Naglaa Mohamed Kamal

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6479859/publications.pdf

Version: 2024-02-01

92 papers 1,055 citations

430442 18 h-index 28 g-index

94 all docs 94 docs citations

times ranked

94

1614 citing authors

#	Article	IF	CITATIONS
1	Infantile Parotid Hemangioma With Diagnostic Dilemma: A Case Report. Clinical Medicine Insights: Case Reports, 2022, 15, 117954762110733.	0.3	O
2	Sexual maturity of children on regular hemodialysis. Medicine (United States), 2022, 101, e28689.	0.4	3
3	Facial Asymmetry in a Newly Born Baby: Diagnostic Challenge!. Clinical Medicine Insights: Case Reports, 2022, 15, 117954762210884.	0.3	O
4	IgA Vasculitis Without Typical Skin Rash Concomitated With c-ANCA Positivity. Clinical Medicine Insights: Case Reports, 2022, 15, 117954762210777.	0.3	1
5	Novel homozygous mutation of PNLIP gene in congenital pancreatic lipase deficiency: an extended family study. Therapeutic Advances in Chronic Disease, 2022, 13, 204062232210787.	1.1	3
6	Novel Melano-Cortin-2-Receptor Gene Mutation Presenting With Infantile Cholestasis: A Case Report. Clinical Medicine Insights: Case Reports, 2022, 15, 117954762210913.	0.3	0
7	Saudi Experts Consensus on Diagnosis and Management of Pediatric Functional Constipation. Pediatric Gastroenterology, Hepatology and Nutrition, 2022, 25, 163.	0.4	4
8	Beta Thalassemia Carrier rate: Problem Burden Among High School Children. Current Pediatric Reviews, 2022, 18, .	0.4	0
9	Serum endocan and endothelial dysfunction in childhood acute lymphoblastic leukemia survivors: a tertiary center experience. Therapeutic Advances in Chronic Disease, 2021, 12, 204062232110159.	1.1	4
10	Huge Non-parasitic Mesothelial Splenic Cyst in a Child: A Case Report and Literature Review. Clinical Medicine Insights Pediatrics, 2021, 15, 117955652110215.	0.7	7
11	Hepatic Injury in Neonates with Perinatal Asphyxia. Global Pediatric Health, 2021, 8, 2333794X2098778.	0.3	5
12	Favorable response to carbamazepine therapy in genetically proven myoclonus-dystonia child. Italian Journal of Pediatrics, 2021, 47, 33.	1.0	1
13	Identification of a Rare Exon 19 Skipping Mutation in ALMS1 Gene in Alström Syndrome Patients From Two Unrelated Saudi Families. Frontiers in Pediatrics, 2021, 9, 652011.	0.9	8
14	CMV, B and C hepatitis among multi-transfused hereditary hemolytic Anemia children: an updated Egyptian experience. Italian Journal of Pediatrics, 2021, 47, 117.	1.0	1
15	Genetic polymorphism of vitamin D receptors and plasminogen activator inhibitorâ€1 and osteonecrosis risk in childhood acute lymphoblastic leukemia. Molecular Genetics & amp; Genomic Medicine, 2021, 9, e1700.	0.6	2
16	Vitamin D status and healthy Egyptian adolescents. Medicine (United States), 2021, 100, e26661.	0.4	5
17	Omega-3 supplementation in children with ADHD and intractable epilepsy. Journal of Clinical Neuroscience, 2021, 94, 237-243.	0.8	2
18	Liver function changes following the introduction of aÂgluten-free diet in patients with celiac disease. Clinical and Experimental Hepatology, 2021, 7, 415-421.	0.6	3

