

Protocol for asymptomatic CSF1R mutation carrier

The first step after receiving your positive CSF1R mutation is to contact Sister's Hope Foundation. They will help connect you to a neurologist that has experience with ALSP. This neurologist will work with a local neurologist, which they will help you find, to manage your screening.

Items to be monitored at least annually:

- 1) Neurological exam
- 2) MRI
- 3) CBC and Metabolic Panel - Bloodwork
- 4) Cognitive/Behavior assessment
 - a) MOCA
 - b) Frontal Behavioral Inventory (FBI)
- 5) If symptoms appear: local neurologist to determine if findings are related to ALSP

*Sisters' Hope Foundation and the ALSP workgroup will update these recommendations as necessary.