

Saleem Ahmed

List of Publications by Year in descending order

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Version: 2024-02-01

8
papers

207
citations

1478505

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1720034

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docs citations

8
times ranked

623
citing authors

#	ARTICLE	IF	CITATIONS
1	A missense mutation in TRAPPC6A leads to build-up of the protein, in patients with a neurodevelopmental syndrome and dysmorphic features. Scientific Reports, 2018, 8, 2053.	3.3	30
2	Mutations in DONSON disrupt replication fork stability and cause microcephalic dwarfism. Nature Genetics, 2017, 49, 537-549.	21.4	81
3	The alkylglycerol monooxygenase (AGMO) gene previously involved in autism also causes a novel syndromic form of primary microcephaly in a consanguineous Saudi family. Journal of the Neurological Sciences, 2016, 363, 240-244.	0.6	18
4	Identification of Two Homozygous Sequence Variants in the <i>COL7A1</i> Gene Underlying Dystrophic Epidermolysis Bullosa by Whole-Exome Analysis in a Consanguineous Family. Annals of Human Genetics, 2015, 79, 350-356.	0.8	3
5	Novel nonsense mutation in the PTRF gene underlies congenital generalized lipodystrophy in a consanguineous Saudi family. European Journal of Medical Genetics, 2015, 58, 216-221.	1.3	31
6	Exome analysis identified a novel missense mutation in the CLPP gene in a consanguineous Saudi family expanding the clinical spectrum of Perrault Syndrome type-3. Journal of the Neurological Sciences, 2015, 353, 149-154.	0.6	37
7	Femoral-facial syndrome in an infant of a diabetic mother. BMJ Case Reports, 2015, 2015, bcr2014208857.	0.5	7
8	Cenani-Lenz syndrome-like limb anomaly with more severe involvement of left side. BMJ Case Reports, 2012, 2012, bcr0120125634-bcr0120125634.	0.5	0