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 health care providers better manage the disease. Here are a few resources to share with your patients:

**National Resource Center on Lupus** – A collection of up-to-date resources and information on lupus. The Resource Center aims to empower, educate, and connect those impacted by this devastating and unpredictable disease. Visit [Lupus.org/Resources](http://Lupus.org/Resources) to learn more.

**National Health Educator Network** – Our nurse and certified health educators are available to answer questions and provide support to people with lupus, their families and caregivers. **Call 1-800-558-0121 or visit Lupus.org/HealthEducator** to speak to a health educator today (in English and Spanish).

**LupusConnect™** – An online lupus community where members can engage with others like them to share experiences and find emotional support. Join the conversation at [Lupus.org/LupusConnect](http://Lupus.org/LupusConnect).

**Research.forME™ Lupus Registry** – The Registry is an online data collection tool for people living with lupus and caregivers who are interested in participation in research. It is designed to accelerate the development of future clinical research and new lupus treatments. To learn more or enroll, visit [Lupus.org/Registry](http://Lupus.org/Registry).

For more information on programs and services offered through the Lupus Foundation of America, visit [Lupus.org](http://Lupus.org).