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TOLL FREE 800-293-6527
www.HemophiliaFlorida.org

Still Time to Register for Camp Spirit

Camp Spirit, a camping experience for children with bleeding disorders ages 7-16, is back this year and will be held at Camp Boggy Creek in Eustis July 14-18, 2022. Campers will enjoy all the activities of camp life in a medically supervised environment. Costs for attending the camp are covered by the HFGF.

To join the fun, go to this link to begin the registration process: www.boggycreek.org/camper-application.

If you want to donate to help make Camp Spirit a reality for a child with a bleeding disorder, go to www.hemophiliaflorida.org/donate/.



HFGF Connections Executive Director's Column

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Greater Florida

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Mission
The mission of the Hemophilia
Foundation of Greater Florida is
dedicated to improving the quality
of life for people with related bleed-
ing disorders and their families
through education, information and
referral services, advocacy and
research.

Disclaimer
The material provided in HFGF
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information only. HFGF does not
give medical advice or engage
in the practice of medicine. The
HFGF does not recommend partic-
ular treatments for specific indi-
viduals and recommends that you
consult your physician or treatment
center before pursuing any course
of treatment.

Dear Friends,

Spring is here and we have so many exciting and fun opportunities for everyone! We have dinners, programs, walks, and Camp Spirit scheduled. It is so nice to get back into the swing of spring.

This was a big year for advocacy. We tried to get the accumulator adjuster bill over the finish line, but we were not able to make it happen. We have a great group of advocacy ambassadors ready to take on the Florida Legislature to get the bill passed in 2023. What is a copay accumulator adjuster program? Thousands of Floridians with chronic health conditions rely on copay assistance to help them pay out-of-pocket costs for expensive specialty medicines. But many insurers have stopped counting copay assistance toward annual deductibles and the out-of-pocket limit, leaving patients with serious illness to foot the bill. These practices harm the most vulnerable Americans—those who need the most help affording their care—and make existing health disparities even worse. When insurers don't count copay assistance toward a patient's deductible or out-of-pocket limit, patients continue to owe that money (even though they've paid it), leaving them facing an enormous bill when they try to refill a prescription or get other health care. That leaves many patients with tough choices between paying for necessities like rent and food or paying for their prescriptions. Too many patients delay care or stop taking their medication, which can lead to serious complications of their illness and even greater health care costs. If you are interested in jumping on board as an advocacy ambassador, email me at Franhaynes@hemophiliaflorida.org.

I look forward to seeing everyone soon. Have a wonderful spring.

Hugs,

Fran



Annual Family Program

June 4, 2022—Ports of Call, Orlando

Tatoos Piercing and Gages

Teen/Father's Day Program

June 19, 2022—Top Golf Tampa

Camp Spirit

July 14-18, 2022—Camp Boggy Creek

Arnold Palmer Hospital Family Program

July 30, 2022—Location TBD

Bleeding Disorder Trivia

Aug. 13, 2022—Adventure Island, Tampa



A photograph of two men sitting in a blue golf cart. The man on the right is wearing a light blue polo shirt, dark shorts, and a dark baseball cap, and is smiling at the camera. The man on the left is wearing a dark blue polo shirt and dark shorts, and is waving his right hand. The golf cart has a tan roof and the number '138' is visible on its side. They are parked on a grassy area next to a parking lot with several cars in the background.

[illegible]

To register for the tournament or sign up for a sponsorship, go to hemophiliaflorida.org/fundraising/golf-tournament/



19th Annual Evening on Broadway Frozen, a Blazing Success!



On Friday, March 4, 2022, a sold-out crowd of HFGF supporters enjoyed orchestra seating for a production of the Broadway hit Frozen, a private reception, and silent auction and raffle at the Dr. Phillips Performing Arts Center in Orlando. The event raised at least \$25,000 to support HFGF's programs including Camp Spirit, the Emergency Assistance Fund, and educational programs on bleeding disorders.

A very special thank-you to our sponsors for helping make the event possible:

Tony Award Winning Sponsor

CSL Behring
Biotherapies for Life®

Program Sponsor

Optum

A big Thank-You to all of the individuals and organizations who contributed goods and services to our silent auction and raffle.

