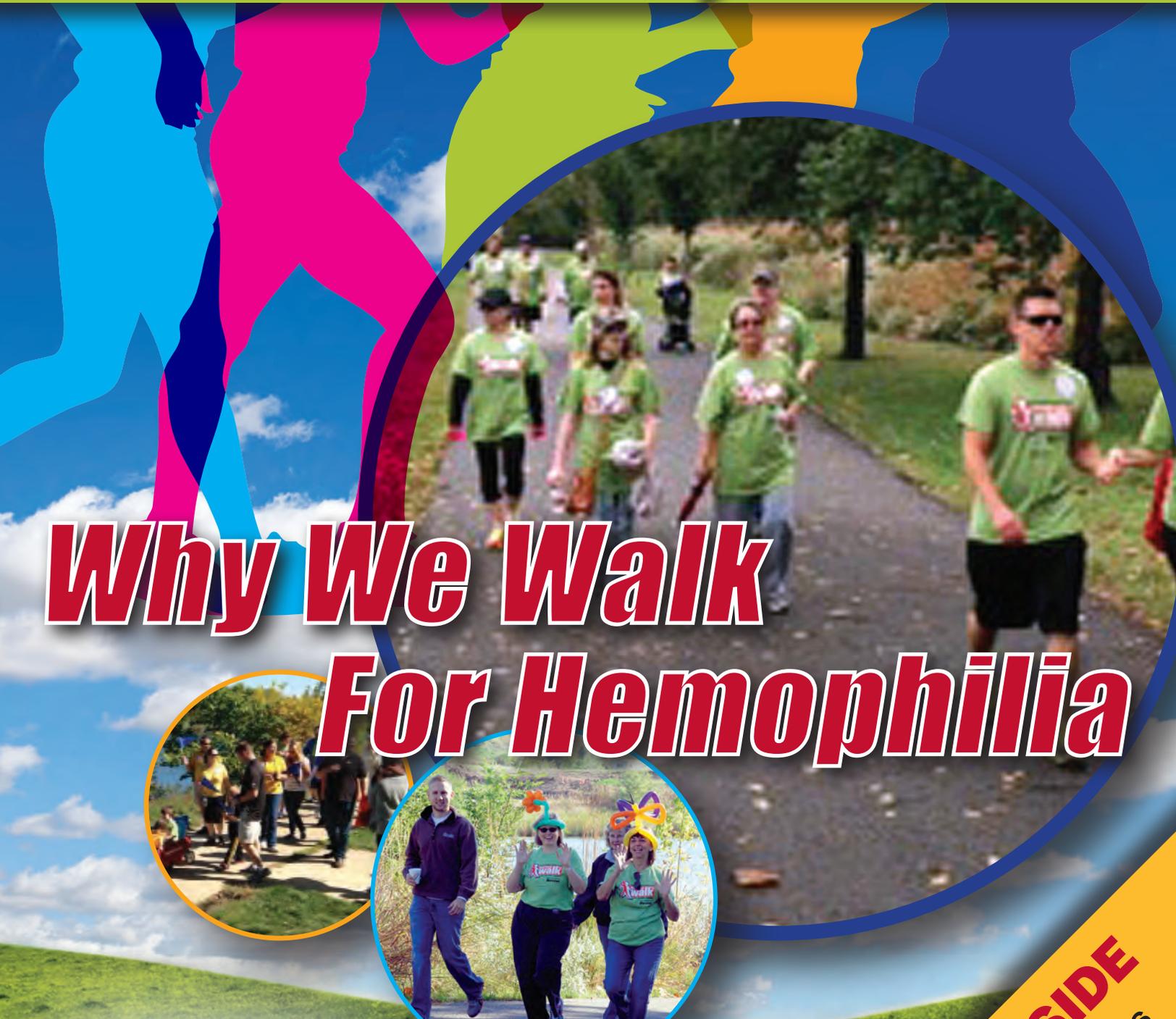


CHAPTER

Factors

SUMMER 2014



Why We Walk For Hemophilia



**Midwest
Hemophilia
Association**

LOOK INSIDE
Hemophilia Walks
Special Insert



Join the 4th Annual Bayer Virtual Walk for Hemophilia!

Help your local hemophilia chapter raise sponsorship funds by participating in the 4th annual Bayer Virtual Walk for Hemophilia.

Spread the word to your friends and family. Every virtual walker brings your chapter one step closer to making a difference in the community!

Virtually walk for a cause TODAY at www.walkforhemophilia.com.



