Maurizio Pocchiari

List of Publications by Year
in descending order

181	12,193	41344 49	27406 106
papers	citations	h-index	g-index

189	189	189	5673
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A new variant of Creutzfeldt-Jakob disease in the UK. Lancet, The, 1996, 347, 921-925.	13.7	2,554
2	Updated clinical diagnostic criteria for sporadic Creutzfeldt-Jakob disease. Brain, 2009, 132, 2659-2668.	7.6	770
3	Analysis of EEG and CSF 14-3-3 proteins as aids to the diagnosis of Creutzfeldtâ€“Jakob disease. Neurology, 2000, 55, 811-815.	1.1	432
4	Genetic prion disease: the EUROCJD experience. Human Genetics, 2005, 118, 166-174.	3.8	391
5	Mortality from Creutzfeldtâ€“Jakob disease and related disorders in Europe, Australia, and Canada. Neurology, 2005, 64, 1586-1591.	1.1	306
6	A Test for Creutzfeldtâ€“Jakob Disease Using Nasal Brushings. New England Journal of Medicine, 2014, 371, 519-529.	27.0	291
7	Determinants of diagnostic investigation sensitivities across the clinical spectrum of sporadic Creutzfeldt-Jakob disease. Brain, 2006, 129, 2278-2287.	7.6	283
8	Iatrogenic Creutzfeldt-Jakob Disease, Final Assessment. Emerging Infectious Diseases, 2012, 18, 901-907.	4.3	280
9	Predictors of survival in sporadic Creutzfeldt-Jakob disease and other human transmissible spongiform encephalopathies. Brain, 2004, 127, 2348-2359.	7.6	244
10	Incidence and spectrum of sporadic Creutzfeldtâ€“Jakob disease variants with mixed phenotype and co-occurrence of PrPSc types: an updated classification. Acta Neuropathologica, 2009, 118, 659-671.	7.7	224
11	Codon 129 prion protein genotype and sporadic Creutzfeldt-Jakob disease. Lancet, The, 1999, 353, 1673-1674.	13.7	203
12	Efficient Transmission and Characterization of Creutzfeldtâ€“Jakob Disease Strains in Bank Voles. PLoS Pathogens, 2006, 2, e12.	4.7	201
13	Phenotypic Variability of Gerstmann-Straussler-Scheinker Disease is Associated with Prion Protein Heterogeneity. Journal of Neuropathology and Experimental Neurology, 1998, 57, 979-988.	1.7	182
14	Diagnosis of Human Prion Disease Using Real-Time Quaking-Induced Conversion Testing of Olfactory Mucosa and Cerebrospinal Fluid Samples. JAMA Neurology, 2017, 74, 155.	9.0	176
15	Amphotericin B treatment dissociates in vivo replication of the scrapie agent from PrP accumulation. Nature, 1992, 356, 598-601.	27.8	166
16	Doxycycline in Creutzfeldt-Jakob disease: a phase 2, randomised, double-blind, placebo-controlled trial. Lancet Neurology, The, 2014, 13, 150-158.	10.2	157
17	Descriptive epidemiology of Creutzfeldtâ€Jakob disease in six european countries, 1993â€“1995. Annals of Neurology, 1998, 43, 763-767.	5.3	154
18	Biomarkers and diagnostic guidelines for sporadic Creutzfeldt-Jakob disease. Lancet Neurology, The, 2021, 20, 235-246.	10.2	151

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19	Detection of Pathologic Prion Protein in the Olfactory Epithelium in Sporadic Creutzfeldtâ€“Jakob Disease. New England Journal of Medicine, 2003, 348, 711-719.	27.0	142
20	Cerebrospinal fluid biomarker supported diagnosis of Creutzfeldtâ€“Jakob disease and rapid dementias: a longitudinal multicentre study over 10 years. Brain, 2012, 135, 3051-3061.	7.6	135
21	Prion-specific and surrogate CSF biomarkers in Creutzfeldt-Jakob disease: diagnostic accuracy in relation to molecular subtypes and analysis of neuropathological correlates of p-tau and AÎ²42 levels. Acta Neuropathologica, 2017, 133, 559-578.	7.7	129
22	Pre-symptomatic detection of prions by cyclic amplification of protein misfolding. FEBS Letters, 2005, 579, 638-642.	2.8	127
23	Cerebrospinal fluid realâ€time quakingâ€induced conversion is a robust and reliable test for sporadic creutzfeldtâ€“jakob disease: An international study. Annals of Neurology, 2016, 80, 160-165.	5.3	107
