Hermann M Schätzl

List of Publications by Year in descending order

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127 papers 13,030 citations

94433 37 h-index 23533 111 g-index

135 all docs 135
docs citations

135 times ranked 23992 citing authors

#	Article	IF	Citations
1	Guidelines for the use and interpretation of assays for monitoring autophagy (3rd edition). Autophagy, 2016, 12, 1-222.	9.1	4,701
2	Guidelines for the use and interpretation of assays for monitoring autophagy. Autophagy, 2012, 8, 445-544.	9.1	3,122
3	Analysis of 27 mammalian and 9 avian PrPs reveals high conservation of flexible regions of the prion protein 1 1Edited by A. R. Fersht. Journal of Molecular Biology, 1999, 289, 1163-1178.	4.2	382
4	Prion Protein Gene Variation Among Primates. Journal of Molecular Biology, 1995, 245, 362-374.	4.2	309
5	The anticancer drug imatinib induces cellular autophagy. Leukemia, 2007, 21, 936-942.	7.2	208
6	In Vitro and In Vivo Neurotoxicity of Prion Protein Oligomers. PLoS Pathogens, 2007, 3, e125.	4.7	201
7	Autophagy induction by trehalose counter-acts cellular prion-infection. Autophagy, 2009, 5, 361-369.	9.1	198
8	PrPC Directly Interacts with Proteins Involved in Signaling Pathways. Journal of Biological Chemistry, 2001, 276, 44604-44612.	3.4	185
9	Lithium induces clearance of protease resistant prion protein in prionâ€infected cells by induction of autophagy. Journal of Neurochemistry, 2009, 109, 25-34.	3.9	169
10	Individuals with antibodies against hepatitis B core antigen as the only serological marker for hepatitis B infection: high percentage of carriers of hepatitis B and C virus. Journal of Hepatology, 1995, 23, 14-20.	3.7	157
11	Intracellular re-routing of prion protein prevents propagation of PrPSc and delays onset of prion disease. EMBO Journal, 2001, 20, 3957-3966.	7.8	147
12	Prion-Protein-Specific Aptamer Reduces PrPSc Formation. ChemBioChem, 2002, 3, 717.	2.6	141
13	Severe Acute Respiratory Syndrome Coronavirus Replication Is Severely Impaired by MG132 due to Proteasome-Independent Inhibition of M-Calpain. Journal of Virology, 2012, 86, 10112-10122.	3.4	130
14	The Tyrosine Kinase Inhibitor STI571 Induces Cellular Clearance of PrPSc in Prion-infected Cells. Journal of Biological Chemistry, 2004, 279, 41918-41927.	3.4	114
15	Humoral immune response to native eukaryotic prion protein correlates with anti-prion protection. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 14670-14676.	7.1	105
16	Polyclonal Anti-PrP Auto-antibodies Induced with Dimeric PrP Interfere Efficiently with PrPSc Propagation in Prion-infected Cells. Journal of Biological Chemistry, 2003, 278, 18524-18531.	3.4	99
17	Essential Role of the Prion Protein N Terminus in Subcellular Trafficking and Half-life of Cellular Prion Protein. Journal of Biological Chemistry, 2003, 278, 3726-3734.	3.4	97
18	Autophagy, prion infection and their mutual interactions. Current Issues in Molecular Biology, 2010, 12, 87-97.	2.4	86

