Naglaa Mohamed Kamal

List of Publications by Year
in descending order

92 papers	1,055 citations	430874 18 h-index	501196 28 g-index
94 all docs	94 docs citations	94 times ranked	1614 citing authors

#	Article	IF	Citations
1	Burden of pediatric hepatitis C. World Journal of Gastroenterology, 2013, 19, 7880.	3.3	102
2	Biomarkers and early detection of late onset anthracycline-induced cardiotoxicity in children. Hematology, 2012, 17, 151-156.	1.5	51
3	Gastrointestinal basidiobolomycosis in children: an overlooked emerging infection?. Journal of Medical Microbiology, 2011, 60, 871-880.	1.8	44
4	Entomophthoromycosis: a challenging emerging disease. Mycoses, 2014, 57, 132-137.	4.0	41
5	Serial serum alkaline phosphatase as an early biomarker for osteopenia of prematurity. Medicine (United States), 2016, 95, e4837.	1.0	41
6	Nutritional Biomarkers in Children and Adolescents with Beta-Thalassemia-Major: An Egyptian Center Experience. BioMed Research International, 2014, 2014, 1-7.	1.9	40
7	Gastrointestinal basidiobolomycosis: an emerging fungal infection causing bowel perforation in a child. Journal of Medical Microbiology, 2011, 60, 1395-1402.	1.8	39
8	Premature atherosclerosis in children with beta-thalassemia major: New diagnostic marker. BMC Pediatrics, 2017, 17, 69.	1.7	32
9	Hodgkin Lymphoma in Childhood. Medicine (United States), 2015, 94, e670.	1.0	31
10	Diagnosis of gastrointestinal basidiobolomycosis: a miniâ€review. Mycoses, 2014, 57, 138-143.	4.0	29
11	NT-proBNP as Early Marker of Subclinical Late Cardiotoxicity after Doxorubicin Therapy and Mediastinal Irradiation in Childhood Cancer Survivors. Disease Markers, 2015, 2015, 1-10.	1.3	28
12	Prevalence and clinical significance of nonorgan specific antibodies in patients with autoimmune thyroiditis as predictor markers for rheumatic diseases. Medicine (United States), 2016, 95, e4336.	1.0	27
13	Extended clinical features associated with novel Glis3 mutation: a case report. BMC Endocrine Disorders, 2017, 17, 14.	2.2	24
14	Cadmium Status Among Pediatric Cancer Patients in Egypt. Medicine (United States), 2015, 94, e740.	1.0	23
15	A pilot single arm observational study of sofosbuvir/ledipasvir (200 + 45 mg) in 6â€•to 12â€•year old children. Alimentary Pharmacology and Therapeutics, 2018, 47, 1699-1704.	3.7	23
16	Diagnosis of spontaneous bacterial peritonitis in infants and children with chronic liver disease: A cohort study. Italian Journal of Pediatrics, 2011, 37, 26.	2.6	20
17	Outcomes of Early Ligation of Patent Ductus Arteriosus in Preterms, Multicenter Experience. Medicine (United States), 2015, 94, e915.	1.0	20
18	Outpatient blind percutaneous liver biopsy in infants and children: Is it safe?. Saudi Journal of Gastroenterology, 2012, 18, 26.	1.1	19

