Adriano Magli

List of Publications by Year
in descending order

61 papers	1,673 citations	567281 15 h-index	315739 38 g-index
61 all docs	61 docs citations	61 times ranked	2598 citing authors

#	Article	IF	Citations
1	A case of retinal vascular involvement in a 6-year-old patient with Covid-19. European Journal of Ophthalmology, 2022, 32, NP1-NP5.	1.3	3
2	Augmented vertical rectus transpositions: Intraoperative measurement of torsion following sequential muscle detachment. European Journal of Ophthalmology, 2021, 31, 2027-2031.	1.3	3
3	Optical coherence tomography angiography for the measurement of optic disc: Macular relationship. European Journal of Ophthalmology, 2021, 31, 543-547.	1.3	1
4	Novel USH1G homozygous variant underlying USH2-like phenotype of Usher syndrome. European Journal of Ophthalmology, 2021, 31, NP18-NP22.	1.3	4
5	Reoperation in esotropic Duane retraction syndrome: Long-term motor outcome of superior rectus transposition. European Journal of Ophthalmology, 2021, 31, 722-726.	1.3	4
6	Optical coherence tomography angiography in healthy children: A comparison of macular structure. European Journal of Ophthalmology, 2021, , 112067212110437.	1.3	2
7	Expanding the Clinical and Genetic Spectrum of RAB28-Related Cone-Rod Dystrophy: Pathogenicity of Novel Variants in Italian Families. International Journal of Molecular Sciences, 2021, 22, 381.	4.1	8
8	Long-Term Effects of Botulinum Toxin in Large-Angle Infantile Esotropia . Clinical Ophthalmology, 2020, Volume 14, 3399-3402.	1.8	3
9	Novel mutations in MFRP and PRSS56 are associated with posterior microphthalmos. Ophthalmic Genetics, 2020, 41, 49-56.	1.2	10
10	Paralog Studies Augment Gene Discovery: DDX and DHX Genes. American Journal of Human Genetics, 2019, 105, 302-316.	6.2	56
11	Antioxidant Saffron and Central Retinal Function in ABCA4-Related Stargardt Macular Dystrophy. Nutrients, 2019, 11, 2461.	4.1	25
12	Pathogenicity of new BEST1 variants identified in Italian patients with best vitelliform macular dystrophy assessed by computational structural biology. Journal of Translational Medicine, 2019, 17, 330.	4.4	2
13	Divergence Excess Intermittent Exotropia: Long-Term Effect of Augmented Bilateral Lateral Rectus Recession. Seminars in Ophthalmology, 2018, 33, 512-516.	1.6	5
14	Long-term development of refractive error in refractive, nonrefractive and partially accommodative esotropia. PLoS ONE, 2018, 13, e0204396.	2.5	9
15	A novel p.(Glu111Val) missense mutation in GUCA1A associated with cone-rod dystrophy leads to impaired calcium sensing and perturbed second messenger homeostasis in photoreceptors. Human Molecular Genetics, 2018, 27, 4204-4217.	2.9	32
16	Multimodal Imaging in Autosomal Dominant Cone-Rod Dystrophy Caused by Novel <i>CRX</i> Variant. Ophthalmic Research, 2018, 60, 169-175.	1.9	4
17	Congenital and Developmental Cataracts: Focus on Strabismus Outcomes at Long-Term Follow-Up. Seminars in Ophthalmology, 2017, 32, 358-362.	1.6	7
18	Essential Infantile Esotropia: Postoperative Sensory Outcomes of Strabismus Surgery. Seminars in Ophthalmology, 2017, 32, 663-671.	1.6	9

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19	Developmental visual deprivation: long term effects on human cone driven retinal function. Graefe's Archive for Clinical and Experimental Ophthalmology, 2017, 255, 2481-2486.	1.9	6
20	Early light deprivation effects on human coneâ€driven retinal function. Acta Ophthalmologica, 2017, 95, 133-139.	1.1	7
21	Long-Term Outcomes of Primary Intraocular Lens Implantation for Unilateral Congenital Cataract. Seminars in Ophthalmology, 2016, 31, 1-6.	1.6	15
22	Infantile esotropia: risk factors associated with reoperation. Clinical Ophthalmology, 2016, Volume 10, 2079-2083.	1.8	6
23	Pediatric keratoconus and iontophoretic corneal crosslinking: refractive and topographic evidence in patients underwent general and topical anesthesia, 18Âmonths of follow-up. International Ophthalmology, 2016, 36, 585-590.	1.4	39
24	Essential infantile esotropia with inferior oblique hyperfunction: long term follow-up of 6 muscles approach. International Journal of Ophthalmology, 2016, 9, 1802-1807.	1.1	0
25	Long-Term Follow-Up after Surgery for Congenital and Developmental Cataracts. Seminars in Ophthalmology, 2014, 31, 1-5.	1.6	3
26	Short-term effects of vision trainer rehabilitation in patients affected by anisometropic amblyopia: electrofunctional evaluation. Documenta Ophthalmologica, 2014, 129, 177-189.	2.2	12
27	Cataract Management in Juvenile Idiopathic Arthritis: Simultaneous versus Secondary Intraocular Lens Implantation. Ocular Immunology and Inflammation, 2014, 22, 133-137.	1.8	24
28	Essential infantile esotropia: postoperative motor outcomes and inferential analysis of strabismus surgery. BMC Ophthalmology, 2014, 14, 35.	1.4	12
29	Refractive Surgery for Accommodative Esotropia: 5-Year Follow-up. Journal of Refractive Surgery, 2014, 30, 116-120.	2.3	9
30	Adalimumab for juvenile idiopathic arthritis-associated uveitis. Graefe's Archive for Clinical and Experimental Ophthalmology, 2013, 251, 1601-1606.	1.9	45
31	Long-term outcome of primary versus secondary intraocular lens implantation after simultaneous removal of bilateral congenital cataract. Graefe's Archive for Clinical and Experimental Ophthalmology, 2013, 251, 309-314.	1.9	42
32	Treatment of Infantile Capillary Hemangioma of the Eyelid with Systemic Propranolol. American Journal of Ophthalmology, 2013, 155, 165-170.e2.	3.3	29
33	Evaluation of retinal nerve fiber layer and ganglion cell complex thickness after ocular blunt trauma. Eye, 2013, 27, 1382-1387.	2.1	5
34	Functional changes after treatment of optic pathway paediatric low-grade gliomas. Eye, 2013, 27, 1288-1292.	2.1	7
35	Epithelium-Off Corneal Collagen Cross-linking Versus Transepithelial Cross-linking for Pediatric Keratoconus. Cornea, 2013, 32, 597-601.	1.7	129
36	Bilateral methicillin-resistant Staphylococcus aureus keratitis following hyperopic photorefractive surgery. International Ophthalmology, 2012, 32, 47-49.	1.4	9

