Negar Khanlou

List of Publications by Citations

22 papers	715 citations	11 h-index	25 g-index
25 ext. papers	873 ext. citations	4.2 avg, IF	3.37 L-index

#	Paper	IF	Citations
22	Tunable-combinatorial mechanisms of acquired resistance limit the efficacy of BRAF/MEK cotargeting but result in melanoma drug addiction. *Cancer Cell*, **2015**, 27, 240-56	24.3	226
21	HIV DNA Is Frequently Present within Pathologic Tissues Evaluated at Autopsy from Combined Antiretroviral Therapy-Treated Patients with Undetectable Viral Loads. *Journal of Virology*, **2016**, 90, 8968-83	6.6	95
20	The pathogenesis of ataxia-telangiectasia. Learning from a Rosetta Stone. *Clinical Reviews in Allergy and Immunology*, **2001**, 20, 87-108	12.3	83
19	Mutations in TFAM, encoding mitochondrial transcription factor A, cause neonatal liver failure associated with mtDNA depletion. *Molecular Genetics and Metabolism*, **2016**, 119, 91-9	3.7	69
18	Increased frequency of alpha-synuclein in the substantia nigra in human immunodeficiency virus infection. *Journal of NeuroVirology*, **2009**, 15, 131-8	3.9	55
17	DMD genotype correlations from the Duchenne Registry: Endogenous exon skipping is a factor in prolonged ambulation for individuals with a defined mutation subtype. *Human Mutation*, **2018**, 39, 1193-1202	4.7	39
16	Incidence, survival, pathology, and genetics of adult Latino Americans with glioblastoma. *Journal of Neuro-Oncology*, **2017**, 132, 351-358	4.8	27
15	Klotho gene silencing promotes pathology in the mdx mouse model of Duchenne muscular dystrophy. *Human Molecular Genetics*, **2016**, 25, 2465-2482	5.6	25
14	Sample Preparation for Transmission Electron Microscopy. *Methods in Molecular Biology*, **2019**, 1897, 417-424	1.4	25
13	Hydroxychloroquine-induced autophagic vacuolar myopathy with mitochondrial abnormalities. *Neuropathology*, **2018**, 38, 646-652	2	20
12	Brain biopsy in neurologic decline of unknown etiology. *Human Pathology*, **2015**, 46, 499-506	3.7	18
11	Safety of focused ultrasound neuromodulation in humans with temporal lobe epilepsy. *Brain Stimulation*, **2021**, 14, 1022-1031	5.1	11
10	Early experience with formalin-fixed paraffin-embedded (FFPE) based commercial clinical genomic profiling of gliomas-robust and informative with caveats. *Experimental and Molecular Pathology*, **2017**, 103, 87-93	4.4	7
9	Cortical dysplasia with prominent Rosenthal fiber formation in a case of intractable pediatric epilepsy. *Human Pathology*, **2009**, 40, 1200-4	3.7	5
8	Cardiac Transplantation in Dermatomyositis: A case report and literature review. *Human Pathology: Case Reports*, **2017**, 8, 55-58	0.2	4
7	Large in-frame 5'deletions in DMD associated with mild Duchenne muscular dystrophy: Two case reports and a review of the literature. *Neuromuscular Disorders*, **2019**, 29, 863-873	2.9	4
6	Evaluation of PAX8 Expression in Brain Tissue and Related Neoplasms. *Applied Immunohistochemistry and Molecular Morphology*, **2016**, 24, 207-9	1.9	2

List of Publications

5	Case Report: Whole Exome Sequencing Identifies Compound Heterozygous Variants in Gene Causing Juvenile Hypertrophic Cardiomyopathy.. *Frontiers in Cardiovascular Medicine*, **2021**, 8, 798985	5.4	0
4	Recessive ciliopathy mutations in primary endocardial fibroelastosis: a rare neonatal cardiomyopathy in a case of Alstrom syndrome. *Journal of Molecular Medicine*, **2021**, 99, 1623-1638	5.5	0
3	61-year-old man with chronic expansile sellar mass. *Brain Pathology*, **2017**, 27, 397-398	6	
2	Vertical asymmetric mitochondrial ophthalmoplegia. *Canadian Journal of Ophthalmology*, **2019**, 54, e230-e232	1.1	
1	A Case of Becker Muscular Dystrophy With Rimmed Vacuoles and Normal Dystrophin. *Journal of Clinical Rheumatology*, **2020**, 26, e307-e308	1.1	