Every BODY has the power to make a difference

Millions of women around the world live with the physically painful and emotionally distressing effects of lipedema, a fat disorder characterized by increased adipose tissue in the limbs that does not respond to diet or exercise. Because little is known about how and why it develops, lipedema remains a difficult disease to diagnose and treat.

The power to change this lies with you!

Join the Lipedema Foundation Registry (LFR) and help forge a path to better treatments from the comfort of your own home. Established by the Lipedema Foundation, the LFR is an online registry designed to help affected individuals and those who care for them better understand and treat lipedema.

Who can join?
Anyone! People who have or may have lipedema. People who do not have lipedema.

What do I need?
A computer with an internet connection and an email address.

What can I expect?
To answer questions about your medical experiences and lifestyle.

Why should I join?
To help researchers better understand lipedema and figure out:
- why lipedema happens
- who/how many are affected
- what treatments are being used
- how it impacts quality of life

Join the Lipedema Foundation Registry at lipedema.org/registry

Questions? Email registry@lipedema.org