Mohammed Zain Seidahmed

List of Publications by Year
in descending order

16	797	932766 10	996533 15
papers	citations	h-index	g-index

16	16	16	2101
all docs	docs citations	times ranked	citing authors

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