24	Advanced tests for early and accurate diagnosis of Creutzfeldtâ€“Jakob disease. Nature Reviews Neurology, 2016, 12, 325-333.	10.1	105
25	A new point mutation of the prion protein gene in Creutzfeldtâ€Jakob disease. Annals of Neurology, 1993, 34, 802-807.	5.3	104
26	Tissue Handling in Suspected Creutzfeldtâ€Jakob Disease (CJD) and Other Human Spongiform Encephalopathies (Prion Diseases). Brain Pathology, 1995, 5, 319-322.	4.1	103
27	Increased Brain Synthesis of Prostaglandin E₂and F₂-Isoprostane in Human and Experimental Transmissible Spongiform Encephalopathies. Journal of Neuropathology and Experimental Neurology, 2000, 59, 866-871.	1.7	96
28	Transmission of Creutzfeldt-Jakob Disease by Dural Cadaveric Graft. Journal of Neurosurgery, 1989, 71, 954-5.	1.6	92
29	Î±â€Synuclein RTâ€QuIC assay in cerebrospinal fluid of patients with dementia with Lewy bodies. Annals of Clinical and Translational Neurology, 2019, 6, 2120-2126.	3.7	87
30	Ancestral Origins and Worldwide Distribution of the PRNP 200K Mutation Causing Familial Creutzfeldt-Jakob Disease. American Journal of Human Genetics, 1999, 64, 1063-1070.	6.2	85
31	Transmission of the 263K scrapie strain by the dental route. Journal of General Virology, 1999, 80, 3043-3047.	2.9	83
32	Ultra-high-pressure inactivation of prion infectivity in processed meat: A practical method to prevent human infection. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 6093-6097.	7.1	82
33	Polymorphisms of the prion protein gene in Italian patients with Creutzfeldt-Jakob disease. Human Genetics, 1994, 94, 375-9.	3.8	80
34	Extended and direct evaluation of <scp>RT</scp>â€Qu<scp>IC</scp> assays for Creutzfeldtâ€Jakob disease diagnosis. Annals of Clinical and Translational Neurology, 2017, 4, 139-144.	3.7	79
35	Iatrogenic Creutzfeldt-Jakob disease with Amyloid-Î² pathology: an international study. Acta Neuropathologica Communications, 2018, 6, 5.	5.2	79
36	Chronic wasting disease and atypical forms of bovine spongiform encephalopathy and scrapie are not transmissible to mice expressing wild-type levels of human prion protein. Journal of General Virology, 2012, 93, 1624-1629.	2.9	78

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37	Cerebrospinal fluid biomarkers in human genetic transmissible spongiform encephalopathies. Journal of Neurology, 2009, 256, 1620-1628.	3.6	77
38	Prions and related neurological diseases. Molecular Aspects of Medicine, 1994, 15, 195-291.	6.4	75
39	Trapping Prion Protein in the Endoplasmic Reticulum Impairs PrPC Maturation and Prevents PrPSc Accumulation. Journal of Biological Chemistry, 2005, 280, 685-694.	3.4	72
40	Influence of timing on CSF tests value for Creutzfeldt-Jakob disease diagnosis. Journal of Neurology, 2007, 254, 901-906.	3.6	72
41	High incidence of genetic human transmissible spongiform encephalopathies in Italy. Neurology, 2005, 64, 1592-1597.	1.1	70
42	The importance of ongoing international surveillance for Creutzfeldtâ€“Jakob disease. Nature Reviews Neurology, 2021, 17, 362-379.	10.1	69
43	Clinical and neuropathological phenotype associated with the novel V189I mutation in the prion protein gene. Acta Neuropathologica Communications, 2019, 7, 1.	5.2	68
44	Revisiting the Heidenhain Variant of Creutzfeldt-Jakob Disease: Evidence for Prion Type Variability Influencing Clinical Course and Laboratory Findings. Journal of Alzheimer's Disease, 2016, 50, 465-476.	2.6	65
45	A novel PSEN2 mutation associated with a peculiar phenotype. Neurology, 2008, 70, 1549-1554.	1.1	62
