

Marcus Fager Ferrari

List of Publications by Year in descending order

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6
papers

27
citations

2258059

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2272923

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#	ARTICLE	IF	CITATIONS
1	Collagen remodelling and plasma ascorbic acid levels in patients suspected of inherited bleeding disorders harbouring germline variants in collagen-related genes. <i>Haemophilia</i> , 2021, 27, e69-e77.	2.1	0
2	A rare case of IgE kappa monoclonal gammopathy of undetermined significance identified in a Swedish female. <i>Scandinavian Journal of Clinical and Laboratory Investigation</i> , 2021, 81, 385-388.	1.2	2
3	Severe Congenital Thrombocytopenia Characterized by Decreased Platelet Sialylation and Moderate Complement Activation Caused by Novel Compound Heterozygous Variants in GNE. <i>Frontiers in Immunology</i> , 2021, 12, 777402.	4.8	7
4	A rare heterozygous variant in FGB (Fibrinogen Merivale) causing hypofibrinogenemia in a Swedish family. <i>Blood Coagulation and Fibrinolysis</i> , 2020, 31, 481-484.	1.0	0
5	Genetic screening of children with suspected inherited bleeding disorders. <i>Haemophilia</i> , 2020, 26, 314-324.	2.1	6
6	Germline heterozygous variants in genes associated with familial hemophagocytic lymphohistiocytosis as a cause of increased bleeding. <i>Platelets</i> , 2018, 29, 56-64.	2.3	12