



NEWS RELEASE

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For Immediate Release:

Nymox NX-1207 Data Presented Today at American Urological Association Meeting in Chicago

NX-1207 Treatment for BPH Enters Phase 3 Development

HASBROUCK HEIGHTS, NJ (September 25, 2008) Nymox Pharmaceutical Corporation (NASDAQ:NYMX) is pleased to announce that data about the progress of Nymox's drug development for NX-1207, the Company's investigational drug for benign prostatic hyperplasia (BPH), were presented this morning at the Annual Meeting of the North Central Section of the American Urological Association being held in Chicago. Neal D. Shore, MD, FACS, of Myrtle Beach, SC made the podium presentation. Dr. Shore is an independent clinical investigator who has participated in four of the NX-1207 clinical trials as well as six follow-up studies of the drug. Dr. Shore serves as an Editorial Consultant for *Urology Times*. NX-1207 is now in Phase 3 development.

Dr. Shore's presentation provided an overview of the clinical trial results to date showing the safety and efficacy of NX-1207 in the treatment of BPH, including data from the recently completed Phase 2 clinical trial. The presentation also reviewed the extensive pre-clinical animal studies of NX-1207, including histopathological studies showing evidence of widespread prostate cell loss one year after a single intraprostatic injection of NX-1207. Reducing the size of the prostate is known to provide symptomatic relief for men suffering from BPH as well as positive long-term healthcare outcomes.

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.

Results of 6 follow-up studies of available subjects from NX-1207 clinical trials have provided evidence of durable benefits from NX-1207 treatment for up to 4½ years from the date of treatment. The Company recently announced statistically significant improvement compared to placebo in a 22 to 33 month follow-up study of 93 patients treated with NX-1207 at 17 U.S. clinical trial sites. Results in that study showed that patients at follow-up without any other treatment for BPH had a mean of 11.3 points BPH Symptom Score reduction, which represents a 47% improvement in symptoms from before 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.

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