Azza Abdel Gawad Tantawy

List of Publications by Year
in descending order

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47 papers	587 citations	623188 14 h-index	676716 22 g-index
48 all docs	48 docs citations	48 times ranked	987 citing authors

#	Article	IF	Citations
1	Circulating platelet and erythrocyte microparticles in young children and adolescents with sickle cell disease: Relation to cardiovascular complications. Platelets, 2013, 24, 605-614.	1.1	71
2	Methylene tetrahydrofolate reductase gene polymorphism in Egyptian children with acute lymphoblastic leukemia. Blood Coagulation and Fibrinolysis, 2010, 21, 28-34.	0.5	48
3	Flow cytometric assessment of circulating platelet and erythrocytes microparticles in young thalassemia major patients: relation to pulmonary hypertension and aortic wall stiffness. European Journal of Haematology, 2013, 90, 508-518.	1.1	36
4	Romiplostim therapy in children with unresponsive chronic immune thrombocytopenia. Platelets, 2012, 23, 264-273.	1.1	34
5	Subclinical Atherosclerosis In Young Î²-thalassemia Major Patients. Hemoglobin, 2009, 33, 463-474.	0.4	32
6	Clinicopathological and Radiological Study of Egyptian Î²-Thalassemia Intermedia and Î²-Thalassemia Major Patients: Relation to Complications and Response to Therapy. Hemoglobin, 2011, 35, 382-405.	0.4	20
7	Outcome of enzyme replacement therapy in children with Gaucher disease: The Egyptian experience. Egyptian Journal of Medical Human Genetics, 2011, 12, 9-14.	0.5	19
8	Pulmonary Complications in Survivors of Childhood Hematological Malignancies: Single-Center Experience. Pediatric Hematology and Oncology, 2011, 28, 403-417.	0.3	19
9	Spinal Cord Compression in Childhood Pediatric Malignancies. Journal of Pediatric Hematology/Oncology, 2013, 35, 232-236.	0.3	19
10	A longitudinal prospective study of bleeding diathesis in Egyptian pediatric patients. Blood Coagulation and Fibrinolysis, 2012, 23, 411-418.	0.5	17
11	Health-Related Quality of Life in Egyptian Children and Adolescents with Hemophilia A. Pediatric Hematology and Oncology, 2011, 28, 222-229.	0.3	16
12	Soluble CD163 in young sickle cell disease patients and their trait siblings. Blood Coagulation and Fibrinolysis, 2012, 23, 640-648.	0.5	16
13	Retinal changes in children and adolescents with sickle cell disease attending a paediatric hospital in Cairo, Egypt: risk factors and relation to ophthalmic and cerebral blood flow. Transactions of the Royal Society of Tropical Medicine and Hygiene, 2013, 107, 205-211.	0.7	16
14	Endothelial nitric oxide synthase gene intron 4 VNTR polymorphism in sickle cell disease: Relation to vasculopathy and disease severity. Pediatric Blood and Cancer, 2015, 62, 389-394.	0.8	16
15	Evoked potentails and neurocognitive functions in pediatric Egyptian Gaucher patients on enzyme replacement therapy: a single center experience. Journal of Inherited Metabolic Disease, 2013, 36, 1025-1037.	1.7	14
16	D-dimer assay in Egyptian patients with Gaucher disease: correlation with bone and lung involvement. Blood Coagulation and Fibrinolysis, 2011, 22, 176-184.	0.5	13
17	Outcome of childhood acute Lymphoblastic leukemia in Egyptian children: A challenge for limited health resource countries. Hematology, 2013, 18, 204-210.	0.7	13
18	Growth differentiation factor-15 in children and adolescents with thalassemia intermedia: Relation to subclinical atherosclerosis and pulmonary vasculopathy. Blood Cells, Molecules, and Diseases, 2015, 55, 144-150.	0.6	13

