

## Sources

### Haemophilia Foundation Australia publications

<[www.haemophilia.org.au/publications](http://www.haemophilia.org.au/publications)>:

Bleeding disorders and hepatitis C: My choice to tell – family, friends, new partners. Melbourne: HFA, 2009

Bleeding disorders and hepatitis C: My choice to tell – at work. Melbourne: HFA, 2009.

A guide for people living with von Willebrand disorder. Melbourne: HFA, 2010

Living with mild haemophilia: a guide. Melbourne: HFA, 2011

The Female Factors: A snapshot of bleeding disorders in females. Melbourne: HFA, 2016

Haemophilia: a guide for parents of a newly diagnosed child. Melbourne: HFA, 2017.

### Other publications:

Australian Haemophilia Centre Directors' Organisation. Guidelines for the management of haemophilia in Australia. Melbourne; Canberra: AHCCDO; National Blood Authority, 2016. <[www.blood.gov.au/haemophilia-guidelines](http://www.blood.gov.au/haemophilia-guidelines)>

Australian Human Rights Commission. Disability discrimination. Sydney: AHRC, 2014. <[www.humanrights.gov.au](http://www.humanrights.gov.au)>

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Hepatitis Australia. Disclosure of hep C status. Canberra: Hepatitis Australia, 2013. <[www.hepatitisaustralia.com](http://www.hepatitisaustralia.com)>

World Federation of Hemophilia. Rare clotting factor deficiencies. Montreal: WFH, 2014. <[www.wfh.org](http://www.wfh.org)>

## Acknowledgements

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## Important note

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