

February 2024

Caring for people with bleeding disorders in developing countries one at a time

There are ways to help you

AFFORD YOUR HEMOPHILIA A MEDICINE.



## The Power of Determination



Subhankar is finding success

Subhankar, from India, received a Micro-Enterprise Grant to start his own blood collection laboratory.

Before he was awarded this grant, his family struggled to maintain income for basic needs. Subhankar's mother is a homemaker and his father is paralyzed. Now, his business is doing well and Subhankar is able to care for his family.



He is also passionate about helping the poor and gives special discounts to those in need. Subhankar's income has doubled so far, and he is confident in expanding his business in the near future!

"Right now, I am doing it for my family, but in the next two years, I will show everyone that you can do anything."

## Save One Life commits to sending sponsorship funds to beneficiaries — even after they lose a sponsor.

1246 children sponsored through our Sponsorship Program
174 covered by the Save One Life Community Fund



Please consider sponsoring **one of the 174 children** available for sponsorship, like brothers Chomroeun and Sothearak and from Cambodia.

Both Chomroeun and Sothearak have moderate hemophilia A. The family's income is less than \$300/month!



Chomroeun is 6 years old and Sothearak is just a year old!

Let's help this family — Sponsor Chomroeun or Sothearak!









MAT-US-2208907-v3.0-07/2023



## Helping hands work together to accomplish great things —

We are thrilled to have Scott, Brandon, Shannon and Ujjwal carrying the torch across the country to continue Barry's legacy. They will be riding a collective 2,938 miles from San Diego, California to St. Augustine, Florida.



Donate to the Team Fundraiser!

Come out to cheer them on at our community rides along the route!



Wheels for the World 2024

Follow us on social media for updates on our relay team and all our rides!











## **Rare Disease Week**

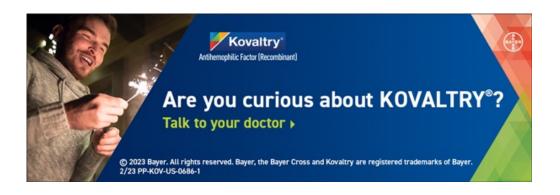
Rare Disease Week on Capitol Hill 2024 took place February 25 - 28 in Washington DC. Hosted by the Rare Disease Legislative Advocates, a program of the EveryLife Foundation for Rare Diseases, this multi-day event brings together rare disease advocates from across the country to make their voices heard by their members of Congress.



As part of the event, on Sunday Feb 25 a screening of Bombardier Blood and panel discussion were held. Executive Director Chris Bombardier had the amazing opportunity to connect with the broader rare disease community. Advocating for change together is a powerful way to start a movement.



 $Thank\ you\ to\ the\ Every Life\ Foundation\ and\ sponsor\ of\ the\ screening,\ Takeda.$ 



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Save One Life, Inc. | 55 Ferncroft Rd., Suite 200, United States, Danvers, MA 01923

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