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19	Risk Factors of Intractable Epilepsy in Children with Cerebral Palsy. Iranian Journal of Child Neurology, 2021, 15, 75-87.	0.2	1
20	Conventional intensive <i>versus</i> LED intensive phototherapy oxidative stress burden in neonatal hyperbilirubinaemia of haemolytic origin. Paediatrics and International Child Health, 2020, 40, 30-34.	0.3	6
21	Whole exome sequencing identifies rare biallelic ALMS1 missense and stop gain mutations in familial Alström syndrome patients. Saudi Journal of Biological Sciences, 2020, 27, 271-278.	1.8	11
22	Vici syndrome with pathogenic homozygous EPG5 gene mutation. Medicine (United States), 2020, 99, e22302.	0.4	8
23	Abdominal mystery in a neonate. Archives of Disease in Childhood: Education and Practice Edition, 2020, 105, 227-229.	0.3	O
24	Factor VIII inhibitor development in Egyptian hemophilia patients: does intron 22 inversion mutation play a role?. Italian Journal of Pediatrics, 2020, 46, 129.	1.0	8
25	Sjogren–Larsson Syndrome: A case series of five members from an extended family with a novel mutation. Molecular Genetics & Genomic Medicine, 2020, 8, e1487.	0.6	2
26	Primary hyperoxaluria Type 1. Medicine (United States), 2020, 99, e20371.	0.4	3
27	Perinatal transmission of hepatitis C virus: an update. Archives of Medical Science, 2020, 16, 1360-1369.	0.4	8
28	Serum levels of interleukin-6 and tumor necrosis factor alpha in children with attention-deficit hyperactivity disorder. Journal of Pediatric Neurosciences, 2020, 15, 402.	0.2	16
29	Acute lymphoblastic leukemia in a β-thalassemia intermedia child: A case report. World Journal of Clinical Pediatrics, 2020, 9, 1-6.	0.6	3
30	Pediatric Liver Disease in theÂAfrican Continent. , 2019, , 699-741.		0
31	Knowledge of Neonatal Hyperbilirubinemia Among Primary Health Care Physicians: A Single-Center Experience. Clinical Medicine Insights Pediatrics, 2019, 13, 117955651882437.	0.7	6
32	Intrafamilial Transmission of Hepatitis C Virus Among Families of Infected Pediatric Oncology Patients. Pediatric Infectious Disease Journal, 2019, 38, 692-697.	1.1	3
33	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.	1.1	1
34	Congenital chloride losing diarrhea. Medicine (United States), 2019, 98, e15928.	0.4	10
35	Effects of dual sofosbuvir/daclatasvir therapy on, chronic hepatitis C infected, survivors of childhood malignancy. World Journal of Clinical Cases, 2019, 7, 2247-2255.	0.3	11
36	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.	0.4	3

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37	Pilot study of classic galactosemia: Neurodevelopmental impact and other complications urge neonatal screening in Egypt. Journal of Advanced Research, 2018, 12, 39-45.	4.4	12
38	A pilot single arm observational study of sofosbuvir/ledipasvir (200 + 45 mg) in 6―to 12―year old children. Alimentary Pharmacology and Therapeutics, 2018, 47, 1699-1704.	1.9	23
39	Sotos syndrome. Medicine (United States), 2018, 97, e12867.	0.4	3
40	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.	0.4	11
41	Lethal neonatal mitochondrial phenotype caused by a novel polymerase subunit gamma mutation. Medicine (United States), 2018, 97, e12591.	0.4	1
42	Extended clinical features associated with novel Glis3 mutation: a case report. BMC Endocrine Disorders, 2017, 17, 14.	0.9	24
43	Premature atherosclerosis in children with beta-thalassemia major: New diagnostic marker. BMC Pediatrics, 2017, 17, 69.	0.7	32
44	Detection of minimal residual disease in childhood B-acute lymphoblastic leukemia by 4-color flowcytometry. International Journal of Hematology, 2017, 105, 784-791.	0.7	2
45	Pediatric sarcoidosis presenting as huge splenomegaly. Pediatrics International, 2017, 59, 366-367.	0.2	6
46	Late presentation of necrotizing enterocolitis associated with rotavirus infection in a term infant with hyperinsulinism on octreotide therapy. Medicine (United States), 2017, 96, e7949.	0.4	7
47	Alström syndrome. Medicine (United States), 2017, 96, e6192.	0.4	9
48	Idiopathic hypoparathyroidism with extensive intracranial calcification in children. Medicine (United) Tj ETQq0 0 0	rgBT /Ove	rlock 10 Tf 5
49	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.	1.0	9
50	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.	0.4	27
51	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.	O.5	1
52	Study of Trace Elements and Role of Zinc Supplementation in Children with Idiopathic Intractable Epilepsy. Journal of Pediatric Epilepsy, 2016, 05, 026-033.	0.1	4
53	A case report of biotin–thiamine-responsive basal ganglia disease in a Saudi child. Medicine (United) Tj ETQq1 1	0.784314 0.4	rgBT /Ov <mark>erl</mark>
54	Serial serum alkaline phosphatase as an early biomarker for osteopenia of prematurity. Medicine (United States), 2016, 95, e4837.	0.4	41

