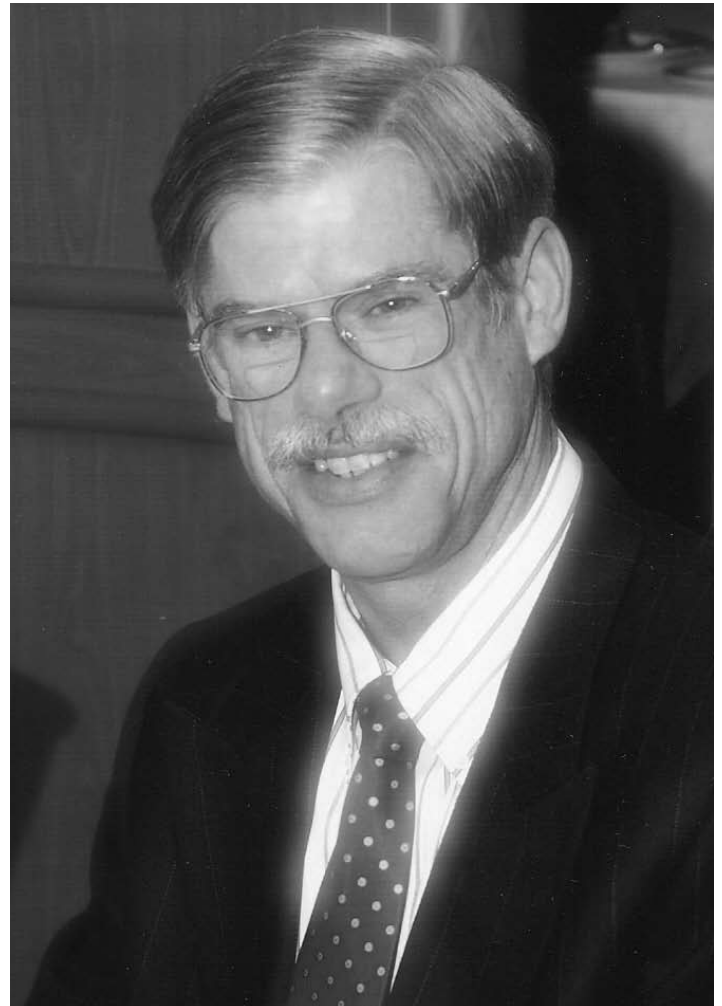
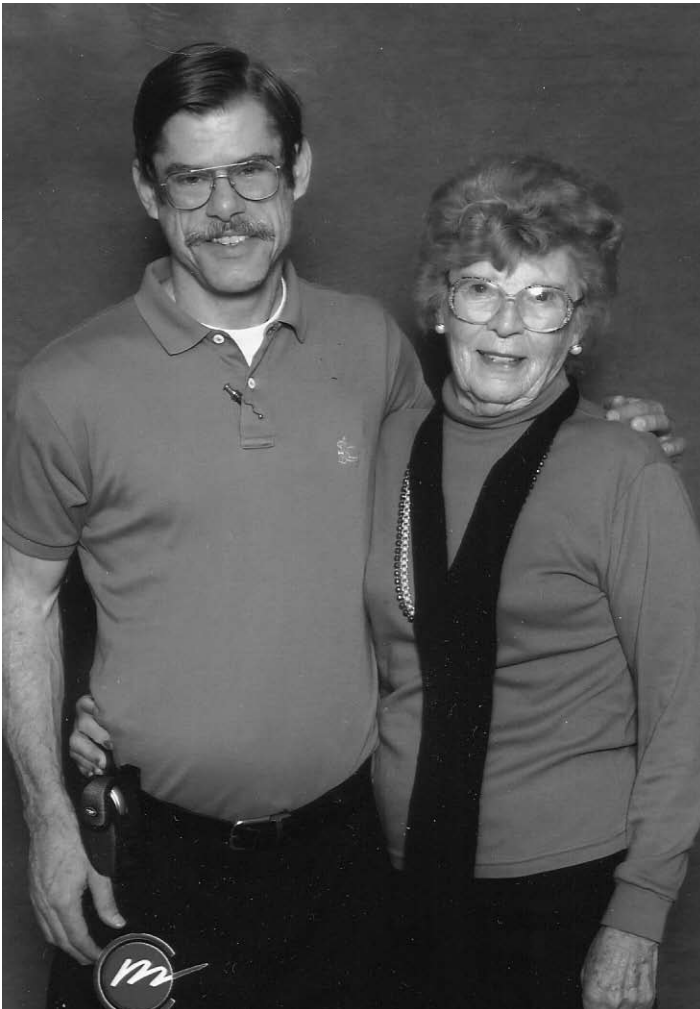


HemophiliACTION

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In Loving Memory - Bob Green



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President's Perspective

At more than halfway through the year, many great events have already come and gone, including Family Information Day in April, regional events in the Inland Empire, High Desert, Los Angeles and Santa Barbara areas from February to August, and our 2008 hemophilia camp session at The Painted Turtle in July! In addition, HFSC has held several entertaining fundraisers, which also serve as opportunities to socialize, including the EWorld red carpet event in February, HFSC On Camera Audiences in May, our first ever Wine Tasting in June,

and the recent Dancing Under the Stars event. The proceeds from these events supplement the many generous grants and donations from individuals, organizations and industry partners that fuel our programs and events, for which we are so thankful. Committed to making 2008 the Year of the Fundraiser, we have more ideas for exciting opportunities to support HFSC and would love to hear about yours.

As promised, in June 2008, HFSC celebrated a historic moment with the kick-off of the new Industry Council (IC) at the San Antonio Winery in Los



Perspectiva del Presidente

A mas de la mitad del año, muchos eventos ya han pasado, incluyendo el Día de Información familiar en abril, eventos regionales en el Inland Empire, el Desierto, Los Ángeles, y las áreas de Santa Bárbara de febrero hasta agosto, y ¡nuestra sesión de campamento del 2008 de Painted Turtle en Julio! Adicionalmente, HFSC ha tenido varios eventos para recolectar fondos, los cuales también han servido como oportunidades de socializar, incluyendo el EWORLDS de alfombra roja en febrero, HFSC Audiencia de Cámaras de HFSC en mayo, nuestra primer Probación de Vinos en junio y recientemente Bailando Bajo las Estrellas. Los fondos de estos eventos suplementaron las muchas becas y donaciones de individuales, organizaciones, y socios de la industria los cuales sirven como combustible de nuestros programas y eventos, de los cuales estamos muy agradecidos. Empeñados en hacer el 2008 el año de Recolección de Fondos, tenemos mas ideas para oportunidades de apoyar HFSC y me encantaría escuchar las tuyas.

