

Barbara Stiller

List of Publications by Year in descending order

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Version: 2024-02-01

8
papers

1,571
citations

1163117

8
h-index

1588992

8
g-index

8
all docs

8
docs citations

8
times ranked

2684
citing authors

#	ARTICLE	IF	CITATIONS
1	<i>Cis</i> -epistasis at the <i>LPA</i> locus and risk of cardiovascular diseases. <i>Cardiovascular Research</i> , 2022, 118, 1088-1102.	3.8	14
2	Interplay of pathogenic forms of human tau with different autophagic pathways. <i>Aging Cell</i> , 2018, 17, e12692.	6.7	148
3	Increased copper toxicity in <i>Saccharomyces cerevisiae</i> lacking <i>VPS35</i> , a component of the retromer and monogenic Parkinson disease gene in humans. <i>Biochemical and Biophysical Research Communications</i> , 2016, 476, 528-533.	2.1	9
4	Structural and Biological Interaction of hsc-70 Protein with Phosphatidylserine in Endosomal Microautophagy. <i>Journal of Biological Chemistry</i> , 2016, 291, 18096-18106.	3.4	52
5	±-Synuclein-Independent Histopathological and Motor Deficits in Mice Lacking the Endolysosomal Parkinsonism Protein <i>Atp13a2</i> . <i>Journal of Neuroscience</i> , 2015, 35, 5724-5742.	3.6	87
6	Pathogenic effects of novel mutations in the P-type ATPase <i>ATP13A2</i> (<i>PARK9</i>) causing Kufor-Rakeb syndrome, a form of early-onset parkinsonism. <i>Human Mutation</i> , 2011, 32, 956-964.	2.5	105
7	Cd ²⁺ , Mn ²⁺ , Ni ²⁺ and Se ²⁺ toxicity to <i>Saccharomyces cerevisiae</i> lacking <i>YPK9p</i> the orthologue of human <i>ATP13A2</i> . <i>Biochemical and Biophysical Research Communications</i> , 2009, 383, 198-202.	2.1	110
8	Hereditary parkinsonism with dementia is caused by mutations in <i>ATP13A2</i> , encoding a lysosomal type 5 P-type ATPase. <i>Nature Genetics</i> , 2006, 38, 1184-1191.	21.4	1,046