



Fostering better education, access and care for the bleeding disorders community.

Our mission at Hemophilia Federation of America, is to assist, educate and advocate for the bleeding disorders community.

As an organization that places individuals, families and community at the heart of all we do, HFA is proud to provide research opportunities, educational tools, assistance and advocacy to anyone in the bleeding disorders community throughout the United States.

Through our resources and initiatives, HFA is well positioned to collaborate with providers, researchers and individuals to create opportunities which benefit everyone.

As we have done for the past 25 years, now, and into the future, we are committed to our mission and bridging the gaps between industry, providers and individuals in the community.

This commitment allows us to move toward better access, education and care for everyone.



Some of HFA's initiatives include:

- **Helping Hands** – Our emergency financial assistance program, which is open to anyone, often by referral from a nurse, social worker, or HTC provider
- **Blood Sisterhood App** – A digital platform to help women with bleeding disorders track their monthly bleeding and turn that data into helpful information for themselves and their medical providers
- **PCORI Research** – Patient-centered research initiatives seeking to investigate and deliver data which can help solve problems and create better solutions for people in the bleeding disorders community
- **Educational Toolkits** – Programming inspired by topics which our community has asked for, these toolkits help deliver online content to patients who can browse content on their own time and at their own speed, while learning about topics and resources which can be helpful to them and their individual circumstances
- **Learning Central** – Our proprietary, medically-vetted learning platform which delivers educational programming and tools to anyone in the bleeding disorders community—from newly diagnosed to veterans; this platform is intended to deliver educational content in a fun, engaging way
- **Symposium** – The second largest national gathering of the bleeding disorders patients and community takes place annually at our conference, which is patient focused, with educational programming, networking and research demonstrations for families, providers and industry professionals
- **Advocacy Fly-In** – Annually in June, HFA provides advocacy training for small groups who take their newly minted skills to Capitol Hill, and meet with Congressional members and their staff to discuss the needs of the bleeding disorders community as they relate to current legislative issues
- **Dateline Federation** – Our quarterly magazine offers news, stories, updates, education and connections for the bleeding disorders community

Interested in learning more or exploring what HFA has to offer? Visit our website for so much more, and to connect with us!

www.hemophiliafed.org



Assisting • Educating • Advocating