Como prometido, en junio 2008, HFSC celebró un momento histórico al empezar el Nuevo Consulado Industrial (IC) en el San Antonio Winery en Los Ángeles. Más de 20 representantes de varios fabricantes, y compañías de cuidado en casa fueron presente en apoyo de HFSC ya que juntos nos embarcamos en un plan para mejorar la comunicación de las necesidades de la comunidad, apoyar participación igual de miembros de la industria y coordinar voluntarios y esfuerzos para recolección de fondos. Con la formación del IC, HFSC ha hecho un cambio de los miembros que estaban en la Junta Directiva de HFSC en acuerdo con la nuevas leyes revisadas. Únase conmigo en agradecerles a la cantidad de miembros de la industria quienes han servido en la Junta Directiva durante todos estos años, Clem Adkins, por sus mas de 15 años de servicio mientras el se mueve de la Junta Directiva y empieza su trabajo como el primer Presidente del Consulado Industrial.

HFSC continúa creando relaciones. En marzo, el Consulado de Hemofilia

Angeles. Over 20 members representing various manufacturer and homecare companies were present in support of HFSC as together we embark on a plan to improve communication of community needs, support equitable participation of industry members and coordinate volunteer and fundraising efforts. With the formation of the IC, HFSC has transitioned the remaining industry members from the HFSC Board of Directors in compliance with our newly revised by-laws. Along with the countless industry members who have served the HFSC Board well over the years, please join me in thanking volunteer extraordinaire, Clem Adkins, for his more than 15 years of service as he moves off the HFSC Board to assume his duties as the first President of the Industry Council.

HFSC continues to build partnership. In March, the Hemophilia Council of California (HCC) facilitated another Chapter Summit meeting of the four California foundation presidents, and our new Advocacy Committee maintains a wide and well-informed perspective through the contributions of representatives of HCC and the Committee of Ten Thousand (COTT) as well as information on the national level from the National Hemophilia Foundation (NHF) and the Hemophilia Federation of America (HFA).

It is hard to believe there is more to look forward to, but we can hardly wait for our Family Retreat at Camp Malibu in September, a regional event in Orange County planned for Fall/Winter, our golf and tennis tournament in October, and, of course,

the HFSC holiday gatherings at the end of the year, which will come all too soon. As we move into the second half of 2008, your HFSC Board, staff, partners and volunteers remain committed to our recently restated mission:

To improve the quality of life and build community for families and individuals living with hemophilia or other bleeding disorders by offering a variety of programs and services that educate, advocate and support the needs of the bleeding disorders community in Southern California.

**Jen Mermmilliod, President
Hemophilia Foundation
Southern California**



Jennifer presents Clem Adkins with a graduation cap as he graduates from the HFSC Board of Directors after over 15 years of service. Thank you, Clem!

de California (HCC) ayudó en otra reunión del Chapter Summit de los cuatro Presidentes de la Fundaciones de California, y nuestro nuevo Comité de Advocación los cuales mantienen una perspectiva de variedad de contribuciones de los representantes de HCC y el Comité de Diez mil (COTT) también como información a un nivel nacional de la Fundación Nacional de Hemofilia (NHF) y la Federación de Hemofilia de America.

Es difícil creer que hay mas que esperar, pero no podemos esperar para los eventos que vienen como el Retiro Familiar en el Campamento Malibu en septiembre., un evento regional en Orange County planeado para el otoño/

invierno, nuestro Torneo de golf y tenis este octubre y claro, las reuniones navideñas de HFSC al final del año, las que vendrán pronto. Ya que nos movemos a la segunda mitad del 2008, su Junta Directiva, los empleados, compañeros, y voluntarios continúan empeñados a nuestra reciente misión revisada:

A mejorar las calidad de vida y crear una comunidad para familias e individuos viviendo con hemofilia y otras enfermedades sanguíneas al ofrecer una variedad de programas y servicios para educar, advocar, y apoyar las necesidades de la comunidad con enfermedades sanguíneas en el Sur de California .

**Jen Mermmilliod, Presidenta
Hemophilia Foundation
Southern California**

Executively Speaking

We dedicate this edition of the HemoAction newsletter to Bob Green, who passed away on June 16th. Both Bob and his mother Gerry Green, have been active members of the hemophilia community, faithfully volunteering at the Foundation office and at our many events throughout the years. His warm smile and dry wit made all of our guests and community feel right at home.

Meeting Bob's family gave me a deeper appreciation for his kind and wonderful personality and actions. The family photographs featured in this newsletter reflect how important Bob's love of family



and his commitment to enjoying each day to its fullest.

The lessons learned from Bob are great. These photos speak volumes on how he loved his family and what his smile and warm welcome can mean to those that need extra reassurance. Although his hardships were great, that was never his message to us. He taught me the equivalent of a graduate degree in humility, kindness, generosity, and appreciation for what is important... your family. And, especially evident was his love for his Mother. These are lessons he learned and put into practice

each and every day.

Please join me in paying tribute to Bob Green and Ivan Escobar at our 29th Ivan Escobar and Bob Green Memorial Golf and Tennis Tournament on Monday, October 6 at Bramaer Country Club in Tarzana. Both men are so greatly missed. The legacy will continue as we follow in their footsteps to love and give unconditionally.

Warmest regards,

Linda Corrente
Executive Director

Hablando Ejecutivamente

Dedicamos esta edición del boletín de noticias, HemoAcción, a Bob Green, que falleció el 16 de junio. Bob Green y su madre, Gerry Green han sido miembros activos de la comunidad de la hemofilia, fielmente ofreciéndose voluntariamente en la oficina de la fundación y en nuestros muchos acontecimientos a través de los años. Su sonrisa cálida e gracia hicieron a nuestra comunidad sentirse en casa.

Haber conocido a la familia de Bob me dio un aprecio más profundo para su personalidad y acciones buenas y maravillosas. Las fotografías de la familia ofrecidas en este boletín de noticias reflejan cómo el amor hacia su familia era importante y de su compromiso de gozar cada día a su más lleno.

Las lecciones aprendidas de Bob son inmensas. Estas fotos habla volúmenes en cómo el amor a su familia y lo que pueden significar su sonrisa y recepción calidad a los que necesiten reaseguro adicional.

Aunque sus dificultades eran grandes, ése nunca era su mensaje a nosotros. Él me enseñó el equivalente de un grado graduado en humildad, amabilidad, generosidad, y el aprecio para cuál es importante... la familia. Y, especialmente evidente era su amor para su madre. Éstas son lecciones que él aprendió y que puso en práctica cada día.

Por favor ayúdeme en pagar tributo a Bob Green y a Ivan Escobar en nuestro 29th Ivan Escobar y Bob Green en el torneo conmemorativo de golf y tenis el lunes 6 de octubre en el club "Bramaer Country Club" en Tarzana. Extrañamos a ambos hombres grandiosamente. La herencia continuará si seguimos en sus pasos de amar y dar incondicionalmente.

Con el respeto más calido,

Linda Corrente
Executive Director



Robert (Bob) Green
September 9, 1952—June 16, 2008

God Saw He Was Getting Tired

God saw he was getting tired,
And a cure was not to be.
So He put His arms around him,
And whispered, "Come with me."

With tearful eyes we watched him suffer,
And saw him fade away.
Although we loved him dearly,
We could not make him stay.

A golden heart stopped beating,
Hard working hands laid to rest.
God broke our hearts to prove to us,
He only takes the best.

