

WHAT SETS THE ALS ASSOCIATION APART

In the competitive community of ALS organizations, The ALS Association stands out as the world's preeminent leader in the fight against amyotrophic lateral sclerosis. ALSA's comprehensive and strategic approach to the disease is founded on the world's finest research enterprise, a national network of chapters that deliver high quality patient services and a highly effective advocacy program that influences change for the betterment of patients, families and caregivers. Because of ALSA's research stature, international influence and patient services network, chapters will experience contacts or overtures from other ALS organizations. Here is what sets ALSA apart from other ALS organizations:

Organizational

- The ALS Association is the only national, not-for-profit voluntary health organization dedicated solely to the fight against ALS through research, patient services, advocacy and public awareness.

Research

- The ALS Association leads the world's premier ALS research enterprise with the largest effort in scope and diversity of projects ever undertaken to unlock the mysteries of ALS.
- The ALS Association directs the largest private research effort into ALS, with a \$25 million investment in research during the last decade - more research funding than any other ALS organization.
- The ALS Association's research program focuses on achieving scientific breakthroughs that will improve the lives of people living with ALS today while searching for a cure for the disease.
- The ALS Association's research program seeks to understand the causes of the disease to someday be able to prevent or eradicate ALS.
- The ALS Association's research program focuses on assay and drug development as a part of its portfolio.
- The ALS Association's research program adheres to the highest scientific standards.

Continued on page 2



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info@alsa-midwest.org with "Dialog" in the subject line.

Include your name, mailing address and your email address.

Continued from page 1 - What Sets The ALS Association Apart

- With a current inventory of approximately 100 funded projects, The ALS Association's research portfolio encompasses every credible hypothesis about the origination and development of the disease.
- Every research project funded by The ALS Association is peer reviewed and evaluated by prestigious scientific panels for credibility and relevance to ALS.
- The ALS Association is recognized by the National Institutes of Health and by scientists worldwide as the primary source of credible scientific information about ALS. NIH routinely consults with The ALS Association.
- The ALS Association is partnering on research into ALS with the National Institute of Neurological Disease and Stroke (NINDS) and the Institute of Environmental Health Sciences.
- The ALS Association is collaborating with the Centers for Disease Control, the U.S. Department of Defense and the Department of Veterans Affairs on research into specific avenues that hold potential in ALS research.
- The ALS Association is the only ALS organization that administers the only post doctoral fellowship for ALS research - ensuring that young scientists join the effort to investigate ALS with fresh insight.
- The ALS Association is the only ALS organization funding clinical management research projects aimed at improving the lives of people with ALS.
- The ALS Association holds and sponsors numerous workshops, meetings and discussions that bring together scientists researching ALS and other neurodegenerative diseases to find commonalities and to generate new research suggestions and fresh insight.

Patient Services

- The ALS Association provides patients, families and caregivers with the most comprehensive and uniform array of services delivered through 39 chapters.
- People with ALS, their families and caregivers receive high quality and diverse services that encompass 12 core programs of excellence.
- Educational
- Authoritative information on research and clinical issues
- ALS-specific support groups
- Caregiver programs and services
- Respite care
- Equipment loan
- Certification of clinical programs with criteria and standards
- Home visits
- Case management
- Clinical management research program
- The ALS Association's educational services are more extensive than other ALS organizations and include the informative "Ask the Experts" series as well as other programs that enable the ALS community to interact with leading researchers, physicians and experts.
- The ALS Association establishes rigorous clinical criteria that set the standard of care for people with ALS
- The ALS Association's 19 certified ALSA Centers across the United States provide state-of-the-art care for people with ALS in a multidisciplinary setting in a supportive, family-oriented environment.
- Patient services staff are health care specialists who provide in-depth expertise to individuals and groups
- The ALS Association produces the widely-recognized "Living with ALS" series of manuals and videos that provide the ALS community with specialized printed and visual materials.

Kansas City Legends

Tom Watson, Len Dawson, Buck O'Neil and George Brett **Headline Joe McGuff ALS Golf Classic**

The ALS Association Keith Worthington Chapter is proud to announce Tom Watson, Len Dawson, Buck O'Neil and George Brett's participation in the 2nd Annual Joe McGuff ALS Golf Classic. The tournament will be held Monday, May 24th at the Nicklaus Golf Club at Lion's Gate in Overland Park, Kansas. For additional information about the Classic please contact Sarah Kerwin at 913.648.2062, ext. 223 or skerwin@alsa-midwest.org.



