



# ALS ROUNDTABLE PROGRAM IMPACTS AND OUTCOMES

## Spring

## Fall

2019

### Setting ALS Association Strategic Priorities

- Key input into [strategic plan](#)
- Topic ideas for future Roundtables and [ALS Focus](#) surveys

### Access to New Therapies

- ALS Focus survey: [Understanding Insurance Needs and Financial Burdens](#)
- Partnership with [Patient Advocate Foundation](#)
- Strategic engagement with the FDA

2020

### Reducing Time to Diagnosis

- [thinkALS](#) tool
- [Educational webinar](#)
- [Consensus statement](#)
- Genetic counseling and testing education campaign

### Decreasing ALS Complications

- Complication tracking for people living with ALS
- [Quality of Care Research Awards](#)

2021

### Access to Essential Care, Services and Supports

- [Patient journey web tool](#)

### Strategic Priorities for ALS

- [Trial Capacity Awards](#)

2022

### Developing Action Plans to Advance Inclusion in ALS Diagnosis, Care and Research

- Refinement of next steps during [Spring 2023 Roundtable](#)

### Increasing the Number of, and Access to High-Quality Clinical Trials in ALS

- Refinement of next steps during [Spring 2023 Roundtable](#)