4 Rivers Smokehouse
American Eagle Outfitters
Busch Gardens Tampa Bay
DoTerra Essential Wellness
Dr. Phillips Center for the Performing Arts
Foxtail Coffee Co.
Kendra Scott
Knitpicks
Lakeridge Winery
Lillian and Chip Hearn
Lucille Coleman
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Mayor Jerry Demmings
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Orlando Magic
Orlando Shakes
Phoenix East Aviation
PRP Wine International
Rifle Paper Company
RocketBook Digital Smart Notebook
San Sebastian Winery
Shake Shack
Solomon Family
The Glass Knife
TopGolf Orlando
Total Wine & More
Trader Joes
Wonder Works



Research Update

NHLBI Grant Supports Scientific Research to Enhance Hemophilia A Gene Therapies

A group of investigators are embarking on a new research program designed to unpack some of the outstanding fundamental questions associated with current approaches to gene therapy for hemophilia A. While multiple gene therapies are currently in various stages of preclinical or clinical studies there remain concerns relevant to the biological nuances affecting long term safety and efficacy.

The new program, which is being supported by a \$12 million grant from the National Heart Lung and Blood Institute (NHLBI), represents a multi-institutional effort to ultimately help improve these therapies. It will be led by Roland Herzog, PhD, at the Indiana University (IU) School of Medicine.

Herzog is the Director of IU's Gene and Cell Therapy Program. He was also a recipient of NHF's Career Development Award from 2000-2003, for his funded project "Immunology of Liver-Derived Expression of Factor IX from AAV Vectors."

"Several companies have taken this forward into clinical trials, and in some of these trials, the patients initially looked like they were cured," said Herzog, who is the Riley Children's Foundation Professor of

Immunology. "But what they all have in common is that they need to deliver a lot of the virus to get the desired results, and over time clotting factor levels started to decline. So, it's clear that we need to further study the biology of this phenomenon."



According to an IU School of Medicine press release, this program will include three major projects with investigators conducting molecular-level analysis of key components in liver-directed gene therapy, including human liver cells and factor VIII (FVIII) viral vectors. Herzog and his colleagues hope to garner new insights that can ultimately lead to lower levels of cellular toxicity and "improved longevity of FVIII production" in individuals who are treated with gene therapy for hemophilia A.

"This is an incredibly significant and urgent medical question, and it requires the synergy of multiple groups with different expertise to come together and solve a problem that they wouldn't be able to solve on their own," said Herzog. "My hope is that our studies will help the field as a whole move toward curing hemophilia A."

Source: Indiana University School of Medicine news release dated March 8, 2022

When you shop with

amazon.com
Smile

they'll donate to

The Hemophilia Foundation of
Greater Florida



DONATE



Hemophilia Treatment Centers Staff

This issue we are featuring Bayview St. Joseph's Center for Bleeding and Clotting Disorders in Tampa, Florida.

Meet Francesca Ferretti



Hello! My name is Francesca Ferretti, and I am the new Hemophilia Treatment Center social worker at St. Joseph's Hospital. Having recently taken over the position, I am excited to introduce myself and share with you my role, and how I will be supporting both pediatric and adult patients at the Center for Bleeding and Clotting Disorders.

I have a degree in Psychology from the University of Florida and a Master of Social Work Degree from the University of Central Florida. In 2018, I became a Licensed Clinical Social Worker.

I bring with me many years of clinical experience in both an inpatient and outpatient medical setting. My role as a medical social worker began at UF Health Shands Hospital where I was a pediatric hematology/oncology social worker for 7 years.

I am excited to join the HTC team here at St. Joseph's Hospital. In my current role, I will be providing crisis intervention, supportive counseling, referrals, assistance with insurance, school and social service issues, as well as assisting with community outreach and program development.

The bleeding disorders community offers a strong network of support, and I'm excited to be a part of it. I look forward to serving the community, and I hope to have the opportunity to meet many of you at future events.

Should you have any questions, I am happy to answer them over e-mail or phone.

Francesca Ferretti, LCSW
Licensed Clinical Social Worker
St. Joseph's Hospital | Center for Bleeding and Clotting Disorders
3001 W. Dr. Martin Luther King Jr. Blvd, Tampa, FL 33607
Phone: (813) 870-4017
Fax: (813) 554-8937
Email: Francesca.Ferretti@baycare.org



Hemophilia Treatment Centers Staff *cont'd*

Meet Johnson Vilayhong



Hello Everyone,

My name is Johnson Vilayhong, and I serve as the Data Coordinator for BayCare – St. Joseph's HTC. Born, bred and raised in sunny St. Petersburg, FL, which is a part of the greater Tampa Bay area. My role is to serve the technical needs of the HTC team with analytical support, database management, IT solutions, project management and process improvement initiatives.

