A Commitment to those Living with ALS:

The ALS Association Rated with High Marks Among Top Charity Watchdogs

Thank you for the outpouring of support.

#ALSiceBucketChallenge

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JGP Wealth Management Group of Wells Fargo Advisors

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We are grateful for our dedicated Board of Directors. These community volunteers generously donate their time, treasure and talents to provide strategic direction and oversight to our chapter.
In large part by the Ice Bucket Challenge, our chapter is thrilled to increase its capacity and its care services to continue to meet the needs of local families living with ALS. Last year, alone, our qualified staff of 10 professionals provided hands-on support and tangible benefits to 410 people living with ALS along with their family members, friends, neighbors and co-workers.

Lance Christian

On behalf of The ALS Association Oregon and SW Washington Chapter, thank you again for taking the Ice Bucket Challenge for ALS last summer, which raised unprecedented levels of awareness and support for the critical fight against this relentless disease. Not only did your participation in this effort allow our organization to triple its commitment to ALS research, it also allowed our local chapter to:

- **Restock** our durable medical equipment, and augmentative and alternative communication loan closets.
- **Add** more home-based assistive technology services across the chapter so that people with ALS do not lose their ability to use computers, communicate and direct their own lives.
- **Create** new programming for children, teens, and young adults with a parent or grandparent with ALS, recognizing that is in not just the person with ALS, but the whole family, who needs support.
- **Increase** our respite and in-home care assistance programs by 30% - adding over 2,500 hours of paid professional care giving hours provided to local ALS families.
- **Start** the effort to provide mental health and counseling services to support people with ALS and their families.
- **Designate** substantial funds for immediate release to fund local ALS research that will lead to a treatment for ALS.
We launched new programs and partnerships to support the children and teens in families living with ALS, supported The Dougy Center in launching a new Pathways Program for children with a parent who has a life-shortening illness, and created an ALS Family Day at the Oregon Zoo where adults gathered for education on communicating with kids and teens about ALS, and children were able to have fun with others who are living with ALS in their families.

This year, our staff:

- averaged 9 home visits per week— 450 in all, distributed respite care grants to 75 families,
- funded 6,500 hours of professional in-home caregiving,
- provided 180 families with durable medical equipment and 150 families with communication-enabling equipment, hosted 132 monthly support groups,
- sponsored 16 local healthcare professionals and staff in attending The National ALS Clinical Conference, where they networked with over 400 other professionals from across the nation, shared best practices and were presented with new and current ways to treat and care for people living with ALS, developed a relationship with Asante to create an ALS multi-disciplinary clinic to serve families in Southern Oregon, and strengthened our support in Multi-Disciplinary Clinic partnerships with The Providence ALS Center, Kaiser Permanente, the Veterans Administration and St. Charles Medical Center.
- A sampling of local care services provided free of charge to people and families living with ALS:

  - On-staff Care Services Coordinators who offer individual support and home visits to assess the needs of those living with ALS, suggest equipment and help families plan for the future.
  - A Medical Equipment Loan Program that shares equipment to aid in daily living activities and with mobility when insurance and other programs cannot fund needed items.
  - A comprehensive Augmentative Communication and Assistive Technology Program to support people living with ALS in their communication and technological needs.
  - Caregiver Support Program to relieve stress and “burnout” which includes family caregiver support groups, Care Connections guidance, In-Home Caregiving financial support and Respite Care grants.
  - Monthly Support Group Program for people living with ALS, family members, caregivers and friends.
  - Support and resources for children who have ALS in their families.
  - Bereavement support for family members and friends who have lost a loved one to ALS.
  - Partnerships in ALS multidisciplinary centers and clinics staffed by professionals with ALS expertise.
  - Outreach to the health care community, home health care agencies and the community with updates on current ALS needs and solutions.

We sponsored a delegation, including Rob Chadwick, who is living with ALS in Scotts Mills, Oregon, and the Feldhaus family, who is living with ALS in Hillsboro, Oregon, to attend the National ALS Advocacy Day in Washington DC to lobby for ALS research.
Program Services Break Down

Outreach and Advocacy: advocacy efforts in Washington DC, Defeat ALS awareness events, National ALS Registry, and research funding.

Independence Programs: Medical Equipment and Assistive Technology Loan Programs

Education and Support Programs: Home visits, monthly support groups, and professional education to community partners

Caregiver Support Programs: In-Home Caregiving program, Respite Care program and Bereavement program.