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19	Autophagy regulates exosomal release of prions in neuronal cells. Journal of Biological Chemistry, 2018, 293, 8956-8968.	3.4	82
20	Endemic chronic wasting disease causes mule deer population decline in Wyoming. PLoS ONE, 2017, 12, e0186512.	2.5	77
21	Chronic Wasting Disease. Topics in Current Chemistry, 2011, 305, 51-77.	4.0	69
22	Early detection of chronic wasting disease prions in urine of pre-symptomatic deer by real-time quaking-induced conversion assay. Prion, 2013, 7, 253-258.	1.8	68
23	The yeast Sup35NM domain propagates as a prion in mammalian cells. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 462-467.	7.1	65
24	Glycosylation Deficiency at Either One of the Two Glycan Attachment Sites of Cellular Prion Protein Preserves Susceptibility to Bovine Spongiform Encephalopathy and Scrapie Infections. Journal of Biological Chemistry, 2004, 279, 53306-53316.	3.4	62
25	The prion protein requires cholesterol for cell surface localization. Molecular and Cellular Neurosciences, 2006, 31, 346-353.	2.2	62
26	Chronic wasting disease: Emerging prions and their potential risk. PLoS Pathogens, 2017, 13, e1006619.	4.7	57
27	Analysis of non-infectious HIV particles produced in presence of HIV proteinase inhibitor. Archives of Virology, 1991, 120, 71-81.	2.1	55
28	Polylactide-Coglycolide Microspheres CoEncapsulating Recombinant Tandem Prion Protein with CpG-Oligonucleotide Break Self-Tolerance to Prion Protein in Wild-Type Mice and Induce CD4 and CD8 T Cell Responses. Journal of Immunology, 2007, 179, 2797-2807.	0.8	50
29	Proteasomal Dysfunction and Endoplasmic Reticulum Stress Enhance Trafficking of Prion Protein Aggregates through the Secretory Pathway and Increase Accumulation of Pathologic Prion Protein. Journal of Biological Chemistry, 2011, 286, 33942-33953.	3.4	50
30	Quantification of hepatitis B virus DNA over a wide range from serum for studying viral replicative activity in response to treatment and in recurrent infection. Hepatology, 1995, 21, 1492-1499.	7. 3	47
31	Is codon 129 of prion protein polymorphic in human beings but not in animals?. Lancet, The, 1997, 349, 1603-1604.	13.7	47
32	Cell Line Dependent RNA Expression Profiles of Prion-infected Mouse Neuronal Cells. Journal of Molecular Biology, 2005, 349, 487-500.	4.2	47
33	Cell-to-cell propagation of infectious cytosolic protein aggregates. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 5951-5956.	7.1	45
34	Prion-like propagation of cytosolic protein aggregates. Prion, 2009, 3, 206-212.	1.8	43
35	Vaccination with prion peptide-displaying papillomavirus-like particles induces autoantibodies to normal prion protein that interfere with pathologic prion protein production in infected cells. FEBS Journal, 2007, 274, 1747-1758.	4.7	42
36	The tyrosine kinase inhibitor imatinib mesylate delays prion neuroinvasion by inhibiting prion propagation in the periphery. Journal of NeuroVirology, 2007, 13, 328-337.	2.1	40

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37	Prion Diseases: From Molecular Biology to Intervention Strategies. ChemBioChem, 2003, 4, 1268-1284.	2.6	37
38	Cell Type-Specific Cleavage of Nucleocapsid Protein by Effector Caspases during SARS Coronavirus Infection. Journal of Molecular Biology, 2008, 376, 23-34.	4.2	37
39	Inhibition of cholesterol recycling impairs cellular PrPSc propagation. Cellular and Molecular Life Sciences, 2009, 66, 3979-3991.	5.4	37
40	Scrapie Infection of Prion Protein-deficient Cell Line upon Ectopic Expression of Mutant Prion Proteins. Journal of Biological Chemistry, 2007, 282, 18702-18710.	3.4	34
41	Concomitant Administration of a Virosome-Adjuvanted Hepatitis A Vaccine With Routine Childhood Vaccines at Age Twelve to Fifteen Months: A Randomized Controlled Trial. Pediatric Infectious Disease Journal, 2007, 26, 787-793.	2.0	34
42	Prion-induced Activation of Cholesterogenic Gene Expression by Srebp2 in Neuronal Cells. Journal of Biological Chemistry, 2009, 284, 31260-31269.	3.4	34
43	Detection of PrPSc in peripheral tissues of clinically affected cattle after oral challenge with bovine spongiform encephalopathy. Journal of General Virology, 2012, 93, 2740-2748.	2.9	34
44	Molecular basis of cerebral neurodegeneration in prion diseases. FEBS Journal, 2007, 274, 606-611.	4.7	33
45	Prion infection impairs lysosomal degradation capacity by interfering with rab7 membrane attachment in neuronal cells. Scientific Reports, 2016, 6, 21658.	3.3	33
46	Charged bipolar suramin derivatives induce aggregation of the prion protein at the cell surface and inhibit PrPSc replication. Journal of Cell Science, 2005, 118, 4959-4973.	2.0	32
47	Prominent Stress Response of Purkinje Cells in Creutzfeldt–Jakob Disease. Neurobiology of Disease, 2001, 8, 881-889.	4.4	29
48	The Novel Sorting Nexin SNX33 Interferes with Cellular PrP ^{Sc} Formation by Modulation of PrP ^c Shedding. Traffic, 2008, 9, 1116-1129.	2.7	29
49	Prion strains depend on different endocytic routes for productive infection. Scientific Reports, 2017, 7, 6923.	3.3	29
50	Cervid Prion Protein Polymorphisms: Role in Chronic Wasting Disease Pathogenesis. International Journal of Molecular Sciences, 2021, 22, 2271.	4.1	29
51	Toxic Effects of Intracerebral PrP Antibody Administration During the Course of BSE Infection in Mice. Prion, 2007, 1, 198-206.	1.8	28
52	Aptamers against prion proteins and prions. Cellular and Molecular Life Sciences, 2009, 66, 2445-2455.	5.4	28
53	Evaluation of Modified Vaccinia Virus Ankara as an Alternative Vaccine against Smallpox in Chronically HIV Type 1-Infected Individuals Undergoing HAART. AIDS Research and Human Retroviruses, 2007, 23, 782-793.	1.1	27
54	Ultra-sensitive detection of prion protein fibrils by flow cytometry in blood from cattle affected with bovine spongiform encephalopathy. BMC Biotechnology, 2005, 5, 26.	3.3	26