*Spread the
word and support
your community
TODAY!*



Chapter Factors is published by:

The Midwest Hemophilia Association

PO Box 412866
Kansas City, MO 64141
816-479-5900
www.midwesthemophilia.org

EXECUTIVE DIRECTOR

MARK COX
913-220-9687
mcox@midwesthemophilia.org

BOARD OF DIRECTORS

2014 OFFICERS

JOHN CARLETON
PRESIDENT
Home: (660) 438-5918
jrcarle@hotmail.com

AIMEE TEMPERA-PARKS
VICE-PRESIDENT
Cell: (316) 518-0524
ajtemperaparks@aol.com

DEBBIE NELSON
SECRETARY
Cell: (816) 456-2097
Work: (816) 802-1268
dlnelson@cmh.edu

MARK DUDLEY
TREASURER
Home: (816) 650-4035
Work: (816) 554-6750
mdudley@blueridgebank.net

2014 BOARD MEMBERS

ANGELA BROWN
(314) 952-6265
intangela@yahoo.com

CHARITY MEADOWS
(417) 353-0931
3meadowsboyz@gmail.com

ELLENE SANDER WHITMORE
(816) 377-7174
ellenesw@gmail.com

JENNIFER RENTSCHLER
(660) 888-1618
jenr@accuraterx.net

This material is provided for your general information only. The Midwest Hemophilia Association does not give medical advice or engage in the practice of medicine. The Midwest Hemophilia Association under no circumstances recommends particular treatments for specific individuals. In all cases, it is recommended that individuals consult a physician or local treatment center before pursuing any course of treatment.

Brand names of treatment products are provided for information only. They are not an endorsement of a particular product or company by MHA. Acceptance of advertising for products and services in Chapter Factors in no way constitutes endorsement by the Midwest Hemophilia Association.

© Copyright 2014 by Midwest Hemophilia Association

FROM THE EXECUTIVE DIRECTOR



News and Notes

The dates for our four Hemophilia Walks are getting closer and it's time to kick the fundraising efforts into high gear. Last year we had a total of 22 teams at all four walks, or about 5 or 6 teams per walk. This year, we're hoping for at least 40 total teams to reach our goal of \$60,000 for all four walks.

To help us achieve our goal, we need you be a Team Captain this year. It's really easy. All you need is three other people to be an official team. You can form a personal/family team or corporate/business team. It really doesn't matter. Walk teams are the most effective method of hitting fundraising goals.

So how do you form a team? It's simple – email or call your family, friends and/or co-workers. Facebook or Twitter are great ways to extend your reach in getting more people to support your team. Included in this issue is a special Walk insert that gives information about the walks plus great tips on fundraising. Pull out this insert to use as a helpful resource during your Walk preparations.

Proceeds from the Walks are crucial to the function of MHA. It allows us to schedule more education events, bring in top-notch speakers, supports our advocacy programs, camp and more.

So very simply, we are asking you to be a Team Captain. Go to www.hemophilia.org/walk, click on MO icon on the right side. Follow the instructions from there. If you have any questions please contact me.

Mark Cox
Executive Director

WICHITA WALK
Sunday, October 12, 2014

KANSAS CITY WALK
Saturday, October 18, 2014

SPRINGFIELD WALK
Saturday, October 18, 2014

COLUMBIA WALK
Sunday, October 19, 2014

2014 MEMBERSHIP APPLICATION

Name: _____

Address: _____

Phone: _____ Email: _____

Individual/Family (\$25) Corporate (\$100)

Additional Contribution: \$ _____

Amount Enclosed: \$ _____

Please mail this form and your membership fee to:
Midwest Hemophilia Association • PO Box 412866 • Kansas City, MO 64141
Thank you for your support!



QUICK NOTES

If It Is Going To Be, It Is Up To Me!

In my former life, before my “make believe” retirement, many of you know I was an Elementary Principal for the Warsaw School District. Early in my career as a principal my two schools became lead schools in the Accelerated Schools Project. In the old administrative model principals gave orders and teachers followed those orders. Teachers hibernated in their rooms with little interaction among themselves and little sharing of best practices. The Accelerated Schools model was built around all members of the educational team having joint responsibility in the decision making, openly sharing best practices and taking responsibility for the success of all decisions made by the group.

At this point you’re probably wondering why the stroll down memory lane. One of the key phrases we



constantly used during our Accelerated Schools journey was “If it is going to be, it is up to me!” The message being that each person has to take personal responsibility for the success of any endeavor the group chooses to pursue. We are gearing up for our 2nd Annual NHF/MHA Hemophilia Walk. We, your organization, need each and every member to contribute in some form. How you ask? You can form a team, join a team, participate in the Walk, raise funds in your neighborhood or place of work. The possibilities and ideas are endless.

Last year I did a letter writing campaign and mailed a personal plea to every family member, friend, business associate, etc. I asked each one to send me \$10, or more, if they were able and felt the cause was worthwhile. I also gave them an easy way out and told them that if they didn’t send anything I still loved them! Pam and I ended up being the second place team overall. I didn’t beat on doors, hound already overburdened businesses or wear out the “locals” of our small town. I also know one of our members in western Kansas did a number of small fund raisers and was very

successful. Our number one fundraiser included a request in all of the billing statements in the family business.

