A warm welcome awaits you this spring in Louisville, Kentucky – host city of the Hemophilia Federation of America’s 2011 Educational Symposium.

We cordially invite you to join us at Symposium, the community-centered meeting that brings together more than 400 community members, over 100 exhibitors, numerous speakers and volunteers from across the country. This year, HFA will award at least 100 scholarships to attendees. Check out how to apply below!

Symposium unites the community with an educational program designed exclusively for those with bleeding disorders. Moms, dads, spouses, young adults, teens, children, and extended family members share information and participate in community fellowship in a friendly, relaxed and supportive environment.

Educational sessions address the latest in research, physical and mental health care, legislative advocacy, social advocacy and leadership. HFA’s newest area of focus – FitFactor – is designed to help everyone maintain a healthy lifestyle. This year we will feature sessions with tips and hands-on advice for maintaining good health.

If you choose to arrive early or stay after the conference, Louisville is a host city with a wide range of activities including the Kentucky Derby Museum at Churchill Downs, the Louisville Slugger Museum and the Muhammad Ali Center. You can be sure that HFA Children’s Programs will include visits to some of these exciting venues.

As an added bonus, the kickoff of the annual Kentucky Derby Festival takes place during this year’s Symposium. Our downtown hotel is in the heart of the festival! Symposium attendees who come to our closing event on Saturday night can head outdoors to the largest annual fireworks display in North America, a part of Thunder Weekend. ✨
HCV AND ACCESS TO THERAPY

In December 2010, the Food and Drug Administration (FDA) published new draft guidance for industry entitled “Codevelopment of Two or More Unmarked Investigational Drugs for Use in Combination” to treat Hepatitis C (HCV). Because the existing developmental and regulatory paradigm focuses primarily on assessment of the effectiveness and safety of a single new investigational drug acting alone, or in combination with an approved drug, FDA published draft guidance to assist sponsors in the codevelopment of two or more unmarked drugs.

Previous draft FDA guidance was weak in its approach to testing combinations of advanced agents. It indicated interest, but recommended the older, slower model for testing new drugs via addition of single agents to the existing Standard of Care cocktail.

The new draft guidance is intended to assist sponsors in the codevelopment of two or more novel (not previously marketed) drugs to be used in combination to treat a disease or condition. This guidance is distinct from FDA draft guidance of September 2010 that addressed similar topics for HCV only, and provides recommendations and advice on how to address certain scientific and regulatory issues that will arise during codevelopment. It is intended to assist sponsors in the codevelopment of two or more novel (not previously marketed) drugs to be used in combination to treat a disease or condition.

For people with bleeding disorders and HCV greater urgency and focus on rapid drug development is critical. After 25 or more years of HCV viremia, many of us will not survive another decade absent rapid availability of effective, low-side-effect therapy.

When final, this guidance may help provide a framework and path to do something about the serious and deteriorating liver health conditions among our endangered community. Combinations of advanced HCV drugs represent our best hopes for better therapy for HCV within our lifetimes.

In recent weeks, Bristol-Myers Squibb and Pharmasset have announced a Clinical Collaboration Agreement for Proof of Concept Combination Study in Patients Chronically Infected with Hepatitis C. This study is the first cross-company collaboration combining two oral, direct-acting antivirals to evaluate the combination with and without ribavirin in treatment-naive patients.

