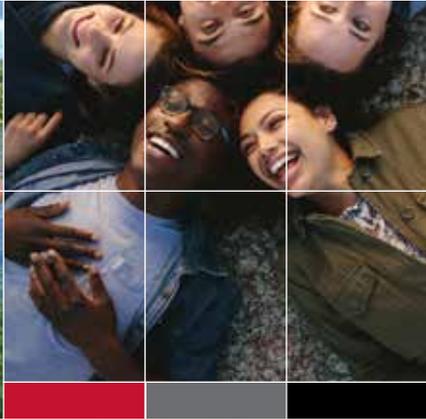




**Midwest
Hemophilia
Association**



2019



**WELCOME!
YOU BELONG HERE**

**27TH ANNUAL
Family Education Weekend
September 14 & 15, 2019**



Table of Contents

Welcome Letter..... 1

Board of Directors..... 2 - 3

Fundraising Events: Unite for Bleeding Disorders Walk-A-Thon..... 4

Fundraising Events: Annual Golf Tournament & Trivia Night..... 5

Educational Events: Summer Camp..... 6

Educational Events: Community Outreach 7

Community Support: Advocacy 8

Community Support: Scholarships 9

2018 Annual Report 10 - 11

Family Education Weekend Sponsors 12 - 13

Family Education Weekend Agenda..... 14 - 15

2019 Donors & Sponsors 16



Hello.

A simple word used globally in a variety of different languages. Bonjour. Hola. Jambo. It is a quick greeting, a salutation at introductions, or a brief conversation.

Yet there is so much potential in the word Hello. It can lead to the beginning of a friendship, a new career, a business partnership, or even a life-long relationship.

It is how Midwest Hemophilia Association would like to start our relationship with you.

When we say Hello to you we look forward to the potential it brings. We want our Hello to go beyond just a quick greeting. We want our Hello to be the start of a life-long relationship that includes you as part of our family.

Inside this brochure is the our way of moving to next step after Hello: Welcome. It is a brief overview of the Midwest Hemophilia Association family, a sort of family album of our advocacy efforts, education programs, fundraising endeavors, and the building of our family through outreach and resources.

Please take look, browse through the pages. It is our hope that you will find your place within the family.

Hello. Welcome to our family.

Sincerely,



Angela Brown
MHA Executive Director





Executive Director

Angela Brown

angela.brown@midwesthemophilia.org

Angela Brown was selected as Executive Director of Midwest Hemophilia Association in July 2019 by the Board and hiring committee. Angela lives in Ozark, Missouri, with her husband, Matt, and four children, one of whom has a bleeding disorder.

Prior to this position, Angeala worked for the midwest division of a specialty pharmacy to develop quality relationships and service with the bleeding disorders community, chapter leaders and other industry representatives. Using her knowledge and experience with IEPs and 504s, she assisted families with their health and school-related needs to improve their experiences and communication with school personnel. Angela has been a member of MHA for over 15 years, working in several different capacities: participant, volunteer, fundraiser, board member, and event planner. She also has background in elementary and special education.

Angela grew up with a brother who has a bleeding disorder, but says she gained a whole new vantage point when her son was diagnosed with the bleeding disorder. Angela explains that her family initially faced many of the common challenges: mis-diagnosis, doctors with little knowledge, long ER visits, and fear of the unknown. They found hope, peace of mind, freedom and independence, however, when they were introduced to the local and national chapters for bleeding disorders and began to attend events provided by MHA. Angela says, "Because of what I learned and experienced through these groups, I knew I wanted to be an active part of the MHA."

Angela strives to pass on the knowledge and resources that benefited her family as well as sharing up-to-date information on product and treatment options, insurance information, and other related health issues. She believes that this information is helpful to newly diagnosed families, imperative for maintaining support networks to experienced families, and essential to sustain our future generations. Angela's goal as Executive Director is to continue to provide the bleeding disorders community with tools through educational events, outreach groups, advocacy leadership, and summer camp.



2019 Board of Directors

BOARD OFFICERS



Nora Ancel
Vice President
nora.e.ancel@gmail.com



Jaci Colter
Treasurer
treasurer@midwesthemophilia.org



Bridget Castro
Secretary
brdgt_c@yahoo.com

BOARD MEMBERS



Jennifer Rentschler
jenr0429@gmail.com



Dakota Rosenfelt
dakotarosenfelt@me.com



Kimberly Rosenfelt
krosenfelt@sbcglobal.net



Kyrie Smith
kyrielynn24@gmail.com



Fundraising Events:

UNITE for Bleeding Disorders Walk-A-Thon

The Unite For Bleeding Disorders Walk-A-Thon is a nationwide event held throughout the year in over 40 cities each year to raise funds and awareness for all bleeding disorders. Participants raise money for crucial research, advocacy and education by joining together in the fun and festive event.