Please, don’t sit on the sideline this year and expect someone else to do the work. We had a very successful first year, but there is a huge potential for growth. MHA is your organization. Get involved at some level and remember: “If it is going to be, it is up to me!”

— John Carleton

MHA 16th Annual Golf Tournament

Join us Friday, September 26th for an awesome day of golf at MHA’s 16th Annual Golf Tournament to be held at the Drumm Farm Golf Club in Independence, Missouri. It is a great course and the staff is extremely supportive of MHA.

A continental breakfast will be available at 7:00 a.m. to all participants prior to the shotgun start at 8:00 a.m. Lunch will be served following the tournament. Both are included in the entry fee.

Prizes will be awarded for longest drive, longest putt, and closest to the pin plus a special award to be announced tournament day for any hole-in-one. Cash awards will be given to the top three four-person teams.





A variety of sponsorship levels are available to help support the golf tournament, such as Driving Range Sponsorship, Golf Cart Sponsorship and Hole Sponsorship. This is a great way for organizations as well as individuals to contribute to MHA. Proceeds help fund MHA educational seminars, support groups and summer camp. Contact Mark Cox at (816) 479-5900 or mcox@midwesthemophilia.org for more information.

Sign up your foursome today to help make this event a success. For more information and registration forms, visit <http://midwesthemophilia.org/events/mha-annual-golf-tournament/>. Hope to see everyone participate this year.

2014 Family Fun Fair

MHA's Family Fun Fair 2014 will be held Saturday, September 27 and Sunday, September 29 at the Hilton Garden Inn in Independence, MO. The cost is \$30 per family for MHA members, or \$55 per family for non-members. Registration fee is non-refundable.

The weekend will feature a full slate of informative speakers and educational topics, plus attendees will enjoy a fun evening at Paradise Park.

Teens ages 13 and up will enjoy an afternoon of fun activities and entertainment. Children under the age of 13 will have quality child care with fun and safe activities provided for them at the Hilton Garden Inn hotel complex.

Once again, numerous exhibitors will be in attendance, giving participants the opportunity to meet representatives from each of the companies and find out what they provide in the way of educational materials, programs, and services for people with bleeding disorders. There will also be information MHA programs like summer camp, the First Step program, legislative work, and walkathons.

Check-in and exhibits open at 9:00 a.m. Space is limited. Reserve your spot early to guarantee a fun weekend for the entire family! Visit www.midwesthemophilia.org for additional details.

2014 Calendar of Events

September 18 – 20	NHF Annual Meeting Washington D.C.	
September 26	16th Annual Golf Tournament Falcon Ridge Golf Course Lenexa, KS	
September 27 – 28	Family Fun Fair Hilton Garden Inn Independence, MO	
October 12, 2014	Wichita Hemophilia Walk Exploration Place Wichita, KS	
October 18, 2014	Springfield Hemophilia Walk Nathanael-Greene Park Springfield, MO	
October 18, 2014	Kansas City Hemophilia Walk Waterfall Park Independence, MO	
October 19, 2014	Columbia Hemophilia Walk Stephens Lake Park Columbia, MO	
November 15	20th Annual Recognition Banquet Kansas City, Missouri	

HELPFUL NUMBERS

NATIONAL HEMOPHILIA FOUNDATION (NHF)