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55	Peptide Aptamers Expressed in the Secretory Pathway Interfere with Cellular PrPSc Formation. Journal of Molecular Biology, 2007, 371, 362-373.	4.2	25
56	Critical Significance of the Region between Helix 1 and 2 for Efficient Dominant-Negative Inhibition by Conversion-Incompetent Prion Protein. PLoS Pathogens, 2013, 9, e1003466.	4.7	25
57	LRP/LR Antibody Mediated Rescuing of Amyloid-β-Induced Cytotoxicity is Dependent on PrPc in Alzheimer's Disease. Journal of Alzheimer's Disease, 2015, 49, 645-657.	2.6	24
58	The celecoxib derivatives AR-12 and AR-14 induce autophagy and clear prion-infected cells from prions. Scientific Reports, 2017, 7, 17565.	3.3	24
59	A prime-boost vaccination protocol optimizes immune responses against the nucleocapsid protein of the SARS coronavirus. Vaccine, 2008, 26, 6678-6684.	3.8	23
60	Autophagy pathways in the treatment of prion diseases. Current Opinion in Pharmacology, 2019, 44, 46-52.	3.5	22
61	The first B/G intersubtype recombinant form of human immunodeficiency virus type 1 (HIV-1) identified in Germany was undetected or underquantitated by some commercial viral load assays. Journal of Medical Virology, 2006, 78, 311-317.	5.0	21
62	CpG and LPS can interfere negatively with prion clearance in macrophage and microglial cells. FEBS Journal, 2007, 274, 5834-5844.	4.7	21
63	From Highâ€Throughput Cell Culture Screening to Mouse Model: Identification of New Inhibitor Classes against Prion Disease. ChemMedChem, 2011, 6, 1928-1937.	3.2	21
64	Combining autophagy stimulators and cellulose ethers for therapy against prion disease. Prion, 2019, 13, 185-196.	1.8	21
65	Neurotrophic factors: ready to go?. Trends in Neurosciences, 1995, 18, 463-464.	8.6	20
66	Vaccination against hepatitis A: comparison of different short-term immunization schedules. Vaccine, 1992, 10, S126-S128.	3.8	19
67	Recognition of Lumenal Prion Protein Aggregates by Post-ER Quality Control Mechanisms Is Mediated by the Preoctarepeat Region of PrP. Traffic, 2004, 5, 300-313.	2.7	19
68	Therapy in Prion Diseases: From Molecular and Cellular Biology to Therapeutic Targets. Infectious Disorders - Drug Targets, 2009, 9, 3-14.	0.8	19
69	Stability and conformational properties of doppel, a prion-like protein, and its single-disulphide mutant. Biochemical Journal, 2003, 373, 485-494.	3.7	18
70	Small-scale Triton X-114 Extraction of Hydrophobic Proteins. Bio-protocol, 2014, 4, .	0.4	18
71	An astrocyte cell line that differentially propagates murine prions. Journal of Biological Chemistry, 2020, 295, 11572-11583.	3.4	18
72	Ligands binding to the prion protein induce its proteolytic release with therapeutic potential in neurodegenerative proteinopathies. Science Advances, 2021, 7, eabj1826.	10.3	18

