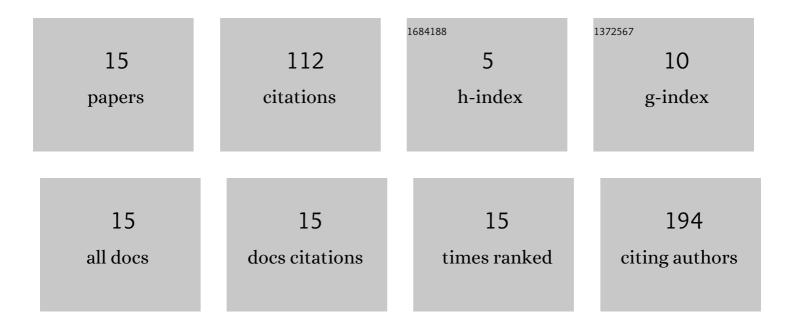
## Sevcan TuÄ BozdoÄän

List of Publications by Year in descending order

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| #  | Article  | IF  | CITATIONS |
|----|--|-----|-----------|
| 1  | 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.  | 3.5 | 7         |
| 2  | Current Status of Genetic Diagnosis Laboratories and Frequency of Genetic Variants Associated with<br>Cystic Fibrosis through a Newborn-Screening Program in Turkey. Genes, 2021, 12, 206.   | 2.4 | 5         |
| 3  | Evaluation of the results of patients who applied to the Çukurova University, Medical Genetics<br>Department for prenatal diagnosis and determination of genetic counseling principles. Turkish<br>Journal of Medical Sciences, 2021, 51, 657-660.                   | 0.9 | Ο         |
| 4  | Vandetanib in a Child Affected by Neurofibromatosis Type 1 and Medullary Thyroid Carcinoma with<br>Both <i>NF1</i> and Homozygous <i>RET</i> Proto-oncogen Germ-line Mutations.<br>JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 342-346. | 0.9 | 3         |
| 5  | Bi-allelic JAM2 Variants Lead to Early-Onset Recessive Primary Familial Brain Calcification. American<br>Journal of Human Genetics, 2020, 106, 412-421.  | 6.2 | 47        |
| 6  | Identification of a novel mutation in GRIN2A gene with global developmental delay and refractory epilepsy. Annals of Indian Academy of Neurology, 2020, 23, 696.   | 0.5 | 1         |
| 7  | BRCAmutation characteristics in a series of index cases of breast cancer selected independent of family history. Breast Journal, 2019, 25, 1029-1033.  | 1.0 | 6         |
| 8  | 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.  | 0.9 | 6         |
| 9  | No Association between Polymorphisms of Vitamin D and Oxytocin Receptor Genes and Autistic<br>Spectrum Disorder in a Sample of Turkish Children. Clinical Psychopharmacology and Neuroscience,<br>2018, 16, 415-421.   | 2.0 | 9         |
| 10 | Shprintzen-Goldberg Syndrome: Case Report. Meandros Medical and Dental Journal, 2018, 19, 175-177.   | 0.2 | 2         |
| 11 | Hereditary Spastic Paraplegia Type 35 with a Novel Mutation in Fatty Acid 2-Hydroxylase Gene and<br>Literature Review of the Clinical Features. Annals of Indian Academy of Neurology, 2018, 21, 335-339.  | 0.5 | 3         |
| 12 | No Association between Polymorphisms of Vitamin D and Oxytocin Receptor Genes and Autistic<br>Spectrum Disorder in a Sample of Turkish Children. Clinical Psychopharmacology and Neuroscience,<br>2018, 16, 415-421.   | 2.0 | 1         |
| 13 | A Rare Double Aneuploidy Case (Down–Klinefelter). Journal of Pediatric Genetics, 2017, 06, 241-243.  | 0.7 | 8         |
| 14 | Alpha-Thalassemia Mutations in Adana Province, Southern Turkey: Genotype-Phenotype Correlation.<br>Indian Journal of Hematology and Blood Transfusion, 2015, 31, 223-228.  | 0.6 | 14        |
| 15 | A Novel Mutation Diagnosing in Allan–Herndon–Dudley's Syndrome. Journal of Pediatric Genetics, 0, ,  | 0.7 | 0         |