Diane L Ritchie

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/4126546/publications.pdf

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		567281	713466
21	2,101	15	21
papers	citations	h-index	g-index

21	21	21	1085
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Preclinical vCJD after blood transfusion in a PRNP codon 129 heterozygous patient. Lancet, The, 2004, 364, 527-529.	13.7	794
2	Prevalence of lymphoreticular prion protein accumulation in UK tissue samples. Journal of Pathology, 2004, 203, 733-739.	4.5	393
3	Peripheral Tissue Involvement in Sporadic, Iatrogenic, and Variant Creutzfeldt-Jakob Disease. American Journal of Pathology, 2004, 164, 143-153.	3.8	158
4	Accumulation of prion protein in tonsil and appendix: review of tissue samples. BMJ: British Medical Journal, 2002, 325, 633-634.	2.3	125
5	Detection of Type 1 Prion Protein in Variant Creutzfeldt-Jakob Disease. American Journal of Pathology, 2006, 168, 151-157.	3.8	111
6	Detection and Localization of PrPSc in the Skeletal Muscle of Patients with Variant, Iatrogenic, and Sporadic Forms of Creutzfeldt-Jakob Disease. American Journal of Pathology, 2006, 168, 927-935.	3.8	100
7	Amyloid-Î² accumulation in the CNS in human growth hormone recipients in the UK. Acta Neuropathologica, 2017, 134, 221-240.	7.7	85
8	Prion diseases. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 145, 393-403.	1.8	68
9	Transmissions of variant Creutzfeldtâ€“Jakob disease from brain and lymphoreticular tissue show uniform and conserved bovine spongiform encephalopathy-related phenotypic properties on primary and secondary passage in wild-type mice. Journal of General Virology, 2009, 90, 3075-3082.	2.9	42
10	UK Iatrogenic Creutzfeldtâ€“Jakob disease: investigating human prion transmission across genotypic barriers using human tissue-based and molecular approaches. Acta Neuropathologica, 2017, 133, 579-595.	7.7	31
11	Sporadic Fatal Insomnia in Europe: Phenotypic Features and Diagnostic Challenges. Annals of Neurology, 2018, 84, 347-360.	5.3	31
12	Clinical, neuropathological and immunohistochemical features of sporadic and variant forms of Creutzfeldtâ€“Jakob disease in the squirrel monkey (Saimiri sciureus). Journal of General Virology, 2007, 88, 688-695.	2.9	30
13	Neuropathology of Human Prion Diseases. Progress in Molecular Biology and Translational Science, 2017, 150, 319-339.	1.7	27
14	Phenotypic diversity of genetic Creutzfeldtâ€“Jakob disease: a histo-molecular-based classification. Acta Neuropathologica, 2021, 142, 707-728.	7.7	24
15	Abnormal prion protein in the pituitary in sporadic and variant Creutzfeldtâ€“Jakob disease. Journal of General Virology, 2007, 88, 1068-1072.	2.9	20
16	Prion Diseases: A Unique Transmissible Agent or a Model for Neurodegenerative Diseases?. Biomolecules, 2021, 11, 207.	4.0	15
17	Variant CJD: Reflections a Quarter of a Century on. Pathogens, 2021, 10, 1413.	2.8	15
18	Blood transmission studies of prion infectivity in the squirrel monkey (<scp>S</scp>aimiri) Tj ETQq0 0 0 rgBT /Overlock 10 Tf	1.6	12

#	Article	IF	Citations
19	A Naturally Occurring Bovine Tauopathy Is Geographically Widespread in the UK. PLoS ONE, 2015, 10, e0129499.	2.5	9
20	Prion strains associated with iatrogenic CJD in French and UK human growth hormone recipients. Acta Neuropathologica Communications, 2021, 9, 145.	5.2	7
21	Renewed assessment of the risk of emergent advanced cell therapies to transmit neuroproteinopathies. Acta Neuropathologica, 2019, 137, 363-377.	7.7	4