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55	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.	0.5	6
56	Arginine dimethylation products in pediatric patients with chronic kidney disease. Annals of Medicine and Surgery, 2016, 9, 22-27.	0.5	14
57	Heart-type fatty acid-binding protein as a predictor of cardiac ischemia in intractable seizures in children. Journal of Pediatric Neurosciences, 2016, 11, 175.	0.2	6
58	Value of electroencephalographic monitoring in newborns with hypoxic-ischemic encephalopathy treated with hypothermia. Journal of Pediatric Neurosciences, 2016, 11, 309.	0.2	10
59	Evaluation oF Epicardial Fat and Carotid Intima-Media Thickness in Obese Children. Iranian Journal of Pediatrics, 2016, 26, e2968.	0.1	13
60	Folic Acid Intake and Neural Tube Defects. Medicine (United States), 2015, 94, e1395.	0.4	1
61	Hodgkin Lymphoma in Childhood. Medicine (United States), 2015, 94, e670.	0.4	31
62	Outcomes of Early Ligation of Patent Ductus Arteriosus in Preterms, Multicenter Experience. Medicine (United States), 2015, 94, e915.	0.4	20
63	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.	0.6	28
64	First Report of Acute Lymphoblastic Leukemia in an Egyptian Child with β-Thalassemia Major. Hemoglobin, 2015, 39, 127-129.	0.4	6
65	Disease patterns of pediatric non-Hodgkin lymphoma: A study from a developing area in Egypt. Molecular and Clinical Oncology, 2015, 3, 139-144.	0.4	14
66	Cadmium Status Among Pediatric Cancer Patients in Egypt. Medicine (United States), 2015, 94, e740.	0.4	23
67	Psychological Impact of Chemotherapy for Childhood Acute Lymphoblastic Leukemia on Patients and Their Parents. Medicine (United States), 2015, 94, e2280.	0.4	15
68	Acute lymphoblastic leukemia: Are Egyptian children adherent to maintenance therapy?. Journal of Cancer Research and Therapeutics, 2015, 11, 54.	0.3	13
69	Cinacalcet in Pediatric and Adolescent Chronic Kidney Disease. Medicine (United States), 2015, 94, e401.	0.4	12
70	Ovarian dysgerminoma with normal serum tumour markers presenting in a child with precocious puberty. Journal of Cancer Research and Therapeutics, 2015, 11, 661.	0.3	7
71	Nutritional Biomarkers in Children and Adolescents with Beta-Thalassemia-Major: An Egyptian Center Experience. BioMed Research International, 2014, 2014, 1-7.	0.9	40
72	Study of non-organ-specific antibodies in Egyptian children with genotype-4 chronic hepatitis C. Egyptian Liver Journal, 2014, 4, 1-7.	0.3	0

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73	Diagnosis of gastrointestinal basidiobolomycosis: a miniâ€review. Mycoses, 2014, 57, 138-143.	1.8	29
74	Entomophthoromycosis: a challenging emerging disease. Mycoses, 2014, 57, 132-137.	1.8	41
7 5	Serum superoxide dismutase activity in acute and chronic paediatric liver diseases. Arab Journal of Gastroenterology, 2014, 15, 72-75.	0.4	5
76	Role of 18f-fdg-pet/ct in assessment of pediatric Hodgkin's lymphoma. Quarterly Journal of Nuclear Medicine and Molecular Imaging, 2014, , .	0.4	1
77	Toe tourniquet syndrome. Journal of King Abdulaziz University, Islamic Economics, 2014, 35, 865-7.	0.5	1
78	Burden of pediatric hepatitis C. World Journal of Gastroenterology, 2013, 19, 7880.	1.4	102
79	Biomarkers and early detection of late onset anthracycline-induced cardiotoxicity in children. Hematology, 2012, 17, 151-156.	0.7	51
80	Outpatient blind percutaneous liver biopsy in infants and children: Is it safe?. Saudi Journal of Gastroenterology, 2012, 18, 26.	0.5	19
81	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.	0.1	1
82	Predictive accuracy of serum hyaluronic acid as a non-invasive marker of fibrosis in a cohort of multi-transfused Egyptian children with \hat{I}^2 -thalassaemia major. Arab Journal of Gastroenterology, 2012, 13, 45-48.	0.4	12
83	Medical Management of Chronic Liver Diseases in Children (Part I). Paediatric Drugs, 2011, 13, 357-370.	1.3	12
84	Medical Management of Chronic Liver Diseases (CLD) in Children (Part II). Paediatric Drugs, 2011, 13, 371-383.	1.3	15
85	Benign intrascrotal lipoblastoma in a 4-month-old infant: a case report and review of literature. Journal of Pediatric Surgery, 2011, 46, e9-e12.	0.8	15
86	Diagnosis of spontaneous bacterial peritonitis in infants and children with chronic liver disease: A cohort study. Italian Journal of Pediatrics, 2011, 37, 26.	1.0	20
87	Colorectal polyps: a frequently-missed cause of rectal bleeding in Egyptian children. Annals of Tropical Paediatrics, 2011, 31, 213-218.	1.0	9
88	Gastrointestinal basidiobolomycosis: an emerging fungal infection causing bowel perforation in a child. Journal of Medical Microbiology, 2011, 60, 1395-1402.	0.7	39
89	Gastrointestinal basidiobolomycosis in children: an overlooked emerging infection?. Journal of Medical Microbiology, 2011, 60, 871-880.	0.7	44
90	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

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91	Intrafamilial Transmission of Hepatitis C Virus Among Families of Infected Pediatric Oncology Patients. SSRN Electronic Journal, 0, , .	0.4	O
92	Persistent Hypertransaminasemia Uncovered Occult Limb-Girdle-Muscle Dystrophy-Type-2C in a Saudi Child. Journal of Pediatric Research, 0, , 57-59.	0.1	0