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19	Serum levels of interleukin-6 and tumor necrosis factor alpha in children with attention-deficit hyperactivity disorder. Journal of Pediatric Neurosciences, 2020, 15, 402.	0.3	16
20	Medical Management of Chronic Liver Diseases (CLD) in Children (Part II). Paediatric Drugs, 2011, 13, 371-383.	3.1	15
21	Benign intrascrotal lipoblastoma in a 4-month-old infant: a case report and review of literature. Journal of Pediatric Surgery, 2011, 46, e9-e12.	1.6	15
22	Psychological Impact of Chemotherapy for Childhood Acute Lymphoblastic Leukemia on Patients and Their Parents. Medicine (United States), 2015, 94, e2280.	1.0	15
23	A case report of biotinâ€“thiamine-responsive basal ganglia disease in a Saudi child. Medicine (United) Tj ETQq1 1 0.784314 rgBT /Ov	1.0	15
24	Disease patterns of pediatric non-Hodgkin lymphoma: A study from a developing area in Egypt. Molecular and Clinical Oncology, 2015, 3, 139-144.	1.0	14
25	Arginine dimethylation products in pediatric patients with chronic kidney disease. Annals of Medicine and Surgery, 2016, 9, 22-27.	1.1	14
26	Acute lymphoblastic leukemia: Are Egyptian children adherent to maintenance therapy?. Journal of Cancer Research and Therapeutics, 2015, 11, 54.	0.9	13
27	Evaluation oF Epicardial Fat and Carotid Intima-Media Thickness in Obese Children. Iranian Journal of Pediatrics, 2016, 26, e2968.	0.3	13
28	Medical Management of Chronic Liver Diseases in Children (Part I). Paediatric Drugs, 2011, 13, 357-370.	3.1	12
29	Predictive accuracy of serum hyaluronic acid as a non-invasive marker of fibrosis in a cohort of multi-transfused Egyptian children with Î²-thalassaemia major. Arab Journal of Gastroenterology, 2012, 13, 45-48.	0.9	12
30	Cinacalcet in Pediatric and Adolescent Chronic Kidney Disease. Medicine (United States), 2015, 94, e401.	1.0	12
31	Pilot study of classic galactosemia: Neurodevelopmental impact and other complications urge neonatal screening in Egypt. Journal of Advanced Research, 2018, 12, 39-45.	9.5	12
32	A Cross-sectional Study of Two Chemotherapy Protocols on Long Term Neurocognitive Functions in Egyptian Children Surviving Acute Lymphoblastic Leukemia. Current Pediatric Reviews, 2018, 14, 253-260.	0.8	11
33	Whole exome sequencing identifies rare biallelic ALMS1 missense and stop gain mutations in familial AlstrÃ¶m syndrome patients. Saudi Journal of Biological Sciences, 2020, 27, 271-278.	3.8	11
34	Effects of dual sofosbuvir/daclatasvir therapy on, chronic hepatitis C infected, survivors of childhood malignancy. World Journal of Clinical Cases, 2019, 7, 2247-2255.	0.8	11
35	Congenital chloride losing diarrhea. Medicine (United States), 2019, 98, e15928.	1.0	10
36	Value of electroencephalographic monitoring in newborns with hypoxic-ischemic encephalopathy treated with hypothermia. Journal of Pediatric Neurosciences, 2016, 11, 309.	0.3	10

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37	Colorectal polyps: a frequently-missed cause of rectal bleeding in Egyptian children. Annals of Tropical Paediatrics, 2011, 31, 213-218.	1.0	9
38	A randomized clinical trial comparing 3 different replacement regimens of vitamin D in clinically asymptomatic pediatrics and adolescents with vitamin D insufficiency. Italian Journal of Pediatrics, 2016, 42, 106.	2.6	9
39	AlstrÃ¶m syndrome. Medicine (United States), 2017, 96, e6192.	1.0	9
40	Vici syndrome with pathogenic homozygous EPG5 gene mutation. Medicine (United States), 2020, 99, e22302.	1.0	8
41	Factor VIII inhibitor development in Egyptian hemophilia patients: does intron 22 inversion mutation play a role?. Italian Journal of Pediatrics, 2020, 46, 129.	2.6	8
42	Perinatal transmission of hepatitis C virus: an update. Archives of Medical Science, 2020, 16, 1360-1369.	0.9	8
43	Identification of a Rare Exon 19 Skipping Mutation in ALMS1 Gene in AlstrÃ¶m Syndrome Patients From Two Unrelated Saudi Families. Frontiers in Pediatrics, 2021, 9, 652011.	1.9	8
44	Late presentation of necrotizing enterocolitis associated with rotavirus infection in a term infant with hyperinsulinism on octreotide therapy. Medicine (United States), 2017, 96, e7949.	1.0	7
45	Huge Non-parasitic Mesothelial Splenic Cyst in a Child: A Case Report and Literature Review. Clinical Medicine Insights Pediatrics, 2021, 15, 117955652110215.	1.4	7
46	Ovarian dysgerminoma with normal serum tumour markers presenting in a child with precocious puberty. Journal of Cancer Research and Therapeutics, 2015, 11, 661.	0.9	7
47	First Report of Acute Lymphoblastic Leukemia in an Egyptian Child with Î²-Thalassemia Major. Hemoglobin, 2015, 39, 127-129.	0.8	6
48	Early detection of myocardial dysfunction in poorly treated pediatric thalassemia children and adolescents: Two Saudi centers experience. Annals of Medicine and Surgery, 2016, 9, 6-11.	1.1	6
49	Pediatric sarcoidosis presenting as huge splenomegaly. Pediatrics International, 2017, 59, 366-367.	0.5	6
50	Knowledge of Neonatal Hyperbilirubinemia Among Primary Health Care Physicians: A Single-Center Experience. Clinical Medicine Insights Pediatrics, 2019, 13, 117955651882437.	1.4	6
51	Conventional intensive versus LED intensive phototherapy oxidative stress burden in neonatal hyperbilirubinaemia of haemolytic origin. Paediatrics and International Child Health, 2020, 40, 30-34.	1.0	6
52	Heart-type fatty acid-binding protein as a predictor of cardiac ischemia in intractable seizures in children. Journal of Pediatric Neurosciences, 2016, 11, 175.	0.3	6
53	Serum superoxide dismutase activity in acute and chronic paediatric liver diseases. Arab Journal of Gastroenterology, 2014, 15, 72-75.	0.9	5
54	Hepatic Injury in Neonates with Perinatal Asphyxia. Global Pediatric Health, 2021, 8, 2333794X2098778.	0.7	5