I came to my current role with extensive experience from ambulatory training and development field. Performing instructional training to our team members on the utilization of practice management and software applications; supporting business initiatives and managing strategic operational projects.

My greatest joy is to bring added value to everyone I meet. As I strongly believe that you should always leave people better than you found them!

I am super excited to be a part of the bleeding disorders support community and look forward to sharing thoughtful conversations with everyone at future events!

Follow/Like HFGF on Social Media!

[Facebook.com/hemophiliafl](https://www.facebook.com/hemophiliafl)
[Twitter.com/hemophiliafl](https://twitter.com/hemophiliafl)
[Instagram.com/hemophiliaflorida](https://www.instagram.com/hemophiliaflorida)



**CHANGE THEIR WORLD. CHANGE YOURS.
THIS CHANGES EVERYTHING.**

[BECOME A VOLUNTEER](#)

OR

[DONATE](#)



2022 Gainesville Walkers and Runners Were Tough as Gators!

More than 125 walkers and runners participated in the 11th annual Gator Clot Trot Saturday, April 9th at Veterans Memorial Park in Gainesville. Cheered on by University of Florida mascots Albert and Alberta, the group raised more than \$31,500 for the HFGF's work with the bleeding disorders community.

The top three fund-raising teams were Team E, UF HTC: The Hemostasis Oasis, and Gator Blood – UF Hematology Fellows. Top honors for individual fundraisers went to Morgan Gillam, Nancy Yardy and Melissa Dolan.



Thanks are due to our sponsors for helping make the event possible:

Champion Sponsors

Bayer Healthcare
CSL Behring
Genentech
Novo Nordisk
Sanofi
Takeda
UF Health

Gold Medal Sponsors

Accredo

In Kind Sponsors

I Am 106.9 Country Radio
ADJE
Kanapaha Botanical Gardens
LifeSouth Community Blood Centers
Piesano's Pizza
Satchel's Pizza
Sun Country Sports Center
The Floating Lotus Spa
Splitz & Skate Station
SunCountry
WinSupply of Gainesville



More Opportunities to Walk for the HFGF!

It's not too late to get into the fun of joining a walk to raise funds to support the work of the HFGF including Camp Spirit, the Emergency Assistance Fund that helps with housing, medical care and other basic life needs, Scholarships, Bleeding Disorders Education Programs, and Legislative Advocacy.

So, choose the walk that's right for you and put on your walking shoes! To register for a walk, go to <https://www.hemophiliaflorida.org/walks-5ks/>.



SuperHero Fun Walk September 24, 2022

Registration starts at
8 am

Vinoy Park
701 Bay Shore Dr. NE
St. Petersburg, FL



Orlando Creepy Crawl Fun Walk or 5K Run October 29, 2022

Registration starts at 4
pm Harbor Park at
Baldwin Park
4990 New Broad Street
Orlando, Florida





Factor Assistance Programs

Information provided below was valid as of 4.08.20

Source: National Hemophilia Foundation for all bleeding disorders.

Bayer

\$0 Product Co-pay Program (Kogenate, Kovaltry, Jivi)

You may be able to receive up to \$12,000 in assistance per year, regardless of income.

Assistance is awarded per patient. Multiple members of the same household can apply.

Enrollment can be started and completed in one short phone call.

Lab Monitoring Co-pay Program (Jivi)

You may be able to receive up to \$250 per year to apply towards out-of-pocket costs for laboratory monitoring of Jivi.

Free Trial Program (Kogenate, Kovaltry, Jivi)

Enroll today for up to 6 free doses of Jivi®, KOVALTRY®, or Kogenate® FS.

Free Trial product is delivered to your home.

Any patient who has never been treated with the product they are receiving through the Free Trial program is able to participate, regardless of type of insurance or if you have insurance.

Loyalty Program (Kogenate, Kovaltry, Jivi)

When you enroll, you get a year's worth of Loyalty points.

Each month you use Jivi®, KOVALTRY®, or Kogenate® FS, you will earn 1 Loyalty Program

point.

You may redeem 3 points for a 1-month supply of Bayer product, up to 12 points at 1 time, to help fill insurance gaps when:

You experience challenges getting insurance coverage for Jivi®, KOVALTRY®, or Kogenate® FS.

