# Oliver Andres

List of Publications by Year
in descending order



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|---|---|---|---|
| 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 | Genotypeâ€phenotype correlation and molecular heterogeneity in pyruvate kinase deficiency. American Journal of Hematology, 2020, 95, 472-482. | 4.1 | 47 |
| 3 | 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 |
| 4 | Platelets in neonates: Central mediators in haemostasis, antimicrobial defence and inflammation. Thrombosis and Haemostasis, 2015, 113, 3-12. | 3.4 | 24 |
| 5 | Diagnosis of platelet function disorders: A standardized, rational, and modular flow cytometric approach. Platelets, 2018, 29, 347-356. | 2.3 | 24 |
| 6 | 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 |
| 7 | 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 |
| 8 | 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 |
| 9 | 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 |
| 10 | 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 |
| 11 | 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 |
| 12 | Novel variants in FERMT3 and RASGRP2â€”Genetic linkage in Glanzmannâ€like bleeding disorders. Pediatric Blood and Cancer, 2020, 67, e28078. | 1.5 | 6 |
| 13 | Functional Classification of Paediatric Patients with Non-syndromic Delta-Storage Pool Deficiency. Hamostaseologie, 2019, 39, 383-391. | 1.9 | 5 |
| 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 | 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 |