For those of you interested, notify your infectious disease/hepatology docs and garner any information. These steps are critical for people with special unmet needs to gain access.
### Thursday, April 14, 2011

<table>
<thead>
<tr>
<th>Time</th>
<th>Event</th>
</tr>
</thead>
<tbody>
<tr>
<td>1:00 PM - 6:00 PM</td>
<td>Registration</td>
</tr>
<tr>
<td>1:00 PM</td>
<td>Exhibitor Set Up</td>
</tr>
<tr>
<td>1:00 PM - 2:30 PM</td>
<td>Nonprofit Best Practices: Facebook, Twitter, LinkedIn and Blogging</td>
</tr>
<tr>
<td>3:00 PM</td>
<td>Young Leaders Kickoff: Today’s Technology, Today’s Advocacy</td>
</tr>
<tr>
<td>5:30 PM - 7:00 PM</td>
<td>Exhibit Hall Opens to Public</td>
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</tbody>
</table>

### Friday, April 15, 2011

<table>
<thead>
<tr>
<th>Time</th>
<th>Event</th>
</tr>
</thead>
<tbody>
<tr>
<td>9:00 AM - 12:00 PM</td>
<td>Registration</td>
</tr>
<tr>
<td>9:30 AM - 11:30 AM</td>
<td>Exhibit Hours</td>
</tr>
<tr>
<td>9:00 AM - 11:30 AM</td>
<td>Young Leaders Session: Today’s Technology, Today’s Advocacy</td>
</tr>
<tr>
<td>11:00 AM - 11:45 AM</td>
<td>First Time Attendee Reception</td>
</tr>
<tr>
<td>11:30 AM - 5:30 PM</td>
<td>Children’s Programming and Childcare (Check-in begins at 11:30am)</td>
</tr>
<tr>
<td>11:30 AM</td>
<td>HFA Teen Programming</td>
</tr>
<tr>
<td>12:15 PM - 1:00 PM</td>
<td>Welcome And Recognition Luncheon</td>
</tr>
<tr>
<td>1:15 PM - 2:15 PM</td>
<td>Keynote Speaker: Clifford C. Kuhn, MD, The Laugh Doctor It All Starts With A SMILE!</td>
</tr>
<tr>
<td>2:30 PM - 3:30 PM</td>
<td>Healthcare Reform: The Law Passed, Now What?</td>
</tr>
<tr>
<td>4:00 PM - 5:30 PM</td>
<td>Exhibit Hall Open</td>
</tr>
<tr>
<td>5:30 PM - 7:30 PM</td>
<td>Industry Dinner hosted by Baxter</td>
</tr>
<tr>
<td>7:30 PM - 10:00 PM</td>
<td>Blood Brotherhood &amp; Dads Poker Tournament</td>
</tr>
</tbody>
</table>

### Breakout Sessions

#### Breakout 1
- **Title:** Ready - Set - Go = A Call to Take Action
- **Details:** Be Effective: Using Legislative Web Tools

