Fahad A Bashiri

List of Publications by Year
in descending order

54 papers	1,942 citations	516215 16 h-index	276539 41 g-index
60 all docs	60 docs citations	60 times ranked	4023 citing authors

#	Article	IF	Citations
1	Childhood absence epilepsy. International Journal of Pediatrics and Adolescent Medicine, 2022, 9, 131-135.	0.5	3
2	Faculty Membersâ€™ Perspective on Virtual Interviews for Medical Residency Matching during the COVID-19 Crisis: A National Survey. Healthcare (Switzerland), 2022, 10, 16.	1.0	4
3	Prevalence of autism spectrum disorder among Saudi children between 2 and 4 years old in Riyadh. Asian Journal of Psychiatry, 2022, 71, 103054.	0.9	13
4	AGREEing on Clinical Practice Guidelines for Autism Spectrum Disorders in Children: A Systematic Review and Quality Assessment. Children, 2022, 9, 1050.	0.6	6
5	Effect of lumbar puncture educational video on parental knowledge and self-reported intended practice. International Journal of Pediatrics and Adolescent Medicine, 2021, 8, 112-116.	0.5	2
6	Phenotypic and molecular spectrum of pyridoxamineâ€5â€²â€phosphate oxidase deficiency: A scoping review of 87 cases of pyridoxamineâ€5â€²â€phosphate oxidase deficiency. Clinical Genetics, 2021, 99, 99-110.	1.0	25
7	Consensus Statement on the Management of Duchenne Muscular Dystrophy in Saudi Arabia During the Coronavirus Disease 2019 Pandemic. Frontiers in Pediatrics, 2021, 9, 629549.	0.9	7
8	Adapting evidence-based clinical practice guidelines for people with attention deficit hyperactivity disorder in Saudi Arabia: process and outputs of a national initiative. Child and Adolescent Psychiatry and Mental Health, 2021, 15, 6.	1.2	6
9	Syncopal attacks in children: Is it cardiac or epilepsy related?. Autonomic Neuroscience: Basic and Clinical, 2021, 231, 102771.	1.4	0
10	Proxy-Reported Quality of Life and Access to Nusinersen Among Patients with Spinal Muscular Atrophy in Saudi Arabia. Patient Preference and Adherence, 2021, Volume 15, 729-739.	0.8	6
11	The Leukodystrophy Spectrum in Saudi Arabia: Epidemiological, Clinical, Radiological, and Genetic Data. Frontiers in Pediatrics, 2021, 9, 633385.	0.9	15
12	2020 Update to Spinal Muscular Atrophy Management in Saudi Arabia. Frontiers in Pediatrics, 2021, 9, 684134.	0.9	2
13	Newborn screening programs for spinal muscular atrophy worldwide: Where we stand and where to go. Neuromuscular Disorders, 2021, 31, 574-582.	0.3	94
14	Medical Facultyâ€™s and Studentsâ€™ Perceptions toward Pediatric Electronic OSCE during the COVID-19 Pandemic in Saudi Arabia. Healthcare (Switzerland), 2021, 9, 950.	1.0	11
15	Remote interviews for medical residency selection during the initial COVID-19 crisis: a national survey. BMC Medical Education, 2021, 21, 462.	1.0	9
16	Parental Attitudes and Hesitancy About COVID-19 vs. Routine Childhood Vaccinations: A National Survey. Frontiers in Public Health, 2021, 9, 752323.	1.3	106
17	Acute Necrotizing Encephalopathy of Childhood: A Multicenter Experience in Saudi Arabia. Frontiers in Pediatrics, 2020, 8, 526.	0.9	20
18	JAK Inhibitor Therapy in a Child with Inherited USP18 Deficiency. New England Journal of Medicine, 2020, 382, 256-265.	13.9	69