ALS Clinics & Certified Center Support: partnerships with Providence ALS Center as well as recognized ALS treatment clinics with The Portland Veteran's Administration Hospital, Kaiser Sunnyside Medical Center, and St. Charles Medical Center.

Statement of Activities
Year ended January 31, 2015 (with comparative totals as of January 31, 2014)

<table>
<thead>
<tr>
<th></th>
<th>Unrestricted</th>
<th>Temporarily Restricted</th>
<th>Total</th>
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<tr>
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<td>$ 1,716,068</td>
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<td>Contributions</td>
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<td>60,000</td>
<td>65,000</td>
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<td>Special event revenue, less direct benefit costs of $67,485 and $84,900, respectively</td>
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<td>(23,000)</td>
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<td>Total revenues released from restrictions</td>
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<td>50,529</td>
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<td>Expenses</td>
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<td>Program services</td>
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<td>Management and general</td>
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<td>567,199</td>
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<td>Net assets, end of year</td>
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<td>$ 50,529</td>
<td>$ 1,955,479</td>
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Statement of Financial Position
Year ended January 31, 2015 (with comparative totals as of January 31, 2014)

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<th>2014</th>
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<td>Revenue sharing refund receivable</td>
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<td>Grants and contributions receivable</td>
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<td>Total assets</td>
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<td>LIABILITIES AND NET ASSETS</td>
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<td>Accounts payable</td>
<td>$ 26,676</td>
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<td>Accrued payroll and related expenses</td>
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<td>Deferred revenue</td>
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<td>Total liabilities</td>
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<td>Unrestricted</td>
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<td>Temporarily restricted net assets</td>
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<tr>
<td>Total net assets</td>
<td>$ 1,955,479</td>
<td>$ 1,337,761</td>
</tr>
<tr>
<td>Total liabilities and net assets</td>
<td>$ 2,077,108</td>
<td>$ 1,591,105</td>
</tr>
</tbody>
</table>
Anonymous
Angeline Adler
Mary Beth Baker
In memory of my husband, Gary.
Jeff & Trina Bandelow
Anita & Dana Cadonau-Huseby
Lance Christian
Leonard Davis*
Barbara C. Deeming
Dr. Kimberly Goslin
Cynthia A. Greene
Joseph P. Gross
In loving memory of my mother, Helen Gross.
Alexis Halmy
Matson Haug & Julie Isaacson
Randy* & Debi Kyte
Michaels L. Lamont*
Al Lee
Lou & Jocelyn Libby
Brian & Aubrey McCauley
Doug & Claudia McClure
Ralph & Susan Nicholson
Fred Noble*
Steve* & Teri Sall
John & Barbara Seibert
Corliss Sherry
Nancy E. Shire
Laurie Speight*
Elisabeth Twist

Members of The Legacy Society have expressed their commitment to The ALS Association Oregon and SW Washington Chapter through a very special and important form of financial support. These donors have named The ALS Association as the ultimate beneficiary of a planned gift. Such gifts might include a bequest and/or charitable income gifts such as gift annuities, remainder trusts, or gifts of life insurance.

Your membership involves no dues, obligations, or solicitations, but it allows us to thank you and recognize you for the plans you have made, and it may inspire generosity in others. Benefits of membership in The Legacy Society also include invitations to special events and seminars and a subscription to our newsletter. The most important benefit, however, you will receive from joining The Legacy Society is the satisfaction derived from making a lasting contribution to our organization’s long-term strength.

If your estate plans include The ALS Association, please let us know by contacting Lance Christian, LMSW, Executive Director, at LanceChristian@alsa-or.org or 800.681.9851 ext 101. We would like to thank you for your generosity by including you in The Legacy Society.

* denotes those no longer with us
We were excited to introduce a new program this year, our Extra Mile Club. This club consists of participants who pledged to “Walk the Extra Mile” and raise $1,000 or more as an individual fundraiser by August 29th.

Thank you for your tremendous efforts this Walk season! If you are interested in joining the club next year, contact walk@alsa-or.org for more information.