46	Levels of infectivity in the blood throughout the incubation period of hamsters peripherally injected with scrapie. Archives of Virology, 1989, 108, 145-149.	2.1	60
47	Failure to Ameliorate Creutzfeldt-Jakob Disease with Amphotericin B Therapy. Journal of Infectious Diseases, 1992, 165, 784-785.	4.0	60
48	Polymorphism at codon 129 or codon 219 of PRNP and clinical heterogeneity in a previously unreported family with Gerstmann-Straussler-Scheinker disease (PrP-P102L mutation). Neurology, 1996, 47, 734-741.	1.1	60
49	Identification of the prion protein allotypes which accumulate in the brain of sporadic and familial Creutzfeldt-Jakob disease patients. Nature Medicine, 1997, 3, 521-525.	30.7	58
50	Regulation of intrinsic prion protein by growth factors and tnf-Î±: the role of intracellular reactive oxygen species. Free Radical Biology and Medicine, 2003, 35, 586-594.	2.9	54
51	Amphotericin B: A Novel Class of Antiscrapie Drugs. Journal of Infectious Diseases, 1989, 160, 795-802.	4.0	53
52	Increased CSF levels of prostaglandin E ₂ in variant Creutzfeldtâ€“Jakob disease. Neurology, 2002, 58, 127-129.	1.1	51
53	Transmission of sporadic Creutzfeldtâ€Jakob disease by blood transfusion: risk factor or possible biases. Transfusion, 2011, 51, 1556-1566.	1.6	51
54	Highly Infectious Purified Preparations of Disease-Specific Amyloid of Transmissible Spongiform Encephalopathies Are Not Devoid of Nucleic Acids of Viral Size. Intervirology, 1997, 40, 238-246.	2.8	49

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55	Sublethal Doses of Î²-Amyloid Peptide Abrogate DNA-dependent Protein Kinase Activity. Journal of Biological Chemistry, 2012, 287, 2618-2631.	3.4	49
56	Proteinase-resistant protein in human neuroblastoma cells infected with brain material from Creutzfeldt-Jakob patient. Lancet, The, 1995, 345, 594-595.	13.7	46
57	Standards for the assay of Creutzfeldtâ€“Jakob disease specimens. Journal of General Virology, 2004, 85, 1777-1784.	2.9	46
58	Combination Ultrafiltration and 6 MUrea Treatment of Human Growth Hormone Effectively Minimizes Risk from Potential Creutzfeldt-Jakob Disease Virus Contamination. Hormone Research, 1991, 35, 161-166.	1.8	44
59	KDEL-tagged anti-prion intrabodies impair PrP lysosomal degradation and inhibit scrapie infectivity. Biochemical and Biophysical Research Communications, 2005, 338, 1791-1797.	2.1	44
60	Proteomic profiling of PrP27â€30â€enriched preparations extracted from the brain of hamsters with experimental scrapie. Proteomics, 2009, 9, 3802-3814.	2.2	43
61	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
62	Identification of the pathological prion protein allotypes in scrapie-infected heterozygous bank voles (Clethrionomys glareolus) by high-performance liquid chromatographyâ€“mass spectrometry. Journal of Chromatography A, 2005, 1081, 122-126.	3.7	41
63	Cyclooxygenaseâ€2, Prostaglandin E2, and Microglial Activation in Prion Diseases. International Review of Neurobiology, 2007, 82, 265-275.	2.0	41
64	Two-dimensional mapping of three phenotype-associated isoforms of the prion protein in sporadic Creutzfeldt-Jakob disease. Electrophoresis, 2002, 23, 347-355.	2.4	40
65	Apolipoprotein E in sporadic and familial Creutzfeldt-Jakob disease. Neuroscience Letters, 1995, 199, 95-98.	2.1	39
66	Migration of dendritic cells into the brain in a mouse model of prion disease. Journal of Neuroimmunology, 2005, 165, 114-120.	2.3	39
67	Molecular diagnostics of transmissible spongiform encephalopathies. Trends in Molecular Medicine, 2002, 8, 273-280.	6.7	37
68	Detection of exosomal prions in blood by immunochemistry techniques. Journal of General Virology, 2015, 96, 1969-1974.	2.9	37
69	Prion protein glycotype analysis in familial and sporadic Creutzfeldt-Jakob disease patients. Brain Research Bulletin, 1999, 49, 429-433.	3.0	36