FUNERAL MASS For BOB GREEN was held
Saturday, June 28, 2008 at 10:00 am
St. John Vianney Catholic Church
1345 Turnbull Canyon Road
Hacienda Heights, CA 91745

In lieu of flowers:
Donations to Hemophilia Foundation of
Southern California are greatly appreciated.
6720 Melrose Avenue,
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(323) 525-0440

- Bob Green -

In Loving Memory Hemophilia Foundation Southern California

I met Bob several years ago. I have known his mother Gerry much longer. Bob was the quiet and serious type until you really got to know him. When I first met Bob I thought he was the son of Ruth Ann and Thurston Levay. Ruth Ann and Thurston are very good friends of Gerry. I really got to know Bob at our Los Angeles Holiday party last year. He and I had been talking for quite a while when we decided to go and get a slice of a really good cheesecake that someone had bought for

our party. There was a long line and by the time Bob and I got to the front there was only one slice left. He looked at me and I looked at him. Then we both said at the same time, "you take it". I said no, look at me, can't you tell that I have had too many slices of cheesecake in my life? He laughed and said let's split it and I said okay. While we were eating our cheesecake, we talked about a lot of old-timers who are no longer with us. We spoke about Vern Hayes and his sons, he also mentioned his

brothers. He really surprised me when he said, Clem, why aren't there more people like you? I jokingly said, do you think the world could tolerate more than one of me? He smiled and said probably not. Gerry joined us and we started to talk about other things. We are going to miss gentle Bob. Mary Long said it best when she said that we lost a true gentleman.

Clem Adkins



Yo conocí a Bob hace varios años. He conocido a su Mama Gerry por mucho mas tiempo. Bob era callado y serio hasta que uno lo conocía bien. Cuando primero lo conocí pensé que era el hijo de Ruth Ann y Thurston Levay. Ruth Ann y Thurston son muy buenos amigos de Gerry. Yo conocí bien a Bob en nuestra fiesta Navideña el año pasado. El y Yo habíamos estado hablando por bastante tiempo cuando el decidió ir y agarrar un pedazo de pastel que alguien

había traído. Había una larga línea y cuando Bob y Yo llegamos al frente solo había un pedazo. Me miró y Yo lo mire. Luego los dos dijimos al mismo tiempo, "tu tómallo." Yo le dije no, mírame no puedes ver que he tenido muchos pedazos de pastel en mi vida? El se rió y dijo, compártamelo y Yo le dije que si. Mientras nos comíamos nuestro pedazo de pastel, hablamos sobre las personas en el pasado quienes ya no están con nosotros. Hablamos sobre Vern Hayes y sus

hijos, el también menciona a sus hermanos. Me sorprendió cuando dijo, Clem por que no hay mas gente como Usted? Yo bromeando le dije, piensas que el mundo toleraría a otro como Yo? El sonrió y dijo, probablemente no. Gerry se acompaño y empezamos a hablar sobre otras cosas. Te extrañaremos Calido Bob. Mary Long lo dijo, que el mundo había perdido a un caballero.

Clem Adkins



HFSC Family Information Day & Annual Meeting

Saturday, April 26, 2008

As families began to arrive at the Los Angeles Police Academy they were greeted by one of our dedicated volunteers Gerry Green to proceed with the check-in process. During the morning adults had the opportunity to visit pharmaceutical and homecare company booths and an insurance information booth staffed by Meredith Zerbe, Baxter Director, Advocacy and Healthcare Initiatives, Andrew Matthews, Bayer, Insurance/Reimbursement Specialist, and Lisa Pullens RN, City Of Hope.

Youths twelve years and older, with bleeding disorders, participated in "Leading Edge", a self-efficacy development program facilitated by Pat Torrey.

Judy Brannan, kids activity room coordinator, facilitated a fun and action packed day that kept our active three to eleven year olds engaged throughout the day. Entertainment included Sunshine Storyteller Ina Buckner-Barnette, who weaves international tales, movement and song to create her participatory "Sunshine Story Shows".

Adults began their educational segment with an opportunity to ask a panel of physical and medical staff their questions. Audience members had the opportunity to ask questions in either English or Spanish on topics ranging from Joint damage/target joints, prophylaxis treatment, arthritis, pain management, and surgical options.

The Ask the Doctor panel included Dr. Laurence Logan, Dr. Doris Quon, Physical Therapist Theresa Wong from Orthopaedic Hemophilia

Treatment Center and Dr. Albert Kheradpour from Loma Linda University Health Care.

At noon families enjoyed a chicken and steak fry buffet style lunch prepared by LA Finest, LAPD Police Academy restaurant staff led by Colleen Zirbes.

After lunch, President Jennifer Mermilliod delivered the President's Annual Report which included the slate of officers for 2008.

The afternoon sessions included an Advocacy Update Panel with Greg Mermilliod as moderator, Dave Cavanaugh, Corey Dubin, Chris Templin, Kim Bernstein, and Meredith Zerbe as advocacy experts.

The day ended with a well received Parents and Teens Panel moderated by Jim Dawdy. Panel members were Greg Mermilliod, Raphael Forbes, Michael Wells, Miriam Pimentel and Rene Pimentel. Topics included life with inhibitors, ports, experiences with high school, college, job seeking, siblings, vWD, mild, moderate and severe Hemophilia.

The Hemophilia Foundation Southern California Family Information Day 2008 concluded with families enjoying a typical southern California evening at a baseball game at Dodger Stadium, courtesy of the LA Dodger's organization.



Families were greeted by Kristen Truccone, Gerry Green, HFSC Lifetime Achievement Honoree, and Linda Corrente



Moms, Marcela Shaughnessy, Judy Brannon, and Kelle McCarthy visit at the Kids activity room at Family Information day



Families at FID registration and the LAPD on horseback, and booth vendors.



HFSC Jornada Informativa de la Familia y Reunión Anual

Sábado 26 de abril de 2008

Ya que las familias empezaban a llegar a la Academia de Policía de Los Angeles, fueron bienvenidos por una de nuestras dedicadas voluntarias Gerry Green para proceder al proceso de llegada. Durante la mañana los adultos tuvieron la oportunidad de visitar las mesas de las compañías farmacéuticas y de cuidado en casa y una mesa con información de seguros la cual estaba ocupada por Meredith Zerbe, Directora de Baxter, Advocación y Cuidado de Salud, Andrew Matthews, de Bayer, Especialista en seguros y Reembolso, y Lisa Pullens RN, de City Of Hope.

Jóvenes de doce años y mayores con enfermedades sanguíneas participaron en "Lider del Borde", un programa de desarrollo y auto eficacia facilitado por Pat Torrey.

Judy Brannan, coordinadora de las actividades de los niños, facilitó un día lleno de actividades el cual mantuvo activos a niños de tres a once años de edad. El entretenimiento incluyó Contadora de Historias de Sunshine, Ina Buckner-Barnette, la cual cuenta historias internacionales, de movimiento y canciones para crear una participación de los "Sunshine Story Shows".