1-800-42-HANDI • www.hemophilia.org

UNIVERSITY OF MISSOURI HOSPITAL AND CLINICS HEMOPHILIA TREATMENT CENTER

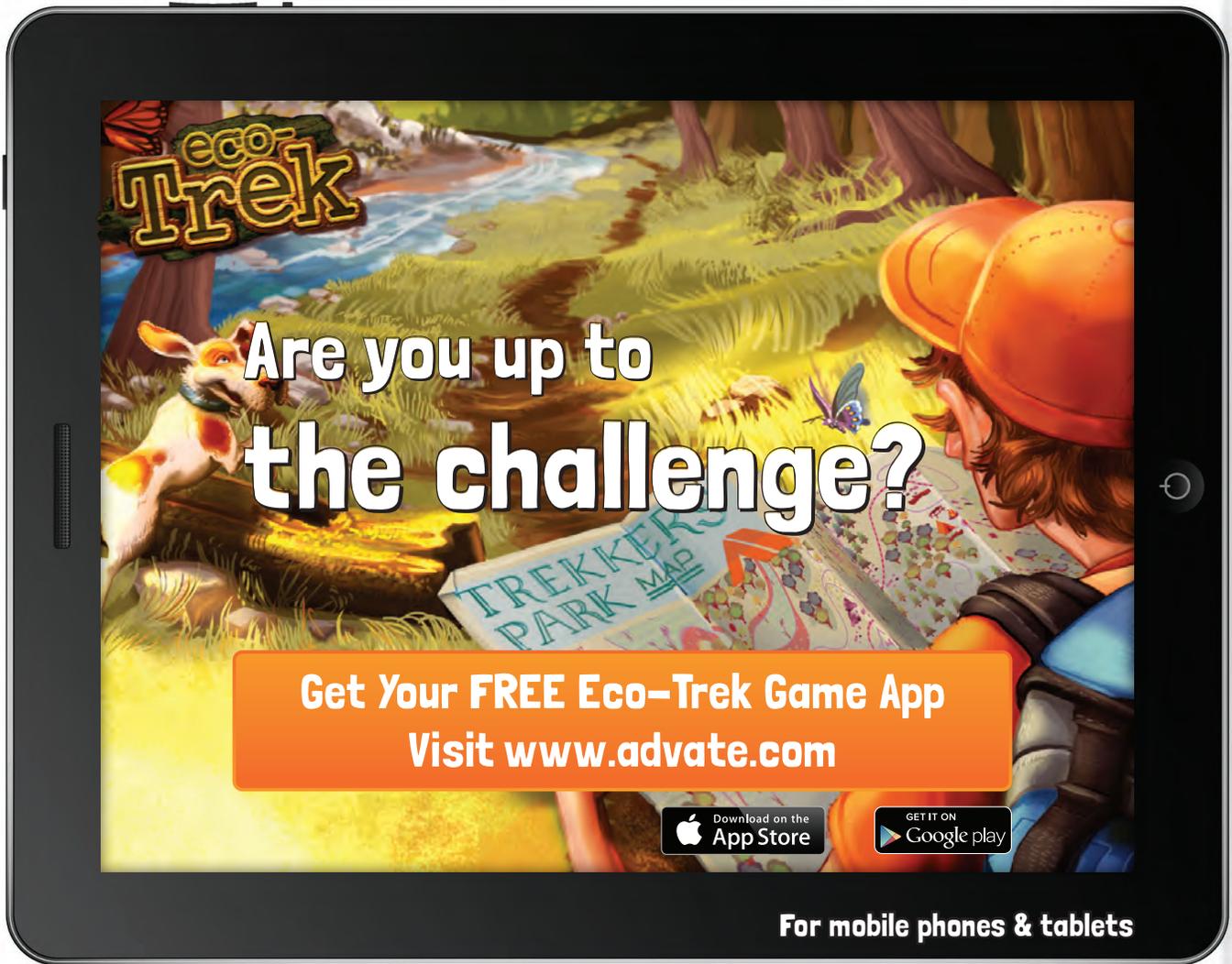
1-573-882-9355

Dr. Barbara Gruner • Dr. Carl Freter • Dr. Tamara Hopkin • Lauren Grana, MSW

KANSAS CITY REGIONALHEMOPHILIA CENTER

1-816-234-3508 • 1-800-236-1713

Dr. Brian Wicklund • Dr. William Jennings • Dr. Shannon Carpenter
Dr. Jill Moormeier • Judy Kauffman, RN, MS, CPNP
Andrew Wilson, RN, CPHON • Katie Foote, LMSW



Eco-Trek is a one-player, multi-level game package that tests your outdoor survival skills

-  3 outdoor adventure games in 1
-  An online leaderboard for nationwide competition
-  Answer educational questions about ADVATE for bonus scoring

Bring your A-game to www.advate.com for your free download!

For more information, contact your Baxter representative today:

David Dale

Phone: (913) 416-0158

Email: david_dale@baxter.com


 [Antihemophilic Factor (Recombinant),
 Plasma/Albumin-Free Method]

Baxter and Advate are registered trademarks of Baxter International Inc.
 All other trademarks, logos, or product brands appearing herein are the property of their respective owners. USBS/34/14-0031

Baxter

...for the human factor®

NATIONAL REACH. LOCAL PRESENCE.

With a nationwide network of pharmacies, Accredo is one of the largest suppliers of clotting factor in the industry. What's more, we have been a part of your local community since 1990, with experienced community advocates who can offer assistance with all aspects of therapy. We deliver what you need, when you need it, wherever you are.

A personal touch from people who know bleeding disorders.

Janie Summers | 1 314 304-6424

accredo®

© 2013 Accredo Health Group, Inc. An Express Scripts Company. All Rights Reserved.

ARJ Infusion SERVICES

Where caring is...



...the ultimate Factor

866.451.8804

www.arjinfusion.com

- Award Winning Nursing
- Licensed Home Health Agency
- Pharmacy Onsite
- 24 Hour On-call Service
- Reimbursement Experts
- Infusion Education for Patients and Families



Specializing In ...

Hemophilia
Bleeding Disorders
Von Willebrand's Disease

Relax...

We take the worry out of living with hemophilia and other bleeding disorders, so you can enjoy life.



Peace of mind for your healthcare needs.

We're looking out for you.

