Janis Blevins

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/11714566/publications.pdf

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11	877	840776	1281871
papers	citations	11 h-index	11 g-index

11	11	11	1119
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Subtype Diagnosis of Sporadic <scp>Creutzfeldtâ€“Jakob</scp> Disease with Diffusion <scp>Magnetic Resonance Imaging</scp>. Annals of Neurology, 2021, 89, 560-572.	5.3	30
2	Identification of novel risk loci and causal insights for sporadic Creutzfeldt-Jakob disease: a genome-wide association study. Lancet Neurology, The, 2020, 19, 840-848.	10.2	42
3	Evaluation of a New Criterion for Detecting Prion Disease With Diffusion Magnetic Resonance Imaging. JAMA Neurology, 2020, 77, 1141.	9.0	46
4	Prion propagation estimated from brain diffusion MRI is subtype dependent in sporadic Creutzfeldtâ€“Jakob disease. Acta Neuropathologica, 2020, 140, 169-181.	7.7	28
5	Diagnosis of prion diseases by RT-QuIC results in improved surveillance. Neurology, 2020, 95, e1017-e1026.	1.1	72
6	Diagnostic and prognostic value of human prion detection in cerebrospinal fluid. Annals of Neurology, 2017, 81, 79-92.	5.3	184
7	Rapidly progressive Alzheimerâ€™s disease features distinct structures of amyloid-Î². Brain, 2015, 138, 1009-1022.	7.6	166
8	Co-existence of Distinct Prion Types Enables Conformational Evolution of Human PrPSc by Competitive Selection. Journal of Biological Chemistry, 2013, 288, 29846-29861.	3.4	45
9	Small Protease Sensitive Oligomers of PrPSc in Distinct Human Prions Determine Conversion Rate of PrPC. PLoS Pathogens, 2012, 8, e1002835.	4.7	72
10	Protease-Sensitive Conformers in Broad Spectrum of Distinct PrPSc Structures in Sporadic Creutzfeldt-Jakob Disease Are Indicator of Progression Rate. PLoS Pathogens, 2011, 7, e1002242.	4.7	66
11	Co-existence of scrapie prion protein types 1 and 2 in sporadic Creutzfeldtâ€“Jakob disease: its effect on the phenotype and prion-type characteristics. Brain, 2009, 132, 2643-2658.	7.6	126