Our chapter holds its walk each fall. The 2019 walk took place on Saturday, August 24 at Shawnee Mission Park in Shawnee, Kansas. Over 100 participants gathered to promote awareness and bring in donations.

The walk-a-thon is the primary fundraising event for MHA. 100% of every dollar raised stays here with our chapter and goes to directly support our advocacy, education and research initiatives. We strongly encourage industry partners to support our local walk and ask our members to seek out sponsorships from local companies.

This event is crucial to our budget and we invite all our members to lace up their shoes and join us every year!



Fundraising Events:

Annual Golf Tournament & Trivia Night

In addition to the UNITE for Bleeding Disorders Walk-A-Thon, MHA hosts two additional fundraising events each year: Trivia Night held in the spring and the annual MHA Golf Tournament held in the fall.

Displays of knowledge and lots of laughter abound as participants battle for prizes – all while raising money for MHA. Trivia Night brings together community and non-community members to enjoy food and fun. This year's event was held in May at Parlor KC restaurant. Funds raised from entry fees and donations benefit our local chapter.

The MHA Annual Golf Tournament is in its 21st year supporting bleeding disorders and offers a variety of sponsorship opportunities to support MHA. The 2019 contest is being held at WinterStone Golf Course in Independence, Missouri. The day starts with a continental breakfast before a shotgun start. Participants compete for team and individual prizes and raffles. An awards luncheon follows at the conclusion of play.





Educational Events:

Summer Camp

MHA's Summer Camp is for kids, ages 7-17, living with hemophilia and other bleeding disorders and traditionally occurs in late summer. Its mission is to provide campers with a fun, safe, and educational experience along with the opportunity to meet other young people with rare bleeding disorders. The 2019 camp was held July 29th through August 2nd at Lake Doniphan Retreat and Conference Center in Excelsior Springs, MO. This year's theme was the Oregon Trail.

In addition to the traditional camp activities, campers receive onsite medical care and education, including self-infusion training. Another educational focus is teaching campers about healthy eating and the benefits of maintaining an active lifestyle.

For most participants, summer camp is the only opportunity to meet peers who also have a bleeding disorder. Camp is that special place where they can discuss their experiences of living with a bleeding disorder, and know that there are others close by that are able to listen and understand.



Educational Events:

Community Outreach

Educational workshops are held each year throughout our coverage area that enable the bleeding disorders community to come together to exchange information on a wide variety of topics. The current year will include sessions held in Kansas City, Missouri; Springfield, Missouri; Dodge City, Kansas; and Wichita, Kansas.

The Springfield workshop was held April 13th. Attendees participated in advocacy bingo and heard from brand sponsors. Following lunch, education sessions and exhibits, participants were given tickets to a Springfield Cardinals baseball game.

The Dodge City education event on June 15th revolved around a Minute to Win It theme. Community members played games, listened to brand sponsor speakers and viewed exhibits. Access to Long Branch Lagoon water park was provided after the presentations.

Wichita will host the final workshop of 2019 on October 19th. National Hemophilia Foundation staff will discuss acronyms associated with bleeding disorders plus talk about choosing a factor provider. The education day will dismiss to the Wichita Zoo following lunch and presentations.





Community Support:

Advocacy

MHA advocates for the unique needs and interests of the people affected by bleeding disorders by educating the general public, elected officials and others in government. In conjunction with the National Hemophilia Foundation (NHF), government funding has been used to improve medical care, services, education, and safety of blood and blood products as a result of these efforts.

MHA partners with Gateway Hemophilia Association in St. Louis, Missouri to form the Missouri Bleeding Disorder Advocacy Coalition. The two organizations come together each year for Missouri Advocacy Day in Jefferson City, Missouri. Also known as “Red Tie Day,” advocates from all over the state meet with state officials to chat and ask questions important to the bleeding disorders community. A training session is offered the night before for attendees to prepare for the event.

Representatives of MHA also advocate on a federal level at NHF’s Washington Days held in the nation’s capitol. The event provides the opportunity to discuss key issues impacting the community with federal officials while putting a face on bleeding disorders.



Community Support:

Scholarship

From our inception as an organization in 1990, MHA has always believed in education. One way we encourage our younger community members to continue their education is by providing scholarships to assist with the costs of secondary education.

The Midwest Hemophilia Association Scholarship was established and has been offered for the past 25+ years. In 2016, the Board authorized a second scholarship, the Mark Dudley Scholarship, in honor of our past board treasurer. These two scholarships together have awarded over \$30,000 in educational funding.

