## **Eman Elghoroury**

## List of Publications by Year in Descending Order

Source: https://exaly.com/author-pdf/4241758/eman-elghoroury-publications-by-year.pdf

Version: 2024-04-20

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

34	320	11	17
papers	citations	h-index	g-index
39	370 ext. citations	1.8	2.82
ext. papers		avg, IF	L-index

#	Paper	IF	Citations
34	Low Dose Iron Therapy in Children with Iron Deficiency: DNA Damage and Oxidant Stress Markers. <i>Indian Journal of Hematology and Blood Transfusion</i> , <b>2021</b> , 37, 287-294	0.7	
33	Clinical and molecular correlation of hepcidin RNA expression in sickle cell patients with iron overload. <i>Journal of the Arab Society for Medical Research</i> , <b>2020</b> , 15, 1	0.4	1
32	The Relation between Serum Hepcidin, Ferritin, Hepcidin: Ferritin Ratio, Hydroxyurea and Splenectomy in Children with Erhalassemia. <i>Open Access Macedonian Journal of Medical Sciences</i> , <b>2019</b> , 7, 2434-2439	1	5
31	Evaluation of serum beta-endorphin and substance P in knee osteoarthritis patients treated by laser acupuncture. <i>Journal of Complementary and Integrative Medicine</i> , <b>2018</b> , 15,	1.5	19
30	Klotho G-395A gene polymorphism: impact on progression of end-stage renal disease and development of cardiovascular complications in children on dialysis. <i>Pediatric Nephrology</i> , <b>2018</b> , 33, 10	19 <del>2</del> 702	7 <sup>10</sup>
29	Association of variable number tandem repeats polymorphism in the IL-4 gene with end-stage renal disease in children. <i>Egyptian Journal of Medical Human Genetics</i> , <b>2018</b> , 19, 191-195	2	2
28	Evaluation of miRNA-21 and miRNA Let-7 as Prognostic Markers in Patients With Breast Cancer. <i>Clinical Breast Cancer</i> , <b>2018</b> , 18, e721-e726	3	38
27	Influence of thiopurine methyltransferase gene polymorphism on Egyptian children with acute lymphoblastic leukaemia. <i>Journal of Genetics</i> , <b>2017</b> , 96, 905-910	1.2	
26	FcRIIa defunctioning polymorphism in paediatric patients with renal allograft. <i>Central-European Journal of Immunology</i> , <b>2017</b> , 42, 363-369	1.6	2
25	Von Willebrand disease among Egyptian women suffering from menorrhagia. <i>Evidence Based Women S Health Journal</i> , <b>2016</b> , 6, 131-133		
24	Glutathione S-transferase gene polymorphism: Relation to cardiac iron overload in Egyptian patients with Beta Thalassemia Major. <i>Hematology</i> , <b>2016</b> , 21, 46-53	2.2	5
23	Advanced glycation end products and soluble receptor as markers of oxidative stress in children on hemodialysis. <i>Renal Failure</i> , <b>2015</b> , 37, 1452-6	2.9	8
22	Bone mass and biochemical markers of bone turnover in children and adolescents with chronic immune thrombocytopenia: relation to corticosteroid therapy and vitamin D receptor gene polymorphisms. <i>Platelets</i> , <b>2013</b> , 24, 282-7	3.6	7
21	Adiponectin: an adipocyte-derived hormone, and its gene encoding in children with chronic kidney disease. <i>BMC Research Notes</i> , <b>2012</b> , 5, 174	2.3	5
20	Endothelial nitric oxide synthase gene intron4 VNTR polymorphism in patients with chronic kidney disease. <i>Blood Coagulation and Fibrinolysis</i> , <b>2011</b> , 22, 487-92	1	15
19	Rationale for an international consortium to study inherited genetic susceptibility to childhood acute lymphoblastic leukemia. <i>Haematologica</i> , <b>2011</b> , 96, 1049-54	6.6	32
18	Genetic polymorphism of ACE and the angiotensin II type1 receptor genes in children with chronic kidney disease. <i>Journal of Inflammation</i> , <b>2011</b> , 8, 20	6.7	17