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55	Vitamin D status and healthy Egyptian adolescents. Medicine (United States), 2021, 100, e26661.	1.0	5
56	Study of Trace Elements and Role of Zinc Supplementation in Children with Idiopathic Intractable Epilepsy. Journal of Pediatric Epilepsy, 2016, 05, 026-033.	0.2	4
57	Serum endocan and endothelial dysfunction in childhood acute lymphoblastic leukemia survivors: a tertiary center experience. Therapeutic Advances in Chronic Disease, 2021, 12, 204062232110159.	2.5	4
58	Saudi Experts Consensus on Diagnosis and Management of Pediatric Functional Constipation. Pediatric Gastroenterology, Hepatology and Nutrition, 2022, 25, 163.	1.2	4
59	Sotos syndrome. Medicine (United States), 2018, 97, e12867.	1.0	3
60	Intrafamilial Transmission of Hepatitis C Virus Among Families of Infected Pediatric Oncology Patients. Pediatric Infectious Disease Journal, 2019, 38, 692-697.	2.0	3
61	Primary hyperoxaluria Type 1. Medicine (United States), 2020, 99, e20371.	1.0	3
62	Serum Neuron-specific Enolase and S100 Calcium-binding Protein B in Pediatric Diabetic Ketoacidosis. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 374-387.	0.9	3
63	Acute lymphoblastic leukemia in a Î²-thalassemia intermedia child: A case report. World Journal of Clinical Pediatrics, 2020, 9, 1-6.	2.1	3
64	Liver function changes following the introduction of aÂgluten-free diet in patients with celiac disease. Clinical and Experimental Hepatology, 2021, 7, 415-421.	1.3	3
65	Sexual maturity of children on regular hemodialysis. Medicine (United States), 2022, 101, e28689.	1.0	3
66	Novel homozygous mutation of PNLIP gene in congenital pancreatic lipase deficiency: an extended family study. Therapeutic Advances in Chronic Disease, 2022, 13, 204062232210787.	2.5	3
67	Detection of minimal residual disease in childhood B-acute lymphoblastic leukemia by 4-color flowcytometry. International Journal of Hematology, 2017, 105, 784-791.	1.6	2
68	Idiopathic hypoparathyroidism with extensive intracranial calcification in children. Medicine (United) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 5	1.0	2
69	Sjogrenâ€“Larsson Syndrome: A case series of five members from an extended family with a novel mutation. Molecular Genetics & Genomic Medicine, 2020, 8, e1487.	1.2	2
70	Genetic polymorphism of vitamin D receptors and plasminogen activator inhibitorâ€1 and osteonecrosis risk in childhood acute lymphoblastic leukemia. Molecular Genetics & Genomic Medicine, 2021, 9, e1700.	1.2	2
71	A comparative study of endoscopic ultrasonography versus endoscopic retrograde cholangiopancreatography in children with chronic liver disease. Indian Journal of Medical Sciences, 2008, 62, 345.	0.1	2
72	Omega-3 supplementation in children with ADHD and intractable epilepsy. Journal of Clinical Neuroscience, 2021, 94, 237-243.	1.5	2

