Deepti M Warad Mbbs

List of Publications by Year
in descending order

38	313	933447	888059
papers	citations	10 h-index	17 g-index

39	39	39	599
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Lemierre Syndrome: A Retrospective Study of the Role of Anticoagulation and Thrombosis Outcomes. Acta Haematologica, 2017, 137, 59-65.	1.4	42
2	Intratumoral CD14+ Cells and Circulating CD14+HLA-DRlo/neg Monocytes Correlate with Decreased Survival in Patients with Clear Cell Renal Cell Carcinoma. Clinical Cancer Research, 2015, 21, 4224-4233.	7.0	33
3	Obesity, sedentary lifestyle, and video games: The new thrombophilia cocktail in adolescents. Pediatric Blood and Cancer, 2018, 65, e27041.	1.5	33
4	Diagnostic laboratory standardization and validation of platelet transmission electron microscopy. Platelets, 2018, 29, 574-582.	2.3	31
5	A prospective, blinded study of a PF4-dependent assay for HIT diagnosis. Blood, 2021, 137, 1082-1089.	1.4	28
6	Comprehensive Platelet Phenotypic Laboratory Testing and Bleeding History Scoring for Diagnosis of Suspected Hereditary Platelet Disorders. American Journal of Clinical Pathology, 2017, 148, 23-32.	0.7	16
7	A Retrospective Analysis of Outcomes of Dalteparin Use in Pediatric Patients: A Single Institution Experience. Thrombosis Research, 2015, 136, 229-233.	1.7	14
8	A retrospective review of pediatric antiphospholipid syndrome and thrombosis outcomes. Blood Coagulation and Fibrinolysis, 2017, 28, 205-210.	1.0	14
9	Pediatric Coagulation Disorders. Pediatrics in Review, 2016, 37, 279-291.	0.4	13
10	Cross-Cultural Care Training for Pediatric Hematology/Oncology Fellows. MedEdPORTAL: the Journal of Teaching and Learning Resources, 2017, 13, 10543.	1.2	13
11	Pediatric Mayâ€Thurner Syndromeâ€”Systematic review and individual patient data metaâ€analysis. Journal of Thrombosis and Haemostasis, 2021, 19, 1283-1293.	3.8	9
12	Cross-Cultural Medical Care Training and Education: a National Survey of Pediatric Hematology/Oncology Fellows-in-Training and Fellowship Program Directors. Journal of Cancer Education, 2019, 34, 478-487.	1.3	8
13	Lemierre's Syndrome: Role of Anticoagulation and Thrombosis Outcomes, a Retrospective Study. Blood, 2015, 126, 2296-2296.	1.4	8
14	EBV-PTLD, Adenovirus, and CMV in Pediatric Allogeneic Transplants With Alemtuzumab as Part of Pretransplant Conditioning: A Retrospective Single Center Study. Journal of Pediatric Hematology/Oncology, 2018, 40, e473-e478.	0.6	7
15	Hermansky-Pudlak syndrome subtype 5 (HPS-5) novel mutation in a 65 year-old with oculocutaneous hypopigmentation and mild bleeding diathesis: The importance of recognizing a subtle phenotype. Platelets, 2018, 29, 91-94.	2.3	7
16	A mutation in SLC37A4 causes a dominantly inherited congenital disorder of glycosylation characterized by liver dysfunction. American Journal of Human Genetics, 2021, 108, 1040-1052.	6.2	7
17	Management of pediatric hepatocellular carcinoma: A multimodal approach. Pediatric Transplantation, 2017, 21, e13007.	1.0	6
18	Clinical Outcomes of Mayâ€“Thurner Syndrome in Pediatric Patients: A Single Institutional Experience. TH Open, 2020, 04, e189-e196.	1.4	6