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19	Expanding the phenotype and the genotype of Stromme syndrome: A novel variant of the CENPF gene and literature review. European Journal of Medical Genetics, 2020, 63, 103844.	0.7	12
20	Predicting future handedness and hemispheric dominance during infancy by analyzing sleep spindles. Journal of King Abdulaziz University, Islamic Economics, 2020, 25, 188-192.	0.5	0
21	Nusinersen in spinal muscular atrophy: Respiratory outcomes at tertiary care centers. Journal of Nature and Science of Medicine, 2020, .	0.1	0
22	A consensus statement on spinal muscular atrophy management in Saudi Arabia in the context of COVID-19. Journal of King Abdulaziz University, Islamic Economics, 2020, 25, 230-237.	0.5	3
23	Clinical spectrum of neurofibromatosis type 1 among children in a tertiary care center. Neurosciences, 2020, 25, 375-379.	0.1	0
24	Clinical spectrum of neurofibromatosis type 1 among children in a tertiary care center. Journal of King Abdulaziz University, Islamic Economics, 2020, 25, 375-379.	0.5	2
25	Appraisal of clinical practice guidelines for the management of attention deficit hyperactivity disorder (ADHD) using the AGREE II Instrument: A systematic review. PLoS ONE, 2019, 14, e0219239.	1.1	9
26	Lessons Learned from Large-Scale, First-Tier Clinical Exome Sequencing in a Highly Consanguineous Population. American Journal of Human Genetics, 2019, 104, 1182-1201.	2.6	184
27	Pyridox(am)ine 5â€²-Phosphate Oxidase Deficiency: Severe Prenatal Presentation with Hypoxic Ischemic Encephalopathy. Journal of Pediatric Epilepsy, 2019, 08, 049-055.	0.1	2
28	Genomic and phenotypic delineation of congenital microcephaly. Genetics in Medicine, 2019, 21, 545-552.	1.1	85
29	Satisfaction and perceived usefulness with newly-implemented Electronic Health Records System among pediatricians at a university hospital. Computer Methods and Programs in Biomedicine, 2019, 169, 51-57.	2.6	26
30	Hypokalemic periodic paralysis due to CACNA1S gene mutation. Journal of King Abdulaziz University, Islamic Economics, 2019, 24, 225-230.	0.5	7
31	Effect of new modalities of treatment on physiciansâ€™ management plan for patients with spinal muscular atrophy. Journal of King Abdulaziz University, Islamic Economics, 2019, 24, 16-21.	0.5	3
32	Physicians' Understanding of Nutritional Factors Determining Brain Development and Cognition in the Middle East and Africa. Pediatric Gastroenterology, Hepatology and Nutrition, 2019, 22, 536.	0.4	1
33	Pediatric intracranial hypertension. Journal of King Abdulaziz University, Islamic Economics, 2019, 24, 257-263.	0.5	1
34	GWAS signals revisited using human knockouts. Genetics in Medicine, 2018, 20, 64-68.	1.1	6
35	A Case of Neonatal Epileptic Encephalopathy due to SCN2A Mutation Responsive to a Ketogenic Diet. Journal of Pediatric Epilepsy, 2018, 07, 148-151.	0.1	3
36	Vitamin D supplementation to prevent vitamin D deficiency for children with epilepsy. Medicine (United) Tj ETQq0 0 0 rgBT /Overlock 10	0.4	14

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37	Assessment of physiciansâ€™ knowledge and attitudes in the management of febrile seizures. Journal of King Abdulaziz University, Islamic Economics, 2018, 23, 314-319.	0.5	5
38	Respiratory support attitudes among pediatric intensive care staff for spinal muscular atrophy patients in Saudi Arabia. Journal of King Abdulaziz University, Islamic Economics, 2018, 23, 62-65.	0.5	2
39	Drowning in the desert: Family denial of brain death. Sudanese Journal of Paediatrics, 2018, 18, 48-52.	0.6	2
40	Auto-immune anti-N-methyl-D-aspartate receptor (anti-NMDAR) encephalitis: three case reports. Paediatrics and International Child Health, 2017, 37, 222-226.	0.3	6
41	Confirming the recessive inheritance of <scp>SCN1B</scp> mutations in developmental epileptic encephalopathy. Clinical Genetics, 2017, 92, 327-331.	1.0	32
42	Mutations in NKX6-2 Cause Progressive Spastic Ataxia and Hypomyelination. American Journal of Human Genetics, 2017, 100, 969-977.	2.6	38
43	The landscape of genetic diseases in Saudi Arabia based on the first 1000 diagnostic panels and exomes. Human Genetics, 2017, 136, 921-939.	1.8	209
44	Clinical genomics expands the morbid genome of intellectual disability and offers a high diagnostic yield. Molecular Psychiatry, 2017, 22, 615-624.	4.1	187
45	Management of convulsive status epilepticus in children: an adapted clinical practice guideline for pediatricians in Saudi Arabia. Journal of King Abdulaziz University, Islamic Economics, 2017, 22, 146-155.	0.5	12
46	Childhood epilepsies: What should a pediatrician know?. Journal of King Abdulaziz University, Islamic Economics, 2017, 22, 14-19.	0.5	6
47	Idiopathic intracranial hypertension in children: Diagnostic and management approach. Sudanese Journal of Paediatrics, 2016, 16, 67-76.	0.6	14
48	Mycoplasma pneumoniae Infection. Global Pediatric Health, 2015, 2, 2333794X1559276.	0.3	2
49	Accelerating Novel Candidate Gene Discovery in Neurogenetic Disorders via Whole-Exome Sequencing of Prescreened Multiplex Consanguineous Families. Cell Reports, 2015, 10, 148-161.	2.9	375
50	ARL6IP6, a susceptibility locus for ischemic stroke, is mutated in a patient with syndromic Cutis Marmorata Telangiectatica Congenita. Human Genetics, 2015, 134, 815-822.	1.8	13
51	The clinical utility of molecular karyotyping for neurocognitive phenotypes in a consanguineous population. Genetics in Medicine, 2015, 17, 719-725.	1.1	19
52	Sturge-Weber syndrome: Continued vigilance is needed. Sudanese Journal of Paediatrics, 2015, 15, 63-70.	0.6	5
53	Biotin-responsive basal ganglia disease should be renamed biotin-thiamine-responsive basal ganglia disease: a retrospective review of the clinical, radiological and molecular findings of 18 new cases. Orphanet Journal of Rare Diseases, 2013, 8, 83.	1.2	123
54	SLC25A22 is a novel gene for migrating partial seizures in infancy. Annals of Neurology, 2013, 74, 873-882.	2.8	102