#### Breakout 2
- **Title:** Getting Personal: Telling My Story

#### Breakout 3
- **Title:** Meet the Press: Getting Noticed!

### Saturday, April 16, 2011

<table>
<thead>
<tr>
<th>Time</th>
<th>Event</th>
</tr>
</thead>
<tbody>
<tr>
<td>7:00 AM - 9:30 AM</td>
<td>Industry Dinner hosted by Bayer</td>
</tr>
<tr>
<td>9:30 AM - 5:30 PM</td>
<td>Childcare &amp; Programming</td>
</tr>
<tr>
<td>9:30 AM - 10:45 AM</td>
<td>Medical Advancement on the Horizon (Panel)</td>
</tr>
<tr>
<td>11:00 AM - 12:00 PM</td>
<td>Universal Data Collection...Public Health</td>
</tr>
<tr>
<td>12:00 PM - 1:30 PM</td>
<td>Exhibit Hall Hours (Exhibit Hall Closes at 1:30)</td>
</tr>
<tr>
<td>1:30 PM - 2:30 PM</td>
<td>Dads In Action: &quot;Parents Working Together to Make it Happen″...Creating Harmony in Families with a Bleeding Disorder (Moms Included)</td>
</tr>
<tr>
<td>1:30 PM - 2:00 PM</td>
<td>Blood Brotherhood: Fifactor, Be Healthy! (Partners/Spouses Welcome)</td>
</tr>
<tr>
<td>2:00 PM - 2:30 PM</td>
<td>Blood Brotherhood: HCV and Access to Therapy (Partners/Spouses Welcome)</td>
</tr>
<tr>
<td>2:30 PM - 3:30 PM</td>
<td>Blood Sisterhood: Beyond My Diagnosis, “Self-Advocating” within the HealthCare System</td>
</tr>
<tr>
<td>2:30 PM - 3:30 PM</td>
<td>Dads in Action: Fifactor...Focusing on Family</td>
</tr>
<tr>
<td>2:30 PM - 3:30 PM</td>
<td>Moms: Rap Session, Partners/Spouses of Blood Brothers: Rap Session</td>
</tr>
<tr>
<td>2:30 PM - 3:30 PM</td>
<td>Blood Brotherhood: Just Click on “IT.” How to connect with your Brothers Whenever, Wherever!</td>
</tr>
<tr>
<td>3:30 PM - 4:30 PM</td>
<td>Women’s Health &amp; Wellness Reception</td>
</tr>
<tr>
<td>3:30 PM - 4:30 PM</td>
<td>Dads in Action: Rap Session, Blood Brotherhood: Rap Session</td>
</tr>
<tr>
<td>5:00 PM</td>
<td>Remembrance Service</td>
</tr>
<tr>
<td>6:30 PM - 9:30 PM</td>
<td>Kentucky Thunder Fun Fest! A final night event filled with food, dancing, entertainment and fun for the entire family!</td>
</tr>
</tbody>
</table>
By Paul Brayshaw, Matthew Compton, Chad Stevens and Tracy Cleghorn

The Hemophilia Federation of America is proud to introduce a new section of the Dateline Federation newsletter, called Truth and Transparency. This section will seek to address various concerns, as well as define issues which stakeholders, including patients, families, medical providers, industry and academia ask regarding access to care, and healthcare delivery.

As the audience, your input is critical. When issues and scenarios are presented, please engage in the discussion and share your comments for future topics. If you have a question or issue that needs to be discussed, please submit them at any time. HFA ensures the privacy of all communications.

HFA does not offer medical advice. Please direct all treatment questions to medical professional.

Question
If I have private insurance, what kind of incentives can I accept?

Answer
There is a big difference in having private and government-based insurance coverage, but if you are enrolled in a government program (i.e. Medicare and/or Medicaid) costs of goods, meals or services cannot be greater than $9.99. For a patient with private insurance the rules are not as clear. Some policies are regulated by federal legislation, depending on the size of an employer, and others are regulated by the state.

Pharmaceutical companies and manufactures are regulated separately and the rules are strict. Unless an educational component is involved in any dinner or meal, manufacturers are unable to provide any remuneration (payment).

For homecare/specialty pharmacy services in the bleeding disorders community, a consumer must assess the quality and value of customer service, and consider the business ethic of the salesperson and the company as a whole. Examples may include expensive meals, sports/entertainment tickets, or anything considered outside of normal and ordinary business practices.

FITNESS FOR ALL AGES!
Introducing HFA's new wellness tool!

Experience FitFactor!
Register and learn details about the program during special sessions at Symposium.

• Discover what a difference 30 minutes a day can make
• Learn simple techniques to stay active
• Uncover nutrition secrets hiding in your own kitchen

Make wellness a priority by making FitFactor part of YOUR routine. Stop by the HFA Booth to learn about FitFactor and register.

Visit us online at http://fitfactor.hemophiliafed.org
Section 1: Payment Information

Payment information must be received before we can process your registration.

