



LIGHTHOUSE LYMPHEDEMA NETWORK  
10240 Crescent Ridge Drive, Roswell, GA 30076  
770-442-1317  
[www.lighthouselymphedema.org](http://www.lighthouselymphedema.org)

May 20, 2019

The Honorable Maria Cantwell  
511 Hart Senate Office Building  
Washington, DC 20510

The Honorable Todd Young  
400 Russell Senate Office Building  
Washington, DC 20510

Dear Senator Cantwell and Senator Young,

On behalf of the lymphedema patients across the United States, the Lighthouse Lymphedema Network thanks you for your leadership in introducing **S.518, the Lymphedema Treatment Act**.

The Lighthouse Lymphedema Network was founded in Roswell, GA, in 1994. It is a non-profit 501(C)(3) organization of individuals who are lymphedema patients, family members, therapists, caregivers, or who have a genuine interest in lymphedema. Our goal is to educate, promote awareness, and provide support for lymphedema patients, the medical community, family and caregivers, insurance companies, the general public, and other lymphedema support groups.

Lymphedema is a medical condition affecting an estimated 1.5 to 3 million Medicare beneficiaries who are not currently receiving treatment from Medicare according to the current medical standard of care. What is worse is that Medicare is spending billions of dollars every year treating largely preventable lymphedema-related cellulitis infections.

Lymphedema is incurable, progressive, and, if left untreated, the swollen area(s) can become fibrotic and prone to serious, debilitating infections. Over time, untreated lymphedema results in disfigurement, disability, and even death.

The Lighthouse Lymphedema Network applauds your effort to close this coverage gap so that lymphedema patients can access the compression treatment supplies needed to manage this chronic condition. Please contact us with questions.

Sincerely,

Joan White, Director  
Lighthouse Lymphedema Network