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NEWS RELEASE

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For Immediate Release:

Nymox Announces Positive Immunogenicity Results for NX-1207 BPH Program

HASBROUCK HEIGHTS, NJ (February 19, 2013) Nymox Pharmaceutical Corporation (NASDAQ: NYMX) is pleased to announce its lead drug candidate, NX-1207, has not shown any evidence of eliciting an immune reaction in men treated with intraprostatic injections of the drug. Extensive clinical immunogenicity testing of men in the Company's pivotal Phase 3 trials (NX02-0017 and NX02-0018) and Phase 3 repeat injection safety study (NX02-0020) showed no evidence of antidrug antibody formation. Periodic safety monitoring reviews of the NX-1207 trials to date have shown no evidence of any allergic reaction to the drug either on first injection or repeat injection.

Positive immunogenicity testing of NX-1207 is an important step in the Company's drug development program. The results are consistent with the pharmacological and pharmacokinetic profile of the drug and augment the extensive safety experience with the drug in men with over 1,000 men having received the drug to date.

The Company will present more detailed scientific data from these studies at upcoming medical conferences. NX-1207 is in late stage Phase 3 development in the U.S. for the treatment of benign prostatic hyperplasia (BPH), a common condition of older men associated with growth in prostate size as men age. The Company's NX02-0017 pivotal Phase 3 trial has completed enrollment. Phase 3 trial activities of NX-1207 for BPH have begun in Europe sponsored by Recordati S.p.A., the company's European licensing partner. In the BPH studies to date, a single dose of NX-1207 has been found to produce symptomatic improvements about double that reported for currently approved BPH drugs without causing the sexual or cardiovascular side effects associated with those drugs.

NX-1207 is also being evaluated for the treatment of low risk localized prostate cancer where NX-1207 is administered directly into the area of the prostate where the cancer was detected. A U.S. Phase 2 study (NX03-0040) for that indication is in progress.

BPH causes progressive difficulties with urination, such as nocturia, urge to void frequently, acute urinary retention and other problems. The condition can seriously impact the health and quality of life of middle aged and older men. It is estimated that 50% of men in their 50s have pathological signs of prostatic hyperplasia and a high proportion of men as they age suffer from moderate to severe urinary problems and symptoms associated with BPH.

More information about Nymox is available at www.nymox.com <http://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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