Check the registration method: _____ $35 Individual  _____ $70 Family (Parents & Children)
Payment Method: _____ Check  _____ Money Order  _____ Credit Card (VISA, MC, AMEX, DISCOVER)
Credit Card Number ____________________________________________ Security Code __________ Expiration Date __________
(Visa | MC | AMEX  Visa/MC on back, AMEX on front)

Name on Credit Card ________________________________ Signature  _________________________________________

Other Payment Arrangements (Explain): ______________________________

Section 2: Information (please print)

Your Name: ________________________________  (Last Name, First Name, MI)
Preferred Name: ________________________________
Address: _____________________________________  City: _______________ State _______ Zip: _________
Cell Phone: (     ) _______________________  Home Phone:  ________________  Pager  ___________________________
Work Phone: (     ) ________________________  Cell Phone: (     ) ________________________
E-mail Address: ____________________________________________

Relation to Hemophilia/Other Bleeding Disorder (Circle)
Factor VIII  |  Factor IX  |  Other Factor
vonWillebrand's  |  Family Member  |  Friend of Family
Carrier  |  Inhibitor  |  Other

Breakout Session Choice for Saturday, April 16
Please Choose one below.

HFA Blood Brotherhood Session ___  HFA Moms ___
HFA Blood Sisterhood ___  HFA Dads in Action ___
HFA Kids/Teen Connection Program ___  HFA Moms ___

Other (explain)  ______________________

Do you or your spouse have health/accident insurance? ____ Yes   ____ No    If yes, please supply the following:
Company Name ____________________________________________
Policy Holder ____________________________________________
Policy Number ____________________________________________

** There is NO on-site registration for the Children/Teen Program **

To register: (Registration Deadline - April 1st, 2011.)
Mail: Hemophilia Federation of America 210 7th Street SE Ste 200B, Washington, DC 20003
Fax: 202.675.6983  Questions 1.800.230.9797 or 202.675.6984

SINGLE ADULT/FAMILY REGISTRATION

Hemophilia Federation of America
Annual Meeting and Symposium
Marriott  |  Downtown - Louisville, KY
April 15 & 16, 2011

** There is NO on-site registration for the Children/Teen Program **

To register: (Registration Deadline - April 1st, 2011.)
Mail: Hemophilia Federation of America 210 7th Street SE Ste 200B, Washington, DC 20003
Fax: 202.675.6983  Questions 1.800.230.9797 or 202.675.6984

To register children 18 and under, complete a SEPARATE Child/Teen registration for each child you have completed THIS form. Feel free to duplicate any form. Young adults (ages 18-21) living at home with parent(s) may be included as part of the family registration.

Section 1: Payment Information

Payment information must be received before we can process your registration.

Check the registration method: _____ $35 Individual  _____ $70 Family (Parents & Children)
Payment Method: _____ Check  _____ Money Order  _____ Credit Card (VISA, MC, AMEX, DISCOVER)
Credit Card Number ____________________________________________ Security Code __________ Expiration Date __________
(Visa | MC | AMEX  Visa/MC on back, AMEX on front)

Name on Credit Card ________________________________ Signature  _________________________________________

Other Payment Arrangements (Explain): ______________________________

Section 2: Information (please print)

Your Name: ____________________________________________
(Spouse's Name: ________________________________  (Last Name, First Name, MI)
Preferred Name: ________________________________
Address: _____________________________________  City: _______________ State _______ Zip: _________
Cell Phone: (     ) _______________________  Home Phone:  ________________  Pager  ___________________________
Work Phone: (     ) ________________________  Cell Phone: (     ) ________________________
E-mail Address: ____________________________________________

Relation to Hemophilia/Other Bleeding Disorder (Circle)
Factor VIII  |  Factor IX  |  Other Factor
vonWillebrand's  |  Family Member  |  Friend of Family
Carrier  |  Inhibitor  |  Other

Breakout Session Choice for Saturday, April 16
Please Choose one below.

HFA Blood Brotherhood Session ___  HFA Moms ___
HFA Blood Sisterhood ___  HFA Dads in Action ___
HFA Kids/Teen Connection Program ___  HFA Moms ___

Other (explain)  ______________________

Do you authorize the use of photographs or videos for HFA use only? _____ Yes   _____ No
Are you registering any children/teens? _____ Yes   _____ No  (If yes, please list below.) You will need to complete a separate youth/young adult for each, and send all with this form.
Children/Teen:

(On the line above, states the child’s/teen’s name and date of birth. A separate form must be filled out for each child/teen.)