MHA also offers the Georgia Northway Scholarship, which is managed by the Truman Heartland Community Foundation. The scholarship was established when MHA was named an estate beneficiary of Georgia Northway from the Wichita, Kansas area. Ms. Northway had two sons that had passed away due to complications from hemophilia. It was her wish that part of her estate be utilized to create a scholarship fund for individuals with bleeding disorders. Truman Heartland received \$423,924.87 in 2003 to start the scholarship. Each year, approximately 5% of the principal balance is available for scholarships, resulting in awards of \$269,188.44 in 108 scholarships to over 38 students. The great part of the Georgia Northway Scholarship is that it is renewable. Recipients can re-apply each year, potentially receiving funding beyond freshman year.

Truman Heartland has done an exceptional job of managing this fund and the increase in the last 17 years of the stock market has helped as well. As of June 30, 2019, the balance in the fund is \$482,477, meaning scholarships will be offered for many years to come. The fund should have \$23,000 to \$25,000 available for grants in the 2020-21 school year.

Deadlines for all three scholarships are in late spring each year. Details and applications can be found on MidwestHemophilia.org.

2019-2020 Georgia Northway Scholarship Recipients:

William Carter, Kolbe Madden, Ramsey Divine, and Elizabeth Hancock

2019-2020 Midwest Hemophilia Association Scholarship Recipient:

Quintin Castro

2019-2020 Mark Dudley Academic Scholarship Recipient:

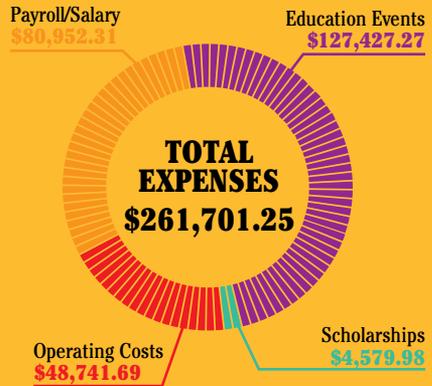
Ramsey Divine

2018 Annual Report

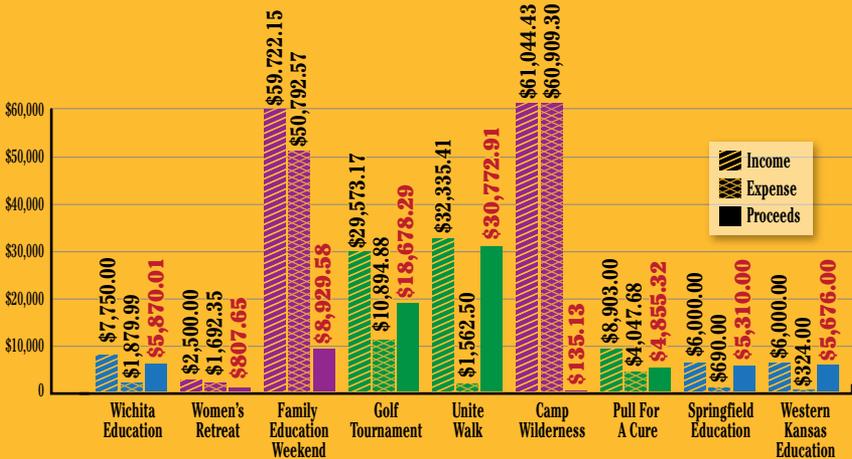
Where the Money Comes From



Where the Money Goes



Event Breakdown by Dollars

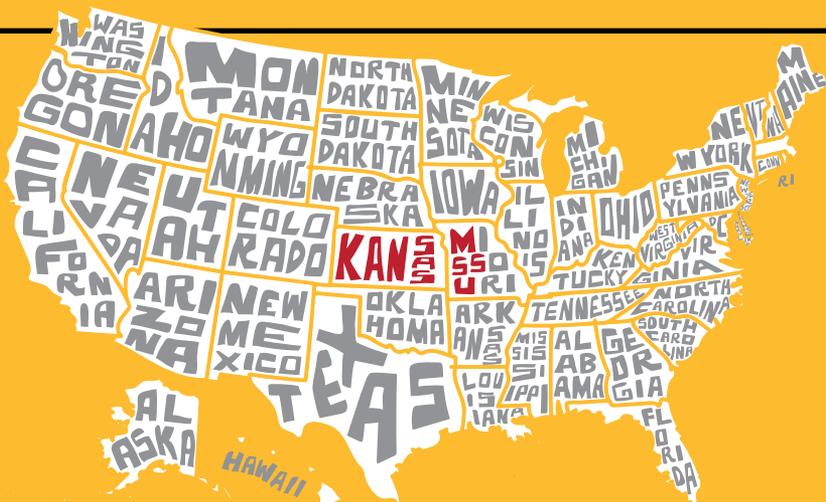


Approximate number of people who attend MHA's events in 2018



Number of events hosted by MHA in 2018 throughout the coverage area





The Midwest Hemophilia Association supports the bleeding disorders communities in **KANSAS** and **WESTERN MISSOURI**.



