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	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
2	Visual Function in Asymptomatic Patients With Homozygous Sickle Cell Disease and Temporal Macular Atrophy. JAMA Ophthalmology, 2017, 135, 1100.	2.5	19
3	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
4	Recent developments in the management of congenital cataract. Annals of Translational Medicine, 2020, 8, 1545-1545.	1.7	13
5	Usefulness of cocaine drops in investigating infant anisocoria. European Journal of Paediatric Neurology, 2017, 21, 852-857.	1.6	12
6	Paramacular temporal atrophy in sickle cell disease occurs early in childhood. British Journal of Ophthalmology, 2019, 103, 906-910.	3.9	12
7	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
8	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
9	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
10	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
11	Early management of sight threatening retinopathy in incontinentia pigmenti. Orphanet Journal of Rare Diseases, 2020, 15, 223.	2.7	8
12	Formulation and Stability of Ataluren Eye Drop Oily Solution for Aniridia. Pharmaceutics, 2021, 13, 7.	4.5	8
13	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
14	RETINAL VASCULAR ABNORMALITIES IN CHILDREN WITH NEUROFIBROMATOSIS TYPE 1. Retina, 2021, 41, 2589-2595.	1.7	7
15	Extraocular muscle positions in anterior plagiocephaly: V-pattern strabismus explained using geometric mophometrics. British Journal of Ophthalmology, 2020, 104, 1156-1160.	3.9	7
16	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
17	Relevance of Identifying JAG1 Mutations in Patients With Isolated Posterior Embryotoxon. Journal of Glaucoma, 2016, 25, 923-925.	1.6	6
18	Functional Vision Analysis in Patients With CHARGE Syndrome. Journal of Pediatric Ophthalmology and Strabismus, 2020, 57, 120-128.	0.7	6

#	Article	IF	Citations
19	Tournay's description of anisocoria on lateral gaze. Neurology, 2014, 82, 452-456.	1.1	4
20	Electroretinogram Findings in Early-Stage Sickle Cell Retinopathy According to Hemoglobin Type. , 2017, 58, 3262.		4
21	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
22	Longâ€term followâ€up of choroidal abnormalities in children with neurofibromatosis type 1. Clinical and Experimental Ophthalmology, 2021, 49, 516-519.	2.6	3
23	Evolution of Retinal Microvascular Abnormalities in Neurofibromatosis Type 1. Ophthalmology, 2020, 127, 1557.	5.2	2
24	Recent developments in the management of congenital cataract. Annals of Translational Medicine, 2020, 8, 1545.	1.7	2
25	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
26	Function of the Retinal Pigment Epithelium in Patients With Neurofibromatosis Type 1. , 2022, 63, 6.		2
27	Spontaneous consecutive esotropia. Eye, 2018, 32, 1197-1200.	2.1	1
28	Intermittent upbeat nystagmus in infancy and antihistamine medications. European Journal of Paediatric Neurology, 2018, 22, 571-572.	1.6	1
29	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
30	Clinical variability and probable founder effect in oculocutaneous albinism type 7. Clinical Genetics, 2020, 97, 527-528.	2.0	1
31	Ocular Motor Apraxia. Contemporary Clinical Neuroscience, 2019, , 451-470.	0.3	1
32	Pseudo-Gaze Deviation Resulting From Positive Angle Kappa and Esotropia. Journal of Neuro-Ophthalmology, 2021, 41, e234-e236.	0.8	1
33	Optic Nerve Abnormalities in Morning Glory Disc Anomaly. Journal of Neuro-Ophthalmology, 2021, Publish Ahead of Print, .	0.8	1
34	Yttrium Aluminum Garnet (YAG) Treatment for Persistent Fetal Vasculature: Nonsurgical Lens-Sparing Procedure to Cut Fibrovascular Stalk. Ophthalmology, 2018, 125, 1353.	5.2	0
35	Extraction of Nystagmus Patterns from Eye-Tracker Data with Convolutional Sparse Coding. , 2020, 2020, 928-931.		0
36	Late Contralateral Recurrence of Unilateral Acute Idiopathic Maculopathy: Adaptive Optics Findings. Ocular Immunology and Inflammation, 2022, 30, 707-712.	1.8	0

#	Article	IF	Citations
37	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