70	Creutzfeldt-Jakob disease associated with the R208H mutation in the prion protein gene. Neurology, 2005, 64, 905-907.	1.1	36
71	Preparation of soluble infectious samples from scrapie-infected brain: a new tool to study the clearance of transmissible spongiform encephalopathy agents during plasma fractionation. Transfusion, 2006, 46, 652-658.	1.6	36
72	Novel Prion Protein Conformation and Glycotype in Creutzfeldt-Jakob Disease. Archives of Neurology, 2007, 64, 595.	4.5	36

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73	Incidence of Creutzfeldt-Jakob disease In Europe in 1993. Lancet, The, 1994, 343, 918.	13.7	34
74	Comparison between plasma and cerebrospinal fluid biomarkers for the early diagnosis and association with survival in prion disease. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 1181-1188.	1.9	34
75	Creutzfeldt-Jakob disease in Europe. Lancet, The, 1995, 346, 898.	13.7	33
76	High incidence of Creutzfeldt-Jakob disease in rural Calabria, Italy. Lancet, The, 1998, 352, 1989-1990.	13.7	33
77	Epidemic of transmissible spongiform encephalopathy in sheep and goats in Italy. Lancet, The, 1999, 353, 560-561.	13.7	33
78	Towards an early clinical diagnosis of sporadic CJD VV2 (ataxic type). Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, 764-772.	1.9	33
79	Creutzfeldt-Jakob disease after non-commercial dura mater graft. Lancet, The, 1992, 340, 614-615.	13.7	32
80	Human transmissible spongiform encephalopathies in eleven countries: diagnostic pattern across time, 1993â€“2002. BMC Public Health, 2006, 6, 278.	2.9	28
81	A Genome Wide Association Study Links Glutamate Receptor Pathway to Sporadic Creutzfeldt-Jakob Disease Risk. PLoS ONE, 2015, 10, e0123654.	2.5	28
82	Validation of Revised International Creutzfeldt-Jakob Disease Surveillance Network Diagnostic Criteria for Sporadic Creutzfeldt-Jakob Disease. JAMA Network Open, 2022, 5, e2146319.	5.9	28
83	Sensitivity to Biases of Case-Control Studies on Medical Procedures, Particularly Surgery and Blood Transfusion, and Risk of Creutzfeldt-Jakob Disease. Neuroepidemiology, 2012, 39, 1-18.	2.3	27
84	Ring trial of 2nd generation RTâ€QuIC diagnostic tests for sporadic CJD. Annals of Clinical and Translational Neurology, 2020, 7, 2262-2271.	3.7	27
85	Pathological prion protein in muscles of hamsters and mice infected with rodent-adapted BSE or vCJD. Journal of General Virology, 2006, 87, 251-254.	2.9	26
86	Detection of proteinase-resistant protein (PrP) in small brain tissue samples from Creutzfeldt-Jakob disease patients. Journal of the Neurological Sciences, 1994, 124, 171-173.	0.6	25
87	Neuroinvasion of the 263K scrapie strain after intranasal administration occurs through olfactory-unrelated pathways. Acta Neuropathologica, 2009, 117, 175-184.	7.7	25
88	Fatal familial insomnia in a new Italian kindred. Neurology, 1998, 51, 1491-1494.	1.1	24
89	Variant Creutzfeldt-Jakob disease in an Italian woman. Lancet, The, 2002, 360, 997-998.	13.7	24
90	Transmission of CJD from nasal brushings but not spinal fluid or RTâ€QuIC product. Annals of Clinical and Translational Neurology, 2020, 7, 932-944.	3.7	23

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91	Assessing Prion Infectivity of Human Urine in Sporadic Creutzfeldt-Jakob Disease. Emerging Infectious Diseases, 2012, 18, 21-28.	4.3	22
92	Mortality trend from sporadic Creutzfeldt-Jakob disease (CJD) in Italy, 1993â€“2000. Journal of Clinical Epidemiology, 2003, 56, 494-499.	5.0	21
93	Variable Phenotype in a P102L Gerstmann-StrÃussler-Scheinker Italian Family. Canadian Journal of Neurological Sciences, 2003, 30, 233-236.	0.5	21
