Hande Kaymakcalan

List of Publications by Year
in descending order

19 papers	1,765 citations	687363 13 h-index	839539 18 g-index
19 all docs	19 docs citations	19 times ranked	4065 citing authors

#	Article	IF	Citations
1	Mutation spectrum of congenital heart disease in a consanguineous Turkish population. Molecular Genetics & Genomic Medicine, 2022, 10, e1944.	1.2	4
2	Prevalence and clinical/molecular characteristics of PTEN mutations in Turkish children with autism spectrum disorders and macrocephaly. Molecular Genetics & Genomic Medicine, 2021, 9, e1739.	1.2	8
3	1q21.1 Deletions and Duplications in 2 Siblings with Psychiatric Problems. Indian Journal of Pediatrics, 2019, 86, 1068-1068.	0.8	3
4	Novel compound heterozygous mutations in GPT2 linked to microcephaly, and intellectual developmental disability with or without spastic paraplegia. American Journal of Medical Genetics, Part A, 2018, 176, 421-425.	1.2	8
5	Biallelic loss of human CTNNA2, encoding Î±N-catenin, leads to ARP2/3 complex overactivity and disordered cortical neuronal migration. Nature Genetics, 2018, 50, 1093-1101.	21.4	70
6	Expanding the Genotypic Spectrum of Bathing Suit Ichthyosis. JAMA Dermatology, 2017, 153, 537.	4.1	17
7	ALPK3 gene mutation in a patient with congenital cardiomyopathy and dysmorphic features. Journal of Physical Education and Sports Management, 2017, 3, a001859.	1.2	20
8	Biallelic Mutations in Citron Kinase Link Mitotic Cytokinesis to Human Primary Microcephaly. American Journal of Human Genetics, 2016, 99, 501-510.	6.2	70
9	Changes in peanut allergy prevalence in different ethnic groups in 2 time periods. Journal of Allergy and Clinical Immunology, 2015, 135, 580-582.	2.9	16
10	NGLY1 mutation causes neuromotor impairment, intellectual disability, and neuropathy. European Journal of Medical Genetics, 2015, 58, 39-43.	1.3	69
11	Brain Malformations Associated With Knobloch Syndromeâ€"Review of Literature, Expanding Clinical Spectrum, and Identification of Novel Mutations. Pediatric Neurology, 2014, 51, 806-813.e8.	2.1	43
12	Mutations in LAMB1 Cause Cobblestone Brain Malformation without Muscular or Ocular Abnormalities. American Journal of Human Genetics, 2013, 92, 468-474.	6.2	96
13	Genomic Analysis of Non- NF2 Meningiomas Reveals Mutations in TRAF7 , KLF4 , AKT1 , and SMO. Science, 2013, 339, 1077-1080.	12.6	714
14	Prenatal Health, Educational Attainment, and Intergenerational Inequality: The Northern Finland Birth Cohort 1966 Study. Demography, 2012, 49, 525-552.	2.5	21
15	Recessive LAMC3 mutations cause malformations of occipital cortical development. Nature Genetics, 2011, 43, 590-594.	21.4	102
16	Whole-exome sequencing identifies recessive WDR62 mutations in severe brain malformations. Nature, 2010, 467, 207-210.	27.8	457
17	Clinical and genomic characterization of distal duplications and deletions of chromosome 4q: Study of two cases and review of the literature. American Journal of Medical Genetics, Part A, 2009, 149A, 2788-2794.	1.2	46
18	Assessment of Nasal Swab Samples, Environmental Microbiological Analyses and Respiratory Function Tests of Bakery Workers Working Under Modern Conditions: A Brief Report. Indoor and Built Environment, 2006, 15, 197-202.	2.8	1

#	Article	IF	Citations
19	Correlation of PAPP-A values with maternal characteristics, biochemical and ultrasonographic markers of pregrancy. Marmara Medical Journal, 0, , .	0.8	0