Sukriye Ayter

List of Publications by Year
in descending order

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

#	Article	IF	Citations
1	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
2	Neurofibromatosisâ€"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
3	Possible modifier genes in the variation of neurofibromatosis type 1 clinical phenotypes. Journal of Neurogenetics, 2018, 32, 65-77.	1.4	18
4	Molecular genetic analyses in neurofibromatosis type 1 patients with tumors. Cancer Genetics and Cytogenetics, 2006, 165, 167-171.	1.0	12
5	Coronin 1A inhibits neurite outgrowth in PC12 cells. Neuroscience Letters, 2014, 582, 38-42.	2.1	12
6	Heightened CXCR4 and CXCL12 expression in NF1-associated neurofibromas. Child's Nervous System, 2018, 34, 877-882.	1.1	11
7	Neurofibromatosis: Novel and Recurrent Mutations in Turkish Patients. Pediatric Neurology, 2007, 37, 421-425.	2.1	9
8	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
9	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
10	Learning disability and oligodendrocyte myelin glycoprotein (OMGP) gene in neurofibromatosis type 1. Turkish Journal of Pediatrics, 2011, 53, 75-8.	0.6	4
11	Clinical findings and mutation analysis of NF1 patients in Turkey. Meta Gene, 2018, 15, 80-83.	0.6	3
12	Gene symbol: NF1. Disease: neurofibromatosis 1. Human Genetics, 2006, 119, 360.	3.8	1
13	NF1 Gene Mutations are the Major Molecular Event in Neurofibromatosis-Noonan Syndrome. Journal of Neurology & Stroke, 2017, 6, .	0.1	0