WORLD HEMOPHILIA DAY 2016
ABOUT THE WFH

The vast majority of people with a bleeding disorder do not know they have one

Treatment for All The Vision of All

For over 50 years, the World Federation of Hemophilia (WFH) has provided global leadership to improve and sustain care for people with inherited bleeding disorders, including hemophilia, von Willebrand disease, rare factor deficiencies, and inherited platelet disorders.

We save and improve lives by:

• training experts in the field to properly diagnose and manage patients;
• advocating for adequate supply of safe treatment products; and
• educating and empowering people with bleeding disorders to help them live healthier, longer lives.

However, much more still remains to be done. The reality is that the majority of people with bleeding disorders still receive very inadequate treatment or no treatment at all. The percentage is even higher for those with von Willebrand disease and rare factor deficiencies. Our vision of Treatment for All is that one day; all people with a bleeding disorder will have proper care, no matter where they live.

Treatment for All means:

• proper diagnosis, management, and care by a multidisciplinary team of trained specialists;
• safe, effective treatment products for all people with bleeding disorders; and
• expanding services beyond hemophilia, to those with von Willebrand disease, rare factor deficiencies, and inherited platelet disorders.

The World Federation of Hemophilia (WFH), an international not-for-profit organization, was established in 1963. It is a global network of patient organizations in 127 countries and has official recognition from the World Health Organization.

For more information go to www.wfh.org
For additional World Hemophilia Day materials, visit www.wfh.org/whd