Juhua Yang

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7911235/publications.pdf

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17	200	1163117 8	1058476 14
papers	citations	h-index	g-index

19	19	19	303
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The Therapeutic Effect of an Anti-TNF-Î±/HSA/IL-6R Triple-Specific Fusion Protein Under Experimental Septic Conditions. Inflammation, 2022, , 1.	3.8	1
2	Long Non-Coding RNA NR-133666 Promotes the Proliferation and Migration of Fibroblast-Like Synoviocytes Through Regulating the miR-133c/MAPK1 Axis. Frontiers in Pharmacology, 2022, 13, 887330.	3.5	3
3	Protective Effects of Nicotinamide Riboside on H₂O₂-induced Oxidative Damage in Lens Epithelial Cells. Current Eye Research, 2021, 46, 961-970.	1.5	6
4	A Dual Target-Directed Single Domain-Based Fusion Protein Against Interleukin-6 Receptor Decelerate Experimental Arthritis Progression Via Modulating JNK Expression. Inflammation, 2021, 44, 1620-1628.	3.8	3
5	Mitochondrial quality control protects photoreceptors against oxidative stress in the H2O2-induced models of retinal degeneration diseases. Cell Death and Disease, 2021, 12, 413.	6.3	20
6	Renoprotective Effect of the Recombinant Anti-IL-6R Fusion Proteins by Inhibiting JAK2/STAT3 Signaling Pathway in Diabetic Nephropathy. Frontiers in Pharmacology, 2021, 12, 681424.	3.5	8
7	The recombinant anti-TNF-Î± fusion protein ameliorates rheumatoid arthritis by the protective role of autophagy. Bioscience Reports, 2020, 40, .	2.4	6
8	Mutation screening of crystallin genes in Chinese families with congenital cataracts. Molecular Vision, 2019, 25, 427-437.	1.1	15
9	Novel mutations in HSF4 cause congenital cataracts in Chinese families. BMC Medical Genetics, 2018, 19, 150.	2.1	10
10	Novel mutations in CRB1 gene identified in a chinese pedigree with retinitis pigmentosa by targeted capture and next generation sequencing. Oncotarget, 2016, 7, 79797-79804.	1.8	3
11	A novel de novo duplication mutation of PAX6 in a Chinese family with aniridia and other ocular abnormalities. Scientific Reports, 2015, 4, 4836.	3.3	4
12	Analysis of TGFBI gene mutations in Chinese patients with corneal dystrophies and review of the literature. Molecular Vision, 2010, 16, 1186-93.	1.1	44
13	Confirmation of the mitochondrial ND1 gene mutation G3635A as a primary LHON mutation. Biochemical and Biophysical Research Communications, 2009, 386, 50-54.	2.1	27
14	Novel A14841G mutation is associated with high penetrance of LHON/C4171A family. Biochemical and Biophysical Research Communications, 2009, 386, 693-696.	2.1	17
15	The novel G10680A mutation is associated with complete penetrance of the LHON/T14484C family. Mitochondrion, 2009, 9, 273-278.	3.4	25
16	Novel human pathological mutations. Gene symbol: RHO. Disease: Retinitis pigmentosa. Human Genetics, 2009, 126, 336.	3.8	0
17	Identification of candidate cancer genes involved in human retinoblastoma by data mining. Child's Nervous System, 2008, 24, 893-900.	1.1	8