**BPN14770**, also known as *zatolmilast*, is a selective PDE4D negative allosteric modulator being developed for the treatment of FXS, Alzheimer’s disease, and other cognitive disorders.

A previous Phase 2a study in FXS (adults males, n=30) comparing BPN14770 25 mg to placebo dosed twice daily showed:

- **Statistically significant changes** in performance-based and parent-rated scales on measures of communication and language
- **A consistent advantage across diverse clinical domains** (including cognitive and behavioral) on parent-rated, clinician-rated and performance measures
- BPN14770 was well-tolerated, with no serious or severe adverse events related to treatment.

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**TETRA’S CURRENT FXS CLINICAL PROGRAM**

**BPN14770-CNS-204**
- Adolescent males 12 to < 18 years
- Double-blind
- Randomized 2 active : 1 placebo
- 13 weeks treatment

**BPN14770-CNS-301**
- Adult males 18-45 years
- Double-blind
- Randomized 2 active : 1 placebo
- 13 weeks treatment

**BPN14770-CNS-302**
- 52-week open label extension
- Subjects who complete -204 or -301 through Week 13

**Primary Eligibility Criteria**
- FXS with a molecular genetic confirmation FMR1 >200 CGG repetitions.
- Current treatment with < 3 psychotropic medications. Anti-epileptics don’t count as psychotropic if used for treatment of seizures.
- Permitted psychotropic must be at a stable dose for at least 4 weeks prior to screening and must remain stable.
- Anti-epileptic medications must be at a stable dose for at least 4 weeks prior to screening and must remain stable.
- Subjects with a history of seizure currently receiving treatment must be seizure-free for 3 months prior to screening, or 2 years if not on treatment.
- Behavioral and non-pharmacological treatments must be stable for 4 weeks prior to screening and must remain stable.
- Subject must be willing to practice barrier methods of contraception or be abstinent.
- Must have consistent caregiver who can attend all visits.

**Overview of Visit Schedules**

**BPN14770-CNS-204/-301 Double-Blind Studies**
- Screening (14 to 28 days before Day 1)
- Day 1 - Baseline assessments
- Week 1 - Telephone call
- Week 4 – Safety assessments
- Week 8 – Safety and Efficacy assessments
- Week 10 – Safety assessments
- Week 13 – Safety and Efficacy assessments
- Weeks 16, 20, 30, 42, 48 – Safety assessments
- Weeks 24, 36, 52 – Safety and Efficacy assessments
- Week 54 – Follow up telephone call

**BPN14770-CNS-302 Open Label Extension**
- Day 1 (same as Week 13 of -204/-301)
- Week 1 - Telephone call
- Week 2, 6, 9 – Safety assessments
- Week 12 Safety and Efficacy assessments
- Weeks 16, 20, 30, 42, 48 – Safety assessments
- Weeks 24, 36, 52 – Safety and Efficacy assessments
- Week 54 – Follow up telephone call
Screening and Day 1 visits and post-treatment visits to assess both safety and efficacy are expected to last 4-6 hours. Visits with only safety assessments are expected to last about 2 hours.

Other than routine blood draws, there are no invasive procedures or scans in these studies. Subjects will complete an IQ test at screening and a portion of the same test at later visits; subjects will also complete cognitive assessments on an iPad.

Caregivers will be asked to assess overall improvement, as well as specific improvement in daily living, language, academic skills (adolescents only), and emotions/behaviors; caregivers will also be asked to complete the Vineland-3, ABC and ADAMS assessment tools.

As clinical sites are activated in each trial, they will be listed on ClinicalTrials.gov. Visit Tetra’s website to find direct links to each study in ClinicalTrials.gov.

www.tetratherapeutics.com