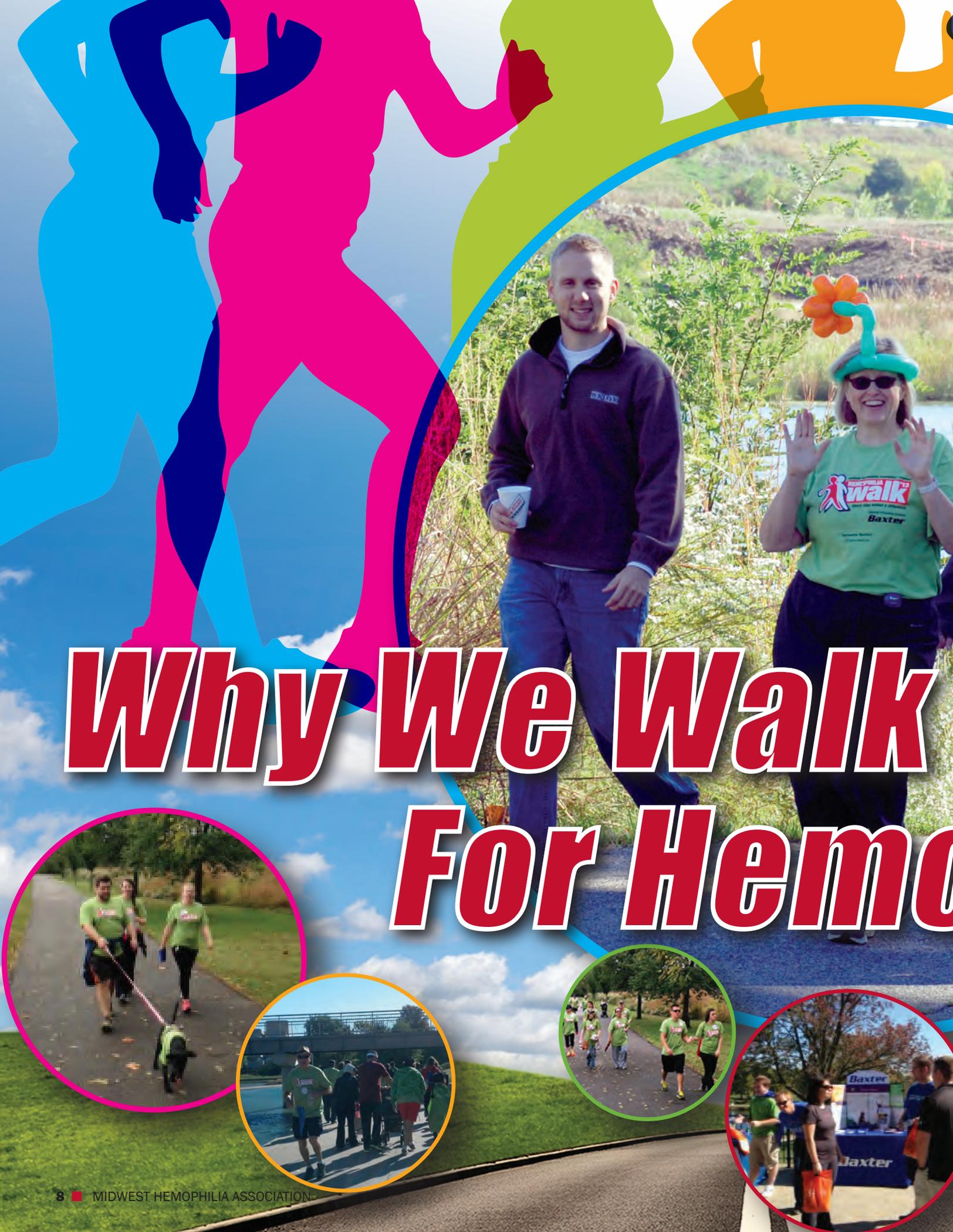
- An individual case member will assist you, advise you and negotiate for you.
- Prescription fulfillment
- Product delivery
- Full line of factor products and custom foil-wrap packaging that maintains proper temperature
- Insurance reimbursement & care provider issues

We're ready to start serving you today.

www.mwcornerstone.com



Mimi Law • Anna MacDonald • Craig Looney • Angela Brown
(877) 646-4529 "Se Habla Español"



Why We Walk For Hemophilia





This is my seventh year to participate in a walk for hemophilia and other bleeding disorders. But the reason why I walk began much earlier.

Hemophilia first entered my life when I was twelve years old. My brother, who was nine months old when he was diagnosed, was the first to have a bleeding disorder in our family. It was scary. We lived in a rural community far from specialized doctors, hemophilia treatment centers, and other families like us. We did not know of education programs, local chapters, or helpful resources.

My parents did the best they could with their given circumstances. My siblings and I watched and learned what we could alongside of them.

Fifteen years later when I delivered our son, it was still a shock hearing the words, “he has hemophilia.” After reliving the same emotions I experienced with my brother’s diagnosis, I determined in my mind that our family goal would to become active members of this community no matter how difficult. We would do it for my brother, our son, and for our family’s future.

