

Help us pass the Lymphedema Treatment Act!

You can support the Lymphedema Advocacy Group's work to ensure that all lymphedema patients have access to their vital compression treatment supplies.

Please help us raise funds for our 2017 Lymphedema Lobby Days by participating in our Walkathon fundraiser on **February 14, 2017**, or simply by donating in support of the millions of men, women, and children affected by this chronic condition.

- ▼ To Donate or Learn More About our Fundraiser:

 https://www.crowdrise.com/love-the-lymphedema-treatment-act-valentines-walk-2017/fundraiser/lagboard
- ▼ To Learn More About the Lymphedema Treatment Act and the Lymphedema Advocacy Group (LAG): http://lymphedematreatmentact.org
- ▼ To Learn More About our Lymphedema Lobby Days: http://lymphedematreatmentact.org/lymphedema-lobby-days/

LAG is an all-volunteer, 501(c)(4) nonprofit organization that engages in lobbying activity strictly for passage of the Lymphedema Treatment Act. Contributions to LAG are not tax deductible.

