Hemophilia Foundation of Greater Florida

HFGF Connections

The Quarterly Newsletter of the HFGF Spring 2020 Volume 32 Issue 54

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

Health and Wellness Update: COVID-19

From Dr. Leonard Valentino, CEO of the National Hemophilia Foundation (NHF)

I know many of you are overwhelmed with often-conflicting information on the COVID-19 virus. As a doctor and researcher, the health of the bleeding disorder community is my highest priority. Every Friday, I will be sending you vetted, trusted information about the pandemic and information you can use to keep you and your family safe. This information will also be posted on the NHF website at www.hemophilia.org.

If you have specific questions about the COVID-19 virus, please email them to communications@hemophilia.org, and I will try to address them in the next email.

What People with Bleeding Disorders Should Know
We have seen no indication that people with bleeding disorders are at an increased risk for contracting COVID-19, and there is no indication that people with bleeding disorders who have no other underlying conditions are more likely to develop severe symptoms. However, the COVID-19 virus is a respiratory illness that may result in severe coughing, and that can increase risk of head bleeds. It is more important than ever to make sure you are adhering to your prophylaxis regimen to decrease the likelihood of all bleeding. Currently, all of our pharmaceutical partners have assured us their supply of medication is unaffected by the outbreak. Do not ration products at this time. Treat as your healthcare provider has prescribed your medications, contact your HTC or healthcare provider with any questions or concerns and above all, stay healthy.

Risk to Pregnant Women and Infants
A small study in Wuhan, China found that pregnant women who tested positive for COVID-19 did not pass the virus to their newborn children.

How Social Distancing Works to Halt the Spread of the Virus
This animated simulation of transmission and recovery shows how social distancing can help contain the spread of the virus.

Health and Exercise
It is important for people with bleeding disorders of all ages to stay active to maintain joint and emotional health. Remember, even during this pandemic it is safe to:

- Let your kids play outside (just no playdates)
- Go for a family bike ride or walk (put those helmets on!)
- Have a family game night
- Exercise as a family

Here are some more resources links to help you stay active during this time:

Coronavirus lockdown: How to stay fit, exercise at home

How We're Staying Fit Through A Quarantine

How to Protect Your Mental Health During a Quarantine

How to Stay Fit While You're Stuck in Quarantine

continued on page 3

HFGF Connections

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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 bleeding disorders and their families through education, information and referral services, advocacy and research.

Disclaimer
The material provided in HFGF Connections is for your general information only. HFGF does not give medical advice or engage in the practice of medicine. The HFGF does not recommend particular treatments for specific individuals and recommends that you consult your physician or treatment center before pursuing any course of treatment.

Executive Director's Column

Dear Friends,

Our hearts go out to everyone who has been impacted by the Coronavirus (COVID-19). The health and safety of everyone in our community comes first, and we wanted to share a few updates as it stands now in this rapidly evolving situation.

The Hemophilia Foundation of Greater Florida is still working to meet our community's needs. We are answering phones, planning programs and here to assist you. To keep you busy while you have to stay home and to promote social distancing we are bringing you innovative, educational webinars and fun activities so check out our Calendar of Events, Social Media and Website.

Unfortunately we have had to postpone the following events, which have been rescheduled:
- The Gainesville Gator Clot Trot has been postponed until December 19, 2020.
- Tampa Walk has been postponed until September 26, 2020.
- Camp Spirit has been postponed – Future Date TBD.

We will continually keep you updated on all the changes occurring.

For more information about the COVID-19 virus, please go to the Centers for Disease Control and Prevention Website:
https://www.cdc.gov/coronavirus/2019-ncov/summary.html
and the National Hemophilia Foundation Website:
https://www.hemophilia.org/.

In times like these, let's continue to support and care about one another.

Take Care

Fran

Being Prepared for an Emergency	7:00 pm April 21, 2020
Step Therapy Advocacy	2:00 pm April 24, 2020
HFGF Board of Directors Meeting	10:00 am April 26, 2020
Mental Health Wokshop	7:00 pm April 30, 2020
Infusion Class	4:00 pm May 1, 2020
Teen Talk	4:00 pm May 7, 2020

RSVP to info@hemophiliaflorida.org for login information

CLICK HERE to subscribe to our email to receive notices and updates of all our upcoming programs and events

Genesis Erickson, Our HFGF NYLI Ambassador

My name is Genesis Erickson and I was selected to be a member of the National Youth Leadership Institute (NYLI). This is a program at the National Hemophilia Foundation (NHF) for 18-24 year olds in the bleeding disorders community. The program provides us leadership opportunities to encourage personal growth, learn how to effect change, and positively influence others. I am excited to learn and share experiences of living with a bleeding disorder while learning how to become a distinguished leader.

