Matthieu P Robert

List of Publications by Year
in descending order

37	249	1163117 8	1058476 14
papers	citations	h-index	g-index

38	38	38	338
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Late Contralateral Recurrence of Unilateral Acute Idiopathic Maculopathy: Adaptive Optics Findings. Ocular Immunology and Inflammation, 2022, 30, 707-712.	1.8	0
2	Neurodevelopmental Disorder in a Child with High Intellectual Potential: Contribution of an Integrative Neuropsychomotor Standardized Assessment, Neuropsychological, and Eye-Tracking in a Single-Case Study. Advances in Neurodevelopmental Disorders, 2022, 6, 84-99.	1.1	0
3	Foveal Hypoplasia Grading in 95 Cases of Congenital Aniridia: Correlation to Phenotype and PAX6 Genotype. American Journal of Ophthalmology, 2022, 237, 122-129.	3.3	8
4	Retinal atrophy and markers of systemic and cerebrovascular severity in homozygous sickle cell disease. European Journal of Ophthalmology, 2022, 32, 3258-3266.	1.3	2
5	The role of irinotecan-bevacizumab as rescue regimen in children with low-grade gliomas: a retrospective nationwide study in 72 patients. Journal of Neuro-Oncology, 2022, 157, 355-364.	2.9	7
6	Function of the Retinal Pigment Epithelium in Patients With Neurofibromatosis Type 1. , 2022, 63, 6.		2
7	Longâ€term followâ€up of choroidal abnormalities in children with neurofibromatosis type 1. Clinical and Experimental Ophthalmology, 2021, 49, 516-519.	2.6	3
8	RETINAL VASCULAR ABNORMALITIES IN CHILDREN WITH NEUROFIBROMATOSIS TYPE 1. Retina, 2021, 41, 2589-2595.	1.7	7
9	Formulation and Stability of Ataluren Eye Drop Oily Solution for Aniridia. Pharmaceutics, 2021, 13, 7.	4.5	8
10	Pseudo-Gaze Deviation Resulting From Positive Angle Kappa and Esotropia. Journal of Neuro-Ophthalmology, 2021, 41, e234-e236.	0.8	1
11	Optic Nerve Abnormalities in Morning Glory Disc Anomaly. Journal of Neuro-Ophthalmology, 2021, Publish Ahead of Print, .	0.8	1
12	Clinical variability and probable founder effect in oculocutaneous albinism type 7. Clinical Genetics, 2020, 97, 527-528.	2.0	1
13	Evolution of Retinal Microvascular Abnormalities in Neurofibromatosis Type 1. Ophthalmology, 2020, 127, 1557.	5.2	2
14	PRPS1 loss-of-function variants, from isolated hearing loss to severe congenital encephalopathy: New cases and literature review. European Journal of Medical Genetics, 2020, 63, 104033.	1.3	11
15	Extraction of Nystagmus Patterns from Eye-Tracker Data with Convolutional Sparse Coding. , 2020, 2020, 928-931.		0
16	An opinion paper on the maintenance of robustness: Towards a multimodal and intergenerational approach using digital twins. Aging Medicine (Milton (N S W)), 2020, 3, 188-194.	2.1	4
17	Focal Areas of High Signal Intensity in Children with Neurofibromatosis Type 1: Expected Evolution on MRI. American Journal of Neuroradiology, 2020, 41, 1733-1739.	2.4	8
18	Early management of sight threatening retinopathy in incontinentia pigmenti. Orphanet Journal of Rare Diseases, 2020, 15, 223.	2.7	8

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19	Functional Vision Analysis in Patients With CHARGE Syndrome. Journal of Pediatric Ophthalmology and Strabismus, 2020, 57, 120-128.	0.7	6
20	Extraocular muscle positions in anterior plagiocephaly: V-pattern strabismus explained using geometric mophometrics. British Journal of Ophthalmology, 2020, 104, 1156-1160.	3.9	7
21	Recent developments in the management of congenital cataract. Annals of Translational Medicine, 2020, 8, 1545.	1.7	2
22	Recent developments in the management of congenital cataract. Annals of Translational Medicine, 2020, 8, 1545-1545.	1.7	13
23	Paramacular temporal atrophy in sickle cell disease occurs early in childhood. British Journal of Ophthalmology, 2019, 103, 906-910.	3.9	12
24	Recent innovations with drugs in clinical trials for neurotrophic keratitis and refractory corneal ulcers. Expert Opinion on Investigational Drugs, 2019, 28, 1013-1020.	4.1	19
25	Sostdc1 is expressed in all major compartments of developing and adult mammalian eyes. Graefe's Archive for Clinical and Experimental Ophthalmology, 2019, 257, 2401-2427.	1.9	1
26	Corpus callosum metrics predict severity of visuospatial and neuromotor dysfunctions in ARID1B mutations with Coffinâ€“Siris syndrome. Psychiatric Genetics, 2019, 29, 237-242.	1.1	8
27	The therapeutic potential of a calorie-restricted ketogenic diet for the management of Leber hereditary optic neuropathy. Nutritional Neuroscience, 2019, 22, 156-164.	3.1	9
28	Ocular Motor Apraxia. Contemporary Clinical Neuroscience, 2019, , 451-470.	0.3	1
29	Spontaneous consecutive esotropia. Eye, 2018, 32, 1197-1200.	2.1	1
30	Intermittent upbeat nystagmus in infancy and antihistamine medications. European Journal of Paediatric Neurology, 2018, 22, 571-572.	1.6	1
31	Yttrium Aluminum Garnet (YAG) Treatment for Persistent Fetal Vasculature: Nonsurgical Lens-Sparing Procedure to Cut Fibrovascular Stalk. Ophthalmology, 2018, 125, 1353.	5.2	0
32	Visual Function in Asymptomatic Patients With Homozygous Sickle Cell Disease and Temporal Macular Atrophy. JAMA Ophthalmology, 2017, 135, 1100.	2.5	19
33	Usefulness of cocaine drops in investigating infant anisocoria. European Journal of Paediatric Neurology, 2017, 21, 852-857.	1.6	12
34	Electroretinogram Findings in Early-Stage Sickle Cell Retinopathy According to Hemoglobin Type. , 2017, 58, 3262.		4
35	Relevance of Identifying JAG1 Mutations in Patients With Isolated Posterior Embryotoxon. Journal of Glaucoma, 2016, 25, 923-925.	1.6	6
36	Tournay's description of anisocoria on lateral gaze. Neurology, 2014, 82, 452-456.	1.1	4

#	Article	IF	Citations
37	Vertical and horizontal smooth pursuit eye movements in children with developmental coordination disorder. Developmental Medicine and Child Neurology, 2014, 56, 595-600.	2.1	51