The 14th International Symposium on ALS/MND

November 17-19, 2003, Milan, Italy

The International Symposium on ALS/MND draws researchers and clinicians together from around the world to learn the newest in ALS research. Approximately 800 clinicians, scientists, allied professionals, people with ALS and their caregivers and members of ALS/MND organizations gathered for this event. It is the premier forum for both biomedical research and clinical investigation and practice. The Motor Neuron Disease Association of England, Wales and Northern Ireland sponsor the Symposium in cooperation with the International Alliance of ALS/MND Association. In 2004, The ALS Association will host the Symposium in Philadelphia.

Excerpts From International ALS/MND Symposium Report

ALS/MND - How do we find the cure?

Presenter: R. Brown Jr. (USA)

"The last decade has witnessed substantial progress in defining factors that contribute to the cause of ALS/MND. Much of the progress in recent years is a consequence of studies of (genetics) in ALS and related motor neuron diseases. There have also been major accomplishments in the epidemiology, biochemistry, immunology and virology of ALS. Taken together, these domains of investigation have provided new insights into (the cause of ALS). ...there have been remarkable technological developments that have facilitated mass-scale analyses of genes, proteins and small molecules in ALS/MND tissues, as well as high-throughput methods for testing new therapies."

Cognitive changes in ALS/MND: An Overview

Presenter: C. Lomen-Hoerth (USA)

Research at the University of California, San Francisco (an ALSA Certified Center) reveals a higher number of people with ALS who also have frontotemporal dementia (FTD) than has been previously reported. ALS with frontotemporal dementia is now being recognized as a sub set of ALS and includes a degeneration of the frontal lobe of the brain with atrophy.

Depressive disorders in late-stage ALS patients and family caregivers

Presenter: S. M. Albert (USA)

The investigators followed 56 patients and their caregivers during the final months of life and evaluated the presence and degree of depression. Findings showed that most patients with ALS were never depressed as they approached the end of their lives.

Intraspinal cord implantation of stem cells

Presenter: L. Mazzini (Italy)

The stem cells were taken from the patient's bone marrow, grown in laboratory culture and then implanted into the patient's spinal cord. Dr. Mazzini reported that the stem cells did develop into neurons and did migrate. Twenty-one months after implantation, there was a trend in some of the seven patients toward improved muscle strength in the lower limbs that was possibly associated with larger numbers of implanted stem cells. "It was not the aim of the study to determine the activity of (the stem cell) implantation in ALS patients; however the clinical results are encouraging." The many unanswered questions about how stem cells work and what promise they may hold for people with ALS raise questions about the advisability of human trials at this time with stem cells. There was no evidence presented that the implanted stem cells replaced diseased motor nerves in the seven people with ALS in this study.

Non-invasive ventilation in ALS

Presenter: N. Leigh (UK)

"Noninvasive ventilation is now an accepted treatment for patients with ALS who have respiratory insufficiency." While there are no published randomized clinical trials (this would require that NIV be withheld from patients in the control group), non-randomized studies have shown that use of NIV improves sleep and quality of life as well as increasing survival. However, studies of caregivers of people who use NIV suggest that the caregivers may have an increase in stress and depression.

Primary placement under radiologic guidance of Entristar™ skin level gastrostomy tubes

Presenter: A. S. Shaw (UK)

Nutritional support for people with ALS is an important consideration to prevent malnourishment, dehydration and further decline in condition. Research studies show that by providing food and fluids through a feeding tube nutrition can be maintained, and some study data suggest that survival is prolonged.

SOAR. BALANCE. HOPE. DREAM. IMAGINE...

This year, “Night of Hope” is flying into fantasy with Neil Goldberg’s *Cirque*. This amazing ensemble cast will exhaust your senses and imagination. Enter into our dreamscape of imagination and be left with a lasting impression of fun and fascination. It’s sure to be a night of delightful adventure, mesmerizing moments and limitless imagination. This magical performance will take place on stage at the Lyric Theatre on Saturday, June 19th. Presale tickets are available starting March 1st at www.alsa-midwest.org. General ticket sales begin March 15th through Ticketmaster. For more information, contact Emily Marsh at 913.648.2062, ext. 221 or emarsh@alsa-midwest.org.



