

Join us

Our members are the heart of the Haemophilia Society, and we would love to have you onboard. Membership is completely free, and gives you access to lots of benefits.

Signing-up online is quick and easy at haemophilia.org.uk/membership

Alternatively, please complete the basic details below, and we will contact you to complete the process and welcome you to the Haemophilia Society.

Title

First name

Surname

Address

.....

Postcode

Telephone number

Email

Data Protection: The Haemophilia Society is registered under the Data Protection Act 2018, which protects personal data of individuals and their privacy. We have requested the information this form in order to function effectively and provide the best possible support. Please tick the box below if are happy to provide us with this information.

I am happy to provide you with the information requested on this form.




Please return your completed form to: FREEPOST RUBC-ASXT-CACH, The Haemophilia Society, 52b Borough High Street, London SE1 1XN



We'd love to hear from you. Whether you want to become a member, need some information, or you want to find out about local groups, please get in touch.

Contact details

The Haemophilia Society
52b Borough High Street
London SE1 1XN
Phone: 020 7939 0780
Email: info@haemophilia.org.uk
Web: haemophilia.org.uk

 [HaemophiliaSocietyUK](https://www.facebook.com/HaemophiliaSocietyUK)
 [HaemoSocUK](https://twitter.com/HaemoSocUK)
 [thehaemophiliasociety](https://www.instagram.com/thehaemophiliasociety)

The Haemophilia Society is a charity. We can only continue to support our members for free thanks to the generosity of our supporters and the community raising money on our behalf.

If you'd like to donate to help to fund our work, however large or small the amount, it would be hugely appreciated. You can find out more at haemophilia.org.uk/get-involved/fundraising/



The UK-wide organisation for you and everyone affected by a bleeding disorder.



Join now for free and find out how we can support you.

The Haemophilia Society

We are the UK-wide charity for everyone affected by a genetic bleeding disorder, and our community is here for you.

Living with a bleeding disorder is never black and white. And it's not all blood-red either.

At the Haemophilia Society (THS) we want to empower everyone affected by a genetic bleeding disorder to live life to the full, whatever your stage in life.

One in 2,000 men, women and children in the UK have a diagnosed bleeding disorder, which are a group of conditions, including haemophilia and von Willebrand disorder, that result when the blood cannot clot properly. A third of all bleeding disorders are the result of a gene mutation with no previous family history, which means that a diagnosis can come completely out of the blue.

Life after diagnosis may start out hard, but we learn by doing, making mistakes, asking questions and surrounding ourselves with people who understand. Then, slowly, it just becomes life again.

Together for life

Become a member of THS and you will be joining a community thousands strong. Membership is free and benefits include:

Events. Our events offer expert knowledge and unique friendships. They include weekends for families with a newly diagnosed child, Youth Camps, and a range of events for people with a bleeding disorder and their family. At every event you can learn more, and meet and share experiences with other people in our community.

Expert resources. For every life stage, we have resources to support you to live well. Visit haemophilia.org.uk/resources/publications or telephone **020 7939 0780**.

News and information. Email updates and your member magazine will keep you in touch with the latest developments in the bleeding disorders community, and news from members across the UK.

“Being a member I’ve made friendships with people who I might never normally meet, people who are facing some of the same challenges as me.” Val






Campaigning.

Together with our members, we raise awareness about bleeding disorders and lobby Government, the NHS and clinicians for the best possible care and equal access to effective treatment

Local connections. THS local groups can be a great source of support and friendship. Activities might include fun days, hosting expert speakers, or local awareness raising. Contact us to find your nearest group on **020 7939 0780** or email services@haemophilia.org.uk

Join the conversation.

Follow the latest news and views and meet other THS members on social media:

 [HaemophiliaSocietyUK](https://www.facebook.com/HaemophiliaSocietyUK)
 [HaemoSocUK](https://twitter.com/HaemoSocUK)
 [thehaemophiliasociety](https://www.instagram.com/thehaemophiliasociety)



“Meeting other families going through the same as us really turned things around. I felt reassured that we could be ‘normal’ again and felt a lot more positive about the future.” Lauren