

Mohammed Zain Seidahmed

List of Publications by Year in descending order

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Version: 2024-02-01

16
papers

797
citations

932766

10
h-index

996533

15
g-index

16
all docs

16
docs citations

16
times ranked

2101
citing authors

#	ARTICLE	IF	CITATIONS
1	Hereditary hyperekplexia in Saudi Arabia. <i>Pediatric Neurology</i> , 2022, , .	1.0	0
2	Ancient founder mutation in RUBCN: a second unrelated family confirms Salih ataxia (SCAR15). <i>BMC Neurology</i> , 2020, 20, 207.	0.8	7
3	Autozygome and high throughput confirmation of disease genes candidacy. <i>Genetics in Medicine</i> , 2019, 21, 736-742.	1.1	81
4	Gonadal mosaicism for <i>ACTA1</i> gene masquerading as autosomal recessive nemaline myopathy. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 2219-2221.	0.7	8
5	Hyperekplexia, microcephaly and simplified gyral pattern caused by novel ASNS mutations, case report. <i>BMC Neurology</i> , 2016, 16, 105.	0.8	32
6	Accelerating Novel Candidate Gene Discovery in Neurogenetic Disorders via Whole-Exome Sequencing of Prescreened Multiplex Consanguineous Families. <i>Cell Reports</i> , 2015, 10, 148-161.	2.9	375
7	Sirenomelia and severe caudal regression syndrome. <i>Journal of King Abdulaziz University, Islamic Economics</i> , 2014, 35 Suppl 1, S36-43.	0.5	3
8	Classification, clinical features, and genetics of neural tube defects. <i>Journal of King Abdulaziz University, Islamic Economics</i> , 2014, 35 Suppl 1, S5-S14.	0.5	15
9	Genetic, chromosomal, and syndromic causes of neural tube defects. <i>Journal of King Abdulaziz University, Islamic Economics</i> , 2014, 35 Suppl 1, S49-56.	0.5	8
10	Epidemiology of neural tube defects. <i>Journal of King Abdulaziz University, Islamic Economics</i> , 2014, 35 Suppl 1, S29-35.	0.5	19
11	Epidemiology, prenatal management, and prevention of neural tube defects. <i>Journal of King Abdulaziz University, Islamic Economics</i> , 2014, 35 Suppl 1, S15-28.	0.5	22
12	Genomic analysis of Meckel-Gruber syndrome in Arabs reveals marked genetic heterogeneity and novel candidate genes. <i>European Journal of Human Genetics</i> , 2013, 21, 762-768.	1.4	56
13	Molecular characterization of Joubert syndrome in Saudi Arabia. <i>Human Mutation</i> , 2012, 33, 1423-1428.	1.1	56
14	A TCTN2 mutation defines a novel Meckel Gruber syndrome locus. <i>Human Mutation</i> , 2011, 32, 573-578.	1.1	72
15	A case of fetal valproate syndrome with new features expanding the phenotype. <i>Journal of King Abdulaziz University, Islamic Economics</i> , 2009, 30, 288-91.	0.5	8
16	Brain Stem and Cerebellar Findings in Joubert Syndrome. <i>Journal of Computer Assisted Tomography</i> , 2006, 30, 116-121.	0.5	35