

ABOUT US

The Louisiana Hemophilia Foundation was established in 1976 to help Louisiana residents with hemophilia, von Willebrand disease, and other bleeding disorders, lead normal and productive lives. We are a 501 (C) 3 Organization (non-profit organization).

CONTACT US!

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LOUISIANA
HEMOPHILIA FOUNDATION

ESTABLISHED IN 1976

OUR MISSION



To improve the quality of life and assist persons affected by inherited bleeding disorders by providing education, advocacy, support services, and by promoting research.

What we offer to our community:

EDUCATION

We host programs across the state that are open to all

PATIENT ASSISTANCE

To help during those tough times

ACCESS TO CARE

For needs that arise from receiving medical care, such as gas & lodging

MEDIC ALERT SCHOLARSHIPS

To provide life-saving services in case of emergency

EDUCATIONAL SCHOLARSHIPS

To help better lives through higher education

SUMMER CAMP

To grow and empower our youth into leaders

ANNUAL MEETING

To build our community and strengthen through knowledge



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