Nymox Announces Prostate Cancer Clinical Trial Will Complete Prospective 18 Month Post-Treatment Assessments

Nymox Pharmaceutical Corp. {NASDAQ: NYMX} have announced the imminent completion of the 18 month trial for prostrate cancer NX03-0040.

Results are expected this quarter.

HASBROUCK HEIGHTS, N.J., Oct. 08, 2015 – Nymox Pharmaceutical Corp. {NASDAQ: NYMX} announced today that that participants in prostate cancer clinical trial NX03-0040 had neared completion of the study's 18 month post-treatment assessments. The results from this 18 month study are expected to be reported this quarter, after the 18 month data has been analyzed. Patients in the prostate cancer study have been followed for up to 39 months after treatment.

Study NX03-0040 is a Phase 2 study of NX-1207 for low grade localized prostate cancer. The study was initiated in 2012. 146 men were randomized to a single injection of NX-1207 at two dosage levels (2.5 mg or 15 mg) or standard of care.

Prostate cancer is the most commonly diagnosed cancer in men, other than skin cancer, and is the second leading cause of cancer death for men. Approximately 50% of prostate cancers are initially considered low risk.

To date, NX-1207 has had an excellent safety profile. NX-1207 has shown safety in 9 clinical trials (BPH and prostate cancer) including repeat injection studies. The drug does not lead to immune responses such as antibody formation which can cause significant drug toxicity and/or limit usage to single treatments due to drug neutralizing effects.

One of the major problems with current prostate treatments for localized prostate cancer (radical prostatectomy, external beam radiation, or brachytherapy) is the relatively high incidence of reported sexual dysfunction post-treatment. In 9 studies, NX-1207 treatment has been shown to have no significant adverse effect post-treatment on sexual function or testosterone levels.

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Forward Looking Statements

To the extent that statements contained in this press release are not descriptions of historical facts regarding Nymox, they are forward-looking statements reflecting the current beliefs and expectations of management made pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995, including statements regarding the need for new

options to treat BPH, the potential of NX-1207 to treat BPH and the estimated timing of further developments for NX-1207. Such forward-looking statements involve substantial risks and uncertainties that could cause our clinical development program, future results, performance or achievements to differ significantly from those expressed or implied by the forward-looking statements. Such risks and uncertainties include, among others, the uncertainties inherent in the clinical drug development process, including the regulatory approval process, the timing of Nymox's regulatory filings, Nymox's substantial dependence on NX-1207, Nymox's commercialization plans and efforts and other matters that could affect the availability or commercial potential of NX-1207. Nymox undertakes no obligation to update or revise any forward-looking statements. For a further description of the risks and uncertainties that could cause actual results to differ from those expressed in these forward-looking statements, as well as risks relating to the business of Nymox in general, see Nymox's current and future reports filed with the U.S. Securities and Exchange Commission, including its Annual Report on Form 20-F for the year ended December 31, 2014, and its Quarterly Reports.

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