We started by educating

ourselves through books, the internet, medical team members, and attending a national annual meeting. We listened to others’ stories and experiences. Then we brought that information back to our local doctors, hospitals, and extended family members. Anyone who would listen, heard our story. Next, we became members of our ‘local’ chapter and began attending educational events. Now, as our son grows, we teach him about his bleeding disorder and send him to hemophilia camp. We continue to be part of the community when and wherever we can.

However, the goal for our family has changed. People before us have sacrificed their time, energy, financial resources, and dare I say, their health, to give us the opportunity to successfully manage our bleeding disorder and gain the freedom and power that education brings with it. Yes, it is important to continue our participation, but it is just as important to give back to the community who has given us so much. It’s just not about us.

So...why do we walk for hemophilia? We walk to honor those who have paved the way before us. We walk to widen the road to include those around us now, in and outside the community. We walk to ensure the road continues and grows easier for those in the future. ■



hemophilia

By Angela Brown



Now Available

A new treatment for hemophilia B

ALPROLIX provides protection* from bleeds starting with at least a week between prophylaxis infusions.

Dosing regimen can be adjusted based on individual response.

*Protection is the prevention of bleeding episodes using a prophylaxis regimen.



To learn more, contact CoRe Manager **Vicki Oberkrom**

E: vicki.oberkrom@biogenidec.com T: 660.281.2447

Indications and Important Safety Information

Indications

ALPROLIX, Coagulation Factor IX (Recombinant), Fc Fusion Protein, is a recombinant DNA derived, coagulation factor IX concentrate indicated in adults and children with hemophilia B for:

- Control and prevention of bleeding episodes
- Perioperative management
- Routine prophylaxis to prevent or reduce the frequency of bleeding episodes

ALPROLIX is not indicated for induction of immune tolerance in patients with hemophilia B.

Important Safety Information

Do not use ALPROLIX if you are allergic to ALPROLIX or any of the other ingredients in ALPROLIX.

Tell your healthcare provider if you have or have had any medical problems, take any medicines, including prescription and non-prescription medicines, supplements, or herbal medicines, have any allergies and all your medical conditions, including if you are pregnant or planning to become pregnant, are breastfeeding, or have been told you have inhibitors (antibodies) to factor IX.

Allergic reactions may occur with ALPROLIX. Call your healthcare provider or get emergency treatment right away if you have any of the following symptoms: difficulty breathing, chest tightness, swelling of the face, rash, or hives.

Your body can also make antibodies called “inhibitors” against ALPROLIX, which may stop ALPROLIX from working properly.

ALPROLIX may increase the risk of formation of abnormal blood clots in your body, especially if you have risk factors for developing clots.

Common side effects of ALPROLIX include headache and abnormal sensation of the mouth. These are not all the possible side effects of ALPROLIX. Talk to your healthcare provider right away about any side effect that bothers you or does not go away, and if bleeding is not controlled using ALPROLIX.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch, or call 1-800-FDA-1088.

Please see **Brief Summary of full Prescribing Information on the next page.** This information is not intended to replace discussions with your healthcare provider.

ALPROLIX [Coagulation Factor IX (Recombinant), Fc Fusion Protein], Lyophilized Powder for Solution For Intravenous Injection.

FDA Approved Patient Information

ALPROLIX™ /all' pro liks/ [Coagulation Factor IX (Recombinant), Fc Fusion Protein]

Please read this Patient Information carefully before using ALPROLIX™ and each time you get a refill, as there may be new information. This Patient Information does not take the place of talking with your healthcare provider about your medical condition or your treatment.

What is ALPROLIX™?

ALPROLIX™ is an injectable medicine that is used to help control and prevent bleeding in people with hemophilia B. Hemophilia B is also called congenital Factor IX deficiency.

Your healthcare provider may give you ALPROLIX™ when you have surgery.

Who should not use ALPROLIX™?

You should not use ALPROLIX™ if you are allergic to ALPROLIX™ or any of the other ingredients in ALPROLIX™. Tell your healthcare provider if you have had an allergic reaction to any Factor IX product prior to using ALPROLIX™.

What should I tell my healthcare provider before using ALPROLIX™?

Tell your healthcare provider about all of the medicines you take, including all prescription and non-prescription medicines, such as over-the-counter medicines, supplements, or herbal medicines.

Tell your doctor about all of your medical conditions, including if you:

- are pregnant or planning to become pregnant. It is not known if ALPROLIX™ may harm your unborn baby.
- are breastfeeding. It is not known if ALPROLIX™ passes into breast milk or if it can harm your baby.
- have been told that you have inhibitors to Factor IX (because ALPROLIX™ may not work for you).

How should I use ALPROLIX™?

