

**The ALS Network**, then ALS Golden West, provides people with ALS and their loved ones compassionate and professional care services, free-of-charge. We offer information, support, and resources proven to increase longevity, enhance quality of life, and help with the challenges of living with ALS.



The ALS Network collaborated with a group of neurologists, researchers, people living with ALS, and their caregivers from around the country, to **launch a baseline study to establish practice parameters and guidelines for in-home ALS care.**

442



**442 people newly diagnosed with ALS connected with the ALS Network** to access our evidence-based wraparound model of care.

31



**Collaborated with 31 ALS clinic partners**, including Certified Treatment Centers of Excellence and Recognized Treatment Centers, to seamlessly provide quality, multidisciplinary care.



In a 2023 survey, **85% of people** with ALS and their caregivers indicated that they **found up-to-date information and resources** first and foremost **through the ALS Network.**

1,763

attendees



The ALS Network **held more than 226 online connection groups** with **1,763 participants.**

7,300+



Since the launch of our “ASK ME” educational webinars, there have been **more than 7,300 views** of the live broadcasts and on-demand recordings.

200



**200 military personnel and veterans with ALS**, as well as their families, received access to vital care services and support offered by the ALS Network.

2,100+



Directly helped clients through **more than 2,172 consultations related to loans of durable medical and communication equipment** and provided **more than \$200K in direct assistance** for equipment needs, transportation support, emergency assistance, and respite care.

2,000



16 care managers **served nearly 2,000 people with ALS** and their loved ones throughout California and Hawaii.



The number of families facing ALS who seek our care services continues to grow every year. On average, the **ALS Network is serving more than 100% of newly diagnosed people with ALS across our service area.**



**95% of 2023 survey respondents** reported being **satisfied/very satisfied with the services and support** provided by their ALS Network care manager.



We offered **unique connection groups**, including meetings for caregivers, for people with ALS/FTD, those interested in adaptive yoga, those seeking Spanish language groups, veterans, LGBTQ+, and newly diagnosed.



37,900+

**Over 37,900 people** viewed our care services webpages or care services-related email communications from the ALS Network.



The ALS Network joined the **Advisory Board for the National Palliative Care working group**, composed of leaders in the ALS field.

**Your support makes a difference. It is only together that we will end ALS.**