Helpful Resources for Lupus and Clinical Trials

LUPUS FOUNDATION OF AMERICA
RESOURCES

Lupus Foundation of America
☞ 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:
☞ LupusFoundationofAmerica
☞ @LupusOrg
☞ @LupusOrg

National Resource Center on Lupus
☞ 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

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
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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.

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FOR MORE INFORMATION, VISIT LUPUS.ORG/IMPACT