

Helpful References for Women with Bleeding Disorders

www.hemophilia.ca - Canadian Hemophilia Society

www.wfh.org - World Federation of Hemophilia

www.coderougewomen.ca - Canadian Hemophilia Society-CodeRouge (When Women Bleed Too Much)

<https://letstalkperiod.ca/> aims to increase awareness of the signs and symptoms of bleeding disorders.

<https://hemaware.org/womens-health/womens-bleeding-disorders> - HemAware for Women

<https://victoryforwomen.org/community-> Victory for Women - National Hemophilia Foundation website for women with bleeding disorders

www.kelleycom.com - LA Kelley Communications

www.ladybugsfoundation.org - LadyBugs Foundation

Talking Red website from the UK:

<http://haemophilia.org.uk/support/talkingred/>

<http://www.pdsa.org/> - Platelet Disorder Support Association (PDSA) - support resource for idiopathic thrombocytopenic purpura (ITP)

<http://www.fwgbd.org/> Foundation for Women and Girls with Blood Disorders

von Willebrand Disease-specific information

<https://www.kelleycom.com/additional-resources/von-willebrand-disease/>- von Willebrand Disease-specific publications from LA Kelley Communications, Inc.

www.allaboutbleeding.com - CLS Behrings VWD site

Memoir

Cheryl Nineff D'Ambrosio- "Pooling Blood: A Journey of Two Girls with Hemophilia and Their Parents' Struggle to Keep Them Alive" A memoir of a family with sisters who have severe factor V deficiency (Amazon.ca)

This page posts your websites, Facebook groups, blogsites and books.

Hemo Friends	This FB group is intended to be a safe and secure (and fun!) environment in which those connected to the bleeding disorders community can freely seek advice, encouragement, support and camaraderie. Membership is open to anyone who has a legitimate, personal interest in seeking and/or providing these things. (Darcy Zwier@facebook)	Secret
Living Legacy League	This is a place for the women of the hemophilia community, impacted by the hemophilia holocaust, to come together to support one another as women: mothers, spouses, sisters and daughters. (Angela Kay Hendrickson or Priscilla Oren @Facebook sites)	Secret
von Willebrand Disease Awareness	von Willebrand Disease: Awareness, Information, Resources, Advice, Support, etc.	Open
Wisconsin Bleeding Disorders Network	The Wisconsin Bleeding Disorders Network is a not-for-profit* agency whose goals are to improve the quality of life for those with bleeding disorders, and their families, through education, networking, advocacy and financial support. https://www.facebook.com/WBDN1/	Open
Women Who Bleed	https://www.womenwhoblead.com/	Open
The Women's Bleeding Disorder Coalition	The WBDC is a division of The Valkyrie Foundation. TVF, as a whole, focuses on unmet needs among specific female populations. TVF exists to serve female veterans and The WBDC continues to focus on women with bleeding disorders (@theWBDC)	Open
Foundation for Women & Girls with Blood Disorders	FWGBD is dedicated to increasing awareness and providing education about blood disorders among women and girls to healthcare providers http://www.fwgbd.org/	Open
Blood Sister to Sister	This group is to support and advocate for all women that have personal ties in the bleeding disorder community. We are not to give or endorse medical advice. BloodSisterToSister@groups.facebook.com	Secret group

	YOUR BLOGSITES
	<p>Laura has a brand blogsite. Be the first to comment and enjoy her writings. http://www.lmuchih.blogspot.com/</p>
	<p>Tiffany is a mother of a daughter with a bleeding disorder and she blogs to provide insight on what it's like to raise a child with a bleeding disorder. Her blog is called: The Art of Lion Taming, or fourlittlelions.com</p>
	<p>Linda, from Florida shares her blogsite called Fibrinogen Free, that tells about what it is like to have been diagnosed with congenital afibrinogenemia. Fibrinogen Free</p>
	<p>Ryanne from Calgary, Canada has created a blogsite about living with factor V deficiency. click this link Hemophilia is for Girls</p>
	<p>Andrea, from The Philippines has a blogsite about her daughter, Star http://fortheloveofstar.com/</p>
	<p>Rosamund, from Kidderminster, Worcestershire UK invites you to her blogsite. It's called FunnyBlood about life with vWD click this link: https://www.funnyblood.co.uk/ Rosamund's Mum has published a book called Funny Blood. Contact Rosamund for details.</p>
	<p>Cheryl, from the Pacific Northwest writes about raising two stepdaughters with severe factor V deficiency blog and "Pooling blood: A Journey of Two Girls with Hemophilia and Their Parents' Struggle to Keep Them Alive" is a memoir of a family with sisters who have severe factor V deficiency (Amazon.ca).</p>
	<p>Laurie, a mother of a son with hemophilia and the Founder of several nonprofits to benefit people with bleeding disorders ("Save one life"), has a blog that is read by people with bleeding disorders all over the world. L A Kelley Communications HemaBlog</p>

YOUR WEBSITES, FACEBOOK SITES, BOOKS

	<p>NHF Hemaware - Women's Stories http://www.hemaware.org/topics/women%27s-bleeding-disorders</p> <p>Hemaware Womens Page: http://www.hemaware.org/women</p>
	<p>Rosamund and Emma area Administrators on the VonWillebrand Disease and Associated Bleeding Disorders site:</p> <p>Von Willebrand Disease & Associated Bleeding Disorders https://www.facebook.com/groups/199489926762054/</p>
	<p>Milena has a facebook for moms called Hemophilia Mother. She supports hemophilia mothers, carriers and women with bleeding disorders: Join in https://www.facebook.com/groups/117299178301385</p>
	<p>Marilyn has a website for her film called BAD BLOOD. She also welcomes you to become a fan of BAD BLOOD on Facebook. Click https://www.facebook.com/Bad-Blood-Documentary-111157142250882/</p>

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