Los Adultos empezaron su segmento educacional con la oportunidad de hacer preguntas a un panel de profesionales médicos de terapia física. Miembros de la audiencia tuvieron la oportunidad de hacer preguntas en ingles o español sobre temas que variaban de deterioro de coyunturas, tratamiento profilaxis, artritis, mantenimiento de dolor, y opciones quirúrgicas.

El panel de Pregúntele a Un Doctor incluyó al Dr., Laurence Logan, Dra. Doris Quon, Terapeuta Física Theresa Wong de Orthopaedic Hemophilia Treatment Center y el Dr. Albert Kheradpour de Loma Linda University Health Care.

A las doce, las familias disfrutaron de un almuerzo estilo buffet de pollo o carne preparado por los mejores de Los Angeles, del Restaurante de la Academia de LAPD empleado por Colleen Zirbes. Después del almuerzo, la Presidenta Jennifer Mermilliod hizo el reporte anual incluyendo al panel de oficiales del 2008.

La sesión de la tarde incluyó el Panel de Advocación con Greg Mermilliod como moderador, Dave Cavanaugh, Corey Dubin, Chris Templin, Kim Bernstein, y Meredith Zerbe como expertos de advocación.

El día terminó con el Panel de Padres y Jóvenes moderado por Jim Dawdy. Miembros del Panel fueron Greg Mermilliod, Raphael Forbes, Michael Wells, Miriam Pimentel y Rene Pimentel. Los temas incluyeron la vida con inhibidores, catéter, experiencia en bachillerato, Universidad, busca de trabajo, hermanos y hermanas, vWd, baja, moderada y hemofilia severa.

El Día de Información de La Fundación de Hemofilia del Sur de California del 2008 concluyó con las familias disfrutando un día en el Sur de California en un juego de béisbol en el Estadio de los Dodger's, cortesía de la organización LA Dodger's.

(HFSC) Camp News 2008

This year, HFSC celebrated its 29th Camp Blood Brothers and Sisters at the Painted Turtle campsite. Since 2004 HFSC has partnered with the Painted Turtle, a Paul Newman's Hole in the Wall camping program.

We recruited 109 youths and are pleased to report that 91 campers enjoyed camp on July 10-15. Eleven campers received the Big Stick award, self infusion without any assistance, and three Big Courage award, attempted self infusion but unsuccessful. Special recognition and accolades went to our youngest seven-year-old Big Stick honoree and our first female camper to be awarded the Big Stick award.

We work diligently with the Painted Turtle to provide a wonderful, life changing, and most of all, a safe camp experience for all of our children. Many individuals are instrumental to provision of a successful and safe camp experience each and every year.

Thank you to our donors for their continued support and financial donations. Our appreciation to the superb medical staff at camp coordinated by Bob Miller for the past 27 years and to all our volunteer camp counselors Jim Dawdy, Edmund Merino, Greg Mermilliod and Linda Reyes.



(HFSC) Noticias del Campamento 2008

Este año HFSC celebró el Campamento Blood Brothers and Sisters número 29 en The Painted Turtle. Desde 2004 HFSC se ha unido con Painted Turtle y el programa de Paul Newman Hole in the Wall.

Reclutamos a 109 jóvenes y estamos contentos en reportar que 91 campistas disfrutaron el campamento en Julio 10-15. Once campistas recibieron el premio Big Stick, por haberse hecho una infusión sin ayuda y tres premios Big Courage, por haber tratado de hacer la infusión pero sin poder haberlo hecho. Enviamos un reconocimiento especial y aprobación a nuestro campista de siete años honrado con el premio Big Stick, y a nuestra primera niña campista a quien se le entregó el premio Big Stick.

Trabajamos diligentemente en Painted Turtle para proveer una fantástica experiencia en el campamento, que les cambia la vida y mas que todo es una experiencia segura para todos los niños. Muchos individuos son instrumentales en proveer una experiencia exitosa y segura cada año.

Gracias a todos los donantes por su continuo apoyo y donaciones financieras. Nuestro aprecio sincero a los empleados médicos en el campamento coordinados por Bob Miller por los últimos 27 años y a todos los voluntarios consejeros Jim Dawdy, Edmund Merino, Greg Mermilliod y Linda Reyes.

Congratulations to All Awardees/ Felicitaciones a todos los Honorados

Big Stick Awards Given at The Painted Turtle (self-infusion without any assistance)/ Premio Big Stick Entregados en The Painted Turtle (Infusión sin ninguna asistencia)

2005 (10)
Jeffrey Pollock
Alfred Coleman
Kevin Fuentes
Eduardo Venegas
Nathaniel Sammiego
Joshua Kim
Vincent Butler
Brandon Weber
Jorge Jimenez

2006 (5)
Rene Macias
Andy Macias
Oscar De Luis
Ivan Rey
Robert Rodriguez

2007 (21)
Austin Blinn
Matthew Porter
Sean Lee
Christopher Diaz
Daniel Cruz
Dustin Huerta
Jacob Bell
Cameron Dutcher
Daniel Webster
Parker Lui
Jonathan Avedano
Diego Cueller
Benjamin Him
Brian Rodriguez
Calvin Dutcher
Austin Reneau
Jarrett Guillow
Dani Arteaga
Keenan Sartain
Francisco Reveles
Lane Higgins

2008 (11)
Leonard Lopez
Thomas Tejada
Jonathan Reynosa
Jacob Powell
Julian Sanchez
Ethan Rucker
Colton Ku
Sean Cato
Bryan Ellsworth
Charlie Inglis
Jocelyn Balbuena

Big Courage Awards given at The Painted Turtle (attempted self-infusion but unsuccessful)/ Premio Big Courage Entregados en The Painted Turtle (Tratar de hacer Infusión pero no fue exitosa)

2008 (3)
Nathaniel Samaniego
Danny Cruz
Joseph Espindola



Camp Visitor's Day - 29th year

Mark Leone Camp Fund

We are pleased to share the news that the Mark Leone Camp Fund has just been established to continue the support of Camp Blood Brothers & Sisters.

Mark Leone was a respected businessman in Orange County. After he passed away, the Leone Family established a Mark Leone Camp Fund in memory of Mark and the memories they had of Mark's independence and positive spirit. Mark's sister Catherine Leone and her husband Keith Lehmann initiated the Fund on May 16 at the Foundation's Leadership Dinner in Orange County. Then Catherine and Mark's parents Aida and Philip Leone attended the Camp's Visitor's Day on July 14 and presented HFSC Executive Director Linda Corrente with another donation.

The Mark Leone Camp Fund joins a distinguished group of Funds from special individuals and their families who have been part of the Hemophilia Foundation family. Thanks to these families and our sponsors, we are able to help many more children and adolescents in the bleeding disorders community.

Camp Blood Brothers & Sisters celebrated its 29th year with 91 campers attending. Our camp expenses for 2009 will be over \$80,000, and we are grateful for the individuals that have donated to help us continue our camp tradition.

Thanks to the donors that have established camp funds and who work to promote the importance of the tradition of Camp Blood Brothers & Sisters remain until we find a cure. Thank you to Verne Hays Camp Fund, Green Family Camp Fund, Steve Emerson Camp Fund and Mark Leone Camp Fund.



