK HEIGHTS, NJ (April 17, 2008) Nymox Pharmaceutical Corporation (NASDAQ: NYMX) today announced that new clinical trial data concerning the safety and efficacy of the Company's NX-1207 for benign prostatic hyperplasia (BPH) will be presented at the South Central Section of the American Urological Association Meeting in San Diego in September. The paper is authored by leading clinical research investigators participating in the U.S. clinical trials of NX-1207.

NX-1207 is Nymox's patented drug candidate for the treatment of BPH, a common affliction of men related to enlarged prostate. Blinded clinical trials to date have shown that men treated with NX-1207 reported statistically significant improvement in BPH symptoms 3 months after a single NX-1207 treatment with no reported serious drug-related side effects, including no (0%) significant sexual side effects. In two multi-center Phase 2 U.S. prospective randomized blinded clinical trials, the aggregated mean improvement in the Primary Endpoint of BPH Symptom Score for 2.5 mg NX-1207 was 10.3 points or a 44% improvement in Symptom Score.

Data from the NX02-0016 study (announced by the Company on February 6 and March 11, 2008) showed that NX-1207 markedly reduced the incidence of nighttime urination (nocturia), a particularly bothersome symptom associated with BPH. After 90 days, subjects treated with a therapeutic dose of NX-1207 had a mean reduction in nocturia symptom score of 41% versus 4% for subjects treated with finasteride, an approved BPH treatment. This improvement was statistically significant ($p < .001$). Having to repeatedly get up in the night to urinate is a common symptom of BPH that can cause chronic sleep loss and, in turn, lead to fatigue, memory deficits, mood changes including depression, and increased risk of long term medical problems.

Results of 5 follow-up studies of available subjects from NX-1207 clinical trials have provided evidence of durable benefits from NX-1207 treatment for up to 3½ years from the date of treatment.

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, and can cause acute urinary retention requiring immediate medical attention.

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. 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.

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