You are between jobs and experience a gap in insurance coverage.

You are uninsured or underinsured.

Open to current and new Jivi®, KOVALTRY®, and Kogenate® FS Patients.

Points are transferrable between Jivi®, KOVALTRY®, and Kogenate® FS.

Live Helpline Support (Kogenate, Kovaltry, Jivi)

Consult with an expert in insurance

Multiple languages, including Spanish, are available

More information:

Phone: 1-800-288-8374 o 8:00 AM–8:00 PM (ET)

Monday–Friday.

Spanish-speaking Case Specialists are also available.

Websites:

www.jivi-us.com/en/copay-support/

www.kovaltry-us.com/access-services-by-bayer

www.kogenatefs.com/accessing-kogenate



Factor Assistance Programs *cont'd*

CSL Behring

“My Access” Program

May cover up to \$12,000 of out-of-pocket (OOP) expenses each year for Afstyl, Idelvion, Humate P, Helixate-FS*

OOP maximum limits for 2017 under the ACA are \$7150 (for individuals) and \$14,300 (for families)

Eligibility Requirements:

Must have private insurance, use a CSL Behring therapy, and be a U.S. resident

Enrollment is not retroactive

Must take a CSL Behring product for the treatment of von Willebrand disease or hemophilia A

Re-enroll every 12 months

Program Information:

My Source: www.mysourcecsl.com
1-800-676-4266

Patient Assistance Program

Provides free product to patients on CSL products for up to 12 months

Eligibility Requirements:

Must be uninsured or underinsured or seeking

insurance (in appeal or delay in coverage decision)

Income is < 250% FPL

Must seek insurance or other form of assistance for up to 3 months after participating for a year

Must participate in insurance counseling

Program Information:

My Source: www.mysourcecsl.com
1-800-676-4266

Assurance Program

Patients earn points for continued use of CSL Behring medication.

The points can be redeemed for free product if a patient has a lapse in insurance coverage.

Eligibility Requirements:

Must have private insurance, use a CSL Behring therapy, and be a U.S. resident.

Must enroll in Assurance program while insured and on product. Enrollment is not retroactive.

Earn 1 point per month of consecutive use of product.

Patient suffers a lapse in coverage and redeems 3 points for 1 month of product supply.

Points are earned monthly and expire 5 years after date earned. (For ex: Point earned in May 2015 expires May 2020)



Factor Assistance Programs *cont'd*

CSL Behring *cont'd*

Program Information:

My Source:
www.mysourcecsl.com or 1-800-676-4266

Genentech **SPECIAL COVID-19 UPDATE**

We realize there is increased anxiety among patients, caregivers, and their families, as resources are stretched due to losses in wages or health insurance. We want to reiterate that we are here to help and our financial assistance programs remain open and operational across all therapeutic areas for both new and existing patients.

Please visit: www.genentech-access.com/patient/brands/hemlibra or call (866) 422-2377.

In addition, we are supporting patients and health care professionals to best navigate the complexities of social distancing while continuing treatment.

The Genentech Patient Foundation, which provides free medicine to qualifying patients, is able to modify shipping locations and frequencies to ensure needed medicine gets to the right place for the patient to be safely treated.

HEMLIBRA Access Solutions

Dedicated case managers to help you understand your insurance coverage and assistance options.

Program information:
www.genentech-access.com/patient/brands/

hemlibrao
(866) 422-2377

HEMLIBRA Co-Pay Program

Helps with up to \$15,000 per year in HEMLIBRA co-pay costs.

Eligibility Requirements

Have been prescribed HEMLIBRA for an FDA-approved indication.

Are 18 years of age or older or have a legal guardian 18 years of age or older to manage the program.

Have commercial (private or non-governmental) insurance. This includes plans available through state and federal health insurance marketplaces.

Do not receive support from the Genentech Patient Foundation or any other independent co-pay assistance foundations for HEMLIBRA.

Are not a government beneficiary and/or participant in a federal or state-funded health insurance program (e.g., Medicare, Medicare Advantage, Medigap, Medicaid, VA, DoD or TRICARE).

Do not reside in a state where the program is prohibited.

Program Info:
hemlibracopay.com/
(844) HEM-COPAY (844-436-2672)



Factor Assistance Programs *cont'd*

Genentech *cont'd*

Genentech Patient Foundation

The Genentech Patient Foundation gives free Genentech medicine to people who don't have insurance coverage or who have financial concerns and to people who meet certain income criteria.