Mark Leone

Fondo de Campamento Mark Leone

Estamos felices en compartir noticias que el Fondo del Campamento en nombre de Mark Leone ha sido establecido y continua apoyando el Campamento Blood Brothers & Sisters.

Mark Leone fue un respetado hombre de negocios de Orange County. Después de haber fallecido, la familia Leone estableció el Fondo de Campamento en nombre de Mark Leone en conmemoración a Mark por su temperamento positivo e independiente. La hermana de Mark, Catherine Leone y sus esposo Keith Lehmann iniciaron el Fondo en mayo 16 en la Comida de Lideres en Orange County hecha por la Fundación. Luego los padres de Catherine y Mark, Aida y Philip Leone, participaron en el Día de visitantes del Campamento en Julio 14, y presentaron a la Directora Ejecutiva Linda Corrente con otra donación.

El Fondo en Nombre de Mark Leone se une a un grupo distinguido de Fondos de individuos y sus familias quienes han tomado parte de la Familia de la Fundación de Hemofilia. Gracias a estas familias y a nuestros patrocinadores quienes han tomado parte del Día Familiar de la Fundación de Hemofilia. Gracias a estas familias y nuestros patrocinadores, podemos ayudar a muchos niños y adolescentes en la comunidad con enfermedades sanguíneas.

Camp Blood Brothers & Sisters celebra su año 29 con 91 campistas participando. Nuestros costos del campamento para el 2009 excederán \$80,000 y estamos agradecidos a los individuos que han donado para ayudarnos y continuar la tradición del campamento.

Gracias a los donantes que han establecido fondos y han trabajado para promover la importancia de la tradición del campamento Camp Blood Brothers & Sisters hasta que encontremos una cura. Gracias al Fondo de Campamento de Verne Hays, el Fondo de Campamento de Steve Emerson y el Fondo de Campamento de Mark Leone.



Linda greets The Leone Family at Camp



Rich Reveles
Cell: 714.272.5560

Cindy Martinez (Habla Español)
Cell: 909.800.2294

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HFSC Regional Meetings

Thank you to Betsy Cook, Candi Nakatani, Mellisa Franzen, and James and Cherie Guillow for their leadership and volunteer efforts in implementing educational and fun meetings throughout the Southern California region. This past year families have benefited from the educational presentations, opportunities to meet new families, and best of all a time to be together. Special appreciation to James and Cherie Guillow for their hospitality and generosity with hosting two Inland Empire meetings in their home.

February 8, 2008

Inland Empire Regional Luncheon at the home of James and Cherie Guillow

March 15, 2008

Ventura/Santa Barbara Luncheon at Coco's Restaurant & Bakery

May 3, 2008

High Dessert Physical Therapy presentation and Roller Skating Party

June 28, 2008

LA Regional Outreach Picnic in the Park

June 28, 2008

Santa Barbara Community Picnic at the Beach Grill, in Carpinteria

August 17, 2008

Inland Empire Pool Party at the Home of James and Cherie Guillow



HFSC Reuniones Regionales

Gracias a Betsy Cook, Candi Nakatani, Mellisa Franzen, y James y Cherie Guillow por su liderazgo y por servir como voluntarios en la implementación de reuniones educativas y divertidas a través del Sur de California. El pasado año las familias han beneficiado de presentaciones educativas y oportunidades de conocer a otras familias y más que todo de reunirse justos. Apreciaciones especiales a James y Cherie Guillow por su hospitalidad y generosidad al ser anfitriones dos veces en las reuniones del Inland Empire en su hogar.

Febrero 8, 2008

Almuerzo Regional en el Inland Empire en la casa de James y Cherie Guillow

Marzo 15, 2008

Almuerzo en Ventura/Santa Barbara en Coco's Restaurant & Bakery

Mayo 3, 2008

High Dessert Presentación de Terapia Física y Fiesta de Patines

Junio 28, 2008

Reunión Regional en LA Regional en el Parque

Junio 28, 2008

Picnic de la Comunidad en Santa Barbara en el Beach Grill, en Carpinteria

Agosto 17, 2008

Fiesta de Piscina en el Inland Empire en al casa de James y Cherie Guillow

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HFSC at the Circus, 2008



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29th Ivan Escobar & Bob Green Memorial Golf and Tennis Tournament Executive Team

Clem Adkins, Tennis Tournament
 Corey Parker & Dan Goodwin, Golf Tournament
 Angie Yanez, Golf Foursomes & Sponsorships
 Randy DeSantis, Regional Corporate Sponsorship
 Karen Arrieta, Debbie Balck, Silent and Live Auction
 Nicole Wruck, Registration
 Linda Corrente, Memorial Reception and Dinner
 Volunteers, Wagner Lemus
 Participants: Premium Gifts, Shirley Bonizan & Linda Reyes



Corey Parker, Randy De Santis, Linda Corrente, Clem Adkins, Debbie Black

HFSC Fundraising: Hollywood Red Carpet E-World Media event



Linda Corrente, HFSC Executive Director, is interviewed.



Red carpet attendees enjoy Play TV



Helena Smith, HFSC office Manager sells raffle tickets at Hollywood Red Carpet event



The HFSC staff promoting our mission and raffle tickets at E-world media event

NHF Leadership Reception at NY Wax Museum



Linda Corrente raffles Guitar donated by Factor Support Network Homecare



Linda tells Joy Philbin about HFSC at recent Community event



Linda at reception for new NHF Ceo Val Bias at the New York Wax Museum, Hollywood Red Carpet EWorld Media Event, February 8



Val Bias and his wife among the celebrities.

Wine Tasting



Orange County Leadership Dinner.



The wine tasting crowd.



A detailed explanation.



Alex Rodriguez supports HFSC wine tasting fundraiser

Ralph's Rewards Card and Community Contributions Fundraiser

Now you can support the HFSC every time you buy groceries. If you provide HFSC with your Ralph's reward number, each time you shop we receive a small % of your shopping total.

Thanks to all of you that have previously joined the Ralph's community contributions program. As some of you may have already discovered, Ralph's recently launched a new Ralph's Rewards program. All customers will be completing a new application and receiving a new card in the Ralph's store.

HOW DOES THIS AFFECT RALPH'S COMMUNITY REWARDS?

Tarjeta de Premios de Ralph's y Contribuciones a Recolectar Fondos para la Comunidad

Ahora usted puede apoyar a HFSC cada vez que compre su comida. Si le provee su número de tarjeta de Ralph's a HFSC, cada vez que usted compre, nosotros recibimos un porcentaje pequeño del total de su compra.

Gracias a todos los que se han unido previamente al programa de Ralph's de contribuciones a la comunidad. Como muchos de ustedes de darían cuenta, Ralph's recientemente empezó un programa

As long as HFSC supporters LINK their previously enrolled Ralph's Club Card with their new Ralph's Rewards Card, their participation with our organization will continue without change. As a confirmation point, everyone will still see the organization contribution note at the bottom of their receipt.