Hemophilia is a rare inherited blood disorder which affects **over 20,000 people** in the United States.



NATIONAL HEMOPHILIA FOUNDATION
for all bleeding disorders

Midwest Hemophilia Association is a **CHAPTER CHARTER MEMBER** of the National Hemophilia Foundation. The **National Hemophilia Foundation**, established in 1948, has chapters throughout the country. It is dedicated to finding cures and treatments for inheritable bleeding disorders and preventing complications through education, advocacy and research.



People in **KANSAS** living with a bleeding disorder



People in **MISSOURI** living with a bleeding disorder



Family Education Weekend Sponsors

DIAMOND

Bayer
CVS Specialty
Genentech
Sanofi Genzyme

PLATINUM

Pfizer
Soleo Health
Takeda

GOLD

HPC Specialty Pharmacy

BRONZE

National Hemophilia Foundation

SILVER

Accredo • Aptevo Therapeutics • ARJ Infusion Services
BioEthics Advantage • Brothers Healthcare
Children's Mercy 340B Factor Program • CSL Behring
Factor One Source • Grifols
Novo Nordisk • Superior Biologics



Family Education Weekend Sponsors

BAYER is dedicated to providing treatment options for people living with rare, serious and life-threatening hematologic diseases. Bayer has been a committed partner to the hemophilia community delivering therapeutic options, educational and support programs and continuing research designed to meet the needs of patients throughout their life journey.

GENENTECH Since 1976, Genentech has stood for scientific innovation that improves the world around us. As part of the Roche Group, we are among the world's leading biotech companies that discovers, develops, manufactures and commercializes medicines to treat patients with serious medical conditions. Our history in hemophilia research dates back to 1984. Today, our community commitment endures, with our Hemophilia community Clinical Educators acting as educational and informational resources for patients with hemophilia A and their caregivers.

SANOFI GENZYME, the specialty care global business unit of Sanofi, focuses on rare diseases, rare blood disorders, multiple sclerosis, oncology, and immunology. We help people with debilitating and complex conditions that are often difficult to diagnose and treat. Our approach is shaped by our experience developing highly specialized treatments and forging close relationships with physician and patient communities. We are dedicated to discovering and advancing new therapies, providing hope to patients and their families around the world.

SOLEO HEALTH is an innovative national provider of home and alternate-site specialty infusion. Our team of experienced clinicians provides exceptional care in managing complex disease states through comprehensive pharmacy, nursing, education, and therapy management programs. Our Mission is to enhance our patients' experience and health through innovative and cost-effective solutions, resulting in exceptional clinical outcomes measured by data-driven quality programs which will afford us the opportunity to be the employer, partner and service provider of choice.

TAKEDA is a global, values-based, R&D-driven biopharmaceutical leader headquartered in Japan, committed to bringing Better Health and a Brighter Future to patients by translating science into highly-innovative medicines. Takeda focuses its R&D efforts on four therapeutic areas: Oncology, Gastroenterology (GI), Neuroscience and Rare Diseases. We also make targeted R&D investments in Plasma-Derived Therapies and Vaccines.



Family Education Weekend Agenda

Saturday, September 14

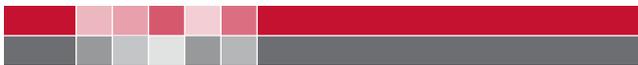
8:00am	Exhibit Hall Open & Event Check-in	
8:30am	Childcare Check-in: Infants to 5yrs	Salon C
	Youth Check-in: Ages K-5th grade	Salon F
9:00am	Teen Check-In: Ages 6th-12th grade	Santa Fe Room
10:00am	Buses Depart for Zoo & Power Play (K-12th)	
11:00am -12:00pm	Opening Ceremony & Lunch <i>Exhibit Hall Closed</i>	
12:10am -1:00pm	Healthy Aging	Salon D
	MyBDC Presentation	Salon E
	Spanish Breakout: Como Abogar Por Si Mismo	Boardroom
1:10-2:00pm	Pain Matters	Salon D
	MyBDC Presentation	Salon E
	An Overview of VWD & Women's Reproductive Issues	Boardroom
2:00-2:30pm	Break & Exhibit Hall Open	
2:30-3:50pm	Women's Breakout: M-Powered	Salon D
	Men's Breakout: Change Can Be Beneficial Hector G.	Salon E
4:00-5:30pm	The Science of Optimism	Salon D
	NHF: Artistically Speaking	Salon E
4:30pm	Buses Return to Hotel K-5th grade	Salon F
	Buses Return to Hotel 6th-12th grade	Santa Fe Room
5:30 pm	Childcare Pick-up: Infants to 5yrs	Salon C
	Youth Pick-up: Ages K-5th grade	Salon F
	Teen Pick-up: Ages 6th-12th grade	Santa Fe Room
6:00pm	Dinner & Evening Activity @ Dave & Busters 6811 W. 135th Street Overland Park, KS 66223	