Program info:

GenentechPatientFoundation.com

My Source:

www.mysourcecsl.com or 1-800-676-4266

Grifols

Factors for Health:

The \$0 Copay Assistance Program, which may cover out-of-pocket expenses not covered or partially covered by insurance.

The Free Trial Program for eligible patients who are new to ALPHANATE.

Benefits investigation and support services to help patients coordinate with their insurer.

The Patient Assistance Program for patients with no coverage or lapsed coverage.

Care Coordination to help patients gain access to and remain on ALPHANATE.

Program information:

844.MY.FACTOR (693.2286)

www.alphanate.com/en/patients/support-and-resources/factors-for-health

Patient Care Programs-Grifols Assurance for Patient (PAP)

Eligibility Requirements:

Temporary lapse in private insurance coverage.

Treated with Grifols products for 3 continuous months prior to lapse in private insurance.

State or federal program recipients not eligible.

Program Information:

www.alphanate.com/en/patients/support-and-resources/factors-for-health

Patient Assistance Program (PAP)

Eligibility Requirements:

Must be uninsured & in temporary need of assistance obtaining Grifols products.

Must be U.S. Citizen or legal resident.

Financial eligibility - below 250% FPL.

Program Information:

www.alphanate.com/en/patients/support-and-resources/factors-for-health



Factor Assistance Programs *cont'd*

Medexus Pharma

XINITY Savings Program

Up to \$12,000 annually.

Eligibility requirements:

Must have a valid prescription for IXINITY.

Must have commercial insurance.

No monthly limits.

No income requirements.

1-855-IXINITY (1-855-494-6489) to sign up.

Co-Pay program may be used retroactively for up to 12 months.

Program information:

IXINITY Patient Assistance Program

Program helps deliver treatment to those in need, even if they do not have insurance. If you are uninsured or experience a lapse in your coverage, this program may cover you.

IXperience Concierge program information:
1-855-IXINITY (1-855-494-6489)

FACTOR IT FORWARD, by IXINITY (New)

This program is accessible through your HTC.

Through Factor It Forward, your HTC may be able to access factor IX for you in times of need. Talk to your HTC to find out if they are part of Factor It Forward, by IXINITY, coagula-

tion factor IX (recombinant).

Program information:

www.ixinity.com/the-ixinity-ixperience/factor-it-forward/

Novo Nordisk

Product Assist Program

Amount of available assistance varies by product.

Eligibility Requirements:

Must be uninsured and currently seeking insurance.

Must be a U.S. Citizen or have legal residence.

Program Information:

www.mynovosecure.com/support/product-assistance-page

1-844-668-6732

Copay Assistance Program

Eligible patients receive up to \$12,000 per year toward their out-of-pocket costs for Novo Nordisk Hemophilia and Rare Bleeding Disorder products.

Eligibility Requirements:

You are not eligible if prescriptions are paid by any state or federally funded programs, including, but not limited to, Medicare, Medicaid, Medigap, VA, DOD, or TRICARE, or where prohibited by law.



Factor Assistance Programs *cont'd*

Offer not valid for prescriptions reimbursed in full by any third-party payor. Offer limited to 1 card per person. Offer void where taxed, restricted, or prohibited. Offer only good in the U.S.A. Eligibility is restricted to individuals; no clubs, groups, or organizations. This savings card is not transferable and is not insurance. When you use the Novo Nordisk Hemophilia & Rare Bleeding Disorders Co-pay Assistance Card, you are certifying that you understand the program rules, terms, and conditions and that you will comply with them. Offer excludes full cash-paying customers. You must be enrolled in a commercial insurance plan.

For a complete set of eligibility criteria, and terms and conditions, please visit

www.mynovosecure.com/copayassistance

NovoSecure Program

NovoSecure™ offers a variety of programs, including reimbursement support, competitive scholarships, lifecoaching with HeroPath™, career counseling, and insurance support. Eligibility for programs vary. Must have hemophilia A, hemophilia A or B with inhibitors, factor VII deficiency, factor XIII deficiency, acquired hemophilia, or Glanzmann's Thrombasthenia to apply.

Program Information:

NovoSecure Program 1-844-668-6732

Octapharma SPECIAL COVID-19 UPDATE

Octapharma is committed to providing support to patients who need our therapies. The COVID-19 pandemic has created unprecedented financial challenges for patients and families at this time. Octapharma is proud to offer relief through our existing and expanded patient support programs. As more families face challenges in the days ahead, we will stand with you to offer the support you need to continue your important therapies.