94	Inactivation of transmissible spongiform encephalopathy agents in food products by ultra high pressureâ€“temperature treatment. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2006, 1764, 558-562.	2.3	21
95	A retrospective study of Creutzfeldt-Jakob disease in Italy (1972?1986). European Journal of Epidemiology, 1988, 4, 482-487.	5.7	20
96	Cerebrospinal Fluid Total Prion Protein in the Spectrum of Prion Diseases. Molecular Neurobiology, 2019, 56, 2811-2821.	4.0	20
97	Small virus-like structure in brains from cases of sporadic and familial Creutzfeldt-Jakob disease. Lancet, The, 1994, 344, 923-924.	13.7	19
98	SPORADIC FATAL INSOMNIA IN A FATAL FAMILIAL INSOMNIA PEDIGREE. Neurology, 2008, 70, 884-885.	1.1	18
99	Codon 129 polymorphism of prion protein gene in sporadic Alzheimerâ€™s disease. European Journal of Neurology, 2008, 15, 173-178.	3.3	17
100	The pathological prion protein forms ionic conductance in lipid bilayer. Neurochemistry International, 2011, 59, 168-174.	3.8	17
101	Neurofilaments in blood is a new promising preclinical biomarker for the screening of natural scrapie in sheep. PLoS ONE, 2019, 14, e0226697.	2.5	17
102	Can potential hazard of Creutzfeldt-Jakob disease infectivity be reduced in the production of human Growth Hormone?. Archives of Virology, 1988, 98, 131-135.	2.1	16
103	Characterisation of antisera raised against species-specific peptide sequences from scrapie-associated fibril protein and their application for post-mortem immunodiagnosis of spongiform encephalopathies. Archives of Virology, 1994, 136, 99-110.	2.1	16
104	Mutation of thePRNP gene at codon 211 in familial Creutzfeldt-Jakob disease. American Journal of Medical Genetics Part A, 2001, 103, 133-137.	2.4	16
105	Quantitative profiling of the pathological prion protein allotypes in bank voles by liquid chromatographyâ€“mass spectrometry. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2007, 849, 302-306.	2.3	16
106	Oral pravastatin prolongs survival time of scrapie-infected mice. Journal of General Virology, 2009, 90, 1775-1780.	2.9	16
107	Diagnostic accuracy of cerebrospinal fluid biomarkers in genetic prion diseases. Brain, 2022, 145, 700-712.	7.6	16
108	Peripheral neuropathy in the course of progressive systemic sclerosis: Light and ultrastructural study. Italian Journal of Neurological Sciences, 1982, 3, 341-348.	0.1	15

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109	Molecular forms of cholinesterases in CSF of Alzheimer's disease/senile dementia of Alzheimer type patients and matched neurological controls. Life Sciences, 1986, 38, 561-567.	4.3	15
110	The scrapie agent and the prion hypothesis. Trends in Biochemical Sciences, 1988, 13, 309-313.	7.5	15
111	Sporadic<scp>C</scp>reutzfeldtâ€“<scp>J</scp>akob disease subtypeâ€specific alterations of the brain proteome: Impact on<scp>R</scp>ab3a recycling. Proteomics, 2012, 12, 3610-3620.	2.2	15
112	Experimental drug treatment of scrapie: A pathogenetic basis for rationale therapeutics. European Journal of Epidemiology, 1991, 7, 556-561.	5.7	14
113	Scrapie infectivity is quickly cleared in tissues of orally-infected farmed fish. BMC Veterinary Research, 2006, 2, 21.	1.9	14
114	Comparison of nanofiltration efficacy in reducing infectivity of centrifuged versus ultracentrifuged 263K scrapieâ€infected brain homogenates in â€œspikedâ€albumin solutions. Transfusion, 2012, 52, 953-962.	1.6	14
115	Prion Strain Characterization of a Novel Subtype of Creutzfeldt-Jakob Disease. Journal of Virology, 2017, 91, .	3.4	14
116	Age at onset of genetic (E200K) and sporadic Creutzfeldt-Jakob diseases is modulated by the CYP4X1 gene. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 1243-1249.	1.9	14