Family Education Weekend Agenda

Sunday, September 15

8:00-8:50am	Breakfast & Exhibit Hall Open <i>Diamond, Platinum & Gold Sponsors Only</i>	
PLEASE EAT BREAKFAST PRIOR TO CHILD CHECK-IN!		
8:00-8:50am	Childcare Check-in: Infants to 5yrs	Salon C
	Youth Check-in: Ages K-5th grade	Salon F
8:45am	Teen Check-In: Ages 6th-12th grade	Salon A
9:00-9:50	Evaluating Your Insurance Options	Salon D/E
	Teen Session 6th-12th grade: The Road to Independence	Salon A
	Youth Session 3-5th grade: Factor is Gold!	Santa Fe Room
10:00-10:50	Gene Therapy with Dr. Brian Wicklund	Salon D/E
	Teen Session 6th-12th grade: Change & Resilience	Salon A
	Youth Session K-2nd grade: Factor is Gold!	Santa Fe Room
PLEASE PICK UP K-12TH GRADE CHILDREN PRIOR TO CLOSING CEREMONY!		
11:00-11:55am	CLOSING CEREMONY: Awards & Recognition <ul style="list-style-type: none"> • <i>Scholarship Winners</i> • <i>UNITE for Bleeding Disorders Top Team & Individual Fundraisers</i> • <i>Golf Tournament Award Winners</i> • <i>Summer Camp Big Stick Awards</i> • <i>2019 Volunteer of the Year</i> • <i>Bleeding Disorder Community Lifetime Service Recognition</i> Raffle & NHF Trip Drawing	Salon D/E
12:00pm	Dismiss & Hotel Check-out	
12:00pm	Childcare Pick-up: Infants to 5yrs	Salon C



2019 Donors & Sponsors

Thank you to our sponsors and donors for your dedication to Midwest Hemophilia Association and your passion for the mission of our organization. Your involvement is very important to the long-term health of the MHA and the fulfillment of its mission.

SPRINGFIELD EDUCATION

Bayer
CSL Behring
Genentech
Takeda
ARJ Infusion Service
BioEthics Advantage

DODGE CITY EDUCATION

CSL Behring
CVS Specialty
Novo Nordisk
BioEthics Advantage
FactorOne Source
Innovative Infusion
Specialists
ARJ Infusion Service
Superior Biologics
Brothers Healthcare

WICHITA EDUCATION

Accurate Rx
Bayer
BioEthics Advantage
Brothers Healthcare
FactorOne Source
Genentech
Sanofi Genzyme
Takeda

SUMMER CAMP

Takeda
Soleo Health
CSL Behring
Novo Nordisk
Bayer
Sanofi Genzyme
R.A. Long Foundation
ARJ Infusion Service

Paragon
Aptevo
Innovative Infusion
Specialists
CVS Specialty
Bob Eschbacher
Superior Biologics
Brothers Healthcare

TRIVIA NIGHT

Central Plains
Cement Company
Sporting KC
Niki Bird
CSL Behring
CVS Specialty
ARJ Infusion Service
BioEthics Advantage

UNITE WALK-A-THON

KC Nannies
Kendra Scott
Flicsha Roberson
Takeda
CVS Specialty
Grifols
Pfizer
Bayer
Briova Rx
Sanofi Genzyme
Accredo
Soleo Health
CSL Behring
ARJ Infusion Services
BioEthics Advantage
Brothers Healthcare
FactorOne Source

GOLF TOURNAMENT

Aptevo
ARJ Infusion Service
Bayer
BioEthics Advantage
Children's Mercy 340B
Factor Program
CSL Behring
Novo Nordisk
Soleo Health
Superior Biologics

We're here for you!

Experts in treating bleeding disorders

ARJ specializes in treating hemophilia, Von Willebrand, and other inherited or acquired complex bleeding disorders.

Rely on ARJ

Dedicated care coordinators are ready to help you and your family with:

- Lifestyle resources
- Financial assistance
- Benefits review



The ARJ ReadyPack®

- Factor and supplies
- Tear-away packaging
- Easy-to-read labels

Learn more

866-451-8804

arjinfusion.com/getready





FEEL EMPOWERED
to step up to the challenge
with Jivi®

Are you ready to see what Jivi offers?
Visit www.explorejivi.com to learn more.

Ask your doctor if Jivi may be right for you.

© 2019 Bayer. All rights reserved. Bayer, the Bayer Cross
and Jivi are registered trademarks of Bayer.