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73	Sequence Note: Phylogenetic Characterization of Simian T Lymphotropic Virus Type I (STLV-I) from the Ethiopian Sacred Baboon (Papio hamadryas). AIDS Research and Human Retroviruses, 1996, 12, 255-258.	1.1	17
74	Prion protein/protein interactions: fusion with yeast Sup35pâ€NM modulates cytosolic PrP aggregation in mammalian cells. FASEB Journal, 2008, 22, 762-773.	0.5	17
75	Gene-edited murine cell lines for propagation of chronic wasting disease prions. Scientific Reports, 2019, 9, 11151.	3.3	17
76	Elevated Epstein–Barr virus loads and lower antibody titers in competitive athletes. Journal of Medical Virology, 2010, 82, 446-451.	5.0	15
77	Modulation of Glycosaminoglycans Affects PrP Sc Metabolism but Does Not Block PrP Sc Uptake. Journal of Virology, 2015, 89, 9853-9864.	3.4	15
78	Promising developments bringing prion diseases closer to therapy and prophylaxis. Trends in Molecular Medicine, 2003, 9, 367-369.	6.7	14
79	Antibodies to a Nonconjugated Prion Protein Peptide 95-123 Interfere with PrP Sc Propagation in Prion-Infected Cells. Cellular and Molecular Neurobiology, 2007, 27, 271-284.	3.3	14
80	Immunization of cervidized transgenic mice with multimeric deer prion protein induces self-antibodies that antagonize chronic wasting disease infectivity in vitro. Scientific Reports, 2017, 7, 10538.	3.3	14
81	Recombinant prion protein vaccination of transgenic elk PrP mice and reindeer overcomes self-tolerance and protects mice against chronic wasting disease. Journal of Biological Chemistry, 2018, 293, 19812-19822.	3.4	14
82	Cellulose ether treatment <i>in vivo</i> generates chronic wasting disease prions with reduced protease resistance and delayed disease progression. Journal of Neurochemistry, 2020, 152, 727-740.	3.9	14
83	Prion infection influences murine endogenous retrovirus expression in neuronal cells. Biochemical and Biophysical Research Communications, 2006, 343, 825-831.	2.1	13
84	Strategies for eliminating PrPc as substrate for prion conversion and for enhancing PrPSc degradation. Veterinary Microbiology, 2007, 123, 377-386.	1.9	13
85	The octarepeat region of prion protein, but not the TM1 domain, is important for the antioxidant effect of prion protein. Free Radical Biology and Medicine, 2008, 45, 1622-1630.	2.9	13
86	A genome-wide survey for prion-regulated miRNAs associated with cholesterol homeostasis. BMC Genomics, 2012, 13, 486.	2.8	13
87	Dimerization of the cellular prion protein inhibits propagation of scrapie prions. Journal of Biological Chemistry, 2018, 293, 8020-8031.	3.4	13
88	Diphenylpyrazole-Derived Compounds Increase Survival Time of Mice after Prion Infection. Antimicrobial Agents and Chemotherapy, 2011, 55, 4774-4781.	3.2	12
89	Failure of Prion Protein Oxidative Folding Guides the Formation of Toxic Transmembrane Forms. Journal of Biological Chemistry, 2012, 287, 36693-36701.	3.4	12
90	Metformin reduces prion infection in neuronal cells by enhancing autophagy. Biochemical and Biophysical Research Communications, 2020, 523, 423-428.	2.1	12