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37	Lower Eyelid Surgery for Lagophthalmos in MÃ¶bius and Poland-MÃ¶bius Syndromes. Journal of Craniofacial Surgery, 2011, 22, e53-e54.	0.7	2
38	Molecular and Clinical Characterization of Albinism in a Large Cohort of Italian Patients. , 2011, 52, 1281.		58
39	Human TUBB3 Mutations Perturb Microtubule Dynamics, Kinesin Interactions, and Axon Guidance. Cell, 2010, 140, 74-87.	28.9	515
40	LASIK and PRK in Refractive Accommodative Esotropia: A Retrospective Study on 20 Adolescent and Adult Patients. European Journal of Ophthalmology, 2009, 19, 188-195.	1.3	24
41	Sonic Hedgehog deletion and distal trisomy 3p in a patient with microphthalmia and microcephaly, lacking cerebral anomalies typical of holoprosencephaly. European Journal of Medical Genetics, 2008, 51, 658-665.	1.3	14
42	Ocular Complications of Eyebrow Piercing. Journal of Pediatric Ophthalmology and Strabismus, 2008, 45, 184-185.	0.7	5
43	Identification ofKIF21AMutations as a Rare Cause of Congenital Fibrosis of the Extraocular Muscles Type 3 (CFEOM3). , 2004, 45, 2218.		83
44	Analysis of nuclear fiber cell compaction in transparent and cataractous diabetic human lenses by scanning electron microscopy. BMC Ophthalmology, 2003, 3, 1.	1.4	23
45	Clinical and surgical data of affected members of a classic CFEOM 1 family. BMC Ophthalmology, 2003, 3, 6.	1.4	11
46	Heterozygous mutations of the kinesin KIF21A in congenital fibrosis of the extraocular muscles type 1 (CFEOM1). Nature Genetics, 2003, 35, 318-321.	21.4	240
47	CFEOM1, the classic familial form of congenital fibrosis of the extraocular muscles, is genetically heterogeneous but does not result from mutations in ARIX. BMC Genetics, 2002, 3, 3.	2.7	42
48	Spontaneous Resolution of Congenital Bilateral Brownâ€™s Syndrome. Ophthalmologica, 2001, 215, 372-375.	1.9	9
49	Measuring contrast sensitivity in aretinopathic patients with insulin dependent diabetes mellitus. Documenta Ophthalmologica, 1997, 93, 199-209.	2.2	5
50	Visual dysfunction in patients with mitochondrial myopathies. Documenta Ophthalmologica, 1995, 89, 211-218.	2.2	9
51	A Case of Parinaudâ€™s Syndrome in a Boy with Delayed Puberty. Ophthalmologica, 1991, 202, 132-137.	1.9	1
52	The Gorlin-Goltz Syndrome: Case Report (With 1 color plate). Ophthalmologica, 1990, 200, 104-106.	1.9	16
53	Ptosis Correction in the Context of the Treatment of External Congenital Ophthalmoplegia. Ophthalmic Plastic and Reconstructive Surgery, 1989, 5, 176-181.	0.8	6
54	The Annette von Droste-Hulshoff syndrome: Pseudostrabismus due to macular ectopia in retinopathy of prematurity. Ophthalmic Paediatrics and Genetics, 1988, 9, 13-16.	0.4	4

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55	Hereditary colobomatous anomalies of the optic nerve head[*]. Ophthalmic Paediatrics and Genetics, 1986, 7, 127-130.	0.4	4
56	Genetic and ultrasound study of hereditary pure microphthalmos. Ophthalmic Paediatrics and Genetics, 1985, 6, 37-42.	0.4	2
57	Genetic and ultrasound study of abnormalities of the optic nerve head. Ophthalmic Paediatrics and Genetics, 1985, 5, 71-78.	0.4	3
58	Coloboma of the lens associated with coloboma of the alar nasal cartilages in a pair of female monozygotic twins: A new syndrome?. Ophthalmic Paediatrics and Genetics, 1983, 2, 83-87.	0.4	0
59	Familial juvenile nephronophthisis and associated ocular anomalies (Senior's syndrome). A study of three families. Ophthalmic Paediatrics and Genetics, 1982, 1, 97-105.	0.4	3
60	Unilateral Inferior Altitudinal Hemianopsia, Argyll Robertson Pupil and Dendritic Keratitis in a Young Patient with Herpes Zoster. Ophthalmologica, 1981, 183, 143-147.	1.9	3
61	Ocular Coloboma with Congenital Heart Disease in the Absence of Chromosomal Abnormalities. Ophthalmologica, 1980, 181, 195-202.	1.9	5