Oliver Andres

List of Publications by Year
in descending order

15 papers	278 citations	1040056 9 h-index	996975 15 g-index
16 all docs	16 docs citations	16 times ranked	491 citing authors

#	Article	IF	Citations
1	Clinical performance evaluation of SARS-CoV-2 rapid antigen testing in point of care usage in comparison to RT-qPCR. EBioMedicine, 2021, 69, 103455.	6.1	63
2	Novel variants in FERMT3 and RASGRP2â€"Genetic linkage in Glanzmannâ€like bleeding disorders. Pediatric Blood and Cancer, 2020, 67, e28078.	1.5	6
3	Genotypeâ€phenotype correlation and molecular heterogeneity in pyruvate kinase deficiency. American Journal of Hematology, 2020, 95, 472-482.	4.1	47
4	Hereditary spherocytosis is associated with decreased pyruvate kinase activity due to impaired structural integrity of the red blood cell membrane. British Journal of Haematology, 2019, 187, 386-395.	2.5	8
5	Functional Classification of Paediatric Patients with Non-syndromic Delta-Storage Pool Deficiency. Hamostaseologie, 2019, 39, 383-391.	1.9	5
6	Urine Proteomic Analysis Reveals Disease-Specific Patterns in Pediatric Patients with Classical Hodgkin's Disease(HD). an Addon Study to the Euronet-PHL-C2 Trial. Blood, 2019, 134, 2804-2804.	1.4	1
7	Diagnosis of platelet function disorders: A standardized, rational, and modular flow cytometric approach. Platelets, 2018, 29, 347-356.	2.3	24
8	Use of Targeted High-Throughput Sequencing for Genetic Classification of Patients with Bleeding Diathesis and Suspected Platelet Disorder. TH Open, 2018, 02, e445-e454.	1.4	18
9	Recessive grey platelet-like syndrome with unaffected erythropoiesis in the absence of the splice isoform GFI1B-p37. Haematologica, 2017, 102, e375-e378.	3.5	16
10	A novel twoâ€nucleotide deletion in HPS6 affects mepacrine uptake and platelet dense granule secretion in a family with Hermanskyâ€"Pudlak syndrome. Pediatric Blood and Cancer, 2017, 64, e26320.	1.5	10
11	Platelets in neonates: Central mediators in haemostasis, antimicrobial defence and inflammation. Thrombosis and Haemostasis, 2015, 113, 3-12.	3.4	24
12	Early postnatal diagnosis of hereditary spherocytosis by combining light microscopy, acidified glycerol lysis test and eosin-5â€²-maleimide binding assay. Annals of Hematology, 2015, 94, 1959-1964.	1.8	10
13	Thrombosis as a complication of central venous access in pediatric patients with malignancies: a 5-year single-center experience. BMC Hematology, 2014, 14, 18.	2.6	31
14	Autosomal dominant overhydrated stomatocytosis associated with the heterozygous Rh<scp>AG</scp> mutation F65S: a case of missed heterozygosity due to allelic dropout. British Journal of Haematology, 2013, 161, 602-604.	2.5	4
15	Even in Pneumococcal Sepsis CD62L Shedding on Granulocytes Proves to be a Reliable Functional Test for the Diagnosis of Interleukin-1 Receptorâ€"associated Kinase-4 Deficiency. Pediatric Infectious Disease Journal, 2013, 32, 1017-1019.	2.0	7