

LE&RN's National Lobby Days brings advocates from around the country to Washington, DC where lawmakers are asked to support increased National Institutes of Health (NIH) lymphatic research funding and insurance coverage for available treatments.

ADVOCACY

LE&RN Chapters now reach across the United States and around the world, bringing the international lymphatic and lymphedema community together as one. Chapters sponsor events such as LE&RN's Walks, educational symposiums, and patient support groups bring the community together and making us stronger.

LE&RN's Lymphatic Disease Diagnosis and Treatment Centers of Excellence standards, as well as our Continuing Medical Education (CME) Seminars are raising the bar in assuring patients optimal, comprehensive care.

To entice the best and brightest researchers into joining our field, LE&RN sponsors up to 25 Travel and Poster Scholarships each year to the most promising young investigators worldwide.

BUILDING COMMUNITY

SETTING PROFESSIONAL **STANDARDS**

INSPIRATION FOR THE NEXT GENERATION

Addressing the lack of certified lymphedema therapists (CLTs) in underserved areas, LE&RN awards ten or more scholarships each year to therapists who enroll in an accredited CLT school.

LE&RN created World Lymphedema Day in 2016, which was officially established by the United States Senate that same year. In 2018, New York State passed a first-in-nation bill, written by LE&RN, that mandates all hospital institutions in the state to provide lymphedema (LE) information packets to all patients at-risk for the disease. LE&RN Chapters will now take this bill to statehouses across the country.

LE&RN hosts monthly Livestream symposia bringing the foremost experts in lymphatics to the medical community, those with LE, and caregivers online. And through our partnership with Harvard Medical Schools, LE&RN's website is host to the presentations delivered at the annual Lymphedema Symposium. For specific questions, LE&RN"s "Ask the **Experts"** provides a direct link to a panel of international experts.

LE&RN knows new treatments

and cures are waiting to be

discovered with research.

We remain steadfastly committed to

funding research fellowships and we

EXPANDING ACCESS TO CARE

LEGISLATIVE ACTION

CUTTING-EDGE ANSWERS

As LE&RN's Spokesperson since 2015, Academy Award-Winning actress Kathy Bates has been a tireless advocate for the lymphedema and lymphatic disease community. From national televised broadcasts, to meeting with politicians and officials at NIH, CDC, AMA, her partnership with LE&RN has exponentially advanced awareness of lymphatic diseases worldwide.

millions of people who live with lymphedema (LE) and lymphatic disease (LD) together in a collective fight to make our fights a global priority. LE&RN's website, social media channels, and e-newsletter ensure that everyone in our community is informed about the issues as well as how they can be an activist in this movement.

LE&RN is committed to bringing the

CREATING A MOVEMENT

actively promote LD/LE research being a top priority at the National Institutes of Health (NIH), the Department of Defense (DoD) and Centers for Disease Control and Prevention (CDC).

RESEARCH FOR NEW TREAMENTS AND A CURE



SPREADING THE WORD