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73	Diarrhea in neutropenic children with cancer: An Egyptian center experience, with emphasis on neutropenic enterocolitis. Indian Journal of Medical and Paediatric Oncology, 2012, 33, 95.	0.2	1
74	Folic Acid Intake and Neural Tube Defects. Medicine (United States), 2015, 94, e1395.	1.0	1
75	First report of co-morbidity of pantothenate kinase-associated neurodegeneration and three types of chronic hemolytic anemias. Annals of Medicine and Surgery, 2016, 5, 11-13.	1.1	1
76	Lethal neonatal mitochondrial phenotype caused by a novel polymerase subunit gamma mutation. Medicine (United States), 2018, 97, e12591.	1.0	1
77	Effects of Dual Sofosbuvir/Daclatasvir Therapy on Weight and Linear Growth in Adolescent Patients with Chronic Hepatitis C Virus Infection. Pediatric Infectious Disease Journal, 2019, 38, e154-e157.	2.0	1
78	Favorable response to carbamazepine therapy in genetically proven myoclonus-dystonia child. Italian Journal of Pediatrics, 2021, 47, 33.	2.6	1
79	CMV, B and C hepatitis among multi-transfused hereditary hemolytic Anemia children: an updated Egyptian experience. Italian Journal of Pediatrics, 2021, 47, 117.	2.6	1
80	Role of 18f-fdg-pet/ct in assessment of pediatric Hodgkin's lymphoma. Quarterly Journal of Nuclear Medicine and Molecular Imaging, 2014, , .	0.7	1
81	Toe tourniquet syndrome. Journal of King Abdulaziz University, Islamic Economics, 2014, 35, 865-7.	1.1	1
82	IgA Vasculitis Without Typical Skin Rash Concomitated With c-ANCA Positivity. Clinical Medicine Insights: Case Reports, 2022, 15, 117954762210777.	0.7	1
83	Risk Factors of Intractable Epilepsy in Children with Cerebral Palsy. Iranian Journal of Child Neurology, 2021, 15, 75-87.	0.3	1
84	Study of non-organ-specific antibodies in Egyptian children with genotype-4 chronic hepatitis C. Egyptian Liver Journal, 2014, 4, 1-7.	0.6	0
85	Pediatric Liver Disease in theÂAfrican Continent. , 2019, , 699-741.		0
86	Abdominal mystery in a neonate. Archives of Disease in Childhood: Education and Practice Edition, 2020, 105, 227-229.	0.5	0
87	Intrafamilial Transmission of Hepatitis C Virus Among Families of Infected Pediatric Oncology Patients. SSRN Electronic Journal, 0, , .	0.4	0
88	Persistent Hypertransaminasemia Uncovered Occult Limb-Girdle-Muscle Dystrophy-Type-2C in a Saudi Child. Journal of Pediatric Research, 0, , 57-59.	0.2	0
89	Infantile Parotid Hemangioma With Diagnostic Dilemma: A Case Report. Clinical Medicine Insights: Case Reports, 2022, 15, 117954762110733.	0.7	0
90	Facial Asymmetry in a Newly Born Baby: Diagnostic Challenge!. Clinical Medicine Insights: Case Reports, 2022, 15, 117954762210884.	0.7	0

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91	Novel Melano-Cortin-2-Receptor Gene Mutation Presenting With Infantile Cholestasis: A Case Report. Clinical Medicine Insights: Case Reports, 2022, 15, 117954762210913.	0.7	0
92	Beta Thalassemia Carrier rate: Problem Burden Among High School Children. Current Pediatric Reviews, 2022, 18, .	0.8	0