This past January, I had the opportunity to attend the NYLI First year leadership training in New York City. At this event, I met the other 8 people in my cohort who come from all over the country. We had public speaking workshops, met with each department of NHF, and worked on self-development as well as emotional intelligence. It was a wonderful introduction to the program, and I am so excited to continue and see where NYLI leads me.

The next month, I was on a plane to our nation's capital for Washington Days! Here, I got to tell my personal story of living with a bleeding disorder to our elected representatives. I was so happy to stand beside other people in the Florida bleeding disorders community as we advocated for a continuation of funding to our HTCs and for access to skilled nursing facilities for people with bleeding disorders. After the day on the hill, NYLI members learned all about intersectionality, and then provided recommendations to the NHF on how we celebrate diversity in the bleeding disorders community and include people of all different backgrounds.

In the future, I will learn how to facilitate workshops and use those skills to deliver workshops to kids and teens all over the country. We will be a part of NHF's annual bleeding disorders conference each year. I will also work closely with HFGF throughout my time in NYLI and after. I encourage any young adults who have an interest in becoming a change maker, especially within the bleeding disorders community, to apply to this program in the future. I am excited to serve our community in the years to come!

Health and Wellness Update: COVID-19 *cont'd*

Mental Health

Coping with stress and managing your mental health is extremely important during these uncertain times. Here are some helpful information links:

Manage Anxiety & Stress
How COVID-19 May Impact Mental Health

Follow/Like HFGF on Social Media!

Facebook.com/hemophiliafl
Twitter.com/hemophiliafl
Instagram.com/hemophiliaflorida

HIGH SCHOOL STUDENTS
EARN COMMUNITY SERVICE HOURS

VOLUNTEER FOR THE HFGF

CONTACT US AT 800-293-6527 or
info@ hemophiliaflorida.org

Factor Assistance Programs

If you are uninsured, underinsured, or experiencing a lapse in insurance coverage, there are assistance programs to provide or navigate access to bleeding disorder prescription products.

Hemophilia/vWD Manufacturer Co-Pay Programs

Company/ Organization	Co-pay/ Co-insurance Program Name & Contact Information	Limit Total	Notes:
Aptevo Therapeutics	IXINITY Savings Program 1-855-494-6489	$12,000 per year	• Must have a valid prescription for IXINITY. • Must have commercial insurance. • No monthly limits unless limit total is reached. • No income requirements. • Co-pay program can be used retroactively for up to 12 months.
Bayer	Bayer Access Services 1-800-288-8374	$12,000 per year Product Copay Program $250 per year Jivi Lab Monitoring	• $0 CoPay Program for Kogenate, Kovaltry, and Jivi regardless of income. • Up to $250 in assistance per year toward out of pocket costs for laboratory monitoring of Jivi. • Assistance is awarded per patient. Multiple members of the same household can apply. • Must have private/commercial insurance.
CSL Behring	MyAccess Program 1-800-676-4266	$12,000 per year	• Must take a CSL Behring eligible product (on-label) for the treatment of von Willebrand disease or hemophilia, including Humate-P, Idelvion, or Afstyla. • Must currently have US-based private insurance that covers your therapy (federally funded program ineligible). • Contact My Source Care Coodinator at the number listed to enroll and to obtain further information about the program. Program benefit DOES NOT apply toward out-of-pocket costs for: • Physician office visit co-pays. • Infusion-related costs or ancillary supplies. • Insurance premiums.