Section 1: Payment Information

Payment information must be received before we can process your registration.

Check the registration method: _____ $35 Individual  _____ $70 Family (Parents & Children)
Payment Method: _____ Check  _____ Money Order  _____ Credit Card (VISA, MC, AMEX, DISCOVER)
Credit Card Number ____________________________________________ Security Code __________ Expiration Date __________
(Visa | MC | AMEX  Visa/MC on back, AMEX on front)

Name on Credit Card ________________________________ Signature  _________________________________________

Other Payment Arrangements (Explain): ______________________________

Section 2: Information (please print)

Your Name: ____________________________________________
(Spouse’s Name: ________________________________  (Last Name, First Name, MI)
Preferred Name: ________________________________
Address: _____________________________________  City: _______________ State _______ Zip: _________
Cell Phone: (     ) _______________________  Home Phone:  ________________  Pager  ___________________________
Work Phone: (     ) ________________________  Cell Phone: (     ) ________________________
E-mail Address: ____________________________________________

Relation to Hemophilia/Other Bleeding Disorder (Circle)
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HFA Blood Sisterhood ___  HFA Dads in Action ___
HFA Kids/Teen Connection Program ___  HFA Moms ___

Other (explain)  ______________________

Do you authorize the use of photographs or videos for HFA use only? _____ Yes   _____ No
Are you registering any children/teens? _____ Yes   _____ No  (If yes, please list below.) You will need to complete a separate youth/young adult for each, and send all with this form.
Children/Teen:

(On the line above, states the child’s/teen’s name and date of birth. A separate form must be filled out for each child/teen.)

Please complete a form for each child attending.

Indicate the type of registration below.

_____ Child (6 months - 2 years)  _____ Child (3 - 4 years)  _____ Child (5 - 8 years)
_____ Child (9 - 12 years)  _____ Teen (13 - 18 years)

Section 1: ALL REGISTRANTS

Parent(s) /  Legal Guardian(s): ________________________________
SYMPOSIUM SCHOLARSHIP APPLICATION

SYMPHOSIUM 2011
Annual Meeting and Symposium
Marriott  |  Downtown - Louisville, KY
April 15 & 16, 2011

To Register: (Registration Deadline - March 15, 2011.)
Completed registration forms and appropriate registration fee must be submitted with this scholarship application and returned to: Hemophilia Federation of America 210 7th Street SE Ste 200B, Washington, DC 20003
Fax: 202.675.6983  Questions call 1.800.230.9797

First Time Attendees are eligible for Scholarships. Scholarships are limited to funding availability. HFA’s Symposium Committee has requested that you pay the registration fee in order for your scholarship to be processed. If your request for a scholarship is not granted and you are unable to attend, your registration payment will be refunded.

PLEASE PRINT:

Name: ____________________________ Home Phone: (   ) - ___
Work Phone: (   ) - ___
Address: __________________________ City, State Zip
E-mail: ____________________________

[ ] Individual    [ ] Family  (include names of each family member attending and ages of children under 18)

Names of Each Family member attending

<table>
<thead>
<tr>
<th>Name</th>
<th>Sex</th>
<th>Age</th>
<th>Relation</th>
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[ ] Individual    [ ] Family  (include names of each family member attending and ages of children under 18)

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</table>

PLEASE CIRCLE TYPE OF ASSISTANCE NEEDED: HOTEL  /  AIRFARE

Departing City & Airport: 1st Choice __________________________ Second Choice __________________________
Preferred time of departure: __________________________

Are you or your spouse employed by a Homecare Company or Manufacturer [ ] Yes  [ ] No
If yes, please identify Company __________________________

Have you received a previous HFA Symposium scholarship [ ] Yes  [ ] No

Relationship to Bleeding Disorder: [ ] Factor VIII [ ] Factor IX [ ] vonWillebrand’s [ ] Other  
[ ] Parent    [ ] Child    [ ] Sibling    [ ] Spouse    [ ] Other __________________________

How did you find out about the HFA Educational Symposium?

