



Join ldconnect.org today!

One of the most important steps towards finding effective treatments for lipodystrophy (LD) is to uncover the ways in which the diseases behave, the symptoms that are commonly seen and how those symptoms progress. This requires the collection of as much information as possible about all people affected by LD. LD Connect will serve as that database—the critical link between the patient community and the scientific community.

Join THE LD CONNECT Registry to:

- Tell us about your experience with LD and learn about the experiences with others
- Share information in a program that protects your privacy
- Provide valuable quality-of-life information that can be used to advocate on behalf of the LD community
- Help the research community launch clinical trials and studies We need to hear YOUR VOICE and we hope you will decide to join LD Connect. Every individual who joins makes a vital contribution toward helping all of those affected by lipodystrophy.



Who Can Participate

- Patients
- Family members
- Healthcare providers
- Investigators & researchers
- Stakeholders interested in Lipodystrophy

Who We Are

As its name implies, Lipodystrophy Connect represents the coming together of a wide range of entities and individuals who recognize the value in pooling their combined expertise and information. The Governing Board of Lipodystrophy Connect comes from leaders in the non-profit sector, government agencies, academic institutions and the pharmaceutical industry. Regardless of their individual backgrounds, the thought leaders of Lipodystrophy Connect share a common goal to increase the community's ability to help one another through improved communication, and by doing so, to help the millions of people affected by Lipodystrophy.