If you did not link your two cards together when you applied for your new Rewards Card or see HFSC at the bottom of your receipt, you should contact the HFSC to enroll your new Ralph's Rewards Card.

nuevo de premios. Todos los clientes tendrán que llenar una aplicación nueva y recibirán una tarjeta nueva en la tienda Ralph's.

¿COMO AFECTA ESTO A LOS PREMIOS DE RALPH'S PARA LA COMUNIDAD ?

Con tal de que los apoyantes de HFSC UNAN la tarjeta previamente inscrita con la tarjeta nueva de Premios de Ralph's, su participación con nuestra

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Email: Your email
Click O.K., click Install, click O.K.

Please note: Upper left corner boomerang icon minimizes menu bar
Upper right corner removes menu bar completely from screen
click on - sign to bring menu back into view

organización continuará sin ningún cambio. Como un punto de confirmación todos todavía podrán ver la organización a la que contribuyen en la parte abajo del recibo.

Si no pudo unir las dos tarjetas cuando aplicó para la nueva tarjeta de Premios, o si no ve HFSC en la parte de abajo del recibo, debe comunicarse con HFSC para inscribir la nueva tarjeta de Premiso de Ralph's.

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For more information call the "Time for ReFacto" Trial Prescription Program administrator at 1.800.710.1379, Monday through Friday 9:00 AM to 5:00 PM eastern time.

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Through Participation

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Through Progress

Looking to the future, Baxter is dedicated to improving current therapies. We will continue to invest in research to offer new and better ways to manage hemophilia A—innovations inspired by listening to you.



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Venous Access, Home and Self-Infusion Kits, Advoy, and www.thereforyou.com



COMMUNITY PROGRAMS

Patient & Community Educational Programs

Facts First, Camp SuperFly, Knights' Crossing, Careers, Education, & Opportunities (CEO), and more



INSURANCE & ADVOCACY

Assistance, Support, & Information

Factor Assist, grants for advocacy training and advocacy projects, and a toll-free assistance hotline 1-888-BAXTER9



CLINICAL INITIATIVES

Support of Medical Education & Clinical Training Programs

Funding training for new hematologists and educational programs for nurses and healthcare partners

For more information on Baxter programs and services, visit www.thereforyou.com.

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Hemophilia Council of California - 2008 Legislative Day

Over 100 people came to the State Capitol on May 16th to participate in the Hemophilia Council of California's Legislative Day. This state-wide annual event is a chance for people to show their support for issues important to people in the bleeding disorders community, meet people from all over the state and brush up on their advocacy skills.

Legislative Day is also the final day of the Future Leader program, an advocacy training program for young people with bleeding disorders and their siblings. Nine young men and women tested their newfound advocacy skills by taking leadership roles during visits. They were also awarded certificates during the luncheon to recognize their completion of the training program and their outstanding efforts on Legislative Day. During the luncheon, Joshua Nuno, a young man from Southern California, was given the Young Advocate award for his exceptional efforts to stop his school from excluding him from field trips. The packed room also wished outgoing HCC Executive Director Val Bias well in his new position as Executive Director of NHF. Val has worked to expand the Future Leader and Legislative Day programs while leading HCC.

Val Bias and HCC's Legislative Advocate Terri Cowger Hill briefed everyone on the issues and then the Legislative Day participants met with their senators and assembly members to oppose changes to Medi-Cal, CCS and GHPP. The issues included opposition to the quarterly renewal applications (QSR's), the elimination of adult dental benefits and the 10% provider rate cuts. Moving from annual to quarterly renewals for Medi-Cal would force many eligible

children and adults off of the program and into emergency rooms with more late applications, resulting in reduced access to medical care. Also, it would add more layers of bureaucracy. Eliminating adult dental services under Medi-Cal would also significantly reduce the availability of this necessary care for people with hemophilia. Finally, HCC Future Leadership and Legislative Day participants lobbied to repeal the 10% cuts to Medi-Cal, CCS and GHPP reimbursements as these cuts negatively impact our physicians, homecare pharmacies and HTC's. These specific cuts have already been adopted and are set to take effect July 1, 2008. These cuts will impact access to providers and the availability of care to people with hemophilia.

Participants also lobbied in support of Hemophilia Standards of Service legislation (SB 1594/Steinberg), a bill sponsored by HCC that would establish standards of service for companies that deliver blood clotting factor and related supplies to ensure a continued high quality of care. Senator Darrell Steinberg is the sponsor of this bill, and he and his staff member Myesha Jackson were recognized with awards at a luncheon ceremony to thank them for their long-standing support of the bleeding disorders community. Senator Steinberg was unable to attend the luncheon, so Ms. Jackson accepted the Senator's award on his behalf.

For more information on the Future Leadership or Legislative Day programs, please contact Robyn Ireland: hccassist@aol.com (916) 498-3780.

Consulado de Hemofilia - Día Legislativo 2008

Mas de 100 personas vinieron a la Capital Estatal en mayo 16 para participar en el Día Legislativo del Consulado de Hemofilia de California. Este evento estatal es la oportunidad para que las personas demuestren su apoyo por la comunidad con enfermedades sanguíneas, para que conozcan a personas alrededor del estado y para que aprendan un poco sobre habilidades de advocación.

El Día Legislativo es también el último día del Programa de Líder del Futuro, y entrenamiento de advocación para personas jóvenes con enfermedades sanguíneas, hermanos y/o hermanas. Estos nueve jóvenes tuvieron la oportunidad de probar sus habilidades de advocación al hablar con líderes durante las visitas. También fueron reconocidos con premios por completar el programa de entrenamiento y por sus grandes esfuerzos en el Día Legislativo.

Adicionalmente, Joshua Nuno y joven del Sur de California, fue quien ganó el premio por su Advocación y por sus grandes esfuerzos al prevenir que sus escuela lo excluyera de viajes escolares. La sala llena de gente, también le deseo bien a Val Bias en su posición como Director Ejecutivo de NHF. Val también ha trabajado en ampliar los programas de Día Legislativo Y Líder del Futuro mientras él ha estado en HCC.

Val Bias y Representante Legislativo de HCC informó sobre los temas y los participantes del Día Legislativo se reunieron con Miembros del Senado y la Asamblea para oponer los cambios a Medi-cal, CCS y GHPP. Los temas incluyeron la oposición de renovación cada tres meses, oposición a la eliminación de beneficios dentales y la oposición de cortes de 10% para proveedores. Al cambiar la renovación de cada tres meses de Medi-cal, forzaría a muchos niños y adultos que califican fuera del programa y a salas de emergencia por causa de aplicaciones que lleguen tarde y capas adicionales de burocracia, resultando en poco acceso a cuidado medico. La eliminación de servicios dentales bajo Medi-cal también disminuye significativamente la habilidad de cuidado necesario par personas con hemofilia. Finalmente, Participantes del Día Legislativo avocaron por la reapelación de los cortes de 10% al reembolso para proveedores de Medi-cal, CCS y GHPP, los cuales tiene un impacto negativo para doctores, farmacias de cuidado en casa y Centros de Tratamiento. Estos cortes específicos ya han sido adoptados para tomar efecto Julio, 1 2008. Estos cortes afectaran el acceso a proveedores y disponibilidad de cuidado para personas con hemofilia.