____________________________________________________________________________________________________
____________________________________________________________________________________________________

HFA use only: Date Received __________________ Referred By: __________________ Date of Notification: __________________
Determining Essential Benefits in Health Care Reform
By Eboni Morris, HFA Policy Analyst

The Institute of Medicine (IOM) is doing a study that will make recommendations on the criteria and methods for determining the essential benefits in health plans that will be offered in health insurance exchanges starting in 2014. As you may recall, healthcare reform mandated that by 2014 adults will need to purchase health insurance or face a tax penalty. To help individuals decide what type of plan to purchase, health insurance exchanges will be set up by states to provide a one point “stop and shop” for the public. The exchange is technically not an actual place but will more likely be a virtual marketplace where individuals can go online and browse insurance offerings based on personal criteria they identify such as, age, health status, and other personal characteristics.

The U.S. Department of Health and Human Services (HHS) will most likely be issuing regulations regarding health insurance or face a tax penalty. To help individuals decide what type of plan to purchase, health insurance exchanges will be set up by states to provide a one point “stop and shop” for the public. The exchange is technically not an actual place but will more likely be a virtual marketplace where individuals can go online and browse insurance offerings based on personal criteria they identify such as, age, health status, and other personal characteristics.

The essential benefits package should always include a process to appeal a claim denial. That process should provide assurance that the insurer has an obligation to first confer with the patient’s physician, hemophilia treatment center (HTC) or the home setting.

The essential benefit package should allow patients access to needed specialists and allow the physician to formulate the best treatment regime for patients at the appropriate site of care whether in the hospital, outpatient clinic, office of the physician, hemophilia treatment center (HTC) or the home setting.

The essential benefit should be designed to encourage patient access and not impede patients from obtaining needed treatments and specialized care. Health plans should look to medical literature and treatment guidelines recommended by medical and patient organizations to determine standards of care for patients with rare and chronic conditions.

HFA submitted written comments to the initial survey conducted by the IOM in December, where several areas of concern were identified:

- Essential benefit packages should be designed to encourage patient access and not impede patients from obtaining needed treatments and specialized care. Health plans should look to medical literature and treatment guidelines recommended by medical and patient organization to determine standards of care for patients with rare and chronic conditions.
- The essential benefit package should allow patients access to needed specialists and allow the physician to formulate the best treatment regime for patients at the appropriate site of care whether in the hospital, outpatient clinic, office of the physician, hemophilia treatment center (HTC) or the home setting.
- The essential benefit package should always include a process to appeal a claim denial. That process should provide assurance that the insurer has an obligation to first confer with the patient’s physician to discuss a possible denial and the grounds for rendering such a decision.
- HFA opposes any additional limits on specific or total benefits in the packages given the statutory restrictions on lifetime and annual benefit limits already authorized in healthcare reform. HFA recommends prohibiting the implementation treatment caps of any kind by providers either by cost or in limits on treatment that has been approved by the United States Food and Drug Administration (FDA).
- State mandates have been invaluable to rare, chronic and high cost disease groups, who might otherwise be excluded from private insurance coverage. Any decision to phase out a given state mandate, where the national program provides similar protection, should be made cognizant of the possible disruption in coverage that might ensue for these particularly vulnerable groups.
- As advances in medical evidence or scientific advancement are being uncovered and approved through the FDA, as well as peer reviewed medical journals stating the improved/decreased health outcome, the national program must put in place expeditious pathways for the inclusion of new therapies and treatments in the essential benefits package when these new products or new information become available.

