NYMOX releases AGM uodate

Nymox Pharmaceutical Corp. {NASDAQ:NYMX} updated shareholders at their Annual General Meeting, held in Nassau, Bahamas.

Nassau is the new registered office of the company.

Nymox Provides Update From Shareholders Annual General Meeting December 16

HASBROUCK HEIGHTS, NJ (December 17, 2015) **Nymox Pharmaceutical Corp. {NASDAQ:NYMX}** is pleased to report an update on the Corporation's business as of December 16, 2015.

At the Company's AGM in Nassau, the following highlights were presented:

• The Company is moving forward with the activities of preparing for filing for approval of fexapotide triflutate (NX-1207) 2.5 mg for the treatment of prostate enlargement (BPH). The Company is pleased with the solid progress that has been made. The date(s) of anticipated filing will be reported in due course.

Dr. Paul Averback, CEO said "*Our management team has over the past several months been very focused on taking the necessary steps that will allow our Company to file for regulatory approvals for fexapotide triflutate in various markets. We*

expect to provide an update in the near future with regard to the filing for approval for our BPH drug fexapotide triflutate.”

Phase 3 trial results for fexapotide showed excellent safety profile and after 3.5 years statistically significant increased improvement in symptom scores vs placebo; statistically significant increases in responder rates vs placebo; statistically significant decrease (after 2 years) in need for surgery vs placebo; and statistically significant better nocturia outcomes vs placebo. Fexapotide is associated with a decrease in prostate cancer risk.

By comparison, some commonly used approved BPH treatments have been linked to increased cancer risk. Fexapotide has enhanced compliance and patient convenience compared to oral medications. Fexapotide is given as a single painless office treatment injectable. Approved oral medications generally involve daily pills intended for the rest of the patient’s life.

• The 18 month low grade localized prostate cancer trial of fexapotide 2.5 mg and 15 mg has been completed, and results are nearing completion of analysis. The timing of reporting of results is on target and top-line data will be presented in the very near future. In corporate developments, Richard Cutler of Houston, TX was appointed to the Board of Directors. Richard Cutler is a graduate of Brigham Young University and Columbia University School of Law. Mr. Cutler has worked at several major national law firms, and in 1996, formed Cutler Law Group in Newport Beach, California and subsequently Atlanta, Georgia and Houston, Texas, a firm which specializes in corporate and securities law, as well as international

business transactions.

Dr. Averback, Randall Lanham, Professor David Morse, and James G. Robinson were re-appointed to the Board of Directors. Paul McDonald has retired from the Board. The Company gratefully acknowledges Mr. McDonald’s steadfast, popular and highly valued solid service on the Board of Directors since 2006.

With reference to the Company’s quarterly financial filings, the international clients of Cutler & Co. have been acquired by transfer of responsibility to Thayer & O’Neal of Houston, Texas. Due to this recent transfer the Company’s quarterly filings have been slightly delayed and will be filed in the very near term.

For more information please contact info@nymox.com

NEWS RELEASE 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 and prostate cancer, the potential of fexapotide to treat BPH and prostate cancer and the estimated timing of further developments for fexapotide. 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 fexapotide, Nymox's commercialization plans and efforts and other matters that could affect the availability or commercial potential of fexapotide. 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.

Contact:

Paul Averback Nymox Pharmaceutical Corporation

www.nymox.com