Utility of clinical comprehensive genomic characterisation for diagnostic categorisation in patients presenting with hypocellular bone marrow failure syndromes

Piers Blombery, Lucy Fox, Georgina L. Ryland, Ella R. Thompson, Jennifer Lickiss, Michelle McBean, Satwica Yerneni, David Hughes, Anthea Greenway, Francoise Mechinaud, Erica M. Wood, Graham J. Lieschke, Jeff Szer, Pasquale Barbaro, John Roy, Joel Wight, Elly Lynch, Melissa Martyn, Clara Gaff, and David Ritchie

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