Factor Assistance Programs

cont'd

Hemophilia/vWD Manufacturer Co-Pay Programs

Company/ Organization	Co-pay/Co-insurance Program Name & Contact Information	Limit Total	Notes:
Genentech	HEMLIBRA Co-Pay Program 1-844-436-2672	$15,000 per year	• No income requirements. • Your enrollment in the program is valid for up to 12 months (You may reapply at the end of 12 months). • The program covers the remaining part of your co-pay, up to $15,000 per year. You may be eligible if you: • Have been prescribed HEMLIBRA for an FDA-approved indication. • Have commercial (private or nongovernmental) insurance. This includes plans available through state and federal health insurance marketplaces. • 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.
Grifols	Factors for Health Co-pay Assistance Program 1-844-693-2286 (844-MY-FACTOR)	In-network claims will be approved up to out-of-pocket maximum. Out-of-network may be denied by the program administrator	• Patients with commercial insurance may pay as little as $0 for ALPHANATE. • No monthly or annual maximums. Individual claims exceeding $2000 will be reviewed for network eligibility. Claims that are in-network will be approved, but those out-of-network may be denied. • Prescription is covered up to the patient's annual out of pocket (OOP) maximum. The 2019 OOP maximum limits under the Affordable Care Act are $7,900 (self-only coverage) and $15,800 (coverage for more than self only). • Benefits investigation and support services help patients coordinate with their insurer.
Kedrion Biopharma	1-855-353-7466	N/A	N/A

Spring Walks Postponed, Mark Your Calendars!
HFGF Families Need Your Help More Than Ever

Due to COVID-19, HFGF spring walks have been postponed. COVID-19 has impacted each and every one of us is many different ways. This is one way that we can truly embrace. With postponement, that means that we can take advantage of the time to tell more people about the walks, which also means more awareness for bleeding disorders and more time to fundraise. With the financial effects of COVID-19, our HFGF families will need us more than ever.

The Gator Clot Trot will now be on December 19 at Depot Park. Online registration is going on now through December 18th at noon. Onsite registration will begin at 11:30 AM on December 12th; 5K Clot Trotters will leave at 12:30 PM, and walkers will leave after. There are still tons of 5K awards up for grabs, as well as spirit and fundraising awards. THIS JUST IN… We are inviting everyone to clot trot in their best holiday gear! We want to see reindeer, elves, or whatever holiday fun you can think of, and we will be ending the day with a holiday party after the 5K and walk. We are talking about a holiday extravaganza! Register now at hemophiliaflorida.org/gainesville.

Don't worry Tampa, we have not forgotten about you! The HFGF Tampa Superhero Fun walk has been postponed until September 26th at Al Lopez Park. Online registration is now open at hemophiliaflorida.org/tampa and will remain open until September 25th at noon. Onsite registration will begin at 9:30 AM, and the walk will begin at 10:30 AM. We are officially putting out a call for all of our superheroes now to battle bleeding disorders in September! We are going to need your super strength help in making this the best walk that Tampa has ever had. Start using your super powers now to gather your teams to fundraise and raise awareness for bleeding disorders.

We anticipate the need for assistance greater than ever this year due to COVID-19. We know that there will be an economic impact, and HFGF will continue to be here for our HFGF families by offering emergency financial assistance, scholarships, and more. HFGF funds initiatives such as these through fundraising events like our walks. When you fundraise at walks like the Gator Clot Trot, Tampa Suphero Fun Walks, and Jacksonville/Orlando Creepy Crawl 5K and Vampire Miles, you are raising money for families who need it. Thank you in advance for your support and love.

Factor Assistance Programs

cont'd

Company/ Organization	Co-pay/Co-insurance Program Name & Contact Information	Limit Total	Notes:
Novo Nordisk	Co-pay Assistance Program 1-844-668-6732 (1-844-NOVOSEC)	$12,000 per year	Apply online. Eligible individuals: • Have hemophilia A and have been prescribed an appropriate Novo Nordisk factor treatment; OR • Have congenital hemophilia A or B with an inhibitor, congenital FVII deficiency, Glanzmann's thrombasthenia with refractoriness to platelet transfusions, or acquired hemophilia and have been prescribed an appropriate Novo Nordisk factor treatment; OR • Have FXIII A-subunit deficiency and have been prescribed an appropriate Novo Nordisk factor treatment; AND • Have private/commercial insurance only (state and federal funded programs ineligible)
Octapharma	Octapharma Co-Pay Program 1-800-554-4440	Up to $12,000 per year	NUWIQ CoPay Program Wilate CoPay Program • Must be receiving treatment from Octapharma or have a prescription to begin treatment. • Must have private/commercial insurance or self-pay. • Co-Pay Assistance Program does not cover costs associated with administration of therapy, such as office visits, infusion costs, or other professional services.
Pfizer	Pfizer Factor Savings Card 1-888-240-9040	$12,000 per year	• Register online to receive a Factor Savings Card or request one from your doctor, or call 1-855-PFZ-HEMO (739-4366). Provides co-pay and co-insurance assistance for a Pfizer factor product. • No financial eligibility requirements. • Must have private/commercial health insurance that covers factor.