Octapharma Expanded Compassionate Care Program

Helps to assist patients using Octapharma Coagulation Therapies who have lost insurance due to the COVID-19 Pandemic by providing access to free factor.

Some restrictions may apply.

Contact the Octapharma Support Center at 1-800-554-4440 for more information

Only applies to patients with private commercial insurance.

NUWIQ Free Trial Program

Allows for up to 6 doses or 20,000 IUs of NUWIQ.

Eligibility Requirements:

Must have a prescription for NUWIQ.

Other restrictions may apply.



Factor Assistance Programs *cont'd*

Octapharma *cont'd*

Program Info:

www.nuwiquisa.com/factor-8-free-trial/#Free-Trial-Program

Octapharma Support Center: 1-800-554-4440

NUWIQ CoPay Program

Assists with copay costs for NUWIQ up to \$12,000 per year.

Eligibility Requirements:

No income requirements.

Only applies to patients with private commercial insurance.

Must have a prescription for NUWIQ.

Other restrictions may apply.

Program Info:

www.nuwiquisa.com/factor-viii-patient-assistance-program/

Octapharma Support Center: 1-800-554-4440

Wilate Free Trial Program

Allows for up to 5,000 IUs of Wilate.

Eligibility Requirements:

Must have a prescription for Wilate.

Other restrictions may apply.

Program Info:

www.wilateusa.com/patient-support/free-trial-program/

Octapharma Support Center: 1-800-554-4440

Wilate CoPay Program

Assists with copay costs for Wilate up to \$12,000 per year.

Eligibility Requirements:

No income requirements.

Only applies to patients with private commercial insurance.

Must have a prescription for Wilate.

Other restrictions may apply.

Program information:

www.hemophiliavillage.com/financial-support

1.844.989.HEMO (4366)

Factor Savings Card

Offers up to \$12,000 in copay, coinsurance and deductible costs associated with Pfizer factor products. Card cannot be combined with any other rebate/coupon, free trial, or similar offer for the specified prescription. The card will be accepted only at participating pharmacies. This coupon is not health insurance. No membership fees.



Factor Assistance Programs *cont'd*

Octapharma *cont'd*

Program Info:

www.wilateusa.com/patient-support/co-pay-assistance-reimbursement/

Octapharma Support Center: 1-800-554-4440

Pfizer

Pfizer Hemophilia Connect

Easy access to all of our hemophilia tools and programs and provides resources to help you connect to the hemophilia community.

Support team is dedicated to working with patients, their caregivers, and health care providers or pharmacies depending on your individual needs.

The Pfizer Patient Assistance Program is a joint program of Pfizer Inc. and the Pfizer Patient Assistance Foundation™. The Pfizer Patient Assistance Foundation is a separate legal entity from Pfizer Inc. with distinct legal restrictions.

Program Info:

www.hemophiliavillage.com/financial-support
(resources and support)

888-240-9040

Eligibility Requirements:

No financial eligibility requirements.

Available to commercially insured patients only; Medicare/Medicaid beneficiaries are not eligible.

Rx Pathways Program

Provides co-pay assistance up to \$10,000 annually, free product, or premium assistance.

Eligibility Requirements:

Must be prescribed a Pfizer specialty drug.

Must have private insurance with prescription drug coverage.

Must meet income eligibility requirements, which are based on the household size and prescription.

Only new patients may use this offer

You are currently covered by a private (commercial) insurance plan

Other restrictions may apply

Program Info:

pfizerhemophiliaresources.com

1.800.710.1379

Free Trial Prescription Program

Eligible patients can receive a one-time, 1-month supply up to 20,000 IU of Pfizer factor product at no cost to you.

For first-time used by commercially insured patients only.



Factor Assistance Programs *cont'd*

Pfizer *cont'd*

Eligibility Requirements:

For a complete list of participating pharmacies, visit PfizerRxPathways.com or call the toll-free number 1.877.744.5675

Program Info:

www.hemophiliavillage.com (resources and support).

Sanofi Genzyme

MyALPROLIX Free Trial Plus Program

Allows for either a 30-day supply of free Alprolix or free

Alprolix for up to 1 year, if needed, until health-care coverage begins.

Eligibility Requirements:

First prescription of ALPROLIX.

You are uninsured or insured by private insurance.