117	Heidenhain variant in two patients with inherited V210I Creutzfeldt-Jakob disease. International Journal of Neuroscience, 2016, 126, 381-3.	1.6	14
118	A rapid and efficient method for the detection of point mutations of the human prion protein gene (PRNP) by direct sequencing. Journal of Neuroscience Methods, 2000, 99, 59-63.	2.5	13
119	Measurement of the concentration of amphotericin B in brain tissue of scrapie-infected hamsters with a simple and sensitive method. Antimicrobial Agents and Chemotherapy, 1991, 35, 1486-1488.	3.2	12
120	<scp>G</scp>erstmannâ€<scp>S</scp>trÃusslerâ€<scp>S</scp>cheinker Syndrome with Variable Phenotype in a New Kindred with <scp>PRNP</scp>â€<scp>P102L</scp> Mutation. Brain Pathology, 2014, 24, 142-147.	4.1	12
121	Synthetic Scrapie Infectivity: Interaction between Recombinant PrP and Scrapie Brain-Derived RNA. Virulence, 2015, 6, 132-144.	4.4	12
122	Diagnosis of Creutzfeldt-Jakob disease. BMJ: British Medical Journal, 1999, 318, 538-538.	2.3	12
123	Choline acetyltransferase activity and [3H]quinuclidinylbenzilate binding in brains of scrapie-infected hamsters. Neuroscience Letters, 1984, 51, 87-92.	2.1	11
124	Expression of wild-type and V210I mutant prion protein in human neuroblastoma cells. Neuroscience Letters, 1999, 270, 41-44.	2.1	11
125	A role for complement in transmissible spongiform encephalopathies. Nature Medicine, 2001, 7, 410-411.	30.7	11
126	Survival in Alzheimerâ€™s Disease Is Shorter in Women Carrying Heterozygosity at Codon 129 of the <i>PRNP</i> Gene and No APOE Îµ4 Allele. Dementia and Geriatric Cognitive Disorders, 2008, 25, 354-358.	1.5	11

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127	Age at Death of Creutzfeldt-Jakob Disease in Subsequent Family Generation Carrying the E200K Mutation of the Prion Protein Gene. PLoS ONE, 2013, 8, e60376.	2.5	11
128	The future for treating Creutzfeldtâ€“Jakob disease. Expert Opinion on Orphan Drugs, 2015, 3, 57-74.	0.8	11
129	Efficacy of phthalocyanine tetrasulfonate against mouse-adapted human prion strains. Archives of Virology, 2009, 154, 1005-1007.	2.1	10
130	Rethinking of doxycycline therapy in Creutzfeldt-Jakob disease. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 705-705.	1.9	10
131	Diagnostic Accuracy of Prion Disease Biomarkers in Iatrogenic Creutzfeldt-Jakob Disease. Biomolecules, 2020, 10, 290.	4.0	10
132	Serotoninergic System in Scrapie-Infected Hamsters. Journal of Neurochemistry, 1985, 44, 862-868.	3.9	9
133	Isonicotinic hydrazide causes seizures in scrapie-infected hamstesr with shorter latency than in control animals: A possible GABAergic defect. Brain Research, 1985, 326, 117-123.	2.2	9
134	Plasma levels of 13,14-dihydro-15-keto PGE2 after vaginal application of a new PGE2 film. Prostaglandins, 1985, 29, 269-272.	1.2	9
135	Alpha1 antichymotrypsin signal peptide polymorphism in sporadic Creutzfeldtâ€“Jakob disease. Neuroscience Letters, 1997, 227, 140-142.	2.1	9
136	THE NUCLEUS BASALIS OF MEYNERT IN PARKINSONISMâ€“DEMENTIA OF GUAM: A MORPHOMETRIC STUDY. Neuropathology and Applied Neurobiology, 1989, 15, 193-206.	3.2	8
137	Recent Italian FFI Cases. Brain Pathology, 1998, 8, 564-566.	4.1	8
138	Creutzfeldt-Jakob disease: hopes for therapy. European Journal of Neurology, 2008, 15, 435-436.	3.3	8
139	Identification of Misfolded Proteins in Body Fluids for the Diagnosis of Prion Diseases. International Journal of Cell Biology, 2013, 2013, 1-10.	2.5	8
140	Subtype-Specific Synaptic Proteome Alterations in Sporadic Creutzfeldt-Jakob Disease. Journal of Alzheimer's Disease, 2013, 37, 51-61.	2.6	8
