Ayse Sahaboglu

List of Publications by Year
in descending order

21 papers	1,275 citations	516710 16 h-index	752698 20 g-index
22 all docs	22 docs citations	22 times ranked	1771 citing authors

#	Article	IF	Citations
1	Drug repurposing studies of PARP inhibitors as a new therapy for inherited retinal degeneration. Cellular and Molecular Life Sciences, 2020, 77, 2199-2216.	5.4	20
2	Poly ADP ribosylation and extracellular vesicle activity in rod photoreceptor degeneration. Scientific Reports, 2019, 9, 3758.	3.3	25
3	Release of Retinal Extracellular Vesicles in aÂModel of Retinitis Pigmentosa. Advances in Experimental Medicine and Biology, 2019, 1185, 431-436.	1.6	8
4	Combination of cGMP analogue and drug delivery system provides functional protection in hereditary retinal degeneration. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E2997-E3006.	7.1	90
5	Temporal progression of PARP activity in the Prph2 mutant rd2 mouse: Neuroprotective effects of the PARP inhibitor PJ34. PLoS ONE, 2017, 12, e0181374.	2.5	23
6	Olaparib significantly delays photoreceptor loss in a model for hereditary retinal degeneration. Scientific Reports, 2016, 6, 39537.	3.3	45
7	HDAC inhibition in thecpfl1mouse protects degenerating cone photoreceptorsin vivo. Human Molecular Genetics, 2016, 25, ddw275.	2.9	39
8	Organotypic retinal explant cultures as in vitro alternative for diabetic retinopathy studies. ALTEX: Alternatives To Animal Experimentation, 2016, 33, 459-464.	1.5	29
9	Deletion of myosin VI causes slow retinal optic neuropathy and age-related macular degeneration (AMD)-relevant retinal phenotype. Cellular and Molecular Life Sciences, 2015, 72, 3953-3969.	5.4	10
10	Retinitis pigmentosa: impact of differentPde6apoint mutations on the disease phenotype. Human Molecular Genetics, 2015, 24, 5486-5499.	2.9	41
11	Identification of a Common Non-Apoptotic Cell Death Mechanism in Hereditary Retinal Degeneration. PLoS ONE, 2014, 9, e112142.	2.5	191
12	Knockout of PARG110 confers resistance to cGMP-induced toxicity in mammalian photoreceptors. Cell Death and Disease, 2014, 5, e1234-e1234.	6.3	13
13	Expression of Poly(ADP-Ribose) Glycohydrolase in Wild-Type and PARG-110 Knock-Out Retina. Advances in Experimental Medicine and Biology, 2014, 801, 463-469.	1.6	2
14	Inhibition of Mitochondrial Pyruvate Transport by Zaprinast Causes Massive Accumulation of Aspartate at the Expense of Glutamate in the Retina. Journal of Biological Chemistry, 2013, 288, 36129-36140.	3.4	72
15	Retinitis pigmentosa: rapid neurodegeneration is governed by slow cell death mechanisms. Cell Death and Disease, 2013, 4, e488-e488.	6.3	67
16	Neuroprotective Strategies for the Treatment of Inherited Photoreceptor Degeneration. Current Molecular Medicine, 2012, 12, 598-612.	1.3	68
17	Calpain and PARP Activation during Photoreceptor Cell Death in P23H and S334ter Rhodopsin Mutant Rats. PLoS ONE, 2011, 6, e22181.	2.5	94
18	Excessive HDAC activation is critical for neurodegeneration in the rd1 mouse. Cell Death and Disease, 2010, 1, e24-e24.	6.3	100

#	Article	IF	Citations
19	PARP1 Gene Knock-Out Increases Resistance to Retinal Degeneration without Affecting Retinal Function. PLoS ONE, 2010, 5, e15495.	2.5	71
20	Spectral Domain Optical Coherence Tomography in Mouse Models of Retinal Degeneration. , 2009, 50, 5888.		193
21	Protective effects of various antioxidants during ischemia-reperfusion in the rat retina. Graefe's Archive for Clinical and Experimental Ophthalmology, 2006, 244, 627-633.	1.9	74