Other restrictions may apply.

Program Information:

www.alprolix.com/resources-and-support/available-financial-support

MyALPROLIX Copay Program

Assists with monthly out-of-pocket costs for

Alprolix up to \$20,000 a year, and helps link patients to coverage resources.

Eligibility Requirements:

No income requirement or cap.

Only applies to private insurance.

Must have a U.S.-based prescriber and pharmacy.

Other limits may apply.

Program Information:

www.alprolix.com/resources-and-support/available-financial-support

MyALPROLIX Factor Access Program

Provides access to Alprolix.

Eligibility Requirements:

Experiencing a gap in insurance coverage, have reached maximum insurance coverage limit, or have no prescription coverage.

Other restrictions may apply.

Program Information:

www.alprolix.com/resources-and-support/available-financial-support

MyEloctate Free Trial Plus Program

Allows for either a 30-day supply of free Eloctate or Eloctate for up to 1 year, if needed, until healthcare coverage begins.



Factor Assistance Programs *cont'd*

Sanofi Genzyme *cont'd*

Eligibility Requirements:

First prescription of Eloctate.

You are uninsured or insured by private insurance.

Other restrictions may apply.

Program Information:

[www.eloctate.com/myeloctate/resources/My-Eloctate Copay Program](http://www.eloctate.com/myeloctate/resources/My-Eloctate-Copay-Program)

Assists with monthly out-of-pocket costs for Eloctate up to \$20,000 a year and helps link patients to coverage resources.

Eligibility Requirements:

No income requirements or caps.

Only applies to private insurance.

Must have a U.S.-based prescriber and pharmacy.

Other restrictions may apply.

Program Information:

www.eloctate.com/myeloctate/resources/

MyEloctate Factor Access Program 10

Provides access to Eloctate.

Eligibility Requirements:

Experiencing a gap in insurance coverage, have reached maximum insurance coverage limit, have no prescription coverage, or meets specific income guidelines adjusted for family size.

Other restrictions may apply.

Program Information:

www.eloctate.com/myeloctate/resources/

Takeda SPECIAL COVID-19 UPDATE:

At Takeda, our highest priority is the patients we serve. Our Hematology Support Center team is available to support patients with their Takeda treatment especially during the COVID-19 pandemic. Our financial assistance programs remain open and operational. Patient Access Managers (PAM) are available to provide insurance and access education and support for patients currently on Takeda products. PAMs can also help to address barriers to treatment access. Patients and caregivers can call the Hematology Support Center for more information: 888-229-8379 Mon-Fri | 8:30 a.m. to 8:00 p.m. ET | www.hematologysupport.com/

Freedom of Choice:

Provides eligible individuals with a free trial of select Takeda products.

Program information:

1-888-229-8379



Factor Assistance Programs *cont'd*

Takeda *cont'd*

Hematology Support Center

Takeda's Hematology Support Center (HSC) is a team dedicated to help patients who have been prescribed Takeda hematology products including:

Reviewing financial assistance options, including co-pay support and emergency access to Takeda Hematology products for existing patients in case of sudden lapse in coverage.

Explaining insurance coverage options and assisting with insurance challenges.

Providing access to educational tools for disease and insurance information.

Connecting patients and caregivers with informational and community resources, including educational programs.

Offering one-on-one health education support in English or Spanish.

Program Information:

1-888-229-8379

www.hematologysupport.com

Copay Assistance Program

Covers an eligible patient's copay, coinsurance, or deductible medication costs, up to \$20,000 a year.

Not valid for prescriptions reimbursed, in whole or in part, by Medicaid, Medicare, Medigap, VA, DoD, TRICARE or any other federal or state healthcare programs, including state pharma-

ceutical assistance programs, and where prohibited by the health insurance provider or by law.

Non-medication expenses, such as ancillary supplies are not eligible.

If patient has payer coverage for administration services, their out-pocket for the product administration may be covered.

Eligibility Requirements:

Must have a current prescription for an eligible Takeda factor or bypass product.

Must have a diagnosis of Hemophilia A or B, or a hemophilia A or B inhibitor.

Must have commercial insurance.

Additional terms and conditions apply.

Program Information:

1-888-229-8379

www.hematologysupport.com

Patient Assistance Program (PAP):

This program is only available to patients without insurance.

Provides free Takeda Hematology products at no cost if patients meet financial eligibility criteria. Other restrictions may apply.

Program information:

1-888-229-8379