Mission Statement: To discover treatments and a cure for ALS, and to serve, advocate for, and empower people affected by ALS to live their lives to the fullest.

Care Services Programs:

- Home visits for people diagnosed with ALS and their family members to assist with access to local, state and federal resources. Telephone consultation can also be requested.
- Educational materials on a variety of topics including booklets for children, teens and young adults
- *(Virtual)* Support and Resource Groups including educational presentations on a variety of topics along with networking opportunities
- Multidisciplinary ALS Clinical Care Centers in ME, NH and VT, access to clinical trials
- Adaptive Equipment Loan Program (Durable Medical Equipment & Communication tools)
- Annual Education Symposium
- Caregiver Lab Program
- Hope Seeley Beyond Diagnosed Program
- Hope & Healing Bereavement Program to support family members
- In-services for professional caregivers
- Paul LaRochelle Quality of Life Grant
- Informational webinars on a variety of topics
- Support of ALS advocacy efforts with federal legislators

*To contact our care services staff directly call **603-226-8855** or email care.services@alsanne.org*

Upcoming Events

2022 Walks to Defeat ALS

Bangor, Maine: Saturday, August 27
Portland, Maine: Saturday, September 10
Vermont: Saturday, September 24
New Hampshire: Saturday, October 10

Please contact Karrie Boskee for Walk Information: karrie.boskee@als.org