

Make Fresh Connections at #BDC2025

[Register Today! >](#)

Celebrating 75 Years

Honoring history, hope, and progress.

Bleeding Disorders

Learn more about hemophilia, VWD, and rare factor disorders.

Research We Fund

A world without bleeding disorders begins with research.

Guidelines on Care

Standards of bleeding disorders care for healthcare providers



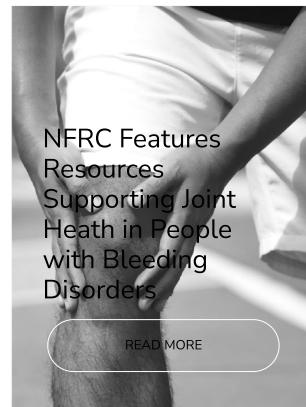
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Add Your Voice

Community Voices in Research (CVR) is a community-powered registry that gathers information through surveys offering researchers a 360-degree view of what it is like to live with a bleeding disorder. Open to all persons affected by an inherited bleeding disorder as well as their non-affected family members (parents, spouses, grandparents, and siblings), it provides all the opportunity to participate in patient-reported outcomes research.

952

Total Enrollment

46%

are Male

54%

are Female

[Enroll Now](#)

Join us at these events



Nov 2, 2025

New York City, New York



15 Oct

[NBDF Research Roundtable](#)

Oct 15, 2025 - Oct 16, 2025

Washington, District of Columbia



21 Aug

[Bleeding Disorders Conference](#)

Aug 21, 2025 - Aug 23, 2025

Aurora, Colorado

[View all events](#)

## You're Not Alone

Connect with other families affected by bleeding disorders in your area through NBDF's chapter network.

[Find a Chapter Near You](#)



## How you can help

"NBDF strengthens our community and gives us a sense that we belong to something bigger than ourselves—that we're all in this together. That's an amazing feeling." — Monica, Community Member

Your generosity has a huge impact!

[Donate Today](#)