Hoda M Hassab

List of Publications by Year
in descending order

24 papers	722 citations	1163117 8 h-index	752698 20 g-index
24 all docs	24 docs citations	24 times ranked	873 citing authors

#	Article	IF	Citations
1	Beta Thalassemia Carrier rate: Problem Burden Among High School Children. Current Pediatric Reviews, 2022, 18, .	0.8	0
2	Voxelotor in adolescents and adults with sickle cell disease (HOPE): long-term follow-up results of an international, randomised, double-blind, placebo-controlled, phase 3 trial. Lancet Haematology,the, 2021, 8, e323-e333.	4.6	61
3	SLC25A38 congenital sideroblastic anemia: Phenotypes and genotypes of 31 individuals from 24 families, including 11 novel mutations, and a review of the literature. Human Mutation, 2021, 42, 1367-1383.	2.5	11
4	Comparing the effect of acupressure and ginger on chemotherapy gastrointestinal side-effects in children with leukemia. Complementary Therapies in Medicine, 2021, 60, 102730.	2.7	8
5	Prevalence of iron deficiency anemia and beta thalassemia carriers among relatives of beta thalassemia patients in Nile Delta region, Egypt: a multicenter study. Journal of the Egyptian Public Health Association, The, 2021, 96, 27.	2.5	5
6	Lowâ€dose immune tolerance induction therapy in children of Arab descent with severe haemophilia A, high inhibitor titres and poor prognostic factors for immune tolerance induction treatment success. Haemophilia, 2021, 28, 65.	2.1	3
7	Evaluation of the efficacy and safety of deferiprone compared with deferasirox in paediatric patients with transfusion-dependent haemoglobinopathies (DEEP-2): a multicentre, randomised, open-label, non-inferiority, phase 3 trial. Lancet Haematology,the, 2020, 7, e469-e478.	4.6	39
8	Efficacy and Safety of Voxelotor in Adolescents and Adults with Sickle Cell Disease: HOPE Trial 72-Week Analysis. Blood, 2020, 136, 19-19.	1.4	3
9	A Phase 3 Randomized Trial of Voxelotor in Sickle Cell Disease. New England Journal of Medicine, 2019, 381, 509-519.	27.0	401
10	Geographic Differences in Phenotype and Treatment of Children with Sickle Cell Anemia from the Multinational DOVE Study. Journal of Clinical Medicine, 2019, 8, 2009.	2.4	8
11	Evaluation of multiplexed biomarkers in assessment of CSF infiltration in pediatric acute lymphoblastic leukemia. International Journal of Hematologic Oncology, 2019, 8, IJH22.	1.6	3
12	Novel findings from the multinational <scp>DOVE</scp> study on geographic and ageâ€related differences in pain perception and analgesic usage in children with sickle cell anaemia. British Journal of Haematology, 2019, 184, 1058-1061.	2.5	13
13	Intracranial Hemorrhage in Egyptian Children with Primary Immune Thrombocytopenia (ITP): Report of 24 Cases in 20 Years. Blood, 2019, 134, 1080-1080.	1.4	1
14	Neutropenia in Children Treated with Deferiprone or Deferasirox: A Report of the Largest Randomized Trial of Oral Chelators in Transfusion-Dependent Pediatric Patients. Blood, 2019, 134, 3552-3552.	1.4	2
15	Deviation from Guidelines in Use of Eltrombopag and Romiplostim in Clinical Practice for Children with Primary Immune Thrombocytopenia; Multi-Center Egyptian Experience. Blood, 2019, 134, 4907-4907.	1.4	0
16	Results from Part A of the Hemoglobin Oxygen Affinity Modulation to Inhibit HbS Polymerization (HOPE) Trial (GBT440-031), a Placebo-Controlled Randomized Study Evaluating Voxelotor (GBT440) in Adults and Adolescents with Sickle Cell Disease. Blood, 2018, 132, 505-505.	1.4	3
17	A Multinational Trial of Prasugrel for Sickle Cell Vaso-Occlusive Events. New England Journal of Medicine, 2016, 374, 625-635.	27.0	117
18	Neurophysiologic Evaluation of Children with Beta-Thalassemia Major. Journal of Pediatric Neurology, 2015, 13, 110-115.	0.2	0

#	ARTICLE	IF	CITATIONS
19	Lymphadenopathy in a Series of Egyptian Pediatric Patients and the Role of Pathology in the Diagnostic Workup. Pediatric and Developmental Pathology, 2014, 17, 344-359.	1.0	11
20	Lymphadenopathy in a Series of Egyptian Pediatric Patients and the Role of Pathology in the Diagnostic Workup.. Pediatric and Developmental Pathology, 2014, , .	1.0	2
21	Intracranial Hemorrhage in Acute and Chronic Childhood Immune Thrombocytopenic Purpura over a Ten-Year Period: An Egyptian Multicenter Study. Acta Haematologica, 2010, 123, 59-63.	1.4	22
22	Phenotypic analysis of bone marrow lymphocytes from children with acute thrombocytopenic purpura. The Egyptian Journal of Immunology / Egyptian Association of Immunologists, 2005, 12, 9-14.	0.4	5
23	Granulocyte-macrophage colony-stimulating factor (GM-CSF) in children with acute immune thrombocytopenic purpura. Medical Science Monitor, 2004, 10, CR330-5.	1.1	3
24	Does TGFBR3 Polymorphism Increase the Risk of Silent Cerebral Infarction in Egyptian Children with Sickle Cell Disease?. Indian Journal of Pediatrics, 0, , .	0.8	1