



# FCS Connect

## Familial Chylomicronemia Syndrome

CONNECTING PATIENTS, HEALTHCARE  
PROFESSIONALS & RESEARCHERS



JOIN  
[FCSCconnect.org](http://FCSCconnect.org)  
TODAY!

## GET CONNECTED.

It takes whole communities to help people living with medical conditions lead healthier and happier lives. The FCS CONNECT Registry brings together individuals affected by lipoprotein lipase deficiency and professionals involved in their medical care and research.

Join **THE FCS CONNECT** Registry to:

- Learn how other patients manage their health
- Take surveys and share your experiences in a program that safeguards your privacy
- Receive information about research opportunities
- Help researchers unlock more effective treatments
- Improve knowledge about FCS among advocacy groups



Encourage everyone  
you know to join today!

### DO YOU HAVE

- Familial chylomicronemia syndrome (FCS)
- Familial LPL deficiency (LPLD)
- Hyperlipoproteinemia Type 1

### Illness is not a personal problem

The more data we collect about an illness, the closer we can get to helping communities find treatments that work. Once registered, view the collected data and learn from others. Your contact information is NEVER shared.

### Putting the patient back into patient data

You are the expert in your own health. By providing your medical information, you're giving researchers a new look at disease. Anyone with FCS can contribute data, and anyone who needs the data can access it. We're making information as useful, comprehensive and safe as it can possibly be.

Registration takes just 10-15 minutes, but the data you provide lasts a lifetime.

**LEARN FROM OTHERS... MAKE A DIFFERENCE.... GET CONNECTED**

[WWW.FCSCONNECT.ORG](http://WWW.FCSCONNECT.ORG)