## Peter K Panegyres

List of Publications by Year in descending order

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DETED K DANECYDES

#	Article	IF	CITATIONS
1	Sigma nonopioid intracellular receptor 1 mutations cause frontotemporal lobar degeneration–motor neuron disease. Annals of Neurology, 2010, 68, 639-649.	5.3	168
2	A review of quality of life after predictive testing for and earlier identification of neurodegenerative diseases. Progress in Neurobiology, 2013, 110, 2-28.	5.7	128
3	The Patterns of Inheritance in Early-Onset Dementia. American Journal of Alzheimer's Disease and Other Dementias, 2015, 30, 299-306.	1.9	66
4	Course and Causes of Suspected Dementia in Young Adults: A Longitudinal Study. American Journal of Alzheimer's Disease and Other Dementias, 2007, 22, 48-56.	1.9	59
5	Early Dementia Screening. Diagnostics, 2016, 6, 6.	2.6	57
6	The Role of Ethnicity in Alzheimer's Disease: Findings From The C-PATH Online Data Repository. Journal of Alzheimer's Disease, 2016, 51, 515-523.	2.6	35
7	Diagnosis and management of Whipple's disease of the brain. Practical Neurology, 2008, 8, 311-317.	1.1	32
8	Prion diseases: immunotargets and therapy. ImmunoTargets and Therapy, 2016, 5, 57.	5.8	27
9	Factors determining recurrence in transient global amnesia. BMC Neurology, 2020, 20, 83.	1.8	22
10	Laryngeal dystonia causing upper airway obstruction in progressive supranuclear palsy. Journal of Clinical Neuroscience, 2007, 14, 380-381.	1.5	19
11	Anti-N-methyl-D-aspartate receptor encephalitis with an imaging-invisible ovarian teratoma: a case report. Journal of Medical Case Reports, 2016, 10, 296.	0.8	18
12	The neurology and natural history of patients with indeterminate CAG repeat length mutations of the Huntington disease gene. Journal of the Neurological Sciences, 2011, 301, 14-20.	0.6	17
13	Exploring the Role of Cognitive Reserve in Early-Onset Dementia. American Journal of Alzheimer's Disease and Other Dementias, 2011, 26, 139-144.	1.9	15
14	The natural history of early-onset dementia: the Artemis Project. BMJ Open, 2012, 2, e001764.	1.9	13
15	A patient with Creutzfeldt-Jakob disease presenting with amyotrophy: a case report. Journal of Medical Case Reports, 2013, 7, 218.	0.8	13
16	Codon 200 mutation of the prion gene: genotype–phenotype correlations. Journal of Neurology, 2012, 259, 2579-2584.	3.6	12
17	The Ancient Greek discovery of the nervous system: Alcmaeon, Praxagoras and Herophilus. Journal of Clinical Neuroscience, 2016, 29, 21-24.	1.5	12
18	Anti-Yo and anti-glutamic acid decarboxylase antibodies presenting in carcinoma of the uterus with paraneoplastic cerebellar degeneration: a case report. Journal of Medical Case Reports, 2012, 6, 155.	0.8	8

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#	Article	IF	CITATIONS
19	Factors influencing the clinical expression of intermediate CAG repeat length mutations of the Huntington's disease gene. Journal of Neurology, 2015, 262, 277-284.	3.6	7
20	Prion disease in Indigenous Australians. Internal Medicine Journal, 2020, 51, 1101-1105.	0.8	7
21	Comparison of two methods for the analysis of CSF Aβ and tau in the diagnosis of Alzheimer's disease. American Journal of Neurodegenerative Disease, 2014, 3, 143-51.	0.1	6
22	Hereditary Neuropathy with Liability to Pressure Palsy Presenting as an Acute Brachial Plexopathy: A Lover's Palsy. Case Reports in Neurology, 2014, 6, 281-286.	0.7	4
23	The Clinical Spectrum of Young Onset Dementia Points to Its Stochastic Origins. Journal of Alzheimer's Disease Reports, 2021, 5, 663-679.	2.2	3
24	Alzheimer's disease, Huntington's disease and cancer. Journal of Clinical Neuroscience, 2021, 93, 103-105.	1.5	3
25	Stochasticity, Entropy and Neurodegeneration. Brain Sciences, 2022, 12, 226.	2.3	2
26	A Caucasian Australian presenting with human T-lymphotropic virus type I associated myelopathy: a case report. Journal of Medical Case Reports, 2014, 8, 382.	0.8	1
27	Antiâ€glial fibrillary acidic protein astrocytopathy. Internal Medicine Journal, 2020, 50, 1017-1019.	0.8	0
28	Guillain-Barre syndrome complicating <i>Neisseria meningitidis</i> infection. Oxford Medical Case Reports, 2022, 2022, .	0.4	0