Yahya Wahba

List of Publications by Year
in descending order

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		1307594	1372567
31	165	7	10
papers	citations	h-index	g-index

33	33	33	197
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Prevalence of congenital heart diseases in children with Down syndrome in Mansoura, Egypt: a retrospective descriptive study. Annals of Saudi Medicine, 2017, 37, 386-392.	1.1	16
2	Rapunzel syndrome (gastric trichobezoar), a rare presentation with generalised oedema: case report and review of the literature. Paediatrics and International Child Health, 2019, 39, 76-78.	1.0	15
3	Novel homozygousSLC29A3mutations among two unrelated Egyptian families with spectral features of H-syndrome. Pediatric Diabetes, 2015, 16, 305-316.	2.9	11
4	Oxidative stress and vitamin D receptor BsmI gene polymorphism in Egyptian children with systemic lupus erythematosus: a single center study. Lupus, 2019, 28, 771-777.	1.6	10
5	Burden of family caregivers of Down syndrome children: a cross-sectional study. Family Practice, 2021, 38, 159-164.	1.9	10
6	Mutation analysis of the GLUT2 gene in three unrelated Egyptian families with Fanconiâ€“Bickel syndrome: revisited gene atlas for renumbering. Clinical and Experimental Nephrology, 2012, 16, 604-610.	1.6	9
7	Lymphocyte subgroups and recurrent infections in children with Down syndrome â€“ a prospective case control study. Central-European Journal of Immunology, 2018, 43, 248-254.	1.2	9
8	Disruptive behavior in Down syndrome children: a cross-sectional comparative study. Annals of Saudi Medicine, 2014, 34, 517-521.	1.1	9
9	Fanconi Bickel Syndrome: Novel Mutations inGLUT 2Gene Causing a Distinguished Form of Renal Tubular Acidosis in Two Unrelated Egyptian Families. Case Reports in Nephrology, 2011, 2011, 1-5.	0.4	8
10	Sporadic Fibrodysplasia Ossificans Progressiva in an Egyptian Infant with c.617G > A Mutation inACVR1Gene: A Case Report and Review of Literature. Case Reports in Genetics, 2013, 2013, 1-8.	0.2	8
11	CYP2C9 (*2&*3) and CYP2C19 (*2&*3) polymorphisms among children with nonlesional epilepsy: a single-center study. Acta Neurologica Belgica, 2021, 121, 1623-1631.	1.1	8
12	The potential impact of CYP2D6 (*2/*4/*10) gene variants among Egyptian epileptic children: A preliminary study. Gene, 2022, 832, 146585.	2.2	8
13	CD40 and CD72 expression and prognostic values among children with systemic lupus erythematosus: a caseâ€“control study. Lupus, 2020, 29, 1270-1276.	1.6	6
14	Polymorphisms of interleukin 4 and interleukin 4 receptor genes and bronchial asthma risk among Egyptian children. Clinical Biochemistry, 2021, 93, 66-72.	1.9	6
15	Novel truncating mutation in the CTNS gene in an Egyptian family with cases of infantile nephropathic cystinosis and congenital heart malformations. Middle East Journal of Medical Genetics, 2012, 1, 71-75.	0.0	4
16	Neurocutaneous melanosis is not always a benign disease. Indian Journal of Dermatology, Venereology and Leprology, 2020, 86, 335.	0.6	4
17	Comparison of Soybean-based Oil and MCT-olive-fish-soy Oil Intravenous Lipid Emulsions on Soluble Adhesion Markers in Preterm Neonates with Sepsis: A Randomized Controlled Trial. Indian Pediatrics, 2019, 56, 841-844.	0.4	3
18	Psychiatric Disorders and Quality of Life in Egyptian Patients with Chronic Immune Thrombocytopenic Purpura: A Single Center Study. Indian Journal of Hematology and Blood Transfusion, 2019, 35, 347-351.	0.6	3

#	Article	IF	Citations
19	Cluster of Differentiation 96 in Children with Acute Leukemia: A Single Center Cohort Study. Indian Journal of Hematology and Blood Transfusion, 2020, 36, 178-182.	0.6	3
20	Comparison of Soybean-based Oil and MCT-olive-fish-soy Oil Intravenous Lipid Emulsions on Soluble Adhesion Markers in Preterm Neonates with Sepsis: A Randomized Controlled Trial. Indian Pediatrics, 2019, 56, 841-844.	0.4	3
21	Camuratiâ€“Engelmann disease: New clinical insights in an Egyptian case report. Journal of Orthopaedic Science, 2020, 25, 529-532.	1.1	2
22	A novel fibrillin-1 mutation in an egyptian marfan family: A proband showing nephrotic syndrome due to focal segmental glomerulosclerosis. Saudi Journal of Kidney Diseases and Transplantation: an Official Publication of the Saudi Center for Organ Transplantation, Saudi Arabia, 2017, 28, 141.	0.3	2
23	Prevalence of Helicobacter pylori infection among children with primary nephrotic syndrome: a cross-sectional study. African Health Sciences, 2020, 20, 1624-31.	0.7	2
24	Prevalence and patterns of chromosomal abnormalities among Egyptian patients with infertility: a single institutionâ€™s 5-year experience. Middle East Fertility Society Journal, 2022, 27, .	1.5	2
25	Genetic and biochemical studies of hepatic carcinoma in the Egyptian population. Journal of Research in Medical Sciences, 2021, 26, 62.	0.9	1
26	Endothelin-1 RS5370 gene polymorphism in primary nephrotic syndrome: A case-control study. Anales De PediatrÃa (English Edition), 2020, 95, 406-412.	0.2	1
27	Effects of Cerebrolysin on the Neurodevelopmental Outcomes in Infants with Down Syndrome: A Randomized Controlled Pilot Trial. Journal of Mental Health Research in Intellectual Disabilities, 0, , 1-16.	2.0	1
28	Pre versus post application of a 0.12% chlorhexidine based oral hygiene protocol in an Egyptian pediatric intensive care unit: Practice and effects. Egyptian Journal of Critical Care Medicine, 2017, 5, 87-91.	0.4	0
29	Iron Deficiency Anemia and Serum Hepcidin Level in Children with Typhoid Fever: A Caseâ€“Control Study. Journal of Pediatric Infectious Diseases, 2020, 15, 288-292.	0.2	0
30	Serum amyloid A versus C-reactive protein in sepsis: new insights in an Egyptian ICU. Research and Opinion in Anesthesia and Intensive Care, 2019, 6, 429.	0.2	0
31	Evaluation of the left ventricular systolic strain in hypocalcemic infants with twoâ€dimensional speckle tracking echocardiography. Echocardiography, 2022, , .	0.9	0