HEMOPHILIA FOUNDATION OF GREATER FLORIDA
CALVIN DAWSON MEMORIAL SCHOLARSHIP

Deadline: April 30

Click here for the 2020 application

Factor Assistance Programs

cont'd

Hemophilia/vWD Manufacturer Co-Pay Programs

Company/ Organization	Co-pay/Co-insurance Program Name & Contact Information	Limit Total	Notes:
Sanofi Genzyme (Formerly Bioverativ)	ALPROLIX Co-pay Program 1-855-692-5776 ELOCTATE Co-pay Program 1-855-693-5628	$20,000 per year	• Available to those who use ALPROLIX or ELOCTATE. • Must have commercial insurance. • Must be treated by licensed doctor in US or Puerto Rico • Have a United States pharmacy. • Enrollment forms available via links provided • Not responsible for costs associated with administration of therapy, such as office visits, infusion costs, or other professional services.
Shire (now Takeda)	Shire's Hematology Support Center CoPay Assistance Program 1-888-229-8379	$12,000 per year	• For Advate, Adynovate, Feiba, Hemofil M, Recombinate, Rixubis, Vonvendi. • Must have commercial insurance. 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 pharmaceutical assistance programs, and where prohibited by health insurance provider or by law. • Provides coverage for medication-related co-payment/co-insurance. Non-medication expenses, such as ancillary

Chart updated 10/25/2019.

Factor Assistance Programs

cont'd

For those in the bleeding disorder community who are uninsured, underinsured, or experiencing lapses in insurance coverage, there are also assistance programs to help provide or navigate access to factor products.

Hemophilia/vWD Manufacturer Product Assistance and Insurance Navigation Programs

Company/Organization	Product Assistance Program Name & Contact Information
Akorn Pharmaceuticals	Patient Assistance Program: This needs-based program assists patients in accessing Amicar. Programs include copay assistance for patients with commercial insurance coverage and a need-based support program for those patients without commercial insurance. 1-844-202-5909
Aptevo Therapeutics	IXINITY Patient Assistance Program: This program helps deliver treatment to those in need, even if they don't have insurance. If you are uninsured or experience a lapse in your coverage, this program may cover you. 1-855-494-6489
Bayer	Loyalty Program allows patients to earn points that can be redeemed for Bayer products (Kogenate FS, Kovaltry, Jivi) in the event of a gap in insurance coverage or challenge with coverage. Patients with government insurance are not eligible. Enroll online or call 1-800-288-8374. Free Trial Program available for Kovaltry and Jivi. Live Helpline Support is available to help with insurance coverage questions in multiple languages including Spanish. 1-800-288-8374
CSL Behring	Patient Assistance Program: To be eligible, patients must be underinsured or uninsured. When enrolled, must actively be seeking insurance. 1-844-PAP-CSLB (727-2752) Assurance Program: Must currently have private health insurance coverage; this is a points program that allows current users to earn points that can be redeemed in the event of a future lapse in private health insurance coverage. 1-866-415-2164 CSL Behring My Source Hotline: Assistance navigating the complexities of insurance approvals, denials and appeals for patients with Coagulation disorders. 1-800-676-4266
Genentech	Genentech Patient Foundation provides free medicines to people who don't have insurance or whose treatment is not covered by insurance (with income less than $150,000), or patients with insurance coverage who are struggling with high out-of-pocket costs and meet certain guidelines. Call 866-422-2377 or see Genentech Access Solutions to find out your options. HEMLIBRA Starter Program helps eligible patients taking HEMLIBRA receive free medicine while awaiting an insurance coverage determination.

