Alexander Howard Peden

List of Publications by Year
in descending order

15	1,566	840776 11	996975 15
papers	citations	h-index	g-index

15	15	15	1163
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	Real time quakingâ€induced conversion analysis of cerebrospinal fluid in sporadic Creutzfeldtâ€“Jakob disease. Annals of Neurology, 2012, 72, 278-285.	5.3	315
3	Sensitive and specific detection of sporadic Creutzfeldtâ€“Jakob disease brain prion protein using real-time quaking-induced conversion. Journal of General Virology, 2012, 93, 438-449.	2.9	111
4	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
5	Amyloid-Î² accumulation in the CNS in human growth hormone recipients in the UK. Acta Neuropathologica, 2017, 134, 221-240.	7.7	85
6	Molecular Pathology in Neurodegenerative Diseases. Current Drug Targets, 2012, 13, 1548-1559.	2.1	31
7	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
8	The prion protein protease sensitivity, stability and seeding activity in variably protease sensitive prionopathy brain tissue suggests molecular overlaps with sporadic Creutzfeldt-Jakob disease. Acta Neuropathologica Communications, 2014, 2, 152.	5.2	23
9	Abnormal prion protein in the pituitary in sporadic and variant Creutzfeldtâ€“Jakob disease. Journal of General Virology, 2007, 88, 1068-1072.	2.9	20
10	Risks of transmission of variant Creutzfeldt-Jakob disease by blood transfusion. Folia Neuropathologica, 2005, 43, 271-8.	1.2	16
11	Variant CJD: Reflections a Quarter of a Century on. Pathogens, 2021, 10, 1413.	2.8	15
12	Advances in the development of a screening test for variant Creutzfeldtâ€“Jakob disease. Expert Opinion on Medical Diagnostics, 2008, 2, 207-219.	1.6	9
13	Review: pathology of variant Creutzfeldt-Jakob disease. Folia Neuropathologica, 2004, 42 Suppl A, 85-91.	1.2	7
14	Study protocol for enhanced CJD surveillance in the 65+ years population group in Scotland: an observational neuropathological screening study of banked brain tissue donations for evidence of prion disease. BMJ Open, 2019, 9, e033744.	1.9	5
15	Epitope mapping of the protease resistant products of RT-QuIC does not allow the discrimination of sCJD subtypes. PLoS ONE, 2019, 14, e0218509.	2.5	4