Aldhalaan Hesham

List of Publications by Year
in descending order

14 papers	825 citations	1307594 7 h-index	1199594 12 g-index
19 all docs	19 docs citations	19 times ranked	2394 citing authors

#	Article	IF	Citations
1	Clinicoâ€radiological features, molecular spectrum, and identification of prognostic factors in developmental and epileptic encephalopathy due to inosine triphosphate pyrophosphatase (ITPase) deficiency. Human Mutation, 2022, 43, 403-419.	2.5	9
2	DNA Methylation Level of Transcription Factor Binding Site in the Promoter Region of Acyl-CoA Synthetase Family Member 3 (ACSF3) in Saudi Autistic Children. Pharmacogenomics and Personalized Medicine, 2022, Volume 15, 131-142.	0.7	0
3	A Novel GEMIN4 Variant in a Consanguineous Family Leads to Neurodevelopmental Impairment with Severe Microcephaly, Spastic Quadriplegia, Epilepsy, and Cataracts. Genes, 2022, 13, 92.	2.4	6
4	Saudi Arabian Consensus Statement on Vagus Nerve Stimulation for Refractory Epilepsy. Saudi Journal of Medicine and Medical Sciences, 2021, 9, 75.	0.8	1
5	Bi-allelic variants in HOPS complex subunit VPS41 cause cerebellar ataxia and abnormal membrane trafficking. Brain, 2021, 144, 769-780.	7.6	33
6	Potential health risks of maternal phthalate exposure during the first trimester - The Saudi Early Autism and Environment Study (SEAES). Environmental Research, 2021, 195, 110882.	7.5	5
7	Genetics of ataxia telangiectasia in a highly consanguineous population. Annals of Human Genetics, 2021, , .	0.8	4
8	Effects of early and recent mercury and lead exposure on the neurodevelopment of children with elevated mercury and/or developmental delays during lactation: A follow-up study. International Journal of Hygiene and Environmental Health, 2020, 230, 113629.	4.3	10
9	Tuberous sclerosis complex: Clinical spectrum and epilepsy: A retrospective chart review study. Translational Neuroscience, 2018, 9, 154-160.	1.4	10
10	The landscape of genetic diseases in Saudi Arabia based on the first 1000 diagnostic panels and exomes. Human Genetics, 2017, 136, 921-939.	3.8	209
11	Expanding the genetic heterogeneity of intellectual disability. Human Genetics, 2017, 136, 1419-1429.	3.8	122
12	Congenital disorders of glycosylation: The Saudi experience. American Journal of Medical Genetics, Part A, 2017, 173, 2614-2621.	1.2	31
13	Accelerating Novel Candidate Gene Discovery in Neurogenetic Disorders via Whole-Exome Sequencing of Prescreened Multiplex Consanguineous Families. Cell Reports, 2015, 10, 148-161.	6.4	375
14	Challenges of Autism Spectrum Disorders Families Towards Oral Health Care in Kingdom of Saudi Arabia. Pesquisa Brasileira Em Odontopediatria E Clinica Integrada, 0, 20, .	0.9	6