Hemophilia/vWD Manufacturer Product Assistance and Insurance Navigation Programs

Company/Organization	Product Assistance Program Name & Contact Information
Grifols	Grifols Patient Assistance Program is for patients using ALPHANATE, AlphaNine SD or ProfilNINE who are uninsured or experiencing a temporary lapse of insurance coverage.
Kedrion Biopharma	1-855-353-7466.
Novo Nordisk	Novo Nordisk Product Assistance Program: Provides medication to qualifying applicants at no charge. Eligible patients must have been prescribed a Novo Nordisk product for an indicated condition (check the website for a complete list of eligible conditions), have no prescription coverage, have a household income at or below 400% the federal poverty level. Patient must be a documented US resident or on a path to documented status with reasonable likelihood of attaining it. Federal government insurance programs are ineligible. 1-844-668-6732
Octapharma	Reimbursement and Claims Support: Contact the Octapharma Support Center for help with reimbursement for your Wilate or NUWIQ prescription. Octapharma representatives are available to provide expert advice and information about insurance matters including individual claims processing reviews, assistance in appeals, insurance investigations into product coverage, and gaining approvals for prior authorizations for Wilate. 1-800-554-4440 or usreimbursement@octapharma.com NUWIQ Free Trial Program allows for up to six (6) doses, or 20,000 IUs, of NUWIQ. A prescription for NUWIQ is required and other restrictions may apply. Wilate Free Trial Program allows for up to 5,000 IUs for Wilate. A prescription for Wilate is required and other restrictions may apply. 1-800-554-4440
Pfizer	Pfizer Hemophilia Connect gives you easy one-stop access to Pfizer tools and programs. 1.844.989.HEMO (4366) Reimbursement Support services for eligible patients, including: • Benefit verifications • Prior authorization assistance • Appeals assistance • Claim denial review and research • Nurse-drafted appeals submitted directly to payer on Pfizer patient's behalf • Timely status follow-up with the payer until the outcome is received Pfizer Trial Prescription Program for patients with commercial insurance prescribed factor products for the first time.

Factor Assistance Programs

cont'd

Hemophilia/vWD Manufacturer Product Assistance and Insurance Navigation Programs

Company/Organization	Product Assistance Program Name & Contact Information
Sanofi Genzyme (Formerly Bioverativ)	Factor Access Program: Patients using ALPROLIX or ELOCTATE who have no prescription coverage, are facing a gap in coverage, or have reached their maximum insurance coverage limit, may be eligible to receive product for free. Those with federal or state government insurance are not eligible. Other restrictions may apply. See the enrollment form or contact the support line.
Shire (now Takeda)	Shire's Hematology Support Center helps educate on insurance options, navigate insurance access issues, and provide access to downloadable tools. Case Managers, Reimbursement and Access Managers, and Healthcare Educators are all part of the team. 888-229-8379

Chart updated 10/25/2019
Information compiled by the Hemophilia Federation of America

SAVE THE DATE!

Hemophilia Foundation of Greater Florida

Teen Talk

HFGF's very own Natalie Rubin will be leading this talk.

Natalie has been part of the HFGF community throughout middle and high school. She gets what it's like to live with a bleeding disorder as a teen.

Thursday May 7, 2020
4:00pm
(Ages 13-19)

Hey Teens! How are you dealing with COVID-19? We want to give you a safe space to chat about how you're feeling through all of this. Do you like distance learning? Do you miss your friends? What do you miss most about life before COVID-19? This talk will be done over the Zoom.

RSVP by emailing info@hemophiliaflorida.org
or call 800-293-6572
You will receive login information the day of the Teen Talk.

By rsvping to this event you hereby grant the Hemophilia Foundation of Greater Florida permission to use your likeness in a photograph, video, or other digital media ("photo") in any and all of its publications, including web-based publications, without payment or other consideration.

We Are Here For You

CHANGE THEIR WORLD. CHANGE YOURS.
THIS CHANGES EVERYTHING.

BECOME A VOLUNTEER OR DONATE

HTC's and COVID19 Changes

By Leanne Buchanan, R.N.

St. Joseph's Hospital Center for Bleeding and Clotting Disorders

1. Staff will screen everyone for illness at the door. Patient and one parent/guardian allowed back for visit with the doctor. No siblings permitted in the clinic.
2. Telemedicine is available with Dr. Cockell. Once you have an appointment, the office staff guides you through signing up on the telemedicine app.
3. No direct admitting to the hospital. You must call ahead to the office to be triaged. This is to rule out the possibility of an appointment in the clinic. If you are sent to the ER, there will be a triage there as well.
4. Limiting all visits in the office to those who have urgent needs.
5. Patients continue to contact the clinic as before. Clinic phones are forwarded to the employees that are working from home. There is always someone available to help you.

Arnold Palmer Hospital

1. Telehealth has been implemented, and they are attempting to do most visits through telehealth. If a virtual visit is not an option, the nurse will go into the clinic. A nurse will call prior to your scheduled appointment to screen for illness.
2. Staff will give a mask to everyone at the door, and you are required to wear the mask the entire time you are in the hospital.
3. If you suspect an ER visit is needed, call first. The triage nurse will determine if an office visit is possible versus a visit to the ER.

