

Part D – National ALS Registry Data only request

The National ALS Registry collects a variety of risk factor data.

Please select from the options below (mark all that apply). Note that not all survey data may be available at this time.

Demographics

Occupational History

Military History

Smoking/Alcohol History

Physical Activity

Disease Progression (ALSFRS-R)

Family History of Neurological Diseases

Clinical Data (e.g. devices used, body onset)

Lifetime Residential History

Lifetime Occupational History

Residential Pesticide Use

Hobbies with Toxicant Exposures

Caffeine Consumption

Reproductive History (women)

Health Insurance Status

Trauma History