Printed in USA 02/19 PP-JIV-US-0467-1

Jivi
antihemophilic factor
rimbunim) PEGylated-aud

LET'S GO



Brothers

HEALTHCARE



Welcome

Mimi, Anna & Noel to BROTHERS HEALTHCARE

MIMI LAW: 314.920.3081 | mimil@brothershealthcare.com

ANNA MACDONALD: 760.540.3118 | annam@brothershealthcare.com

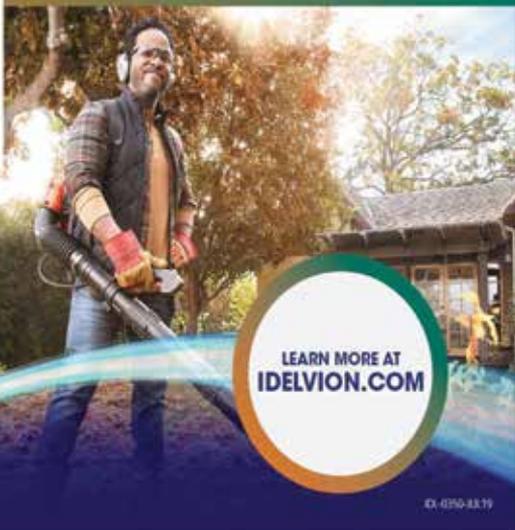
NOEL MINOR, RN, BSN: 316.866.0114 | noelm@brothershealthcare.com

BONDED BY BLOOD

BROTHERSHEALTHCARE.COM | 800.291.1089



IDELVION
Coagulation Factor IX (Recombinant), Albumin Fusion Protein



LEARN MORE AT
IDELVION.COM

ID-0390-83.19

Biologics for Life™ **CSL Behring**



Learn more at
AFSTYLA.COM

AFSTYLA
Antihemophilic Factor
(Recombinant), Single Chain

AFSTYLA (AF016)

Do you have a Severe Factor Deficiency?

Specifically searching for V, VII, VIII, IX, and XI

Join our paid plasma donation program! Make a difference! George King Bio-Medical, Inc. is searching for persons who have a Severe Factor Deficiency.

- Your health and safety are our primary concern.
- Upon acceptance into our program, donors are generously compensated for their plasma donation.
- All associated expenses are paid.
- Visit our website at www.kingbiomed.com to learn more about us.
- Call 800-255-5108 or email plasma@kingbiomed.com if you are interested.

Your plasma will be used in laboratories to aid in the diagnosis of others. It may also be used in research labs seeking improved tests and better treatments. Human plasma is the best source for this critical laboratory product. George King Bio-Medical, Inc. has been a leader in the clinical community since 1973.

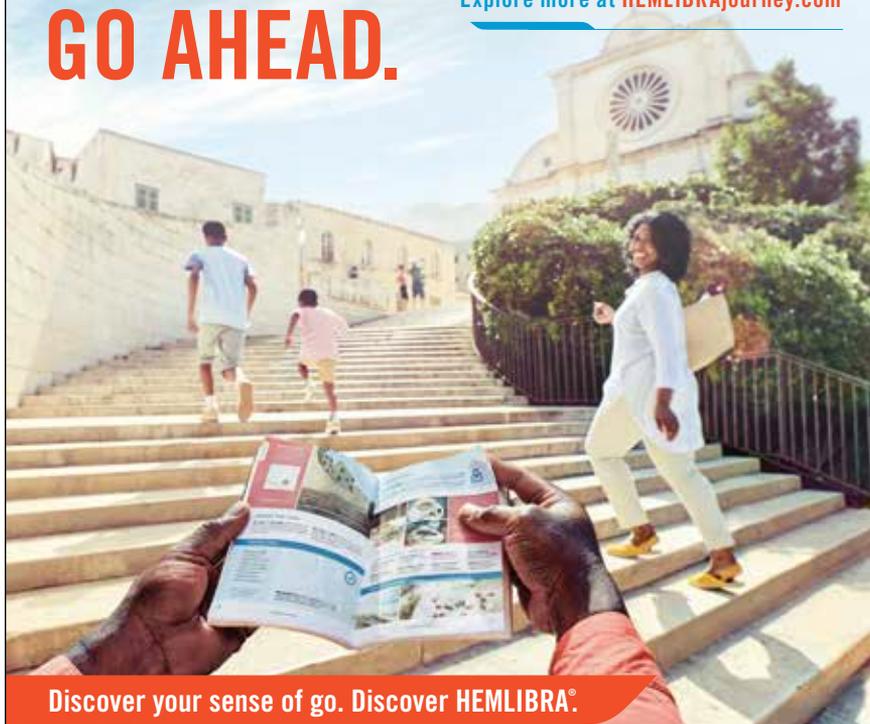


11771 W 112th Street Overland Park, KS 66210 www.kingbiomed.com 800-255-5108



GO SEEK. GO EXPLORE.
GO AHEAD.