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91	From Seeds to Fibrils and Back: Fragmentation as an Overlooked Step in the Propagation of Prions and Prion-Like Proteins. Biomolecules, 2020, 10, 1305.	4.0	12
92	Astrocyte in prion disease: a double-edged sword. Neural Regeneration Research, 2022, 17, 1659.	3.0	12
93	Targeting prion proteins in neurodegenerative disease. Expert Opinion on Biological Therapy, 2008, 8, 923-940.	3.1	11
94	Overexpression of quality control proteins reduces prion conversion in prion-infected cells. Journal of Biological Chemistry, 2018, 293, 16069-16082.	3.4	11
95	Disulfide-crosslink scanning reveals prion–induced conformational changes and prion strain–specific structures of the pathological prion protein PrPSc. Journal of Biological Chemistry, 2018, 293, 12730-12740.	3.4	11
96	Sephin1 Reduces Prion Infection in Prion-Infected Cells and Animal Model. Molecular Neurobiology, 2020, 57, 2206-2219.	4.0	11
97	Introducing a Rigid Loop Structure from Deer into Mouse Prion Protein Increases Its Propensity for Misfolding In Vitro. PLoS ONE, 2013, 8, e66715.	2.5	11
98	Dynamic interactions of Sup35p and PrP prion protein domains modulate aggregate nucleation and seeding. Prion, 2008, 2, 99-106.	1.8	10
99	Genomic Characterization of a Novel HIV Type 1 B/G Intersubtype Recombinant Strain from an Injecting Drug User in Germany. AIDS Research and Human Retroviruses, 2005, 21, 654-660.	1.1	9
100	Therapeutic vaccination reduces HIV sequence variability. FASEB Journal, 2008, 22, 437-444.	0.5	9
101	Isolated norovirus GII.7 strain within an extended GII.4 outbreak. Journal of Medical Virology, 2010, 82, 1058-1064.	5.0	9
102	Piperazine derivatives inhibit PrP/PrPres propagation in vitro and in vivo. Biochemical and Biophysical Research Communications, 2014, 445, 23-29.	2.1	9
103	Early detection of prion protein aggregation with a fluorescent pentameric oligothiophene probe using spectral confocal microscopy. Journal of Neurochemistry, 2021, 156, 1033-1048.	3.9	9
104	Variability of the hepatitis B surface protein in HBV-infected liver transplant recipients. Journal of Biomedical Science, 1997, 4, 146-154.	7.0	8
105	Fatal Epstein-Barr virus-associated lymphoproliferative disorder following treatment with a novel mTOR Inhibitor for relapsed chronic lymphocytic leukemia. Haematologica, 2007, 92, 1282-1283.	3.5	8
106	A Seroepidemiological Survey of Antibodies to HTLV-I/ HTLV-II in Selected Population Groups in Paraguay. Scandinavian Journal of Infectious Diseases, 1992, 24, 397-398.	1.5	7
107	Inhibition of Prion Amplification by Expression of Dominant Inhibitory Mutants - A Systematic Insertion Mutagenesis Study. Infectious Disorders - Drug Targets, 2009, 9, 40-47.	0.8	7
108	Preparation and Characterization of Cellulose Ether Liposomes for the Inhibition of Prion Formation in Prion-Infected Cells. Journal of Pharmaceutical Sciences, 2019, 108, 2814-2820.	3.3	7

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109	Prevalence of human T-Cell lymphotropie virus infections in Germany. Journal of Medical Virology, 1994, 43, 159-160.	5.0	6
110	Neuroendocrine cultured cells counteract persistent prion infection by down-regulation of PrPc. Molecular and Cellular Neurosciences, 2008, 38, 98-109.	2.2	6
111	Conditional Modulation of Membrane Protein Expression in Cultured Cells Mediated by Prion Protein Recognition of Short Phosphorothioate Oligodeoxynucleotides. Journal of Biological Chemistry, 2011, 286, 6911-6917.	3.4	6
112	Clearance of variant Creutzfeldt–Jakob disease prions <i>in vivo</i> by the Hsp70 disaggregase system. Brain, 2022, 145, 3236-3249.	7.6	6
113	The immune response to different doses of inactivated hepatitis A vaccine. Journal of Hepatology, 1993, 18, S38-S40.	3.7	5
114	Identifying critical sites of PrP ^c -PrP ^{Sc} interaction in prion-infected cells by dominant-negative inhibition. Prion, 2013, 7, 452-456.	1.8	5
115	Assessing Proteinase K Resistance of Fish Prion Proteins in a Scrapie-Infected Mouse Neuroblastoma Cell Line. Viruses, 2014, 6, 4398-4421.	3.3	5
116	Modulation of Host Cell Death by SARS Coronavirus Proteins. , 2010, , 231-245.		5
117	GABAA receptor subunit β1 is involved in the formation of proteaseâ€resistant prion protein in prionâ€infected neuroblastoma cells. FEBS Letters, 2010, 584, 1193-1198.	2.8	4
118	Polymorphisms in glia maturation factor \hat{l}^2 gene are markers of cellulose ether effectiveness in prion-infected mice. Biochemical and Biophysical Research Communications, 2021, 560, 105-111.	2.1	4
119	Small-scale Subcellular Fractionation with Sucrose Step Gradient. Bio-protocol, 2014, 4, .	0.4	2
120	Cellular Mechanisms of Propagation and Clearance. , 2013, , 147-160.		1
121	Variability of the Hepatitis B Surface Protein in HBV-Infected Liver T ransplant Recipients. Journal of Biomedical Science, 1997, 4, 146-154.	7. O	0
122	Parvovirus B19 and necrotizing enterocolitis in neonates. Journal of Pediatrics, 2012, 160, 887.	1.8	0
123	Prionen. , 2010, , 667-685.		0
124	Prions., 2013,, 919-947.		0
125	The autophagy inducers ARâ€12 and ARâ€14 control prion infection. FASEB Journal, 2018, 32, 795.5.	0.5	0
126	Insights into the Cellular Trafficking of Prion Proteins. , 2005, , 379-405.		0

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127	Gene-Edited Cell Models to Study Chronic Wasting Disease. Viruses, 2022, 14, 609.	3.3	0