## LIST OF PUBLICATIONS

17	A comparison of obese and nonobese Egyptian children with asthma and exploring serum eotaxin level as a link between obesity and asthma. <i>Medical Research Journal</i> , <b>2011</b> , 10, 63-68		1
16	Predictors of bone disease in Egyptian prepubertal children with Ethalassaemia major. <i>Archives of Medical Science</i> , <b>2010</b> , 6, 584-91	2.9	7
15	Methylene tetrahydrofolate reductase gene polymorphism in Egyptian children with acute lymphoblastic leukemia. <i>Blood Coagulation and Fibrinolysis</i> , <b>2010</b> , 21, 28-34	1	41
14	Oxidative stress markers and C-reactive protein in pediatric patients on hemodialysis. <i>Annals of Nutrition and Metabolism</i> , <b>2009</b> , 55, 309-16	4.5	15
13	Malondialdehyde and coenzyme Q10 in platelets and serum in type 2 diabetes mellitus: correlation with glycemic control. <i>Blood Coagulation and Fibrinolysis</i> , <b>2009</b> , 20, 248-51	1	23
12	Intradialytic and postdialytic platelet activation, increased platelet phosphatidylserine exposure and ultrastructural changes in platelets in children with chronic uremia. <i>Blood Coagulation and Fibrinolysis</i> , <b>2009</b> , 20, 230-9	1	12
11	Anti-beta2-glycoprotein I in childhood immune thrombocytopenic purpura. <i>Blood Coagulation and Fibrinolysis</i> , <b>2008</b> , 19, 26-31	1	10
10	Study of factor VII, tissue factor pathway inhibitor and monocyte tissue factor in noninsulin-dependent diabetes mellitus. <i>Blood Coagulation and Fibrinolysis</i> , <b>2008</b> , 19, 7-13	1	9
9	The spectrum of inherited bleeding disorders in pediatrics. <i>Blood Coagulation and Fibrinolysis</i> , <b>2008</b> , 19, 771-5	1	15
8	Study of genetic polymorphism of xenobiotic enzymes in acute leukemia. <i>Blood Coagulation and Fibrinolysis</i> , <b>2007</b> , 18, 489-95	1	18
7	Role of Thrombopoietin in Megakaryopoiesis and Thrombopoiesis with Relation to Platelets Ultrastructure. <i>Journal of Medical Sciences (Faisalabad, Pakistan)</i> , <b>2007</b> , 7, 179-186	0.5	
6	Study of Genetic Polymorphism of Xenobiotic Enzymes in Acute Leukemia. <i>Journal of Medical Sciences (Faisalabad, Pakistan)</i> , <b>2007</b> , 7, 354-360	0.5	
5	Intra-and Postdialytic Platelet Activation, Increased Platelet Phosphatidylserine Exposure and Ultrastructural Changes in Platelets in Children with Chronic Uremia. <i>Journal of Medical Sciences</i> (Faisalabad, Pakistan), <b>2007</b> , 7, 319-329	0.5	2
4	Tumor Necrosis Factor-IGene Polymorphism in Hemodialysis Pediatric Patients: Association with Comorbidity, Functionality and Serum Albumin. <i>Journal of Medical Sciences (Faisalabad, Pakistan)</i> , <b>2007</b> , 7, 776-782	0.5	1
3	Anti- Glycoprotein I in Childhood Immune Thrombocytopenic Purpura. <i>Journal of Medical Sciences</i> (Faisalabad, Pakistan), <b>2006</b> , 6, 279-285	0.5	
2	Plasma Nitric Oxide Level in Myocardial Disorders with Left Ventricular Diastolic Dysfunction. Journal of Medical Sciences (Faisalabad, Pakistan), <b>2006</b> , 6, 439-444	0.5	
1	Antithrombin-III in Pediatric Systemic Lupus Erythematosus. <i>Journal of Medical Sciences (Faisalabad, Pakistan)</i> , <b>2006</b> , 6, 779-785	0.5	