Senate Bill 0800 ALS/Lou Gehrig Disease Fund

Summary: S.B.0800

This act creates a one-dollar check-off on the Missouri income tax return. The money designated by the check off will be deposited into the ALS Lou Gehrig's Disease Fund, to be distributed to the Missouri ALS Associations for the St. Louis and Kansas City regions. The check-off of one dollar is primarily for taxpayers who are to receive a refund. However, taxpayers who owe taxes may also contribute to the fund and any taxpayer may elect to contribute more than one dollar.

Sponsored by: Senator Sarah Steelman-Sponsor
Assigned to Committee: Financial and Governmental Organization, Veteran's Affairs and Elections
Testimony Heard: January 26, 2004
Committee Members:

Anita Yeckel, <i>Chairperson</i>	1 st	314-729-0541	or	573-751-2887
Chuck Gross, <i>Vice Chair</i>	23 rd	636-949-2323	or	573-751-8635
Matt Bartle	8 th	573-751-1464		
Jon Dolan	2 nd	573-751-4964		
Bill Foster	25 th	573-751-3859		
Delbert Scott	28 th	573-751-8793		
Mary Bland	9 th	816-444-1347	or	573-751-2770
Maida Coleman	5 th	573-751-2606		
Rita Days	14 th	573-751-4106		
Jim Mathewson	21 st	573-751-4771		

To find out additional information and/or to contact the above members of the committee via the Internet, follow the steps below:

1. Go to <http://www.state.mo.us>
2. Click on **Legislative Branch** v (a drop down screen appears)
3. Click on **Missouri Senate** from the drop down screen.
4. A new page appears with the heading **Missouri State Senate**.
5. Click on **Committees** (in green lettering).
6. Under the heading Standing Committees, click on **Financial and Governmental Organizations, Veterans Affairs and Elections-Assigned Bills**.
7. You will see the members of the Committee who will determine if S.B. 0800 gets out of committee and if it will be brought to vote. (You should see the above named members.)
8. On the bottom of this page you will also see the line **Senate Hearing Schedule**. If you click on this line and then scroll down the new page, you will see **S.B.800**, if you click on this line it will give full details of the bill if you are so interested.
9. Click on the committee members name and their office information including phone numbers, fax numbers, email line, etc. **Please email, call, fax, or send a letter asking for their support in bringing S.B. 0800 out of committee for a full vote.**

Thank you for your support!

Support Group News

Kansas City

Approximately 29 were in attendance at the February 2nd Kansas City support group meeting. The speaker was Mike Dorris with American Mobility Solutions who talked about conversion vans and cars, adaptation to motor vehicles, internal/external lifts, modular ramps, Blue Button Life Lines and more.

*** Due to timing and/or inclement weather cancellations, the Wichita, Hutchinson, Springfield, Omaha and Lincoln support group meeting reports were unavailable. Look for them in next month's issue ***

3rd Annual Bob Hohn Memorial Golf Classic for ALS

Bob Hohn's unconquerable courage and heroism while he confronted ALS will always be an inspiration to all who knew him. He lost his battle with ALS on November 27th, 2003. The golf tournament Bob championed and lent his name to will continue on in his memory.

The 3rd Annual Bob Hohn Memorial Golf Classic for ALS in Nebraska will feature Honorary Celebrity Gale Sayers, Nebraska sports legends and other hometown



celebrities just like Bob. The festivities begin Friday evening, June 25th, with an autograph signing session open to the general public. A Pairings Party (location to be determined) will follow the autograph session.

On Saturday, June 26th, teams will gather at the spectacular Wilderness Ridge Championship golf course for one of

two shotguns, morning or afternoon. Each team will consist of three players, plus a celebrity. At the end of play there will be an awards ceremony at the clubhouse.

The Bob Hohn ALS Foundation is committed to establishing an equipment loan closet and providing an income assistance program to help those with ALS. The programs and research funded by the ALS Association Keith Worthington Chapter are a vital link for patients and families to education, information and services to help them live with ALS. Together, both organizations make a difference in the lives of those living with ALS. This is a terrible and terrifying disease.

For more information about the Bob Hohn Memorial Golf Classic for ALS, contact Ric Miller in Omaha at 866-762-6361 or rmiller@alsa-midwest.org.

Help Network

For many of us, there are not enough hours in a day to complete the tasks we all face daily, weekly, monthly. This is especially true for families dealing with ALS. From minor jobs such as walking the dog, sweeping the garage or raking the yard to the larger more time-consuming projects such as painting a room, building a ramp or enlarging a doorway - these are things that need to be done.

