

Azza Abdel Gawad Tantawy

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8059574/publications.pdf>

Version: 2024-02-01

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 | Endothelial specific isoform of type XVIII collagen (COL α 18N): A marker of vascular integrity in haemophilic arthropathy. <i>Haemophilia</i> , 2022, 28, 849-856. | 1.0 | 2 |
| 2 | Joint health state in relation to different clinical, laboratory, and radiological variables of hemophilia A patients. <i>QJM - Monthly Journal of the Association of Physicians</i> , 2021, 114, . | 0.2 | 0 |
| 3 | Abdominal lymphadenopathy in children with Gaucher disease: Relation to disease severity and glucosylsphingosine. <i>Pediatric Hematology and Oncology</i> , 2021, , 1-14. | 0.3 | 1 |
| 4 | Pulmonary manifestations in young Gaucher disease patients: Phenotype \leftrightarrow genotype correlation and radiological findings. <i>Pediatric Pulmonology</i> , 2020, 55, 441-448. | 1.0 | 7 |
| 5 | Serum progranulin levels in paediatric patients with Gaucher disease; relation to disease severity and liver stiffness by transient elastography. <i>Liver International</i> , 2020, 40, 3051-3060. | 1.9 | 1 |
| 6 | Inner Ear Complications in Children and Adolescents with Sickle Cell Disease. <i>Hemoglobin</i> , 2020, 44, 411-417. | 0.4 | 2 |
| 7 | Ganglion Cell Complex Thinning in Young Gaucher Patients: Relation to Prodromal Parkinsonian Markers. <i>Movement Disorders</i> , 2020, 35, 2211-2219. | 2.2 | 2 |
| 8 | Cognitive decline and depressive symptoms: early non-motor presentations of parkinsonism among Egyptian Gaucher patients. <i>Neurogenetics</i> , 2020, 21, 159-167. | 0.7 | 7 |
| 9 | Silent bleeding in children and adolescents with immune thrombocytopenia: relation to laboratory parameters and health related quality of life. <i>Journal of Thrombosis and Thrombolysis</i> , 2020, 50, 258-266. | 1.0 | 3 |
| 10 | Surfactant protein D as a marker for pulmonary complications in pediatric patients with sickle cell disease: Relation to lung function tests. <i>Pediatric Pulmonology</i> , 2019, 54, 610-619. | 1.0 | 3 |
| 11 | Psychiatric manifestations in Egyptian Gaucher patients on enzyme replacement therapy. <i>Journal of Psychosomatic Research</i> , 2019, 122, 75-81. | 1.2 | 3 |
| 12 | Evaluation of continuous glucose monitoring system for detection of alterations in glucose homeostasis in pediatric patients with β -thalassemia major. <i>Pediatric Diabetes</i> , 2019, 20, 65-72. | 1.2 | 10 |
| 13 | Tartrate-Resistant Acid Phosphatase 5b in Young Patients With Sickle Cell Disease and Trait Siblings. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2017, 23, 64-71. | 0.7 | 7 |
| 14 | Soluble fms-Like Tyrosine Kinase 1 as a Link Between Angiogenesis and Endothelial Dysfunction in Pediatric Patients With β -Thalassemia Intermedia. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2017, 23, 943-950. | 0.7 | 3 |
| 15 | Influence of thiopurine methyltransferase gene polymorphism on Egyptian children with acute lymphoblastic leukaemia. <i>Journal of Genetics</i> , 2017, 96, 905-910. | 0.4 | 0 |
| 16 | Clinical Predictive Value of Cystatin C in Pediatric Sickle Cell Disease: A Marker of Disease Severity and Subclinical Cardiovascular Dysfunction. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2017, 23, 1010-1017. | 0.7 | 13 |
| 17 | Development of anti-velaglycerase alfa antibodies in clinical trial-treated patients with Gaucher disease. <i>Blood Cells, Molecules, and Diseases</i> , 2016, 59, 37-43. | 0.6 | 9 |
| 18 | Endothelial nitric oxide synthase gene intron 4 variable number tandem repeat polymorphism in β -thalassemia major. <i>Blood Coagulation and Fibrinolysis</i> , 2015, 26, 419-425. | 0.5 | 6 |

| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 19 | 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 |
| 20 | 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 |
| 21 | 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 |
| 22 | 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 |
| 23 | The scope of clinical morbidity in sickle cell trait. Egyptian Journal of Medical Human Genetics, 2014, 15, 319-326. | 0.5 | 8 |
| 24 | 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 |
| 25 | 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 |
| 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 | 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 |
| 28 | Evoked potentials 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 |
| 29 | Outcome of childhood acute Lymphoblastic leukemia in Egyptian children: A challenge for limited health resource countries. Hematology, 2013, 18, 204-210. | 0.7 | 13 |
| 30 | 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 |
| 31 | Spinal Cord Compression in Childhood Pediatric Malignancies. Journal of Pediatric Hematology/Oncology, 2013, 35, 232-236. | 0.3 | 19 |
| 32 | Soluble CD163 in young sickle cell disease patients and their trait siblings. Blood Coagulation and Fibrinolysis, 2012, 23, 640-648. | 0.5 | 16 |
| 33 | A longitudinal prospective study of bleeding diathesis in Egyptian pediatric patients. Blood Coagulation and Fibrinolysis, 2012, 23, 411-418. | 0.5 | 17 |
| 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 | Romiplostim therapy in children with unresponsive chronic immune thrombocytopenia. Platelets, 2012, 23, 264-273. | 1.1 | 34 |
| 36 | Prevalence of XmnI G ¹³ polymorphism in Egyptian patients with β^2 -thalassemia major. Annals of Saudi Medicine, 2012, 32, 487-491. | 0.5 | 3 |

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 37 | Clinicopathological and Radiological Study of Egyptian $\hat{\alpha}^2$ -Thalassemia Intermedia and $\hat{\alpha}^2$ -Thalassemia Major Patients: Relation to Complications and Response to Therapy. Hemoglobin, 2011, 35, 382-405. | 0.4 | 20 |
| 38 | Health-Related Quality of Life in Egyptian Children and Adolescents with Hemophilia A. Pediatric Hematology and Oncology, 2011, 28, 222-229. | 0.3 | 16 |
| 39 | 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 |
| 40 | 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 |
| 41 | Radionuclide Ventriculography Detects Early Anthracycline Cardiotoxicity in Children With Hodgkin Lymphoma. Journal of Pediatric Hematology/Oncology, 2011, 33, e132-e137. | 0.3 | 11 |
| 42 | Pulmonary Complications in Survivors of Childhood Hematological Malignancies: Single-Center Experience. Pediatric Hematology and Oncology, 2011, 28, 403-417. | 0.3 | 19 |
| 43 | Methylene tetrahydrofolate reductase gene polymorphism in Egyptian children with acute lymphoblastic leukemia. Blood Coagulation and Fibrinolysis, 2010, 21, 28-34. | 0.5 | 48 |
| 44 | Predictors of bone disease in Egyptian prepubertal children with $\hat{\alpha}^2$ -thalassaemia major. Archives of Medical Science, 2010, 4, 584-591. | 0.4 | 8 |
| 45 | 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 |
| 46 | Spinal Cord Compression And Extramedullary Hematopoiesis in Young Egyptian $\hat{\alpha}^2$ -thalassemia Patients. Hemoglobin, 2009, 33, 448-462. | 0.4 | 5 |
| 47 | Subclinical Atherosclerosis In Young $\hat{\alpha}^2$ -thalassemia Major Patients. Hemoglobin, 2009, 33, 463-474. | 0.4 | 32 |