Meral Topcu

List of Publications by Year
in descending order

19	391	1040056 9	839539 18
papers	citations	h-index	g-index

19	19	19	1289
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The Complex Genetic Landscape of Hereditary Ataxias in Turkey and Implications in Clinical Practice. Movement Disorders, 2021, 36, 1676-1688.	3.9	9
2	Unraveling neuronal ceroid lipofuscinosis type 2 (CLN2) disease: A tertiary center experience for determinants of diagnostic delay. European Journal of Paediatric Neurology, 2021, 33, 94-98.	1.6	4
3	Management of CLN1 Disease: International Clinical Consensus. Pediatric Neurology, 2021, 120, 38-51.	2.1	10
4	Genetic and phenotypic features of patients with childhood ataxias diagnosed by next-generation sequencing gene panel. Brain and Development, 2020, 42, 6-18.	1.1	12
5	MINPP1 prevents intracellular accumulation of the chelator inositol hexakisphosphate and is mutated in Pontocerebellar Hypoplasia. Nature Communications, 2020, 11, 6087.	12.8	28
6	Long-term effects of vagus nerve stimulation in refractory pediatric epilepsy: A single-center experience. Epilepsy and Behavior, 2020, 110, 107147.	1.7	13
7	Comprehensive clinical, biochemical, radiological and genetic analysis of 28 Turkish cases with suspected metachromatic leukodystrophy and their relatives. Molecular Genetics and Metabolism Reports, 2020, 25, 100688.	1.1	4
8	Prenatal enzymatic diagnosis of lysosomal storage diseases using cultured amniotic cells, uncultured chorionic villus samples, and fetal blood cells: Hacettepe experience. Prenatal Diagnosis, 2019, 39, 1080-1085.	2.3	1
9	Clinical outcomes of two patients with a novel pathogenic variant in ASNS: response to asparagine supplementation and review of the literature. Human Genome Variation, 2019, 6, 24.	0.7	31
10	Neurologic Involvement in Primary Immunodeficiency Disorders. Journal of Child Neurology, 2018, 33, 320-328.	1.4	12
11	Mystery Case: Pontine tegmental cap dysplasia in a neonate. Neurology, 2018, 91, e2100-e2101.	1.1	2
12	Management Strategies for CLN2 Disease. Pediatric Neurology, 2017, 69, 102-112.	2.1	80
13	Electrical status epilepticus during sleep: A study of 22 patients. Brain and Development, 2015, 37, 250-264.	1.1	13
14	Etiological yield of SNP microarrays in idiopathic intellectual disability. European Journal of Paediatric Neurology, 2014, 18, 327-337.	1.6	26
15	Developmental abnormalities and mental retardation. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2013, 111, 211-217.	1.8	5
16	Mutations in CLN7/MFSD8 are a common cause of variant late-infantile neuronal ceroid lipofuscinosis. Brain, 2009, 132, 810-819.	7.6	116
17	Effects of Johnstone pressure splints combined with neurodevelopmental therapy on spasticity and cutaneous sensory inputs in spastic cerebral palsy. Developmental Medicine and Child Neurology, 2001, 43, 307-313.	2.1	0
18	Letters to the editor. Muscle and Nerve, 1996, 19, 675-681.	2.2	1

#	Article	IF	Citations
19	Griscelli syndrome: Report of Three Cases. Pediatric Pathology & Laboratory Medicine: Journal of the Society for Pediatric Pathology, Affiliated With the International Paediatric Pathology Association, 1995, 15, 309-319.	0.3	24