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19	Clinical Predictive Value of Cystatin C in Pediatric Sickle Cell Disease: A Marker of Disease Severity and Subclinical Cardiovascular Dysfunction. Clinical and Applied Thrombosis/Hemostasis, 2017, 23, 1010-1017.	0.7	13
20	Radionuclide Ventriculography Detects Early Anthracycline Cardiotoxity in Children With Hodgkin Lymphoma. Journal of Pediatric Hematology/Oncology, 2011, 33, e132-e137.	0.3	11
21	Cytokines in Gaucher disease: Role in the pathogenesis of bone and pulmonary disease. Egyptian Journal of Medical Human Genetics, 2015, 16, 207-213.	0.5	11
22	Growth differentiation factor-15 in young sickle cell disease patients: Relation to hemolysis, iron overload and vascular complications. Blood Cells, Molecules, and Diseases, 2014, 53, 189-193.	0.6	10
23	Evaluation of continuous glucose monitoring system for detection of alterations in glucose homeostasis in pediatric patients with Î²-thalassemia major. Pediatric Diabetes, 2019, 20, 65-72.	1.2	10
24	Development of anti-velaglucerase alfa antibodies in clinical trial-treated patients with Gaucher disease. Blood Cells, Molecules, and Diseases, 2016, 59, 37-43.	0.6	9
25	Predictors of bone disease in Egyptian prepubertal children with Î²-thalassaemia major. Archives of Medical Science, 2010, 4, 584-591.	0.4	8
26	Pathogenesis and Prognosis of Neutropenia in Infants and Children Admitted in a University Children Hospital in Egypt. Pediatric Hematology and Oncology, 2013, 30, 51-59.	0.3	8
27	The scope of clinical morbidity in sickle cell trait. Egyptian Journal of Medical Human Genetics, 2014, 15, 319-326.	0.5	8
28	Tartrate-Resistant Acid Phosphatase 5b in Young Patients With Sickle Cell Disease and Trait Siblings. Clinical and Applied Thrombosis/Hemostasis, 2017, 23, 64-71.	0.7	7
29	Pulmonary manifestations in young Gaucher disease patients: Phenotypeâ€genotype correlation and radiological findings. Pediatric Pulmonology, 2020, 55, 441-448.	1.0	7
30	Cognitive decline and depressive symptoms: early non-motor presentations of parkinsonism among Egyptian Gaucher patients. Neurogenetics, 2020, 21, 159-167.	0.7	7
31	Endothelial nitric oxide synthase gene intron 4 variable number tandem repeat polymorphism in Î²-thalassemia major. Blood Coagulation and Fibrinolysis, 2015, 26, 419-425.	0.5	6
32	One-Year Prospective Study of Community Acquired Influenza and Parainfluenza Viral Infections in Hospitalized Egyptian Children with Malignancy: Single Center Experience. Pediatric Hematology and Oncology, 2015, 32, 304-314.	0.3	6
33	Spinal Cord Compression And Extramedullary Hematopoiesis in Young Egyptian Î²-thalassemia Patients. Hemoglobin, 2009, 33, 448-462.	0.4	5
34	Haemophilia A patients are not at increased risk of hepatitis A virus infection: An Egyptian experience. Egyptian Journal of Medical Human Genetics, 2012, 13, 93-97.	0.5	3
35	Soluble fms-Like Tyrosine Kinase 1 as a Link Between Angiogenesis and Endothelial Dysfunction in Pediatric Patients With Î²-Thalassemia Intermedia. Clinical and Applied Thrombosis/Hemostasis, 2017, 23, 943-950.	0.7	3
36	Surfactant protein D as a marker for pulmonary complications in pediatric patients with sickle cell disease: Relation to lung function tests. Pediatric Pulmonology, 2019, 54, 610-619.	1.0	3

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37	Psychiatric manifestations in Egyptian Gaucher patients on enzyme replacement therapy. Journal of Psychosomatic Research, 2019, 122, 75-81.	1.2	3
38	Silent bleeding in children and adolescents with immune thrombocytopenia: relation to laboratory parameters and health related quality of life. Journal of Thrombosis and Thrombolysis, 2020, 50, 258-266.	1.0	3
39	Prevalence of XmnI Gî³ polymorphism in Egyptian patients with Î²-thalassemia major. Annals of Saudi Medicine, 2012, 32, 487-491.	0.5	3
40	Inner Ear Complications in Children and Adolescents with Sickle Cell Disease. Hemoglobin, 2020, 44, 411-417.	0.4	2
41	Ganglion Cell Complex Thinning in Young Gaucher Patients: Relation to Prodromal Parkinsonian Markers. Movement Disorders, 2020, 35, 2211-2219.	2.2	2
42	Somatosensory Evoked Potential for Detection of Subclinical Neuropathy in Egyptian Children with Acute Lymphoblastic Leukaemia. Pakistan Journal of Biological Sciences, 2010, 13, 527-536.	0.2	2
43	Endothelial specific isoform of type XVIII collagen (COLâ€18N): A marker of vascular integrity in haemophilic arthropathy. Haemophilia, 2022, 28, 849-856.	1.0	2
44	Serum progranulin levels in paediatric patients with Gaucher disease; relation to disease severity and liver stiffness by transient elastography. Liver International, 2020, 40, 3051-3060.	1.9	1
45	Abdominal lymphadenopathy in children with Gaucher disease: Relation to disease severity and glucosylsphingosine. Pediatric Hematology and Oncology, 2021, , 1-14.	0.3	1
46	Influence of thiopurine methyltransferase gene polymorphism on Egyptian children with acute lymphoblastic leukaemia. Journal of Genetics, 2017, 96, 905-910.	0.4	0
47	Joint health state in relation to different clinical, laboratory, and radiological variables of hemophilia A patients. QJM - Monthly Journal of the Association of Physicians, 2021, 114, .	0.2	0