ALPROLIX™ should be administered as ordered by your healthcare provider. You should be trained on how to do infusions by your healthcare provider. Many people with hemophilia B learn to infuse their ALPROLIX™ by themselves or with the help of a family member.

See the Instructions for Use for directions on infusing ALPROLIX™. The steps in the Instructions for Use are general guidelines for using ALPROLIX™. Always follow any specific instructions from your healthcare provider. If you are unsure of the procedure, please ask your healthcare provider.

Do not use ALPROLIX™ as a continuous intravenous infusion.

Contact your healthcare provider immediately if bleeding is not controlled after using ALPROLIX™.

What are the possible side effects of ALPROLIX™?

Common side effects of ALPROLIX™ include headache and abnormal sensation in the mouth.

Allergic reactions may occur. Call your healthcare provider or get emergency treatment right away if you have any of the following symptoms: hives, chest tightness, wheezing, difficulty breathing, or swelling of the face.

ALPROLIX™ may increase the risk of forming abnormal blood clots in your body, especially if you have risk factors for developing blood clots.

Your body can also make antibodies called, "inhibitors," against ALPROLIX™, which may stop ALPROLIX™ from working properly. Your healthcare provider may need to test your blood for inhibitors from time to time.

These are not all the possible side effects of ALPROLIX™. Talk to your healthcare provider about any side effect that bothers you or that does not go away.

How should I store ALPROLIX™?

Store ALPROLIX™ vials at 2°C to 8°C (36°F to 46°F). Do not freeze.

ALPROLIX™ vials may also be stored at room temperature up to 30°C (86°F) for a single 6 month period.

If you choose to store ALPROLIX™ at room temperature:

- Note on the carton the date on which the product was removed from refrigeration.
- Use the product before the end of this 6 month period or discard it. Do not return the product to the refrigerator.
- Do not use product or diluent after the expiration date printed on the carton, vial or syringe.

After Reconstitution:

- Use the reconstituted product as soon as possible; however, you may store the reconstituted product at room temperature up to 30°C (86°F) for up to 3 hours. Protect the reconstituted product from direct sunlight. Discard any product not used within 3 hours after reconstitution.
- Do not use ALPROLIX™ if the reconstituted solution is cloudy, contains particles or is not colorless.

What else should I know about ALPROLIX™?

Medicines are sometimes prescribed for purposes other than those listed here. Do not use ALPROLIX™ for a condition for which it was not prescribed. Do not share ALPROLIX™ with other people, even if they have the same symptoms that you have.

Manufactured by
Biogen Idec Inc.
14 Cambridge Center
Cambridge, MA 02142
U.S. License #1697

A visit to WFHF 2014

By Judy Kauffman, RN, CPNP
Kansas City Regional Hemophilia Center

I had the amazing opportunity to travel to Melbourne, Australia recently to attend my very first World Federation of Hemophilia Congress. A number of things had to come together to get me to Australia and I am really glad that I got the chance to do it. When people ask me how it was in Australia, I answer, "I learned A LOT!" Here is a partial list of what I learned and experienced:

A 15-hour plane ride is LONG! I thought it would never end! Total travel time to Melbourne was 25 hours. Whoa! Talk about jet lag!

Convention centers everywhere are the same – big and challenging to navigate!

Big cities are the same everywhere (sort of). I found that Australian English is very different than American English. I laughed every time I turned on the TV at some of the ways things are said there!

People with bleeding disorders



Photos courtesy of The Congress Daily: wfhcongressdaily.org

Welcome Reception at the World Federation of Hemophilia Congress held May 11-15 in Melbourne, Australia.

from other countries are the same AND different! At the WFH Congress, there is a treatment room where people can get evaluation and factor treatment for bleeding. There were people who had the money to travel to Australia at the meeting, but limited access to factor in the countries. Some of those took advantage of the chance to get factor and PT care in the treatment room.

Hemophilia doctors, nurses, physical therapists (they're called "physios" in many countries) are the same except for their accents. They care about providing the best treatment to their patients. Providers that I met were very focused on learning all they could at the meeting.

I went to many great sessions at the Congress and learned a lot myself. Some of what I learned was about new factor products, inhibitors, lab testing and von Willebrand disease.

I learned that before getting on a bus in a new city that is very far away from home, one should ask when the bus quits for the day, because the one I was on just dropped us off (where, we weren't sure) saying "that's all for today." That was a bit of an adventure trying to find my way back to the hotel.

I learned that Australia has these really cute little penguins, sometimes called "fairly penguins," that live on their shore. People can watch them every day as they march up the beach to their burrows on land. They were cute and it was fun to watch them. We also got to see some koalas, kangaroos and wallabys.

