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A message for ALS
caregivers

Care, Support \u0026amp;

ALS How ALS clinics
also support caregivers

~~THE HARD TRUTH~~

~~-DITL of a~~

~~CAREGIVER || This~~

~~is ALS~~ How ALS can
affect your relationship

Respiratory Care in

ALS - \"Your Decisions

for Your Quality of

Read Book Als Patient And

Life\" ABC's of ALS
Part 3 Sandra's Story:
An Artist Living with
ALS How An ALS
Patient Battled With
Choosing His Last Day
| NBC News

ABC's of ALS (Part 3)
An ALS patient and
Her Desire to Help
Others

A qualitative analysis of
the impact of caregiving
on ALS carer's lives

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Chris And Kelsie Snow
Still Full Of Hope In
Spite Of ALS Diagnosis

ALS Patient Robin
Mower's Caregiver
Wants Help with a

\ "Therapeutic" \

Activity Breakthrough
ALS Treatment

Promises New Hope |
TODAY Caregiving:

How to Survive and
Stay Sane NUR 211

AMYOTROPHIC

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~~Manual~~
SCLEROSIS ALS--

Ford Keep Memory

Alive - Frontotemporal

Dementias ~~Living Life~~

~~to the Fullest with ALS:~~

~~The Role of~~

Occupational Therapy

National Amyotrophic

lateral sclerosis (ALS)

Registry -- Impact,

Challenges, and Future

Directions ~~Als Patient~~

~~And Caregiver~~

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Caregivers Resources As

a service to the ALS

community, we ' re

providing online access

to a variety of resources,

including publications,

videos, books, and

informative websites

that provide a wealth of

easy-to-access

information on

important topics for

ALS patients and their

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Caregivers. Find a
support group in your
area.

Resource Manual

~~Caregivers Resources |~~
~~The ALS Association~~
For Caregivers Being
the primary caregiver
for a loved one with
ALS can be stressful —
not only are you on call
24/7, but you may have
additional outside
responsibilities, like

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working a full- or part-time job. Often, caregivers are so busy supporting their loved ones that their own physical or mental health takes second priority.

~~For Caregivers | The
ALS Association~~

Resources As a service to the ALS community, we have assembled

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Original Resource Manual
various resources,
including publications
and videos produced by
The Association, other
related books, fact
sheets, as well as
resource guides so that
you can quickly and
easily access information
pertaining to a variety of
subjects related to ALS.

~~Resources | The ALS
Association~~

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RESOURCES FOR CAREGIVERS

CareGuide Website:
www.careguide.com

ElderCare On-line:
Website: [www.ec-](http://www.ec-online.net)

[www.ec-](http://www.ec-online.net)
[online.net](http://www.ec-online.net) Empowering
Caregivers Website:

www.care-givers.com
Phone/fax: (212)

807-1204 Today ' s
Caregiver On-line

Website:

www.caregiver.com The

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ALS Association 1275 K
Street NW, Suite 250
Washington, DC 20005
Toll-free Information
and Referral Service:
(800) 782-4747

~~Caregiving COPING
WITH ALS CARING
FOR THE
CAREGIVER~~

ALS Caregiver ' s
Guide. This
comprehensive guide to

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Caring for someone with ALS is published by MDA. You can also order a free print copy from your local MDA office. National Alliance for Caregiving. The National Alliance for Caregiving is a nonprofit coalition of national organizations focusing on issues of family caregiving with the goal of improving

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Quality of life for families
and care recipients.

~~Resources for People
Caring for Those with
ALS~~

A good resource for
building a circle of help
and care is the Care
Connection program.

Be Aware of Depression
Depression can strike
anyone and caregivers
are especially vulnerable

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~~Caregiver~~
Resource
Manual

-it is the most common
healthcare condition
reported by family
caregivers. Be aware of
the early signs of
depression

~~Caregiving Tips and
Hints | The ALS
Association~~

What Is ALS?

Amyotrophic Lateral
Sclerosis (ALS), also
called Lou Gehrig ' s

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Manual

disease or motor neuron disease, is a progressive, neuromuscular disease that attacks nerve cells and pathways in the brain and spinal cord.

~~Amyotrophic Lateral Sclerosis (ALS) | Family Caregiver ...~~

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Home care is non-
medical care provided

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in the home, either by loved ones or paid caregivers. Home care can include assistance with dressing, bathing, eating, getting to the toilet, and moving around. Providing 24/7 care can be challenging for family members, no matter how much love and energy you have to offer.

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~~Home Care - Facts,
Tips, Resources for
People with ALS ...~~

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Each episode explores a
new and interesting
subject: learn more
about the science of
ALS, how to prepare for
and manage symptoms,

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and tips for both patients and caregivers. The ALS Pathways Podcasts are free audio files that you can stream or download using your smartphone, tablet device, or desktop computer. Each episode in this series is about 10 minutes.

~~ALS Patient Resources~~

~~— ALS Podcast | ALS~~

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Patient And

Pathways

PATIENT AND

CAREGIVER

RESOURCES

Caregiving

Administration on

Community Living

alzheimers.gov Supports

individuals living with

Alzheimer ' s disease or

other dementias and

their caregivers by

increasing access to

community resources.

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Aging Life Care

Association

aginglifecare.org Locate

a geriatric care

manager.

ALZConnected®

alzconnected.org

~~Patient and Caregiver~~

~~Resources - Alzheimer's~~

~~Association~~

What Does Being an

ALS Caregiver Mean?

Caregivers are

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individuals who provide physical and emotional support to help loved ones manage ALS.

Anyone can be a caregiver, but the role is usually undertaken by family members, such as a spouse, partner, sibling, parent, or adult child.

~~ALS Caregiver~~ | ~~ALS
Equipment~~ | ~~ALS~~

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Caregiver

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Challenges in limitation
and restriction – The
unending and often time-
delimited course of ALS
is mirrored in providing

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Care as being a caregiver can be restrictive in time and place. The increased demands on time, render it a limited and restricted resource, time is taken from the caregiver and given to the patient.

~~Disease Burden in
Caregivers of ALS
Patients ...~~

Resources for

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Caregivers As a service to the ALS community, we ' re providing online access to a variety of resources, including publications, videos, books, and informative websites that provide a wealth of easy-to-access information on important topics for ALS patients and their caregivers. Find support groups in your area.

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~~Resources for
Caregivers - ALS Texas
Manual~~
Family Caregiver

Alliance (FCA) seeks to improve the quality of life for caregivers through education, services, research and advocacy. FCA's National Center on Caregiving offers information on current social, public policy and

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Caregiving issues and provides assistance in the development of public and private programs for caregivers.

~~ALS | Family Caregiver Alliance~~

Patient and Caregiver Resource Center Since 1983, NORD has been the primary source of support and information for patients and families

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affected by rare diseases.

This Resource Center provides patients and caregivers with free webinars, fact sheets, infographics and other helpful materials to guide you on your journey with a rare disease.

~~Patient and Caregiver
Resource Center
NORD (National ...~~

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The Joe Martin ALS Foundation is a North Carolina-based organization that provides many support services at no cost, including caregiver training, 3-6 hours of in-home care per week, home assessments, wheelchair-accessible transportation to appointments and events, support groups

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and workshops, a camp for children whose lives have been affected by ALS, and a network of volunteers who can help families with meals, errands, and home projects.

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