Los Participantes también avocaron por el Apoyo de la Legislación de Reglas de Servicio para Hemofilia (SB 1594/Steinberg), una ley patrocinada por HCC la cual establecería reglas de servicio para compañías que entregan factores coagulantes y auxiliares relacionados, para asegurar cuidado de alta calidad. El Senador Darrell Steinberg es el patrocinador de esta ley, y el y Myesha Jackson fueron reconocidos con premios en nuestro almuerzo ceremonial para agradecerles por su apoyo de tanto tiempo a la comunidad con enfermedades sanguíneas. El Senador Steinberg no pudo participar en el almuerzo, así que Ms. Jackson acepto el premio del Senador.

Para más información sobre el Programa de Líderes del Futuro o el Día Legislativo, por favor comuníquese con Robyn Ireland: hccassist@aol.com (916) 498-3780

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Dutch Study Measures Aerobic Capacity in Boys with Hemophilia

Researchers from the Netherlands recently conducted a small study to assess whether aerobic capacity differs significantly between unaffected boys vs. those with hemophilia. Aerobic capacity is the maximum amount of oxygen the body can take in and use to generate energy. The study, led by Raoul H.H. Engelbert, PT, PCS, PhD, showed that there was a disparity between the children with hemophilia and those without it.

The study comprised 47 boys with hemophilia from the Van Creveldkliniek Department of Haematology at the University Medical Center in Utrecht. The test subjects ranged in age from 8.2-17.4 years, with an average age of 12.9 years old. Investigators evaluated anthropometry (measurement of human body) in addition to testing muscle strength, joint impairment, functional ability and aerobic capacity. Even though all the affected boys performed the fitness tests at maximum or near-maximum levels and none reported bleeds or other adverse events, an assessment of their aerobic capacity pointed to poorer performance.

Specifically, relative peak oxygen, peak heart rate and peak working capacity were considerably lower among hemophilia patients when compared with unaffected control subjects. Total muscle strength, however, was normal.

Further, test results of joint health status showed only marginal impairments. Although researchers also stated that a "substantial proportion" of Dutch children with hemophilia were overweight, there was no reported difference in the amount of physical activity in which they engaged. It should be noted that these rates of physical activity were self-reported.

The study "Aerobic Capacity in Children with Hemophilia," was published in the June 2008 issue of The Journal of Pediatrics.

Source: Reuters Health Medical News, July 16, 2008

Academic Scholarships

Academic Scholarships for Students with Bleeding Disorders

For U.S. residents:

Beth Carew Memorial Scholarship

Amount: Ten \$2,000 college awards

Candidate: Person or family member with hemophilia or other bleeding disorder, enrolled full-time in an accredited college

Deadline: April 4, 2008

Contact: Colburn-Keenan Foundation, Inc.

Phone: (800) 966-2431

www.colburnkeenan.org

Bill McAdam Scholarship Fund

Amount: one \$2,000 award

Candidate: A person with hemophilia, VWD or other hereditary bleeding disorder or the person's spouse, partner, child or sibling planning to attend an accredited college or university or certified training program.

Deadline: May 15, 2008

Contact: Cathy McAdam Scholarship Fund

22226 Doxtator, Dearborn, MI 48128

Phone: (313) 563-1412

mcmadam@comcast.net

Download Application: <http://www.hemophilia.org/NHFWeb/Resource/StaticPages/menu0/menu1/menu53/BillMcAdamScholarshipApplication.doc>

BioRx/Hemophilia of North Carolina Educational Scholarships

Amount: Three \$2,000 scholarships

Candidate: intended for caregivers of children affected with bleeding disorders, a person who has been diagnosed with hemophilia, or a sibling of a person diagnosed with hemophilia. Preference will be given to a hemophilia community member, parent and family members seeking healthcare related education from an accredited college or university or certified training program.

Deadline: May 1, 2008

Contact: BioRx

Phone: 866-442-4679

www.biorx.net

cbarnes@biorx.net

Christopher Pitkin Memorial Scholarship

Amount: amounts of scholarships vary from \$500 - \$1,000

Who can apply: All members of the hemophilia and bleeding disorders community, including spouses, siblings and children

Deadline: July 25, 2008

Contact: Hemophilia Foundation of Southern California

6720 Melrose Avenue, Hollywood, CA 90038

Phone: (323) 525-0440

www.hemosocal.org

CoaguLife Education Scholarships

Amount: several scholarships up to \$5,000 each

Candidate: Person with hemophilia, von Willebrand disease or other inherited bleeding disorder

Deadline: July 25, 2008

Contact: CoaguLife

Phone: (866)-858-9200

www.coagulife.com

"Education is Power" Scholarship

Amount: A minimum of twenty-five scholarships will be awarded, ranging from \$500-\$2,500

Candidate: Individuals living with hemophilia and von Willebrand Disease entering or attending a community college, junior college, four-year college, university, or vocational school.

Important note: Prior applicants and winners are encouraged to reapply each year

Deadline: May 1st, 2008

Contact: MedPro Rx, Inc.

Phone: (866) 528-4963

www.medprorx.com

educationispower@medprorx.com

Edu-Grants (for K-12 learning assistance)

Amount: Up to \$500 financial award per child per year to help pay for tutors or learning assistance when students (K-12) miss a significant number of school days due to hemophilia

Candidate: people with hemophilia diagnosed with an inhibitor or individuals with factor VII deficiency

Deadline: Ongoing

Contact: Novo Nordisk

Phone: (856) 573-9400

www.novoseven-us.com

Eric Delson Memorial Scholarship

Amount: One \$1,500 award per year, renewable up to four years, for private high school. Three \$2,500 award per year, renewable up to four years, for college or trade school.

Candidate: students clinically diagnosed with hemophilia or von Willebrand disease

Deadline: July 1, 2008

Contact: Caremark

Phone: (866) 792-2731

www.caremark.com

Eric Dostie Memorial College Scholarship

Amount: Ten \$1,000 awards

Candidate: Student with hemophilia or other related bleeding disorder, or a family member. Must be a US citizen and enrolled full-time in an accredited two or four year college program

Deadline: March 1, 2008

Contact: NuFACTOR

Phone: (800) 323-6832 ext 1300

www.nufactor.com

Factor Foundation Scholarship Program

Amount: several scholarships awarded for Continued Education and Creative Arts

Candidate: individual with a congenital bleeding disorder and their siblings or parents

Deadline: not yet posted

Contact: Factor Foundation of America

Phone: (866) 843-3362

www.factorfoundation.org

Hemophilia Federation of America

Amount: Several \$1,500 awards

Candidate: Person with hemophilia or VWD, including parents and siblings attending any accredited two- or four-year college, university or vocational/technical school in the US. Also include awards for completing a visual/performing art project

Deadline: April 1, 2008

Contact: Hemophilia Federation of America

1405 West Pinhook, Suite 101, Lafayette, LA 70503

Phone: (800)-230-9797

www.hemophiliafed.org

info@hemophiliafed.org

Hemophilia Health Services Memorial Scholarship Program

Amount: several \$1,500 and more awards a full academic year.

