



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**



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.

442



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



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

31



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



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.**



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.**



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

1,763 attendees



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



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.

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.



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

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.



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.