141	Assessment of prion reduction filters in decreasing infectivity of ultracentrifuged 263<scp>K</scp> scrapieâ€infected brain homogenates in â€œspikedâ€•human blood and red blood cells. Transfusion, 2014, 54, 990-995.	1.6	8
142	Evaluation of Human Cerebrospinal Fluid Malate Dehydrogenase 1 as a Marker in Genetic Prion Disease Patients. Biomolecules, 2019, 9, 800.	4.0	8
143	Creutzfeldt-Jakob Disease Mortality in Italy, 1982â€“1996. Neuroepidemiology, 1999, 18, 92-100.	2.3	7
144	Prion protein allotype profiling by mass spectrometry. Pure and Applied Chemistry, 2003, 75, 317-323.	1.9	7

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145	Prion (PrPres) Allotypes Profiling: New Perspectives from Mass Spectrometry. European Journal of Mass Spectrometry, 2004, 10, 371-382.	1.0	7
146	Prevalence of variant CJD in the UK. BMJ: British Medical Journal, 2009, 338, b435-b435.	2.3	7
147	Increased levels of acute-phase inflammatory proteins in plasma of patients with sporadic CJD. Neurology, 2012, 79, 1012-1018.	1.1	7
148	Pathogenesis and Transmission of Classical and Atypical BSE in Cattle. Food Safety (Tokyo, Japan), 2016, 4, 130-134.	1.8	7
149	Codon 200 mutation in a new family of Chilean origin with Creutzfeldt-Jakob disease.. Journal of Neurology, Neurosurgery and Psychiatry, 1996, 61, 111-112.	1.9	6
150	Genomic and post-genomic analyses of human prion diseases. Genome Medicine, 2009, 1, 63.	8.2	6
151	Role of proteomics in understanding prion infection. Expert Review of Proteomics, 2012, 9, 649-666.	3.0	6
152	Mutant PrPCJD prevails over wild-type PrPCJD in the brain of V210I and R208H genetic Creutzfeldtâ€“Jakob disease patients. Biochemical and Biophysical Research Communications, 2014, 454, 289-294.	2.1	6
153	Spatial Epidemiology of Sporadic Creutzfeldt-Jakob Disease in Apulia, Italy. Neuroepidemiology, 2020, 54, 83-90.	2.3	6
154	Ultrastructural studies in synaptic formations in dissociated fetal mouse brain cultures. Neuroscience Letters, 1983, 43, 127-130.	2.1	5
155	Progressive Dementia in a Young Patient with a Homozygous Deletion of the PrP Gene.. Annals of the New York Academy of Sciences, 1994, 724, 358-360.	3.8	5
156	BSE and variant Creutzfeldt-Jakob disease: never say never. Acta Neuropathologica, 2002, 103, 627-628.	7.7	5
157	Patient with rapidly evolving neurological disease with neuropathological lesions of Creutzfeldtâ€Jakob disease, Lewy body dementia, chronic subcortical vascular encephalopathy and meningothelial meningioma. Neuropathology, 2017, 37, 110-115.	1.2	5
158	Creutzfeldt-Jakob disease in the province of Siena: Two cases transmitted to monkeys. Italian Journal of Neurological Sciences, 1983, 4, 61-64.	0.1	4
159	Immunodiagnosis of bovine spongiform encephalopathy. Livestock Science, 1994, 38, 41-46.	1.2	4
160	Mortality from Human Transmissible Spongiform Encephalopathies: A Record Linkage Study. Neuroepidemiology, 2005, 24, 214-220.	2.3	4
161	Clinical trials and methodological problems in prion diseases. Lancet Neurology, The, 2009, 8, 782.	10.2	3
162	Clinicopathological features of the rare form of Creutzfeldt-Jakob disease in R208H-V129V PRNP carrier. Acta Neuropathologica Communications, 2019, 7, 47.	5.2	3

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163	Need to improve clinical trials in rare neurodegenerative disorders. Annali Dell'Istituto Superiore Di Sanita, 2011, 47, 55-9.	0.4	3
164	PrPTSE in muscle-associated lymphatic tissue during the preclinical stage of mice infected orally with bovine spongiform encephalopathy. Journal of General Virology, 2009, 90, 2563-2568.	2.9	2
165	Management and prevention of human prion diseases. Current Neurology and Neuroscience Reports, 2009, 9, 423-429.	4.2	2
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