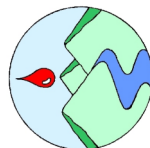


WHAT DO WE DO?

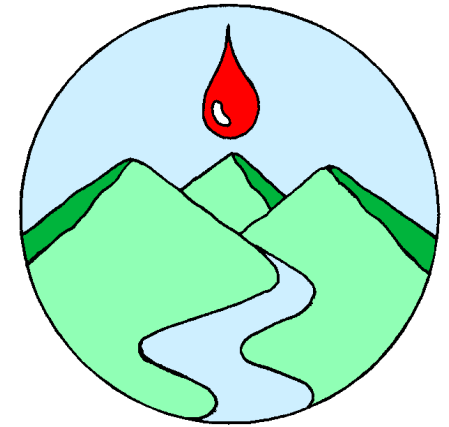
We advocate for the bleeding disorders community by:

1. Legislative lobbying at the national level (*e.g.* Ricky Ray Hemophilia Relief Fund Act of 1998) and the state level (Access to Justice legislation).
2. Camperships
3. Emergency funds
4. Educational scholarships
5. Clinic Travel Assistance Program (CTAP)
6. Community Education
7. Fundraisers - 100% of all monies raised are returned to the community
8. Monitoring of health issues with federal and state agencies, and non-profit organizations
9. Association Activities
 - Educational Retreats
 - Newsletter/Website
 - Annual Picnic/Business Meeting
 - Golf Tournament
 - Bowl-a-thon
 - Cabin Fever Party



Bleeding Disorders Association
of Northeastern New York, Inc.
P.O. Box 947
Rensselaer, NY 12144

Bleeding Disorders Association of Northeastern New York



“We are Here for You”

www.BDANENY.org

P.O. Box 947
Rensselaer, NY 12144
(518) 729-3577

Bleeding Disorders Association of Northeastern New York, Inc. MEMBERSHIP/SUPPORT

Name: _____ E-mail: _____

Address: _____ Home Phone: _____

City: _____ State: _____ Zip Code: _____ County: _____

INDIVIDUAL \$10. SPONSOR \$150.

FAMILY \$20. PATRON \$250.

BENEFACTOR \$50. PRESIDENT'S CLUB \$500.

January 31, 2011

Your contribution is TAX DEDUCTIBLE. Please make all checks payable to:

BLEEDING DISORDERS ASSOCIATION OF NORTHEASTERN NEW YORK, INC.
(BDANENY, Inc.)

No one will be denied membership because of financial hardship.

Please return this form to :

Bleeding Disorder Association of NorthEastern New York, Inc

P.O. Box 947

Rensselaer, New York 12144

E-mail Address: memberservices@bdaneny.org

www.BDANENY.org

Phone: (518) 729-3577

WHO ARE WE?

A chapter of the National Hemophilia Foundation, since 1968. We are a volunteer, not-for-profit organization that serves the congenital bleeding disorders community within North-eastern New York and adjacent areas.

We also reach out to the general public for education about bleeding disorders.

OUR MISSION

The BDANENY is dedicated to volunteer service on behalf of persons with hemophilia, von Willebrand Disease (vWD) and other genetic bleeding disorders.

The organization provides services and referrals to members of the community and encourages them to advocate for their own healthcare. Furthermore, the organization supports research and programs that will improve the quality of life for persons afflicted with genetic bleeding disorders and their associated complications.