NYMOX announces PH3 studies for NX-1207

NYMOX Pharmaceutical (NASDAQ: NYMX} rather surprisingly announced today that despite previous setbacks, they are continuing with the phase 3 trial from their prostrate enlargement drug NX-1207.

CEO Paul Averback commented that management are “still confident that NX-1207 has long term potential in the treatment of BPH”.

NEWS RELEASE

Nymox Announces Phase 3 BPH Studies

HASBROUCK HEIGHTS, NJ – **Nymox Pharmaceutical Corporation {NASDAQ: NYMX}** announced today that the Company is undertaking further analyses of its pivotal U.S. Phase 3 studies of NX-1207 for prostate enlargement (BPH).

This will include new long-term data from Studies NX02-0017 and NX02-0018. The
Company expects to provide these new pivotal Phase 3 study results in Q2 or early Q3 this year.

The pivotal U.S. studies NX02-0017 and NX02-0018 were initiated in 2009. Enrollment of NX02-0017 (499 patients randomized) was completed in 2012; enrollment of NX02-0018 (498 patients randomized) was completed in 2013. 973 patients were injected with either NX-1207 2.5 mg (n=582) or saline vehicle alone as control (n=391). At 12 months post-treatment there was no overall top-line statistical significance for the efficacy of treatment in terms of BPH Symptom Score improvement vs controls. The safety profile of NX-1207 was excellent.

Dr. Paul Averback, CEO of Nymox said, “Despite the setback of top-line results not initially beating controls statistically at 12 months post-treatment in these large studies, we continue to believe that NX-1207 has enormous potential for long-term management of BPH. Additional new blinded protocol data from the same pivotal studies is being prospectively captured in order to assess long-term results in patients up to 5 years after
a single injection of NX-1207 2.5 mg vs placebo.”

NX-1207 is also in late-stage development for low grade localized prostate cancer. In 2014 the Company reported 8 month efficacy results showing statistically significant reduced cancer progression in patients who received NX-1207 compared to standard of care.

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