# Sukriye Ayter

## List of Publications by Year
in descending order

| | | 1307594 | 1281871 |
|---|---|---|---|
| **13** papers | **118** citations | **7** h-index | **11** g-index |
| **13** all docs | **13** docs citations | **13** times ranked | **181** citing authors |

| # | Article | IF | Citations |
|---|---------|----|-----------|
| 1 | Heightened CXCR4 and CXCL12 expression in NF1-associated neurofibromas. Child's Nervous System, 2018, 34, 877-882. | 1.1 | 11 |
| 2 | Possible modifier genes in the variation of neurofibromatosis type 1 clinical phenotypes. Journal of Neurogenetics, 2018, 32, 65-77. | 1.4 | 18 |
| 3 | Clinical findings and mutation analysis of NF1 patients in Turkey. Meta Gene, 2018, 15, 80-83. | 0.6 | 3 |
| 4 | NF1 Gene Mutations are the Major Molecular Event in Neurofibromatosis-Noonan Syndrome. Journal of Neurology & Stroke, 2017, 6, . | 0.1 | 0 |
| 5 | Coronin 1A inhibits neurite outgrowth in PC12 cells. Neuroscience Letters, 2014, 582, 38-42. | 2.1 | 12 |
| 6 | Two pathogenic NF1 gene mutations identified in DNA from a child with mild phenotype. Child's Nervous System, 2012, 28, 943-946. | 1.1 | 4 |
| 7 | Absence of exon 17 c.2970-2872delAAT mutation in Turkish NF1 patients with mild phenotype. Child's Nervous System, 2011, 27, 2113-2116. | 1.1 | 4 |
| 8 | Learning disability and oligodendrocyte myelin glycoprotein (OMGP) gene in neurofibromatosis type 1. Turkish Journal of Pediatrics, 2011, 53, 75-8. | 0.6 | 4 |
| 9 | Neurofibromatosis: Novel and Recurrent Mutations in Turkish Patients. Pediatric Neurology, 2007, 37, 421-425. | 2.1 | 9 |
| 10 | Two neurofibromatosis type 1 cases associated with rhabdomyosarcoma of bladder, one with a large deletion in the NF1 gene. Cancer Genetics and Cytogenetics, 2006, 164, 159-163. | 1.0 | 22 |
| 11 | Molecular genetic analyses in neurofibromatosis type 1 patients with tumors. Cancer Genetics and Cytogenetics, 2006, 165, 167-171. | 1.0 | 12 |
| 12 | Gene symbol: NF1. Disease: neurofibromatosis 1. Human Genetics, 2006, 119, 360. | 3.8 | 1 |
| 13 | Neurofibromatosis—Noonan's Syndrome With Associated Rhabdomyosarcoma of the Urinary Bladder in an Infant: Case Report. Journal of Child Neurology, 2003, 18, 68-72. | 1.4 | 18 |