

Mark Kindy

List of Publications by Year in descending order

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Version: 2024-02-01

15
papers

2,789
citations

567281

15
h-index

996975

15
g-index

15
all docs

15
docs citations

15
times ranked

3686
citing authors

#	ARTICLE	IF	CITATIONS
1	RAGE mediates amyloid- β peptide transport across the blood-brain barrier and accumulation in brain. <i>Nature Medicine</i> , 2003, 9, 907-913.	30.7	1,277
2	RAGE potentiates $\text{A}\beta$ -induced perturbation of neuronal function in transgenic mice. <i>EMBO Journal</i> , 2004, 23, 4096-4105.	7.8	311
3	Receptor-dependent cell stress and amyloid accumulation in systemic amyloidosis. <i>Nature Medicine</i> , 2000, 6, 643-651.	30.7	303
4	Inhibitors of Cathepsin B Improve Memory and Reduce β -Amyloid in Transgenic Alzheimer Disease Mice Expressing the Wild-type, but Not the Swedish Mutant, β -Secretase Site of the Amyloid Precursor Protein. <i>Journal of Biological Chemistry</i> , 2008, 283, 7745-7753.	3.4	185
5	Neprilysin Regulates Amyloid β Peptide Levels. <i>Journal of Molecular Neuroscience</i> , 2004, 22, 5-12.	2.3	141
6	Genetic cathepsin B deficiency reduces β -amyloid in transgenic mice expressing human wild-type amyloid precursor protein. <i>Biochemical and Biophysical Research Communications</i> , 2009, 386, 284-288.	2.1	97
7	The Cysteine Protease Inhibitor, E64d, Reduces Brain Amyloid- β and Improves Memory Deficits in Alzheimer's Disease Animal Models by Inhibiting Cathepsin B, but not BACE1, β -Secretase Activity. <i>Journal of Alzheimer's Disease</i> , 2011, 26, 387-408.	2.6	92
8	Cathepsin B is a New Drug Target for Traumatic Brain Injury Therapeutics: Evidence for E64d as a Promising Lead Drug Candidate. <i>Frontiers in Neurology</i> , 2015, 6, 178.	2.4	76
9	Brain Pyroglutamate Amyloid- β is Produced by Cathepsin B and is Reduced by the Cysteine Protease Inhibitor E64d, Representing a Potential Alzheimer's Disease Therapeutic. <i>Journal of Alzheimer's Disease</i> , 2014, 41, 129-149.	2.6	73
10	Cysteine Cathepsins in the secretory vesicle produce active peptides: Cathepsin L generates peptide neurotransmitters and cathepsin B produces beta-amyloid of Alzheimer's disease. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2012, 1824, 89-104.	2.3	67
11	Cysteine protease inhibitors effectively reduce in vivo levels of brain β -amyloid related to Alzheimer's disease. <i>Biological Chemistry</i> , 2007, 388, 247-52.	2.5	43
12	Pharmacogenetic features of cathepsin B inhibitors that improve memory deficit and reduce β -amyloid related to Alzheimer's disease. <i>Biological Chemistry</i> , 2010, 391, 861-72.	2.5	42
13	Cysteine protease inhibitors reduce brain β -amyloid and β -secretase activity <i>in vivo</i> and are potential Alzheimer's disease therapeutics. <i>Biological Chemistry</i> , 2007, 388, 979-983.	2.5	38
14	Cathepsin B Gene Knockout Improves Behavioral Deficits and Reduces Pathology in Models of Neurologic Disorders. <i>Pharmacological Reviews</i> , 2022, 74, 600-629.	16.0	29
15	Key signaling pathways regulate the biological activities and accumulation of amyloid- β . <i>Neurobiology of Aging</i> , 2001, 22, 967-973.	3.1	15