Zeliha Gormez

List of Publications by Year
in descending order

18	150	1307594 7	1199594 12
papers	citations	h-index	g-index

18	18	18	492
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Expansion of the phenotypic spectrum of SMC1A nonsense variants: a patient with cerebellar atrophy and review of the literature. Clinical Dysmorphology, 2020, 29, 217-223.	0.3	1
2	A Novel and Mosaic WDR45 Nonsense Variant Causes Beta-Propeller Protein-Associated Neurodegeneration Identified Through Whole Exome Sequencing and X chromosome Heterozygosity Analysis. NeuroMolecular Medicine, 2019, 21, 54-59.	3.4	5
3	A novel truncating mutation of DOCK7 gene with an early-onset non-encephalopathic epilepsy. Seizure: the Journal of the British Epilepsy Association, 2019, 66, 12-14.	2.0	4
4	A further family of Stromme syndrome carrying CENPF mutation. American Journal of Medical Genetics, Part A, 2017, 173, 1668-1672.	1.2	12
5	Myophosphorylase (PYGM) mutations determined by next generation sequencing in a cohort from Turkey with McArdle disease. Neuromuscular Disorders, 2017, 27, 997-1008.	0.6	11
6	SYNE1 related cerebellar ataxia presents with variable phenotypes in a consanguineous family from Turkey. Neurological Sciences, 2017, 38, 2203-2207.	1.9	8
7	Exome sequencing identifies a novel homozygous CLN8 mutation in a Turkish family with Northern epilepsy. Acta Neurologica Belgica, 2017, 117, 159-167.	1.1	5
8	Coffin-Siris syndrome with cafÃ©-au-lait spots, obesity and hyperinsulinism caused by a mutation in the <i>ARID1B</i> gene. Intractable and Rare Diseases Research, 2016, 5, 222-226.	0.9	14
9	An Aggregated Cross-Validation Framework for Computational Discovery of Disease-Associative Genes. IFMBE Proceedings, 2016, , 489-494.	0.3	1
10	Loss-of-Function Mutations in ELMO2 Cause Intraosseous Vascular Malformation by Impeding RAC1 Signaling. American Journal of Human Genetics, 2016, 99, 299-317.	6.2	23
11	FMFilter: A fast model based variant filtering tool. Journal of Biomedical Informatics, 2016, 60, 319-327.	4.3	5
12	A clinical variant in SCN1A inherited from a mosaic father cosegregates with a novel variant to cause Dravet syndrome in a consanguineous family. Epilepsy Research, 2015, 113, 5-10.	1.6	8
13	Hypertension prediction by multi-objective optimization methods. , 2014, , .		0
14	HomSI: a homozygous stretch identifier from next-generation sequencing data. Bioinformatics, 2014, 30, 445-447.	4.1	45
15	Multi objective SNP selection using pareto optimality. Computational Biology and Chemistry, 2013, 43, 23-28.	2.3	5
16	Comparison of aggregators for multi-objective SNP selection. , 2013, 2013, 3062-5.		0
17	Statistical bias and variance of gene selection and cross validation methods: A case study on hypertension prediction. , 2012, , .		1
18	Evaluation of the Concatenative Turkish Text-to-Speech System. , 2009, , .		2