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19	A Severe Case of Congenital Thrombotic Thrombocytopenia Purpura Resulting From Compound Heterozygosity Involving a Novel ADAMTS13 Pathogenic Variant. Journal of Pediatric Hematology/Oncology, 2018, 40, 60-62.	0.6	4
20	Acute appendicitis in acute leukemia and the potential role of decitabine in the critically ill patient. Leukemia Research Reports, 2015, 4, 21-23.	0.4	3
21	An Age Distribution of D-Dimer Values in Normal Healthy Donor Population: An Indirect Verification of the Age-Adjusted D-Dimer Cutoffs for VTE Exclusion. Blood, 2016, 128, 1432-1432.	1.4	3
22	Injury to Insult: Infarction After Radiotherapy inÂtheÂTreatmentÂof Pediatric Brain Tumor. Pediatric Neurology, 2015, 52, 552-553.	2.1	1
23	Crossâ€cultural communication in pediatric oncology: Catch them when they're young. Cancer, 2018, 124, 1516-1517.	4.1	1
24	Case Report: Development of Factor VIII Inhibitor in a Patient with an Uncommon de novo Mutation in the Factor VIII Gene. Acta Haematologica, 2019, 141, 129-134.	1.4	1
25	Value of Platelet Esoteric Testing in Laboratory Diagnosis of Platelet Disorders: A Single Center Experience. Blood, 2015, 126, 1061-1061.	1.4	1
26	A Premature Girl with Pallor and Rash. Pediatric Annals, 2011, 40, 296-298.	0.8	1
27	Pediatric Anti-Phospholipid Syndrome: Thromboses Outcomes, Mayo Clinic Children's Center Experience. Blood, 2014, 124, 4258-4258.	1.4	1
28	The Platelet Factor-4 (PF4)-Dependent p-Selectin Expression Assay (PEA) Is Highly Accurate for the Detection of Pathogenic HIT Antibodies: Results of a 440-Sample Prospective Blinded Study. Blood, 2018, 132, 416-416.	1.4	1
29	Pediatric Myxopapillary Ependymomas: A Clinicopathologic Evaluation. Journal of Pediatric Hematology/Oncology, 2021, 43, e1194-e1200.	0.6	1
30	Immune Profiling to Predict Treatment Response from Extracorporeal Photopheresis in Graft-Versus-Host Disease. Biology of Blood and Marrow Transplantation, 2014, 20, S263.	2.0	0
31	Cerebral sinovenous thrombosis in pediatric hemolytic uremic syndrome. Research and Practice in Thrombosis and Haemostasis, 2020, 4, 659-665.	2.3	0
32	Risk of perinatal intracranial hemorrhage and role of prenatal genetic testing in individuals with type 3 von Willebrand disease. Journal of Thrombosis and Haemostasis, 2020, 18, 2779-2780.	3.8	0
33	Computerâ€based simulation to reduce EHRâ€related chemotherapy ordering errors. Cancer Medicine, 2020, 9, 8844-8851.	2.8	0
34	Healthcare Utilization and Costs associated with Hereditary Hemorrhagic Telangiectasia Patients in a Large US Claims Database. Mayo Clinic Proceedings Innovations, Quality & Outcomes, 2021, 5, 55-64.	2.4	0
35	Acute thrombosis of a giant perimedullary arteriovenous fistula in a pediatric HHT patient. Interventional Neuroradiology, 2021, , 159101992110224.	1.1	0
36	Hemorrhagic Complications With Adenotonsillectomy In Children and Young Adults With Bleeding Disorders. Blood, 2013, 122, 3611-3611.	1.4	0

#	Article	IF	Citations
37	Utility of Various Von Willebrant Factor Laboratory Tests in Assessment of Acquired Von Willebrant Syndrome in Patients with Aortic Stenosis. Blood, 2014, 124, 1517-1517.	1.4	0
38	Platelet Transmission Electron Microscopy and Flow Cytometry in the Diagnosis of Congenital/Hereditary Qualitative or Quantitative Platelet Disorders. Blood, 2015, 126, 3476-3476.	1.4	0