NEWS RELEASE

For Further Information Contact:
Erik Danielsen
Nymox Pharmaceutical Corporation
1-800-93NYMOX
www.nymox.com

For Immediate Release:

Nymox Announces Important New Patent Developments

HASBROUCK HEIGHTS, NJ (April 21, 2020) Nymox Pharmaceutical Corporation (NASDAQ: NYMX) is pleased to announce the recent allowances of 4 new different US and international patents concerning the Company's prostate enlargement and prostate cancer treatments. The new US and international patents that have been allowed are further expansions of Nymox's intellectual property covering the Company's prostate treatment technologies.

Nymox CEO, Dr Paul Averback stated, “Intellectual property is one of the backbones of the biopharmaceutical industry. We are therefore very pleased to report that another 4 patents have recently been allowed. During the past year a total of 10 new patents have been allowed which adds to the Company’s very strong intellectual property position for the future. Nymox continues with intensive work to maximize our IP portfolio for the long-term value and strength of the Company's proprietary foundations.”

Dr. Averback added, “Operationally at the present time, our employees are doing very well and are following the prescribed guidelines in their area, including working remotely, where necessary. On a different note, we are also extremely pleased with the reactions and feedback we are receiving from multiple sources with regard to the recent (Feb 2020) peer review journal article in the *World Journal of Urology (*https://link.springer.com/article/10.1007/s00345-020-03127-w) documenting the very promising clinical results of Fexapotide Triflutate in treating men with early prostate cancer. We look forward to advancing this program into pivotal registrational trials.”

The recent peer review publication is entitled "Prospective Evaluation of Fexapotide Triflutate Injection Treatment of Grade Group 1 Prostate Cancer: Four Year Results”. The authors are Neal Shore, Myrtle Beach, SC; Steven A. Kaplan, New York, NY; Ronald Tutrone, Baltimore, MD; Richard Levin, Towson, MD; James Bailen, Louisville, KY; Alan Hay, Salem, OR; Susan Kalota, Tucson, AZ; Mohamed Bidair, San Diego, CA; Sheldon Freedman, Las Vegas, NV; Kenneth Goldberg, Lewisville, TX; Frederick Snoy, Albuquerque, NM; Jonathan I. Epstein, Baltimore, MD.

The Fexapotide (FT) prostate cancer study was started in 2012 and enrolled 147 men with localized Gleason Grade 6 T1c prostate cancer at 28 U.S. clinical investigation sites. Patients were followed with clinical and laboratory evaluations and regular periodic prostate biopsies for up to 5 years. Patients randomized to FT were treated with a single one-time targeted injection of FT, either 2.5mg or 15mg. Statistical comparisons were made over time of the proportions of subjects and untreated controls who progressed to higher Gleason grade and/or invasive treatments instituted with prostatectomy, radiotherapy, or chemotherapy. Important clinical highlights from the study include:

- FT treatment reduced cancer progression (-67.7%) compared to controls (3 years, FT 15mg, $p<.02$, pooled FT $p=.0265$).

- FT treatment group had reduced (-54.7%) incidence of surgery, radiotherapy or chemotherapy (4 years, FT 15mg p<.02; pooled FT p=.0374).
- At 4 years the incidence of surgery, radiotherapy or chemotherapy with increased Gleason grade was significantly reduced (FT 15mg -73.3% p=.0059, pooled FT p=.0064).
- Results for the high dose (FT 15mg) were superior to the lower dose (FT 2.5mg). Safety data showed no serious adverse events related to FT during the study.

FT is a pro-apoptotic proprietary drug which promotes natural programmed cell death (apoptosis) in prostatic glandular cells which compose the prostate cancer. FT has been safely administered to men in clinical trials in the U.S. involving over 1700 patients and controls treated for BPH or prostate cancer. FT has completed Phase 3 studies for BPH and further studies of FT for prostate cancer are planned in the near future.

For more information please contact info@nymox.com or 800-936-9669.

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, 2019, and its Quarterly Reports.