



Appreciating Lupus Caregivers

by NEWLIFEOUTLOOK TEAM

Avoiding Lupus Caregiver Burnout

Lupus is an unpredictable illness that can be as much of an emotional rollercoaster for the caregiver as it is for the person with the diagnosis. The physical and emotional impacts of caring for someone with lupus can take their toll; it's important to do all you can to avoid caregiver burnout.

On top of the tips listed below for avoiding burnout, remember that talking to someone about your experience as a caregiver can lift a huge weight off your shoulders. Whether you talk to a therapist, a fellow lupus caregiver, or someone from a support organization, make sure you express your feelings of loss, frustration, resentment, anger or guilt with someone — keeping them bottled up can be damaging to your mental health, and your ability to provide the best care you can.

Also remember to look for the positives. Perhaps caring for your loved one has strengthened the bond the two of you have, or helped you to see what's really important in life. Every cloud has a silver lining.

Read our lupus caregivers infographic below to learn more about caregiving and avoiding burnout.



90 MILLION
Americans are caregivers for loved ones who are sick or disabled in some way.¹

1 MILLION AMERICANS
are also caring for veterans from the IRAQ+AFGHANISTAN wars who have visible injuries and invisible wounds.

True Dedication
A caregiver may help their loved one with **washing, dressing, eating** and other daily activities, as well as taking **medications**, getting to doctor's **appointments**, and so much more.

Handle with CARE
Each member of our family has the ability to help a loved one with a chronic condition, illness, or disability that may cause a barrier to healthy care options.

- Multiple sclerosis
- Lupus
- Parkinson's
- Crohn's disease
- Diabetes
- Alzheimer's
- Autism
- Cancer
- HIV/AIDS
- Depression
- Dementia
- Heart disease
- Stroke
- Trauma
- Vision impairment

To young and YOUNG AT HEART
Because of America's aging population, the number of people needing care has been rising, and is expected to keep going up. By 2020, people age 65 and older are expected to make up around **19%** of the American population, up from **14.8%** in 2008.²

However, parents caring for children with disabilities are also in the family caregiver ranks. Children with conditions like **autism, muscular dystrophy** and **down's syndrome** all require a caregiver, often through into adulthood.

CAREGIVER BURNOUT
Caregiver burnout is when a caregiver becomes emotionally and physically exhausted under the strain of caring for another person. This can manifest as:

- Irritability and feeling blue
- Fatigue and changes in sleep patterns
- Withdrawing from friends and other loved ones
- Abandoning activities you used to enjoy
- Changes in appetite and/or weight
- Depression and/or anxiety

Six tips to **AVOID BURNOUT**SM

Stay HEALTHY
It's easy to spend so much time thinking about your loved one's health that you can completely forget about your own. Make sure you take care of yourself, too – not work, exercise and get enough sleep.

Pursue your INTERESTS
Giving up on your hobbies and interests because of your caregiving duties can leave you feeling empty. Try to find time to do the things you love to do. You'll be able to enjoy your life more.

Take time to DE-STRESS
Consider taking up a stress-reducing activity like yoga, meditation or deep breathing techniques that will help you let go of the stress of being a caregiver.

Take A BREAK
Being a caregiver, although a labor of love, can be emotionally and physically draining. Give yourself time to be a caregiver every now and then to rest and recharge.

Connect with other CAREGIVERS
Joining a support group or an online forum for caregivers can help you to feel less alone in your struggles and provide a sense of community.

Seek and ACCEPT HELP
There's no shame in admitting you need help. Asking for help, or accepting it when it's offered, can lighten your load and make you better able to cope.

