Jan StÃ¶hr

List of Publications by Year
in descending order

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29	2,514	331670 21	477307 29
papers	citations	h-index	g-index

29	29	29	3582
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	AÎ² and tau prion-like activities decline with longevity in the Alzheimerâ€™s disease human brain. Science Translational Medicine, 2019, 11, .	12.4	96
2	Structural heterogeneity and intersubject variability of AÎ² in familial and sporadic Alzheimerâ€™s disease. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E782-E791.	7.1	105
3	A longâ€lived AÎ² oligomer resistant to fibrillization. Biopolymers, 2018, 109, e23096.	2.4	26
4	Zinc-binding structure of a catalytic amyloid from solid-state NMR. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 6191-6196.	7.1	102
5	A 31-residue peptide induces aggregation of tau's microtubule-binding region in cells. Nature Chemistry, 2017, 9, 874-881.	13.6	67
6	Structural Biology of PrP Prions. Cold Spring Harbor Perspectives in Medicine, 2017, 7, a024455.	6.2	2
7	Structural Polymorphism of Alzheimerâ€™s Î²-Amyloid Fibrils as Controlled by an E22 Switch: A Solid-State NMR Study. Journal of the American Chemical Society, 2016, 138, 9840-9852.	13.7	79
8	Mechanism of Scrapie Prion Precipitation with Phosphotungstate Anions. ACS Chemical Biology, 2015, 10, 1269-1277.	3.4	33
9	Structural Studies of Truncated Forms of the Prion Protein PrP. Biophysical Journal, 2015, 108, 1548-1554.	0.5	25
10	Prion Proteinâ€”Antibody Complexes Characterized by Chromatography-Coupled Small-Angle X-Ray Scattering. Biophysical Journal, 2015, 109, 793-805.	0.5	33
11	Propagation of prions causing synucleinopathies in cultured cells. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, E4949-58.	7.1	191
12	Truncated forms of the prion protein PrP demonstrate the need for complexity in prion structure. Prion, 2015, 9, 333-338.	1.8	2
13	Short peptides self-assemble to produce catalytic amyloids. Nature Chemistry, 2014, 6, 303-309.	13.6	510
14	Distinct synthetic AÎ² prion strains producing different amyloid deposits in bigenic mice. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 10329-10334.	7.1	140
15	Serial propagation of distinct strains of AÎ² prions from Alzheimerâ€™s disease patients. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 10323-10328.	7.1	247
16	Seeded Fibrillation as Molecular Basis of the Species Barrier in Human Prion Diseases. PLoS ONE, 2013, 8, e72623.	2.5	18
17	Prion Protein Aggregation and Fibrillogenesisln Vitro. Sub-Cellular Biochemistry, 2012, 65, 91-108.	2.4	5
18	Degradation of Fungal Prion HET-s(218-289) Induces Formation ofÂa Generic Amyloid Fold. Biophysical Journal, 2012, 102, 2339-2344.	0.5	24

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19	Spontaneous generation of rapidly transmissible prions in transgenic mice expressing wild-type bank vole prion protein. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 3498-3503.	7.1	65
20	Purified and synthetic Alzheimerâ€™s amyloid beta (AÎ²) prions. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 11025-11030.	7.1	327
21	In vitro conversion and seeded fibrillization of posttranslationally modified prion protein. Biological Chemistry, 2011, 392, 415-21.	2.5	6
22	Spontaneous generation of anchorless prions in transgenic mice. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 21223-21228.	7.1	68
23	Protease-Resistant Prions Selectively Decrease Shadoo Protein. PLoS Pathogens, 2011, 7, e1002382.	4.7	39
24	Molecular Interactions between Prions as Seeds and Recombinant Prion Proteins as Substrates Resemble the Biological Interspecies Barrier In Vitro. PLoS ONE, 2010, 5, e14283.	2.5	10
25	Spontaneous and BSE-prion-seeded amyloid formation of full length recombinant bovine prion protein. Biochemical and Biophysical Research Communications, 2008, 373, 493-497.	2.1	19
26	Mechanisms of prion protein assembly into amyloid. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 2409-2414.	7.1	127
27	Structural changes of membrane-anchored native PrP ^C. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 10815-10819.	7.1	83
28	Aggregation and Amyloid Fibril Formation of the Prion Protein Is Accelerated in the Presence of Glycogen. Rejuvenation Research, 2008, 11, 365-369.	1.8	6
29	Assembly of natural and recombinant prion protein into fibrils. Biological Chemistry, 2005, 386, 569-580.	2.5	59