HFA has entered the next phase of this process and will be holding a series of public meetings to hear from various stakeholders regarding essential benefits. The first meeting took place during the week of January 12-14, 2011 and covered numerous topics within essential health benefits including: the original legislative intent of health care reform regarding essential health benefits, stakeholder recommendations for defining health insurance packages, issues regarding medical necessity, how insurers determine benefit design and coverage and various perspectives from patients and health advocacy groups.

Some common themes emerged from the presentations:

- Insurers argue that essential benefits should not be heavily prescribed through regulation to allow for the market to determine which benefits provide the best coverage options. They also cautioned against adopting state mandates because of the tens of thousands already in existence across the country. They argue that adopting a multitude of mandates on the national level will bring a dramatic rise in health insurance premiums.
- Patient and health advocacy groups argued that patients with chronic disease need access to treatment and providers of their choice. Essential benefits must be designed so that this access is not denied. They urged the panel to ensure that the benefits reflect the diverse healthcare needs of the U.S. population and are based on appropriate standards of care and practice. According to the advocacy groups, the inclusions of some state mandates are necessary to maintain comprehensive care for those with complex health needs.

HFA staff will be attending these public meetings in Washington D.C. and will be monitoring how the process unfolds. Be sure to monitor healthcare reform action in your state. States will be holding public hearings and introducing and passing legislation to set up various aspects of health reform.

Resources to Keep You Informed

- www.healthcare.gov
- www.kff.org
- http://www.iom.edu/Activities/HealthServices/FunctionalAreas/EssentialHealthBenefits.aspx

“Making a difference today, for your future!”

toll free: 877-376-4968
www.FactorSupport.com
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BEYOND PHARMACY SERVICES.

With Alphanate® you have a choice!

Available in the following potencies and packaged with Mix2Vial® Filter Transfer Set

<table>
<thead>
<tr>
<th>Potency</th>
<th>Diluent Size</th>
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<td>250 IU FVIII Range</td>
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<td>10 mL</td>
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<tr>
<td>1500 IU FVIII Range</td>
<td>10 mL</td>
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A personal touch from people who know bleeding disorders.

1 800 800-6606

Hemophilia Health Services

...for the human factor®

I joined FactorPlus℠, the Helixate® FS free trial program, because...

I'm Unique. Like Everyone Else.

Ask your doctor if you should join the FactorPlus program

Everyone has unique needs. If you or your child are currently using another product, you may want to consider Helixate FS. With FactorPlus, the Helixate FS free trial program, you can receive a supply of 5 doses, or up to 20,000 IU, of Helixate FS shipped directly to you. You’ll also receive a Welcome Kit, which includes helpful advice, patient resources, and ways to connect with others like you.

Just 3 easy steps to enroll

2. Share the form with your doctor and discuss whether Helixate FS is right for you or your child.
3. Ask your doctor to complete the form and send it in for processing.

For more information and to get started, visit www.FactorPlusprogram.com.

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

Helixate FS is a registered trademark and FactorPlus is a service mark of CSL Behring AG. Alphanate and Mix2Vial are registered trademarks of Grifols Biologicals Inc.
Symposium 2011 details inside!

Calling Dads AND Moms!

Join us for
“Parents Working Together to Make it Happen...”
Creating Harmony in Families with a Bleeding Disorder with Dr. Dave Robinson

Dr. Robinson has a PhD in marriage and family therapy. He is a licensed and practicing marriage and family therapist and an associate professor at the University of Nebraska Medical Center. He and his wife Jamie have five children (three boys and two girls) ages 17 to 5. The oldest and youngest both have moderate factor IX hemophilia.

Followed by
Rap Sessions for Dads
Co-facilitated by Dr. Dave Robinson and Mark Zatyrka

and for Moms
Facilitated by Nikita Murry, RN

Join us for this and many more programs at Symposium
April 14-16, 2011 | Marriott Downtown | Louisville, KY

Racing Ahead, A Community United!