

THE COALITION FOR HEMOPHILIA



ABOUT US

A 30 year-old national non-profit organization, we are a primary source of information nationally for individuals and families with hemophilia B. We provide networking support and information with regard to their physical, emotional and psychological care. We empower our members to advocate for themselves. We spend a large amount of time speaking one-on-one with patients and their caregivers about the importance of adhering to treatment regimens.



OUR MISSION

To make quality of life the focal point of treatment for individuals with hemophilia B; to empower and educate our members to act as advocates on their own behalf; and to ensure patients have timely information about their healthcare so they can have the highest quality of life possible.

OUR GOALS

To empower individuals and families with hemophilia B to advocate on their own behalf, to offer education, and to ensure access to the highest standard of care. We are focused on tackling the unmet needs of the hemophilia B community.

OUR PROGRAMS AND SERVICES

- Annual Educational Symposium
- *On the Road* family meetings and workshops
- Spring and Fall Men's retreats
- Spring and Fall Women's retreats
- Support groups
- Webinars and video series
- BVoice Advocacy Program
- *The Beats* Music Program
- BCares Patient Assistance Program
- *Gen IX* Advocacy, Leadership, and Teen/Mentoring Programs
- William N. Drohan Educational Scholarship
- *Factor Nine News* - Offered in English and Spanish

GET INVOLVED. STAY INVOLVED.

VISIT OUR SOCIAL MEDIA SITES

Website: www.hemob.org

Facebook: www.facebook.com/HemophiliaB/

Twitter: <https://twitter.com/coalitionhemob>

Instagram: www.instagram.com/coalitionforhemophiliab

LinkedIn: <https://www.linkedin.com/company/coalition-for-hemophilia-b/>



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