Where can a person with ALS, a caregiver or family member get help when help is needed?

The answer is in our communities - friends, schools, neighborhoods, churches, businesses, civic groups and organizations. The answer is a "Help Network" - an idea being developed by some of our friends in Springfield, Missouri and our Chapter. With a little help from you, we are ready to make it a reality.

Here is the plan:

- "Help Network" volunteers will contact various resources in their community that might be able to lend a hand to our ALS families. Scout groups, church groups, high school community service clubs, local business community service coordinators, area volunteer agencies, and ALS survivors and friends are just a few of the many possible resources that will be contacted.
- A network of "Helpers" will be created and documented.
- The Chapter will publicize the "Help Network" program on our web site and in our newsletter and will provide a phone number for people to call when they need help.
- Those with ALS, their caregivers or family members will be able to call the number to get in touch with a "Help Network" volunteer, and be matched with a "Helper" or "Group of Helpers."

Right now "Help Network" volunteers are setting up the program in the Springfield area. If you would like to help them, or have another contact for them, please call Jim Barb at 417.582.0505 or email jbarb@centurytel.net.

If you are an individual with ALS, an ALS caregiver or family member in the Springfield area and you need help, please call Jim Barb at 417.582.0505 or email jbarb@centurytel.net.

If you are interested in starting a "Help Network" in your community, please call Sally Dwyer, Program Director, at 800.878.2062, ext. 212.

FOR SALE

HOME

For Sale By Owner - 2709 NE 78th St. Custom built wheelchair accessible home in upper class neighborhood. 5-BR, 3-Gar, 1 1/2 story on treed lot. Separate living area in finished basement for caregiver, etc. w/office. No-Step entrances oversized "roll-in" shower; whirlpool tub w/Hoyer lift, hardwood & ceramic tile floors. \$379,500. Call 816-436-3141.

VANS

1996 Ford Econoline with lift, extended roof. 50,000 miles. Call Joan @ 913-385-1259.

1997 Ford Pinnacle Motor Home Ford F53 Chassi V8 7.5 liter gas engine, double rear end axle, new batteries, good generator, 22,250 miles, 33 ft. long. Blue interior with queen size bed. Call Chuck after 6 p.m. @ 785-459-2281.

1998 Dodge Caravan 12,000 miles, kneeling option, wheelchair ramp, floor-lock wheelchair tie-down clamps, 7 passenger with seats, removable seats. Excellent condition. Asking \$25,000. Call 913-831-4977 or 913-789-7786.

1999 Ford F150 V8 Raised Roof Van w/ wheelchair tie downs. Remote for doors and lift, loaded. 39,000 miles. Call 913-287-7751 or 816-507-3728.

2000 Dodge Caravan IMS Rampvan 18,500 miles, fully loaded, lock down for wheelchair, still under warranty. Contact Jeaneene @ 417-466-2907, 417-466-1471 or by email @ GRANEENE1@aol.com.

2000 Chevy Conversion Van w/ Braun wheelchair lift (up to 800 lbs.), 13,100 miles, TV, VCR. 4 captains chairs, 2 removed to accommodate the wheelchair. Asking \$25,000 or best offer. Call Marilyn @ 785-389-3861.

WHEELCHAIRS

2002 Ranger X Electric Tilt Wheelchair joystick control, several speeds, neck support, carrier on back, barely used. Owner's manual included. Call 712-527-5475.

Quantum Jazzy 1400 Power Wheelchair 4 months old, great condition. \$3,000 or best offer. Call 417-865-2911.

Pride Jazzy Electric Wheelchair with 2 batteries and charger on board. Good condition. \$1,000. Call 816-318-8667 and leave a message.

Invacare R2 Mid-wheel Drive Power Wheelchair 20" seat width, pressure relief seat cushion. Very tight turning radius. \$3,000 includes delivery to location in KC area. Call 913-631-5959.

Invacare Storm Wheelchair headrest, tilt, recline, a gel cushion seat, an oxygen holder and is set up for a vent tray. Contact Jeaneene @ 417-466-2907, 417-466-1471 or by email @ graneene1@aol.com.

Electric Wheelchair 2 yrs. old. Like new. Asking \$1,000. Call 913-362-2354.

Pride Jet 7 Motorized Wheelchair never used. 16.5" turning radius. Call 913-287-7751 or 816-507-3728.