Candidate: A US citizen with hemophilia, VWD or other bleeding disorder. Must be a high school student about to attend college, a college student, a college senior planning to attend graduate school or graduate school student.

Deadline: May 1, 2008

Contact: Hemophilia Health Services

Phone: (800) 289-6501 ext 5175

www.HemophiliaHealth.com access key is HEMO

Joshua Gomes Memorial Scholarship Fund

Amount: several \$1,000 awards

Candidate: Individuals living with HIV/AIDS accepted or enrolled in college in the US

Deadline: July 15, 2008

Contact: Joshua Gomes Memorial Scholarship Fund

2700 South Emerson Street, Englewood, CO 80113-1737

Phone: (303) 761-3055

www.joshuagomes.org



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Academic Scholarships (cont'd)

Kevin Child Scholarship
Amount: one \$1,000 award
Candidate: Individual diagnosed with hemophilia or von Willebrand disease attending college, university or vocational school in the US
Deadline: Friday, June 27, 2008
Contact: National Hemophilia Foundation
116 West 32nd Street, 11th Floor, New York, NY 10001
Phone: (800) 42-HANDI ext. 3700 (ask for Renee)
www.hemophilia.org

Lawrence Madeiros Scholarship
Amount: several \$1,000 or more awards
Candidate: Student with an inherited bleeding disorder or other chronic disorder attending an accredited college or university
Deadline: June 1, 2008
Contact: Positudes
The Lawrence Madeiros Scholarship
Phone: (518) 661-6005
www.adirondackspintacular.com

Michael Bendix Sutton Foundation
Amount: two \$2,000 scholarships
Candidate: Student with hemophilia pursuing pre-law study
Deadline: write to address below
Contact: Michael Bendix Sutton Foundation
c/o Marion B Sutton
300 Maritime Avenue, White Plains, NY 10601

Mike Hylton and Ron Niederman Memorial Scholarships
Amount: Five \$1,000 scholarships
Candidate: A person with hemophilia or von Willebrand disease and their immediate family members
Deadline: April 30, 2008
Contact: Factor Support Network Pharmacy
Phone: (877) 376-4968
www.factorsupport.com
scholarships@factorsupport.com

Millie Gonzalez Memorial Scholarship
Amount: Two \$1000 awards
Candidate: Women diagnosed with hemophilia or von Willebrand disease entering or attending college or vocational school
Deadline: April 30, 2008
Contact: Factor Support Network Pharmacy
Phone: (877) 376-4968
www.factorsupport.com
scholarships@factorsupport.com

National Cornerstone Healthcare Services
Amount: several \$500 to \$1,000 awards per a year
Candidate: A person diagnosed with a bleeding disorder, parent, spouse, partner, child or sibling of a person diagnosed with a bleeding disorder.
Deadline: April 1, 2008
National Cornerstone Healthcare Services
24747 Redlands Blvd, Suite B, Loma Linda, CA 92354
Phone: (877) 616-6247
www.nc-hs.com

Project Red Flag Academic Scholarship for Women with Bleeding Disorders
Amount: two awards of \$2,500
Candidate: Applicants must be female residents of the US and have a bleeding disorder (includes von Willebrand disease, hemophilia carrier or other clotting factor deficiencies)
Deadline: May 16, 2008
Contact: National Hemophilia Foundation
116 West 32nd Street, 11th Floor, New York, NY 10001
Phone: (800) 424-2634 ext. 3700 (ask for Renee)
www.projectredflag.org

Rachel Warner Scholarship
Amount: Funds are varied and limited
Candidate: Person with bleeding disorder
Deadline: May 1, 2008
Contact: The Committee of Ten Thousand
c/o Rachel Warner Scholarship
236 Massachusetts Avenue, Suite 609, Washington, DC 20002
Phone: (800) 488-2688
www.cott1.org
cott-dc@earthlink.net

Salvatore E Quinci Foundation Scholarship
Amount: Two \$2,000 scholarships
Candidate: Person diagnosed with hemophilia or other bleeding disorder accepted into an accredited university, college or vocational/technical school
Deadline: April 4, 2008
Contact: S.E.Q Foundation, Inc
178 Florence Street, Melrose, MA 02176
Phone: (781) 760-7138
www.seqfoundation.org

Soozie Courter Sharing a Brighter Tomorrow Hemophilia Scholarship Program
Amount: Two \$7,500 graduate study awards, sixteen \$5,000 college scholarships and two \$2,500 vocational scholarships to applicants with hemophilia A or B.
Candidate: A student with hemophilia A or B who is a high school senior (or has graduate equivalency diploma), college student (undergraduate or graduate) or vocational school student.
Deadline: April 4, 2008
Contact: Wyeth/Genetics Institute, Inc.
Phone: (888) 999-2349
www.hemophiliavillage.com

Professor Ulla Hedner Scholarships
Amount: from \$2000 to \$7000 per person per year
Candidate: Having either hemophilia with an inhibitor or FVII deficiency . Must be a high school senior, college or vocational student under the age of 23
Deadline: April 30, 2008
Contact: Novo Nordisk
Phone: (877) 668-6777
www.novoseven-us.com

For clarification on any of the scholarships listed above, please contact the individual sponsor for each respective program.

This list was compiled by HANDI and revised January 2008. To request a Scholarship Resource Packet, please call HANDI: 1-800-424-2634 or e-mail: handi@hemophilia.org.

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Bob Green family photo

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2008 Calendar of Events

August 17	Pool Party, Inland Empire
August 27	HFSC Board Meeting Conference Call
Sept. 12-14	HFSC Retreat, Camp Malibu
Sept. 24	HFSC Board Meeting
Oct. 6	Golf & Tennis Tournament, Braemar Country Club
Oct. 22	HFSC Board Meeting Conference Call
Nov. 13-15	NHF Convention, Denver, CO
November 26	HFSC Board Meeting
December	Holiday Parties
December 24	Board Meeting Conference Call

***Dates & locations are subject to change

HFSC phone # 1-800-371-4123 or 323-525-0440

Calendario de Eventos 2008

Agosto 17	Fiesta de Piscina, Inland Empire
Agosto 27	Llamada conferencia de la Junta Directiva de HFSC
Sept. 12-14	Retiro de HFSC, Campamento Malibu
Sept. 24	Reunión de la Junta Directiva HFSC
Oct. 6	Torneo de Golf & Tennis, Braemar Country Club
Oct. 22	Llamada Conferencia de la Junta Directiva de HFSC
Nov. 13-15	Reunión de NHF, Denver, CO
Nov. 26	Reunión de la Junta Directiva de HFSC
Diciembre	Fiestas Navideñas
Diciembre 24	Llamada Conferencia de la Junta Directiva de HFSC

***FECHAS Y LUGARES PUEDEN CAMBIAR .

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