Marcus Fager Ferrari

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/1749486/publications.pdf

Version: 2024-02-01

6	27	2258059 3	2272923 4
papers	citations	h-index	g-index

6	6	6	45
all docs	docs citations	times ranked	citing authors

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