Invacare Ranger X Wheelchair 18" seat with Ishdish pad, power reclining back & legs, 2 new batteries, charger & additional 16" molded back & seat. \$5,000. Call Bob @ 573-592-4013 or 573-659-6692 or email btphillips1@mchsi.com.

Ranger II Electric Wheelchair fully adjustable. As new. Appointment only, 913-469-4188. Invacare Solara Wheelchair adjustable headrest, arm rests & leg supports. Tilt positioning chair with Avanti Curve Back & air inflated seats. Used only 6 weeks. Asking \$1,600 or best offer. Contact Don @ 402-488-5387 or by email @ dwoodbu@tps.org.

Invacare Reclining Manual Wheelchair with 18" seat width. Folding chair with removable headrest, flat free tires. \$600 or best offer. Call 913-631-5959.

Bruno Power Chair - PWC 2300 350 pound weight capacity, drop down controller, long trailing arm suspension, carry-along charger, 90 degree swivel seat, adjustable armrests, security key lock system, curb climbing height: 1:25", turning radius: 22". Used for only 3 months. Asking \$4,000. Contact Jeff Blair @ 402-614-5516.

1998 Electric Wheelchair Invacare Action Ranger II Storm Series loaded with gel seat remote programmer, joystick, switches, indicator, anti-tippers, adjustable high back/headrest recliner, standard and deluxe foot/leg rests and mirrors. 2 battery chargers, 1 used 24 volt, Action-Model18350 Invacare and one new/unused 24 volt dual mode automatic, Lester Electrical, Model-18350. Manuals included. Best offer. Call 785-272-3349.

Model 9000 Action Power Electric Wheelchair complete with two batteries and charger. Includes Owner's Operation and Maintenance Manual. Call 816-246-7761 or 816-564-4841. Jazzy Red Electric Wheelchair can be disassembled and put in car. One year old, valued at \$6,000. Best offer. 913-772-7071.

Invacare Power Wheelchair, Storm TDX5, headrest, tilt, recline, gel cushion seat. Brand new, never used. \$5,000. Call Kim @ 913-385-7952 after 4pm.

SCOOTERS

Deluxe Rascal Scooter used only 6 times, \$2,000. Call 316-321-3494.

Rascal 200 Three-Wheeled Scooter red, slightly used. Swivel seat and flip up armrests. Battery operated. Retailer new for \$1,500. Asking price \$600 or best offer. Call 913-341-5174.

Three-Wheeled Little Rascal battery operated, goes up to 5 mph, has horn, \$1,900. Call 402-371-3763 and leave a message.

LIFTS

Ricon S-Series Wheelchair Lift for van rear access. Load capacity 800 lbs. Weight 325 lbs. Remote control with manual backup. Call 417-865-1977.

Silver Glide Stair Lift with battery pack back up. 18 foot track. Manufacturer - American Access Ind. Like new. Asking \$1,200. 816-943-8334.

MATTRESSES & CUSHIONS

Therapeutic Select Air Mattress has bed controls, barely used. \$1,500 or best offer. Call Milli @ 316-794-8834. (Goodard, KS)

BEDS

Hospital Bed, Serenity Home Care Bed by Medline 2 years old. Used only one year. Semi-electric. Paid \$1,800, asking \$500. Call Wendy @ 913-371-2238.

2 Adjustable Beds. Massage with timer, knee and head adjusts. Good condition. \$1,200 or best offer. Will sell together or separate. Call Judy Cervantes @ 816-455-0105.

CHAIR GLIDES

2 Chair Glides for 12-13 steps. Manufacturer will re-install in the purchaser's home for approx. \$300. One and a half years old. Best offer. 913-772-7071.

TALKING MACHINES

Lightwriter SL 25 Talking Machine. 8 voice choices, dual visual display, 7K memory. New \$3130. Asking \$1500. Call 402-721-4626.

INCLUSION IN THE DIALOG IS NOT AN ENDORSEMENT FOR THESE PRODUCTS AND SERVICES.

Please notify Suzanne at 913-648-2062, ext. 209 or sfrye@alsa-midwest.org if your item has been sold or should be removed.

TRIBUTES

My grandmother was one of the strongest women I knew and seeing her go through ALS and keeping a smile on her face just proved her inner strength to me. My grandmother will always have a part of my heart. She was my best friend; she was everything to me. Now she is gone and I miss her dearly. I will be forever changed by the wonderful woman who would sing, "You are my sunshine" to me. My grandmother taught me to always smile and that things aren't always as bad as they seem to be. My grandma is my sunshine.

