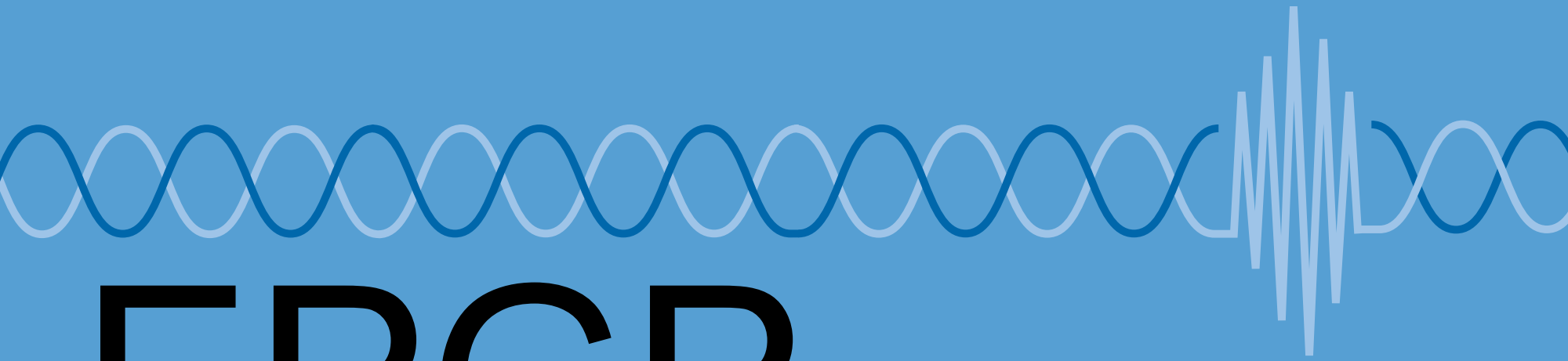

EPGP

EPILEPSY PHENOME/GENOME PROJECT

A consortium of epilepsy centers and the NINDS working to identify genes that influence the development of epilepsy and pharmacoresponsiveness.

Questions? Referrals?

www.epgp.org

info@epgp.org

(888) 279-EPGP

IGE / LRE Pairs

- ☑ Diagnosis of idiopathic generalized epilepsy (IGE) or localization-related epilepsy (LRE)
- ☑ Full-blooded **sibling, parent, or child** with idiopathic IGE or LRE (do not need to be concordant)
- ☒ Siblings are not identical twins (fraternal OK)
- ☒ No head injury, infection, stroke, other *acquired* causes, or *known* genetic mutation associated with epilepsy
- ☒ No significant developmental delay prior to seizure onset

Triads

- ☑ Diagnosis of cryptogenic:
  - infantile spasms
  - Lennox-Gastaut Syndrome
  - polymicrogyria
  - periventricular heterotopia
- ☑ Both biological **parents** willing to participate
- ☒ Neither parent has epilepsy
- ☒ No *known* genetic mutation associated with epilepsy
- ☒ No history of head trauma, hypoxic encephalopathy, meningitis/encephalitis, or other *acquired* causes

Sponsored by the National Institute of Neurological Disorders and Stroke