PEOPLE LIKE YOU. STORIES LIKE YOURS.
Explore more at HEMLIBRAjourney.com



Discover your sense of go. Discover **HEMLIBRA**®.

What is **HEMLIBRA**?

HEMLIBRA is a prescription medicine used for routine prophylaxis to prevent or reduce the frequency of bleeding episodes in adults and children, ages newborn and older, with hemophilia A with or without factor VIII inhibitors.

What is the most important information I should know about **HEMLIBRA**?

HEMLIBRA increases the potential for your blood to clot. Carefully follow your healthcare provider's instructions regarding when to use an on-demand bypassing agent or factor VIII, and the dose and schedule to use for breakthrough bleed treatment. **HEMLIBRA** may cause serious side effects when used with activated prothrombin complex concentrate (aPCC; **FEIBA**®), including thrombotic microangiopathy (TMA), and blood clots (thrombotic events). If aPCC (**FEIBA**®) is needed, talk to your healthcare provider in case you feel you need more than 100 U/kg of aPCC (**FEIBA**®) total.

Please see Brief Summary of Medication Guide on following page for Important Safety Information, including **Serious Side Effects**.



Medication Guide HEMLIBRA® (hem-lee-bruh) (emicizumab-kxwh) injection, for subcutaneous use

What is the most important information I should know about HEMLIBRA?

HEMLIBRA increases the potential for your blood to clot. Carefully follow your healthcare provider's instructions regarding when to use an on-demand bypassing agent or factor VIII (FVIII) and the recommended dose and schedule to use for breakthrough bleed treatment.

HEMLIBRA may cause the following serious side effects when used with activated prothrombin complex concentrate (aPCC; FEIBA®), including:

- **Thrombotic microangiopathy (TMA).** This is a condition involving blood clots and injury to small blood vessels that may cause harm to your kidneys, brain, and other organs. Get medical help right away if you have any of the following signs or symptoms during or after treatment with HEMLIBRA:
 - confusion
 - weakness
 - swelling of arms and legs
 - yellowing of skin and eyes
 - stomach (abdomen) or back pain
 - nausea or vomiting
 - feeling sick
 - decreased urination
- **Blood clots (thrombotic events).** Blood clots may form in blood vessels in your arm, leg, lung, or head. Get medical help right away if you have any of these signs or symptoms of blood clots during or after treatment with HEMLIBRA:
 - swelling in arms or legs
 - pain or redness in your arms or legs
 - shortness of breath
 - chest pain or tightness
 - fast heart rate
 - cough up blood
 - feel faint
 - headache
 - numbness in your face
 - eye pain or swelling
 - trouble seeing

If aPCC (FEIBA®) is needed, talk to your healthcare provider in case you feel you need more than 100 U/kg of aPCC (FEIBA®) total.

See "What are the possible side effects of HEMLIBRA?" for more information about side effects.

What is HEMLIBRA?

HEMLIBRA is a prescription medicine used for routine prophylaxis to prevent or reduce the frequency of bleeding episodes in adults and children, ages newborn and older, with hemophilia A with or without factor VIII inhibitors.

Hemophilia A is a bleeding condition people can be born with where a missing or faulty blood clotting factor (factor VIII) prevents blood from clotting normally.

HEMLIBRA is a therapeutic antibody that bridges clotting factors to help your blood clot.

Before using HEMLIBRA, tell your healthcare provider about all of your medical conditions, including if you:

- are pregnant or plan to become pregnant. It is not known if HEMLIBRA may harm your unborn baby. Females who are able to become pregnant should use birth control (contraception) during treatment with HEMLIBRA.
- are breastfeeding or plan to breastfeed. It is not known if HEMLIBRA passes into your breast milk.

Tell your healthcare provider about all the medicines you take, including prescription medicines, over-the-counter medicines, vitamins, or herbal supplements. Keep a list of them to show your healthcare provider and pharmacist when you get a new medicine.

How should I use HEMLIBRA?

See the detailed "Instructions for Use" that comes with your HEMLIBRA for information on how to prepare and inject a dose of HEMLIBRA, and how to properly throw away (dispose of) used needles and syringes.

- Use HEMLIBRA exactly as prescribed by your healthcare provider.
- **Stop (discontinue) prophylactic use of bypassing agents the day before starting HEMLIBRA prophylaxis.**
- **You may continue prophylactic use of FVIII for the first week of HEMLIBRA prophylaxis.**
- HEMLIBRA is given as an injection under your skin (subcutaneous injection) by you or a caregiver.
- Your healthcare provider should show you or your caregiver how to prepare, measure, and inject your dose of HEMLIBRA before you inject yourself for the first time.
- Do not attempt to inject yourself or another person unless you have been taught how to do so by a healthcare provider.