One of the reasons that I was able to attend the WFH Congress is to prepare for the fact that in 2016 the National Hemophilia Foundation (NHF) will host the WFH meeting in Orlando, Florida! That represents an amazing opportunity for Americans with bleeding disorders to go to a WFH meeting without spending an arm and a leg on airfare. The meeting will be in July 2016 and I encourage everyone to consider saving your nickels, dimes and pennies so that you might be able to attend. Meeting people from all over the world with bleeding disorders is a great experience and one that I really recommend. ■





RIXUBIS

[COAGULATION FACTOR IX
(RECOMBINANT)]

**For more information, contact your
Baxter representative today:**

Jana Zens

Phone: (913) 558-4913

E-mail: jana_zens@baxter.com

To learn more, visit www.RIXUBIS.com.



Inhibitor Study Stresses Screening Importance

Results from a six-year study of patients with hemophilia A and B produced interesting findings. The Hemophilia Inhibitor Research Study (HIRS) enrolled 1,163 patients from 17 federally funded hemophilia treatment centers (HTCs). One of the goals was to predict

which patients were at highest risk for development of inhibitors, antibodies to infused factor.

“A Study of Prospective Surveillance for Inhibitors Among Persons with Haemophilia in the United States,” was published in the March 2014 issue of *Haemophilia*. The lead investigator was Michael Soucie, PhD, Division of Blood Disorders, National Center on Birth Defects and Developmental Disabilities, Centers for Disease Control and Prevention (CDC) in Atlanta.

A central laboratory

performed periodic inhibitor tests using blood samples and genotyped the subjects. In all, 3048 inhibitor tests (some patients were screened more than once) were conducted. The main findings were:

- All people with hemophilia are at risk for developing inhibitors
- One-third of newly developed inhibitors were found in people with non-severe hemophilia
- One-half were older than 5 years old
- Six out of 10 people with hemophilia with an inhibitor had no symptoms
- 23 new FVIII inhibitors were identified
- 431 distinct mutations were genotyped, 151 of which had not previously been reported

HIRS investigators and CDC researchers determined that individuals with hemophilia of all ages were at risk for developing an inhibitor. Further, CDC now estimates that approximately 60% of people with an inhibitor have no symptoms. Without regular screening, a significant number of these patients may not be aware of it until they experience severe bleeding.

The CDC concluded that patients with hemophilia receiving care in federally funded HTCs will be tested yearly for an inhibitor by the CDC Division of Blood Disorders laboratory as part of Community Counts, its new blood monitoring program. ■

— Source: CDC



Your family deserves an
upgrade in service.



Call our 24/7 hotline to experience the
"No Phone Tree" Guarantee

888-335-4279

www accuraterx.net

intake@accuraterx.net

[@AccurateRx](https://twitter.com/AccurateRx)

[/AccurateRxPharmacy](https://www.facebook.com/AccurateRxPharmacy)

ACCURATE Rx
PHARMACY
SPECIALIZING IN NURSING & INFUSION SERVICES

CVS Specialty
CAREMARK Pharmacy

Trust the Experience

At **CVS Caremark**, we've been helping families like yours for more than 30 years. Our caring patient support helps ensure safety, convenient access and satisfaction.

Stephanie Lasister, Regional Sales Manager
1-816-668-5200
stephanie.lasister@cvscaremark.com

www.CVSCaremarkSpecialtyRx.com



©2014 Caremark. All rights reserved.
106-30296a 030514

LIMITLESS.

The possibilities when you're well cared for.



Participating Tricare Hemophilia Provider

At Biomed, our commitment to the bleeding disorder community doesn't end with **skilled pharmacists**, **experienced infusion nurses**, or **dedicated customer service associates**. It continues through education, training, and involvement at the local and national level by Biomed's team of advocates who have extensive experience in the community. Improving the lives of patients and caregivers for a future of unlimited possibilities is our passion.

To learn more about Biomed bleeding disorder programs, contact **John Kowalski** or **Ed Chomyak** at **866-661-0110**.

BIOMED
PHARMACEUTICALS

www.biomed-rx.com
866-661-0110
Follow us: [in](#) [f](#) [t](#)

MOVING? New Address

Name

Address

City

State

ZIP

(Area Code) Phone Number

E-Mail Address

MAIL TO:

MHA • PO Box 412866 • Kansas City, MO 64141



Midwest Hemophilia Association

PO Box 412866
Kansas City, MO 64141

IT'S TIME TO WALK!

WALK LOCATIONS AND DATES

Wichita, KS: Exploration Place
Sunday, October 12, 2014

Kansas City, MO: Waterfall Park
Saturday, October 18, 2014

Springfield, MO: Botanical Garden
Saturday, October 18, 2014

Columbia, MO: Stephens Lake Park
Sunday, October 19, 2014

Join us to raise funds and awareness
for the bleeding disorder community.
Your support is vital to our success.

To become a Team Captain, or to volunteer in your area
contact Mark Cox, Executive Director at **816-479-5900** or
email at events@midwesthemophilia.org. Please visit
www.hemophilia.org/walk to register yourself or your team.