**Cumulative Gifts Donated to Our Chapter in 2014 – 15**

<table>
<thead>
<tr>
<th>Our Hope</th>
<th>$25,000 – $99,999</th>
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<tbody>
<tr>
<td>Alpenrose Dairy</td>
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<td>Reser's Fine Foods, Inc</td>
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<td>Certified Languages International, LLC</td>
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<td>Les Schwab Warehouse Center, Inc.</td>
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<td>Ken and Kathy Fink</td>
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<td>Harold and Arlene Schnitzer CARE Foundation</td>
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<td>Joe Gross and Shelley Noyes</td>
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<td>Shirley Lefkowitz</td>
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<td></td>
<td>Mayer and Janet Schwartz Fund of The OCF</td>
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<td>Jim and Sally McAfee</td>
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<td>Mikatomi Holdings LLC</td>
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<td>Ray Pretti</td>
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<td></td>
<td>The Peterson Family Gift Fund</td>
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<td>Wells Fargo Advisors, LLC</td>
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<tr>
<td>Our Courage</td>
<td>$15,000 – $24,999</td>
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<td>John and Barb Deeming</td>
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<td>Helen John Foundation</td>
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<td>Kevin McGee and Gary Moe</td>
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<td></td>
<td>B.P., Lester and Regina John Foundation</td>
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<td></td>
<td>Leonard Dieterich</td>
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<td></td>
<td>Providence Brain and Spine Institute</td>
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<td></td>
<td>The Anjulicia Foundation and Hayden Homes</td>
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<tr>
<td>Our Strength</td>
<td>$10,000 – $14,999</td>
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<td>Nike Employee Matching Gift Program</td>
<td>Our Determination</td>
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<td>James Flynn</td>
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<td>Fred Meyer Stores</td>
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<td>John and Kathy Connors</td>
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<td>Lou and Jocelyn Libby</td>
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<td>Susie Oswald</td>
<td>Eberhard's Dairy Products</td>
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<tr>
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<td>OCF Joseph E. Weston Public Foundation</td>
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<td>Ken and Debby Willett</td>
<td>Our Resilience</td>
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<td>Jerry Nudelman</td>
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<td>Microsoft Matchings Gifts Program</td>
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<td>Kalan Morinaka Memorial Golf Tournament</td>
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<td>Teri Sall</td>
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<td>Battle Ground School District No. 119</td>
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<td></td>
<td>The Peterson Family Gift Fund</td>
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</tbody>
</table>

Many thanks to the following local business members who each raised $2,500 or more and worked together to raise over $111,000 towards our 2014 Walk to Defeat ALS®.

**Gifts**

<table>
<thead>
<tr>
<th>Gifts</th>
<th>Cumulative Gifts Donated to Our Chapter in 2014 – 15</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mary Beth Baker</td>
<td>Doug and Claudia McClure</td>
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<tr>
<td>Fred Meyer</td>
<td>Jerry Nudelman</td>
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<td>Doug Greenberg</td>
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<td>Hill, Row</td>
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<tr>
<td>Lance Christian</td>
<td>John and Peg Esple</td>
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<tr>
<td>Teri Sall</td>
<td>Shirley Lefkowitz</td>
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<td>Dennis and Ruth Young</td>
<td>Mayer and Janet Schwartz Fund of The OCF</td>
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<td>Battle Ground School District No. 119</td>
<td>Jim and Sally McAfee</td>
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<td>Mikatomi Holdings LLC</td>
<td>Ray Pretti</td>
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<tr>
<td>The Peterson Family Gift Fund</td>
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<tr>
<td>Wells Fargo Advisors, LLC</td>
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For more information, contact walk@alsa-or.org.
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Please accept our apologies for any inaccuracies and contact us at
800.681.9851 ext 105 or AudreeMcCaney@alsa-or.org. Only monetary donations given in this fiscal year have been listed. List is in order of total donations, not alphabetical.
We remember the courage, strength and remarkable spirits of those who lost their lives to ALS.

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Kathy Bennett
Dan Berry
Charles Brown
Georgette Brown
Lois Burns
Doug Byers
Kenneth Byrne
Nicholas Carter
Sue Case
Dolores Cassinelli
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Karen Chinn
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OUR VISION
Create a world without ALS.

OUR MISSION
Leading the fight to treat and cure ALS through global research and nationwide advocacy while also empowering people with Lou Gehrig’s Disease and their families to live fuller lives by providing them with compassionate care and support.

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Southern Oregon: 2305 Ashland St., Ste. C, Box #458, Ashland, Oregon 97520
Willamette Valley: PO Box 22653, Eugene, Oregon 97402

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