Written by Ashley Cable, 12 years old

We are looking for "Living with ALS" articles and TRIBUTES for future issues of the Dialog. Please send your stories to:

The ALS Association - Attn: Sally Dwyer
8340 Mission Road, Suite B-4
Prairie Village, KS 66206

or

Email: sdwyer@alsa-midwest.org

Damage Control

My name is Victoria Wasserman and I reside in Rochester, NY. My mother-in-law and my aunt both lost their lives to ALS, so finding a cure is something that is near and dear to my heart. In light of that, I am donating half the royalties from the sale of my book to ALS.

The book itself, *Damage Control*, actually has nothing to do with ALS. It is a murder mystery that centers around a medical malpractice cover-up and is intended for an adult audience. After it entered the publication process, my husband and I decided that we wanted to donate half the monies from its sale. My mother-in-law loved to read and often asked me how my writing was going. Unfortunately, she never got to see this book in its completed form. Donating the money from *Damage Control* is my way of paying tribute to my mother-in-law and my aunt as well as my attempt to make a difference.

I have a publicist who has included the fact that I am donating money to ALS in the book's press release. However, I thought that one of the best ways to get the word out would be to contact those who care the most.

The book came out in August. I know that people have gone into Barnes and Noble and ordered copies. It is also available online at barnesandnoble.com and amazon.com. Should I get any book signings, I will donate all the profit from those sales to ALS.

I hope that I can raise awareness as well as money. If you have any questions, you can reach me at 585-256-2599.

Thank you for your time.
Victoria Wasserman

March Birthdays

March 1	Charlene Parker	March 8	Tyra Kahre
March 1	Marilyn Cunningham	March 9	Connie Russell
March 1	Karlton Bohrn	March 10	Ann McLain
March 2	Allen Baker	March 17	William Westfall
March 2	Margaret Dodge	March 20	John Calver
March 3	Bertie Hood	March 21	Annette Schotte
March 4	Melvin Hoss	March 23	William Belz
March 4	Dick Franklin	March 23	William Barnes
March 5	Dennis Clark	March 23	Margie Yoakum
March 6	Christopher Batrano	March 28	Lee Barnhardt
March 6	Robert Landman		

Memorials

Thanks to the families of the following for designating our
Chapter for donations:

Kathleen Deardorff
Pat McNeese

Clyde Smith
Philip Wheatley

In Memorium

We send our sympathy & support to the families & friends of
those who have recently died after battling ALS.

Georgia Bourland
Kathleen Deardorff

Clyde Smith
Philip Wheatley

EMAIL ADDRESSES AND PHONE EXTENSIONS TO BETTER SERVE YOU:

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THE DIALOG

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March Support Group Dates

2	KC Night	7:00 p.m.
3	Hutchinson	2:00 p.m.
4	Wichita	7:00 p.m.
9	Springfield	7:00 p.m.
11	Omaha	7:00 p.m.
12	KC Caregivers	12:30 p.m.
17	KC Day	2:00 p.m.
17	Lincoln	6:30 p.m.
20	Wichita Caregivers	11:00 a.m.
22	Topeka	3:00 p.m.
	Joplin - No meeting this month.	
	Salina - No meeting this month.	

Support Group Locations

KC Night SG 2 nd Presbyterian Church 55 th & Oak	Topeka, KS SG Topeka & Shawnee County Public Library 1515 SW 10 th Avenue
KC Day SG Village Presbyterian Church 66 th & Mission Road	Lincoln, NE SG Madonna Rehab Hospital 5401 South Street Sheridan C Room
KC Caregivers SG ALS Office - Must RSVP 8340 Mission Rd., Ste. B-4	Omaha, NE SG St. Pius X Parish Center 6905 Blondo Street
Wichita, KS SG & Caregivers SG Grace Presbyterian Church 5002 East Douglas	Springfield, MO SG Cox Medical Center South 3801 South National Ave.
Hutchinson, KS SG Grace Episcopal Church 20 th & Main	Joplin, MO SG YMCA - 3404 W. McIntosh Circle Room A, South Branch
Salina, KS SG Christ the King Lutheran 111 W. Magnolia Street	

With Offices in the Following Cities, Contact the Office Nearest You!

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