



# THE HAEMOPHILIA SOCIETY



For everybody affected by  
an inherited bleeding disorder

**We're the UK-wide charity for everyone living with an  
Membership is free and open to everyone.  
We're here to help you live life well, providing you and  
services and support from birth to older age.**



## **What we offer**

*The latest information on care and treatment, so you can make informed choices about your treatment and care, and inspire others to do the same.*

- Services and events where you will hear from expert speakers and meet other members of our community.
- Regular email updates and mailings on the issues affecting you and your family: services, opportunities, campaigns and the changing NHS
- Our twice-yearly members' magazine, HQ, packed with new insights, inspiring stories and experiences
- Publications to help you understand more about life with a bleeding disorder
- Practical information, for example where to find help with benefits.

**inherited bleeding disorder, their families and carers.**

**and your family with practical information,**



*Support, friendships and a listening ear when needed: with more than 25,000 people diagnosed across the UK, no-one should have to feel isolated or alone. Support from our member community is available online, face-to-face and over the phone.*

*We're also building specific support networks for:*

- People living with an inhibitor
- Women with a bleeding disorder via our Talking Red activities
- Carriers of an affected gene
- Single parents raising a child / children with a bleeding disorder.

*'The Haemophilia Society has been with us every step of the way'*

*'A massive weight has been lifted. To be with other people who actually 'get it' made the world of difference. I cannot recommend this enough...'*



# About you

Title: ..... First name: ..... Surname: .....

Address: .....

.....

Postcode: .....

Email address\*: .....

Telephone number: .....

Date of birth: .....

Haemophilia centre: .....

\* Whenever possible we try to use email to communicate with our members.

By providing your email address you're helping us to keep our costs to a minimum.

However, we understand email isn't suitable for everyone. If you would prefer to

receive information from us by post, please tick here

Please tick one of the following statements that best describes you:

I am:

- A person with a bleeding disorder (including symptomatic carrier)
- A partner / carer / parent of an adult with a bleeding disorder
- A carer / parent of a child (under 16) with a bleeding disorder
- A relative of someone who died from a bleeding disorder or its treatment
- A family member of person with a bleeding disorder
- A friend of a person with a bleeding disorder
- A healthcare professional
- A member of the general public
- A volunteer with The Haemophilia Society
- Other (please specify)

Please let us know which bleeding disorder is relevant to you:

- Haemophilia A
- Haemophilia B
- von Willebrand Disease
- Another factor deficiency
- A platelet disorder
- Don't know
- Other

### Children and young people (under 16)

If you're a parent / guardian of any children or young people under the age of 16, please provide us with their details below – including any siblings who don't have a bleeding disorder. Providing us with this information will help us to send you relevant information to your child's age-group and to let you know about children's and young people's activities and events they might like to take part in.

Name	Bleeding disorder	Newly diagnosed Y/N	Age	Name of sibling/s without a bleeding disorder	Age

Data Protection: The Haemophilia Society is registered under the Data Protection Act 1998, which protects the personal data of individuals and their privacy. We have requested the information on this form in order to function effectively and provide the best possible support.

Please tick the box below if you 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 RTSK-HLKS-SXLR, The Haemophilia Society, Willcox House, 140 –148 Borough High Street, London, SE1 1LB**

The Haemophilia Society, Willcox House, 140-148 Borough High Street, London SE1 1LB

Freephone: 0800 018 6068, Reception: 020 7939 0780

Email: [info@haemophilia.org.uk](mailto:info@haemophilia.org.uk)

 [HaemophiliaSocietyUK](https://www.facebook.com/HaemophiliaSocietyUK)

 [HaemoSocUK](https://twitter.com/HaemoSocUK)

[www.haemophilia.org.uk](http://www.haemophilia.org.uk)

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*Opportunities to share your views and experiences to help raise UK-wide awareness of the needs – and the capabilities – of people affected by a bleeding disorder.*

- Regular surveys on important issues to shape our work and influence future care and treatment.
- Vote at our AGM and elect our trustees – or even become a trustee yourself.



We work on your behalf, and with our local groups and partner organisations. We also campaign for improvements in treatment and support for our community.

To find out more and join our growing community of individuals and families, supporters and healthcare professionals, visit [www.haemophilia.org.uk](http://www.haemophilia.org.uk) or call 020 7939 0780.

You can also get in touch on Facebook (HaemophiliaSocietyUK) or Twitter (@HaemoSocUK).

*'The support gained from knowing you're not alone in the tough times, and that you will come out even stronger on the other side, was worth its weight in gold.'*

*'Through The Haemophilia Society I found someone else with the same bleeding disorder. This helped me, and I felt less isolated.'*