- Your healthcare provider will prescribe your dose based on your weight. If your weight changes, tell your healthcare provider.
- You will receive HEMLIBRA 1 time a week for the first four weeks. Then you will receive a maintenance dose as prescribed by your healthcare provider.
- If you miss a dose of HEMLIBRA on your scheduled day, you should give the dose as soon as you remember. You must give the missed dose as soon as possible before the next scheduled dose, and then continue with your normal dosing schedule. **Do not** give two doses on the same day to make up for a missed dose.
- HEMLIBRA may interfere with laboratory tests that measure how well your blood is clotting and may cause a false reading. Talk to your healthcare provider about how this may affect your care.

What are the possible side effects of HEMLIBRA?

• See "What is the most important information I should know about HEMLIBRA?"

The most common side effects of HEMLIBRA include:

- redness, tenderness, warmth, or itching at the site of injection
- headache
- joint pain

These are not all of the possible side effects of HEMLIBRA.

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

How should I store HEMLIBRA?

- Store HEMLIBRA in the refrigerator at 36°F to 46°F (2°C to 8°C). Do not freeze.
- Store HEMLIBRA in the original carton to protect the vials from light.
- Do not shake HEMLIBRA.
- If needed, unopened vials of HEMLIBRA can be stored out of the refrigerator and then returned to the refrigerator. HEMLIBRA should not be stored out of the refrigerator for more than a total of 7 days or at a temperature greater than 86°F (30°C).
- After HEMLIBRA is transferred from the vial to the syringe, HEMLIBRA should be used right away.
- Throw away (dispose of) any unused HEMLIBRA left in the vial.

Keep HEMLIBRA and all medicines out of the reach of children.

General information about the safe and effective use of HEMLIBRA.

Medicines are sometimes prescribed for purposes other than those listed in a Medication Guide. Do not use HEMLIBRA for a condition for which it was not prescribed. Do not give HEMLIBRA to other people, even if they have the same symptoms that you have. It may harm them. You can ask your pharmacist or healthcare provider for information about HEMLIBRA that is written for health professionals.

What are the ingredients in HEMLIBRA?

Active ingredient: emicizumab-kxwh

Inactive ingredients: L-arginine, L-histidine, poloxamer 188, and L-aspartic acid.

Manufactured by: Genentech, Inc., A Member of the Roche Group,
1 DNA Way, South San Francisco, CA 94080-4990
U.S. License No. 1048

HEMLIBRA® is a registered trademark of Chugai Pharmaceutical Co., Ltd., Tokyo, Japan
©2018 Genentech, Inc. All rights reserved.

For more information, go to www.HEMLIBRA.com or call 1-866-HEMLIBRA.
This Medication Guide has been approved by the U.S. Food and Drug Administration
Revised: 10/2018



© 2019 Genentech USA, Inc. All rights reserved. EMI/061818/0106w/2
HEMLIBRA® is a registered trademark of Chugai Pharmaceutical Co., Ltd., Tokyo, Japan.
The HEMLIBRA logo is a trademark of Chugai Pharmaceutical Co., Ltd., Tokyo, Japan.
The Genentech logo is a registered trademark of Genentech, Inc.
All other trademarks are the property of their respective owners.

Genentech
A Member of the Roche Group

HEMOPHILIA SOLUTIONS



OUR FIRST PRIORITY is to listen, and continuously develop a better understanding of your needs. At Paragon, we work together closely to provide tools that allow you to customize your care, and a comprehensive system that focuses on all areas of your bleeding disorder!

YOUR HEMOPHILIA TEAM



Provides 24 hour on-call pharmacy support, 365 days a year



Completes all insurance verifications, pre-authorizations, and billing



Offers ongoing patient education
Provides injection training if needed



Communicates regularly with each patient and referring physician for their safety and peace of mind.



Offers free overnight shipping

COLBY DAVIS - 214.435.0929
YOUR LOCAL KANSAS CITY REP!

PARAGON HEMOPHILIA

P: 833.862.4559 • F: 855.862.4373

PARAGONSPECIALTY.COM

We're Listening



At Pfizer Hemophilia, we have always been deeply committed to listening to what you have to say. Our programs and resources are all designed in response to the needs of the hemophilia community.

We are grateful for having the chance to partner with you.

—Your Pfizer Hemophilia Team

P.O. Box 412866 • Kansas City, MO 64141
info@midwesthemophilia.org

MidwestHemophilia.org



Follow us on Facebook

MidwestHemophiliaAssociation



Follow us on Twitter

MidwestHemophi1



Follow us on Instagram

MidwestHemophiliaAssociation

