KornÃ©lia Tripolszki

List of Publications by Year
in descending order

11	170	1307594 7	1281871 11
papers	citations	h-index	g-index

11	11	11	371
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Passive Transfer of Blood Sera from ALS Patients with Identified Mutations Results in Elevated Motoneuronal Calcium Level and Loss of Motor Neurons in the Spinal Cord of Mice. International Journal of Molecular Sciences, 2021, 22, 9994.	4.1	4
2	Passive Transfer of Sera from ALS Patients with Identified Mutations Evokes an Increased Synaptic Vesicle Number and Elevation of Calcium Levels in Motor Axon Terminals, Similar to Sera from Sporadic Patients. International Journal of Molecular Sciences, 2020, 21, 5566.	4.1	3
3	Comprehensive Genetic Analysis of a Hungarian Amyotrophic Lateral Sclerosis Cohort. Frontiers in Genetics, 2019, 10, 732.	2.3	31
4	Experimental Motor Neuron Disease Induced in Mice with Long-Term Repeated Intraperitoneal Injections of Serum from ALS Patients. International Journal of Molecular Sciences, 2019, 20, 2573.	4.1	11
5	Angiogenin mutations in Hungarian patients with amyotrophic lateral sclerosis: Clinical, genetic, computational, and functional analyses. Brain and Behavior, 2019, 9, e01293.	2.2	10
6	Genetic analysis of the SOD1 and C9ORF72 genes in Hungarian patients with amyotrophic lateral sclerosis. Neurobiology of Aging, 2017, 53, 195.e1-195.e5.	3.1	17
7	Identification of two novel mutations in the SLC45A2 gene in a Hungarian pedigree affected by unusual OCA type 4. BMC Medical Genetics, 2017, 18, 27.	2.1	8
8	Delineating the genetic heterogeneity of OCA in Hungarian patients. European Journal of Medical Research, 2017, 22, 20.	2.2	4
9	Somatic mosaicism of the PIK3CA gene identified in a Hungarian girl with macrodactyly and syndactyly. European Journal of Medical Genetics, 2016, 59, 223-226.	1.3	16
10	Phenotypical diversity of patients with LEOPARD syndrome carrying the worldwide recurrent p.Tyr279Cys PTPN11 mutation. Archives of Dermatological Research, 2015, 307, 891-895.	1.9	10
11	<scp>CTSC</scp> and Papillonâ€“LefÃ¨vre syndrome: detection of recurrent mutations in <scp>H</scp>ungarian patients, a review of published variants and database update. Molecular Genetics & Genomic Medicine, 2014, 2, 217-228.	1.2	56