

AFFILIATED COMPREHENSIVE HEMOPHILIA TREATMENT CENTERS

Children's Hospitals and Clinics of MN Hemophilia and Thrombosis Center

2525 Chicago Ave S,
CSC – 175
Minneapolis, MN 55404
Phone | 612.813.5940
Toll Free | 888.811.5940

Mayo Clinic Comprehensive Hemophilia Center

Mayo 10-55E
200 First Street SW
Rochester, MN 55905
Phone | 507.284.8634
Toll Free | 800.344.7726

M Health Center for Bleeding and Clotting Disorders

2512 South 7th Street, Suite 105
Minneapolis, MN 55454
Phone | 612.273.5005
Toll Free | 800-688-5252, ext. 66455

Sanford Health South Dakota Center for Bleeding Disorders

1600 W 22nd St, PO Box 5039
Sioux Falls, SD 57117
Phone | 605.312.1000
Toll Free | 800.850.0064

COMPREHENSIVE CLINICS VISITS

Clinic visits at a hemophilia treatment center (HTC) are encouraged on a regular basis (at least once a year) to review and evaluate the status of a person with bleeding disorders. Goals of the Comprehensive clinic visits are to assess health concerns since the last visit, evaluate current treatment plans, provide educational opportunities, and plan for the treatment on any unmet needs, changes or future needs. Studies have shown that regular attendees at HTC Comprehensive Clinics have better health outcomes.



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**HEMOPHILIA
FOUNDATION**
OF MINNESOTA/DAKOTAS

*We dedicate ourselves to advancing the
quality of life of individuals and families
affected by hemophilia and other bleeding
disorders by providing a broad range of
programs and services.*

E*stablished in 1965,*

the Hemophilia Foundation of Minnesota/Dakotas (HFMD) is an independent 501 (c)(3) non-profit organization and the only resource of its kind in the region. We work to provide education, support, and advocacy for our community and strive to help those affected by a bleeding disorder to become healthier and more independent.

We partner with the medical staff from four Hemophilia Treatment Centers in the region to carry out our unique programs and services and provide valuable information, advocacy, and support for our bleeding disorders community.



Annual Meeting: An educational weekend for patients and families featuring quality programs from leading medical experts who treat hemophilia.



Family Education Events: We feature the latest in medical information and the opportunity for families to learn, network, share experiences and have fun.



Women's Education Program: Annual program for women with bleeding disorders featuring informative presentations and interactive learning. In partnership with a program sponsor.



Group Fitness Events: Events providing safe and fun cardiovascular activities the whole family can enjoy.



Winter Teen Retreat: Connect with other teens while enjoying fun indoor and outdoor winter activities while gaining independence.



Blood Brotherhood: A program exclusively for adult men with bleeding disorders.



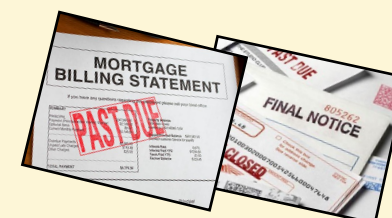
Summer Camp Not-a-Clot: A 6-day summer traditional camp experience for campers, ages 8-17, who have a bleeding disorder. The mission of camp is to foster community among bleeding disorder patients and promote a transition to independence.

Advocate:

For Access to Healthcare



Post-Secondary Scholarships:
For eligible students with hemophilia or related bleeding disorders.



Emergency Assistance: For patients and families experiencing financial hardships.