Shuo Yang

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7592511/publications.pdf

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13	352	1307594	1474206
papers	citations	7	9
		h-index	g-index

14	14	14	404
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Efficacy and Safety of rAAV2-ND4 Treatment for Leberâ€™s Hereditary Optic Neuropathy. Scientific Reports, 2016, 6, 21587.	3.3	131
2	Long-term outcomes of gene therapy for the treatment of Leber's hereditary optic neuropathy. EBioMedicine, 2016, 10, 258-268.	6.1	106
3	Seven-Year Follow-up of Gene Therapy for Leberâ€™s Hereditary Optic Neuropathy. Ophthalmology, 2020, 127, 1125-1127.	5.2	40
4	Tissue Distribution oftrans-Resveratrol and Its Metabolites after Oral Administration in Human Eyes. Journal of Ophthalmology, 2017, 2017, 1-12.	1.3	27
5	Evaluation of Leber's hereditary optic neuropathy patients prior to a gene therapy clinical trial. Medicine (United States), 2016, 95, e5110.	1.0	17
6	Chemical and material communication between the optic nerves in rats. Clinical and Experimental Ophthalmology, 2015, 43, 742-748.	2.6	16
7	Ocular Surface Ion-Channels Are Closely Related to Dry Eye: Key Research Focus on Innovative Drugs for Dry Eye. Frontiers in Medicine, 2022, 9, 830853.	2.6	9
8	Multilocus Mitochondrial Mutations Do Not Directly Affect the Efficacy of Gene Therapy for Leber Hereditary Optic Neuropathy. Journal of Neuro-Ophthalmology, 2020, 40, 22-29.	0.8	4
9	Sema-3A Regulates the Expression of B7-H3 and B7-H4 on Murine DCs.. Transplantation, 2014, 98, 684.	1.0	1
10	Co-Blocking TLR and CD154 Leads Profound Protection Against Renal Ischemia-Reperfusion Injury in Mice. Transplantation, 2012, 94, 1140.	1.0	0
11	A Robust Proof for the Essential Role of MyD88 in DCs in Transplant Responses. Transplantation, 2012, 94, 444.	1.0	0
12	A Meta-Analysis: Comparative Efficacy and Safety of Mizoribine With Mycophenolate Mofetil for Asian Renal Transplant Recipients.. Transplantation, 2014, 98, 585.	1.0	0
13	A novel CEP290 disease-causing variant identified in a patient with leber congenital amaurosis using a medical diagnostic panel sequencing. Ophthalmic Genetics, 2022, 43, 97-103.	1.2	0