Simon Dulz

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5695113/publications.pdf

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17 papers	166 citations	1307594 7 h-index	1199594 12 g-index
18 all docs	18 docs citations	18 times ranked	329 citing authors

#	Article	IF	Citations
1	Retinal Degeneration in Mice Deficient in the Lysosomal Membrane Protein CLN7. , 2016, 57, 4989.		26
2	Retinal toxicity after cisplatin-based chemotherapy in patients with germ cell cancer. Journal of Cancer Research and Clinical Oncology, 2017, 143, 1319-1325.	2.5	26
3	The Ocular Phenotype in Primary Hyperoxaluria Type 1. American Journal of Ophthalmology, 2019, 206, 184-191.	3.3	21
4	Intravitreal Co-Administration of GDNF and CNTF Confers Synergistic and Long-Lasting Protection against Injury-Induced Cell Death of Retinal Ganglion Cells in Mice. Cells, 2020, 9, 2082.	4.1	18
5	Reduced-dose abdominopelvic CT using hybrid iterative reconstruction in suspected left-sided colonic diverticulitis. European Radiology, 2016, 26, 216-224.	4.5	14
6	Retinal hyperreflective foci in Fabry disease. Orphanet Journal of Rare Diseases, 2019, 14, 296.	2.7	10
7	An Ophthalmic Rating Scale to Assess Ocular Involvement in Juvenile CLN3 Disease. American Journal of Ophthalmology, 2020, 220, 64-71.	3.3	10
8	A Case of a Bilateral Cicatricial Upper Eyelid Entropion After Hematopoietic Stem Cell Transplantation in Mucopolysaccharidosis Type I. Ophthalmic Plastic and Reconstructive Surgery, 2017, 33, S75-S77.	0.8	8
9	A comparison of the lateral tarsal strip with everting sutures and the Quickert procedure for involutional entropion. Acta Ophthalmologica, 2019, 97, e933-e936.	1.1	8
10	Oxalate retinopathy is irreversible despite early combined liver-kidney transplantation in primary hyperoxaluria type 1. American Journal of Transplantation, 2019, 19, 3328-3334.	4.7	7
11	Retinal vessel tortuosity as a prognostic marker for disease severity in Fabry disease. Orphanet Journal of Rare Diseases, 2021, 16, 485.	2.7	6
12	Isolated juvenile macular dystrophy without posterior column ataxia associated with FLVCR1 mutation. Ophthalmic Genetics, 2021, 42, 784-786.	1.2	3
13	Ongoing retinal degeneration despite intraventricular enzyme replacement therapy with cerliponase alfa in late-infantile neuronal ceroid lipofuscinosis type 2 (CLN2 disease). British Journal of Ophthalmology, 2023, 107, 1478-1483.	3.9	3
14	Retained visual function in a subset of patients with long-chain 3-hydroxyacyl-CoA dehydrogenase deficiency (LCHADD). Ophthalmic Genetics, 2021, 42, 23-27.	1.2	2
15	Reliability of the ocular trauma score for the predictability of traumatic and post-traumatic retinal detachment after open globe injury. International Journal of Ophthalmology, 2021, 14, 1589-1594.	1.1	2
16	Retinal Tacks for Complicated Retinal Detachment: Retinal Tacks in the Times of Modern Small-Gauge Vitrectomy. Journal of Ophthalmology, 2022, 2022, 1-9.	1.3	0
17	Visual perception and macular integrity in non-classical CLN2 disease. Graefe's Archive for Clinical and Experimental Ophthalmology, 0, , .	1.9	0