Sevcan TuÄ BozdoÄan

List of Publications by Year
in descending order

15 papers	112 citations	1683934 5 h-index	1372474 10 g-index
15 all docs	15 docs citations	15 times ranked	194 citing authors

#	Article	IF	Citations
1	Bi-allelic JAM2 Variants Lead to Early-Onset Recessive Primary Familial Brain Calcification. American Journal of Human Genetics, 2020, 106, 412-421.	2.6	47
2	Alpha-Thalassemia Mutations in Adana Province, Southern Turkey: Genotype-Phenotype Correlation. Indian Journal of Hematology and Blood Transfusion, 2015, 31, 223-228.	0.3	14
3	No Association between Polymorphisms of Vitamin D and Oxytocin Receptor Genes and Autistic Spectrum Disorder in a Sample of Turkish Children. Clinical Psychopharmacology and Neuroscience, 2018, 16, 415-421.	0.9	9
4	A Rare Double Aneuploidy Case (Downâ€“Klinefelter). Journal of Pediatric Genetics, 2017, 06, 241-243.	0.3	8
5	Clinical and molecular evaluation of MEFV gene variants in the Turkish population: a study by the National Genetics Consortium. Functional and Integrative Genomics, 2022, 22, 291-315.	1.4	7
6	BRCAmutation characteristics in a series of index cases of breast cancer selected independent of family history. Breast Journal, 2019, 25, 1029-1033.	0.4	6
7	IL-1RN VNTR, IL-2(-330), and IL-4 VNTR gene polymorphisms in patients with chronic rhinosinusitis with sinonasal polyposis. Turkish Journal of Medical Sciences, 2019, 49, 1411-1417.	0.4	6
8	Current Status of Genetic Diagnosis Laboratories and Frequency of Genetic Variants Associated with Cystic Fibrosis through a Newborn-Screening Program in Turkey. Genes, 2021, 12, 206.	1.0	5
9	Vandetanib in a Child Affected by Neurofibromatosis Type 1 and Medullary Thyroid Carcinoma with Both <i>NF1</i> and Homozygous <i>RET</i> Proto-oncogen Germ-line Mutations. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 342-346.	0.4	3
10	Hereditary Spastic Paraplegia Type 35 with a Novel Mutation in Fatty Acid 2-Hydroxylase Gene and Literature Review of the Clinical Features. Annals of Indian Academy of Neurology, 2018, 21, 335-339.	0.2	3
11	Shprintzen-Goldberg Syndrome: Case Report. Meandros Medical and Dental Journal, 2018, 19, 175-177.	0.1	2
12	Identification of a novel mutation in GRIN2A gene with global developmental delay and refractory epilepsy. Annals of Indian Academy of Neurology, 2020, 23, 696.	0.2	1
13	No Association between Polymorphisms of Vitamin D and Oxytocin Receptor Genes and Autistic Spectrum Disorder in a Sample of Turkish Children. Clinical Psychopharmacology and Neuroscience, 2018, 16, 415-421.	0.9	1
14	Evaluation of the results of patients who applied to the Ã‡ukurova University, Medical Genetics Department for prenatal diagnosis and determination of genetic counseling principles. Turkish Journal of Medical Sciences, 2021, 51, 657-660.	0.4	0
15	A Novel Mutation Diagnosing in Allanâ€“Herndonâ€“Dudley's Syndrome. Journal of Pediatric Genetics, 0, , .	0.3	0