Shands Hospital and UF health

1. Telehealth is up and running. They are seeing most patients virtually. Nurses are scheduling appointments in the clinic if there is an issue.
2. Clinic in Tallahassee is canceled for the months of March, April and May.
3. At the hospital entrance, staff will triage everyone that comes into the hospital. The hospital is limiting who can enter the hospital. No one under 18 is allowed unless they are the patient. One parent/guardian may accompany the minor patient.
4. Once you get to the clinic, a nurse will screen again outside the clinic doors. If you are experiencing any symptoms, a mask will be issued.
5. For an ER visit, call first. A nurse will triage the call and if possible, the patient will see the doctor in clinic. Staff will call the ER if needed.

Johns Hopkins All Children's Hospital

1. Telehealth has been up for about 2 weeks. Dr. Ayalas takes all the telehealth appointments. The appointments are not time limited. Other staff will follow up after the doctor visit as needed.
2. If seen in the clinic, appointments are scattered to decrease wait time.
3. Outpatient lab is open for essential lab work.
4. Staff will screen everyone at the entrance every day. Staff will give you a mask, and everyone must wear the mask the entire time they are in the hospital.
5. One parent/guardian for each patient at the bedside. No siblings allowed in the clinic or hospital.
6. Infusion room is open.
7. For an ER visit, you must call first.

HTC's and COVID19 Changes

Nemours Jacksonville and Orlando

1. Telehealth visits are up and running.
2. Urgent issues will be given in-clinic appointments.
3. The hospital has only 2 entrances during this time.
4. Staff will screen everyone at the entrance every day. A temperature reading is required for admittance into the hospital. Masks are given to anyone with a cough.
5. The infusion room is open and seeing patients.
6. For an ER visit, you must call first.
7. One parent/guardian for each patient at the bedside. No siblings allowed in the clinic or hospital.

University of Miami HTC

1. They are officially live with telehealth as of this week. At UM, the procedure for all of the adult and pediatric patients is that they need to have access to their uhealth chart account to be able to have the telehealth visit scheduled. Registration helps grant our patients access, or the parents of our pediatric patients to have proxy access in order to fill out the appropriate consents so they can continue with their scheduled visit. We have been enjoying the cool experience of seeing our patient families virtually.

2. We are only seeing urgent cases in clinics, which can include post-hospitalization/severe bleeding episodes that we will need to be able to see up close so that Kelli, HTC PT, has access to perform the Ultrasound.

3. Our pediatric clinic only allows one parent/guardian into clinic. There are no siblings or other family members allowed. For the adults, only the patient is allowed unless they need assistance such as wheelchair/elderly.

4. At UM, we have screening tents outside the main door of Sylvester where both the adult and pediatric patients have clinic. They will ask everyone whether they have a fever, loss of taste/smell, and about traveling outside of Miami while also taking their temperatures prior to entering the clinic.

5. Call first for all ER visits.

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The Hemophilia Foundation of Greater Florida

DONATE

HFGF LOVEGRAMS

Support Through Social Distancing

Due to the financial effects of COVID-19, HFGF anticipates an increase in the need for emergency financial assistance from the Florida bleeding disorders community this year. Here's an easy way for you to help those in need AND send some virtual love to your friends and family who need it right now.

Here's how:

- Visit hemophiliaflorida.org/HFGFLovegram
- Pick out the Lovegram(s) you'd like to send (Send as many as you'd like!)
- Fill out form for each Lovegram
- Make payment - Dollars raised support HFGF patients who need it!
- Your loved one(s) receive HFGF designed Lovegram(s) privately via email and on HFGF social media outlets

Bear Hug - $1000
3 exclusive emails sent to HFGF community
3 social media posts (Choice of Facebook/Twitter/Instagram)
Company Logo & Link

Love Bug Hug - $500
2 exclusive emails sent to HFGF community
2 social media posts (Choice of Facebook/Twitter/Instagram)
Company Logo included

Huggie - $250
1 exclusive email sent to HFGF community
1 social media post (Choice of Facebook/Twitter/Instagram)
Company Name Included

Hugs & Kisses - $100
1 social media post (Choice of Facebook/Twitter/Instagram)
1 personal email

I Miss You Hug - $25
1 social media post (Choice of Facebook/Twitter/Instagram)

Hemophilia Foundation of Greater Florida

MORE INFO: LEAHNASH@HEMOPHILIAFLORIDA.ORG
THANK YOU FOR YOUR SUPPORT!