Tabitha Haw

List of Publications by Year in descending order

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ΤΛΡΙΤΗΛ ΗΛΊΑΙ

#	Article	IF	CITATIONS
1	Genotype-phenotype correlation in combined deficiency of factor V and factor VIII. Blood, 2008, 111, 5592-5600.	1.4	63
2	Identification ofP gene mutations in individuals with oculocutaneous albinism in sub-Saharan Africa. Human Mutation, 2000, 15, 166-172.	2.5	43
3	The Utilization and Outcome of Diagnostic, Predictive, and Prenatal Genetic Testing for Huntington Disease in Johannesburg, South Africa. Genetic Testing and Molecular Biomarkers, 2012, 16, 58-62.	0.7	20
4	Phenotypic consequences in black South African Fanconi anemia patients homozygous for a founder mutation. Genetics in Medicine, 2014, 16, 400-406.	2.4	11
5	Genetic Counseling for Fetal Abnormalities in a South African Community. Journal of Genetic Counseling, 2010, 19, 247-254.	1.6	9
6	Breast cancer in high-risk Afrikaner families: Is BRCA founder mutation testing sufficient?. South African Medical Journal, 2016, 106, 264.	0.6	9
7	Hematological consequences of a FANCG founder mutation in Black South African patients with Fanconi anemia. Blood Cells, Molecules, and Diseases, 2015, 54, 270-274.	1.4	8
8	Exploring how mothers of a child with a genetic disorder experience their couple relationship in a low socioâ€economic setting. Journal of Genetic Counseling, 2021, 30, 885-899.	1.6	4
9	Awareness of genetic counseling services among allied healthcare professionals in South Africa. Journal of Genetic Counseling, 2021, 30, 1649-1657.	1.6	1