Manisha Gadgeel

List of Publications by Year
in descending order

21 papers	102 citations	1684188 5 h-index	1474206 9 g-index
21 all docs	21 docs citations	21 times ranked	248 citing authors

#	Article	IF	Citations
1	Lymphocyte HLA-DR/CD-38 co-expression correlates with Hodgkin lymphoma cell cytotoxicity inÂvitro independent of PD-1/PD1-L pathway. Leukemia and Lymphoma, 2022, , 1-8.	1.3	0
2	Persistent pseudo-Pelger-HuÃ«t anomaly. Annals of Hematology, 2021, 100, 2661-2663.	1.8	0
3	Characteristic flow cytometric profile of ectopic intraâ€thyroidal thymic tissue in children. Cytometry Part B - Clinical Cytometry, 2021, 100, 457-459.	1.5	2
4	Distinctive phenotypes in two children with novel germline RUNX1 mutations - one with myeloid malignancy and increased fetal hemoglobin. Pediatric Hematology and Oncology, 2021, 38, 65-79.	0.8	9
5	Aberrant myelomonocytic CD56 expression in Down syndrome is frequent and not associated with leukemogenesis. Annals of Hematology, 2021, 100, 1695-1700.	1.8	5
6	CD20+ T Cells in Primary Mediastinal Large B Cell Lymphoma Microenvironment. Cytometry Part B - Clinical Cytometry, 2020, 98, 16-18.	1.5	3
7	CD14/16 monocyte profiling in juvenile myelomonocytic leukemia. Pediatric Blood and Cancer, 2020, 67, e28555.	1.5	3
8	Differing reflections of paediatric classical Hodgkinâ€™s lymphoma on local and distant immunological microenvironments: a flow cytometric study. Journal of Clinical Pathology, 2020, 73, 176-179.	2.0	1
9	Clonal Tâ€cell large granular lymphocyte proliferations in childhood and young adult immune dysregulation conditions. Pediatric Blood and Cancer, 2020, 67, e28231.	1.5	4
10	Severe macrothrombocytopenia with platelet CD9 deficiency responsive to romiplostim. British Journal of Haematology, 2020, 190, e239-e242.	2.5	1
11	Mild erythrocytosis as a presenting manifestation ofPIEZO1associated erythrocyte volume disorders. Pediatric Hematology and Oncology, 2019, 36, 317-326.	0.8	13
12	Compound heterozygosity in PKLR gene for a previously unrecognized intronic polymorphism and a rare missense mutation as a novel cause of severe pyruvate kinase deficiency. Haematologica, 2019, 104, e428-e431.	3.5	8
13	Different Clonal T-Large Granular Lymphocyte Proliferations in SCID. Journal of Clinical Immunology, 2019, 39, 245-248.	3.8	3
14	Oryzocytosis: A Novel Morphological Variant of Hereditary Elliptocytosis Associated with a Novel Mutation in Î²-Spectrin (SPTB c154 C>T p.Arg52Trp). Blood, 2019, 134, 3509-3509.	1.4	0
15	KLF1 E325K-associated Congenital Dyserythropoietic Anemia Type IV: Insights Into the Variable Clinical Severity. Journal of Pediatric Hematology/Oncology, 2018, 40, e405-e409.	0.6	33
16	Flow cytometric false myeloperoxidaseâ€positive childhood Bâ€lineage acute lymphoblastic leukemia. Cytometry Part B - Clinical Cytometry, 2018, 94, 477-483.	1.5	11
17	RUNX1 associated Familial Platelet Disorder with Myeloid Malignancy (FPD-MM) in Children: A Novel New Phenotype with Juvenile and Chronic Myelomonocytic Leukemia (JMML/CMML) Characteristics. Blood, 2018, 132, 5504-5504.	1.4	1
18	Lymphocyte Co-Expression of CD38/HLA-DR Is a Marker for Anti-Tumor Activity in Hodgkin Lymphoma in Vitro: Evidence for a PD1-Independent Mechanism. Blood, 2018, 132, 4129-4129.	1.4	0

#	Article	IF	Citations
19	Clonal T-Large Granular Lymphocyte Proliferations in Childhood: Friend or Foe?. Blood, 2018, 132, 3719-3719.	1.4	0
20	Methimazole Induced Total Myeloid Aplasia with Delayed Recovery Despite Granulocyte Colony Stimulating Factor (G-CSF): Marrow Progenitor Recovery Kinetics. Indian Journal of Hematology and Blood Transfusion, 2016, 32, 214-218.	0.6	4
21	Red Cell Band 3 Content Evaluation By Eosin Maleimide (EMA) Fluorescence: Beyond Diagnosis of Dominant Hereditary Spherocytosis (HS). Blood, 2015, 126, 3343-3343.	1.4	1