

Helpful Resources for Lupus and Clinical Trials

LUPUS FOUNDATION OF AMERICA RESOURCES

Lupus Foundation of America

🔗 [Lupus.org](https://www.lupus.org)

Lupus Foundation of America is the only national force devoted to solving the mystery of lupus, one of the world's cruelest, most unpredictable, and devastating diseases, while giving caring support to those who suffer from its brutal impact. About 1 in 250 African American women will develop lupus. Lupus is not only more common in African Americans, it is typically more serious. 9 out of 10 people with lupus are women, but men and kids get lupus too.

At the **Lupus Foundation of America**, we understand the physical suffering, emotional turmoil, and economic hardships caused by lupus. We collaborate with lupus medical experts to produce useful and relevant resources, programs and services to directly help people with lupus and their doctors better manage the disease.

Visit **Lupus.org** to learn more.

For lupus news and information follow us here:

f [LupusFoundationofAmerica](https://www.facebook.com/LupusFoundationofAmerica)

t [@LupusOrg](https://twitter.com/LupusOrg)

ig [@LupusOrg](https://www.instagram.com/LupusOrg)

National Resource Center on Lupus

🔗 [Resources.Lupus.org](https://www.Resources.Lupus.org)

Developed by Lupus Foundation of America, the **National Resource Center on Lupus** is a one-stop resource for all things lupus, from treatment to living and relationships. The Resource Center is updated on a regular basis to ensure that you are up-to-date and well-prepared to manage your lupus.

The Resource Center aims to empower, educate and connect people with lupus through reliable



resources that provide emotional support and clear, accurate health information.

Anyone affected by lupus, including people diagnosed with lupus, children and teens, caregivers, health care professionals and the general public can find tailored information on the Resource Center.

Visit **Resources.Lupus.org** to learn more.

National Health Educator Network

🔗 [Lupus.org/HealthEducator](https://www.Lupus.org/HealthEducator)

Lupus Foundation of America's health educators are available to answer questions and provide caring support to people with lupus, their families and their caregivers.

Call 1-800-558-0121 or visit **Lupus.org/HealthEducator** to contact a health educator today. *(Questions are answered in English and Spanish.)*

LUPUS CLINICAL TRIALS DATABASES

National Resource Center on Lupus

🔗 Resources.Lupus.org

For more information on Clinical Trials in your area, visit Resources.Lupus.org/Entry/Search-for-Clinical-Trials

For a listing of Frequently Asked Questions on Clinical Trials, visit Resources.Lupus.org/Entry/Frequently-Asked-Questions-Clinical-Trials

Research.forMe™ Lupus Registry

🔗 Resources.Lupus.org/Registry

A registry is a place to store detailed information about individuals with a specific disease and their families. The **Research.forMe™ Lupus Registry** is for individuals with a diagnosis of lupus. This registry will help researchers and clinicians better understand the disease, and registry participants will receive tailored information on available studies and research opportunities.

Visit Resources.Lupus.org/Registry to learn more.

The Center for Information & Study on Clinical Research Participation (CISCRP)

🔗 CISCRP.org

The CISCRP Education Center offers individuals information and resources on clinical research participation.

Visit CISCRP.org/Education-Center to learn more.



Search Clinical Trials

🔗 SearchClinicalTrials.org

Search Clinical Trials is a public service that compiles clinical trial listings from multiple sources. You can also request a free search for clinical trials in your area.

Visit SearchClinicalTrials.org or call 1-877-MED-HERO to learn more.

ClinicalTrials.gov

🔗 ClinicalTrials.gov

ClinicalTrials.gov is a registry and database of publicly and privately supported clinical studies conducted around the world. ClinicalTrials.gov provides in-depth information on clinical studies including relevant history, policies, and laws.

Visit ClinicalTrials.gov to learn more.

This project is supported by Grant Number 1 CPIMP161122-01-00 from the U.S. Department of Health and Human Services Office of Minority Health. Its contents are solely the responsibility of the authors and do not necessarily represent the official views of the HHS, OMH.

FOR MORE INFORMATION, VISIT LUPUS.ORG/IMPACT

Help Us Solve
The Cruel Mystery
LUPUS[™]
FOUNDATION OF AMERICA
LUPUS.ORG