**Non-Arteritic Anterior Ischemic Optic Neuropathy**

**About the Study:**
Sponsored by Quark Pharmaceuticals, the purpose of this study, called QRK, is to test whether an investigational drug called QPI-1007 helps prevent loss of visual acuity, in participants with Non-Arteritic Anterior Ischemic Optic Neuropathy (NAION). This study will assess the safety and tolerability of QPI-1007.

All participants will be randomized to one of five study groups:

- **Group A**—One 1.5 mg of QPI-1007 administered by intravitreal injection followed by 2 sham injections
- **Group B**—One 3 mg of QPI-1007 administered by intravitreal injection followed by 2 sham injections
- **Group C**—1.5 mg of QPI-1007 administered by intravitreal injection every 2 months, 3 injections total
- **Group D**—3 mg of QPI-1007 administered by intravitreal injection every 2 months, 3 injections total
- **Group E**—3 Sham injections

Participants participating in this study will make up to 8 scheduled study visits over a period of 12 months. Each visit will last anywhere from 2-5 hours.

**Key Inclusion Criteria:**
- NAION in the study eye, with symptom onset within 14 days
- Age 50-80 years

**Key Exclusion Criteria:**
- Present use or history or any treatment for the current episode of NAION
- Prior episode of NAION in the study eye only
- Bilateral (simultaneous) NAION
- Macular disease, proliferative diabetic retinopathy or other eye disease limiting visual acuity in study eye only
- Pain on eye movement in either eye
- Glaucoma or OHT in the study eye

**For Additional Information or To Refer a Patient, Contact:**
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