Dario Pasquale Mucciolo

List of Publications by Year
in descending order

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51	334	840728 11	996954 15
papers	citations	h-index	g-index

51	51	51	455
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	EDI-OCT evaluation of choroidal thickness in retinitis pigmentosa. European Journal of Ophthalmology, 2018, 28, 52-57.	1.3	29
2	Reading Ability and Quality of Life in Stargardt Disease. European Journal of Ophthalmology, 2017, 27, 740-745.	1.3	24
3	Subthreshold yellow micropulse laser for treatment of diabetic macular edema: Comparison between fixed and variable treatment regimen. European Journal of Ophthalmology, 2021, 31, 1254-1260.	1.3	19
4	Dietary profile of patients with Stargardtâ€™s disease and Retinitis Pigmentosa: is there a role for a nutritional approach?. BMC Ophthalmology, 2016, 16, 13.	1.4	17
5	Retinal dystrophy and subretinal drusenoid deposits in female choroideremia carriers. Graefe's Archive for Clinical and Experimental Ophthalmology, 2017, 255, 2099-2111.	1.9	17
6	Optical Coherence Tomography Angiography (OCT-A) in young choroideremia (CHM) patients. Ophthalmic Genetics, 2019, 40, 201-206.	1.2	16
7	Retinal capillaritis in a CRB1-associated retinal dystrophy. Ophthalmic Genetics, 2017, 38, 555-558.	1.2	15
8	Peripapillary comet lesions and comet rain in PXE-related retinopathy. Graefe's Archive for Clinical and Experimental Ophthalmology, 2018, 256, 1605-1614.	1.9	14
9	Long-term follow-up of a CRB1-associated maculopathy. Ophthalmic Genetics, 2018, 39, 522-525.	1.2	13
10	Novel clinical findings in autosomal recessive NR2E3-related retinal dystrophy. Graefe's Archive for Clinical and Experimental Ophthalmology, 2019, 257, 9-22.	1.9	13
11	En face OCT in Stargardt disease. Graefe's Archive for Clinical and Experimental Ophthalmology, 2016, 254, 1669-1679.	1.9	12
12	Case report of an atypical early onset X-linked retinoschisis in monozygotic twins. BMC Ophthalmology, 2017, 17, 19.	1.4	12
13	Coquille dâ€™oeuf in young patients affected with Pseudoxantoma elasticum. Ophthalmic Genetics, 2019, 40, 242-246.	1.2	11
14	Phenotypic Features and Genetic Findings in a Cohort of Italian Pseudoxanthoma Elasticum Patients and Update of the Ophthalmologic Evaluation Score. Journal of Clinical Medicine, 2021, 10, 2710.	2.4	8
15	A novel GRK1 mutation in an Italian patient with Oguchi disease. Ophthalmic Genetics, 2018, 39, 137-138.	1.2	7
16	Pattern dystrophy-like changes and coquille dâ€™oeuf atrophy in elderly patients affected by pseudoxanthoma elasticum. Graefe's Archive for Clinical and Experimental Ophthalmology, 2020, 258, 1881-1892.	1.9	7
17	Computer-Assisted Evaluation of Retinal Vessel Diameter in Retinitis Pigmentosa. Ophthalmic Research, 2016, 56, 139-144.	1.9	6
18	Fundus phenotype in retinitis pigmentosa associated with EYS mutations. Ophthalmic Genetics, 2018, 39, 589-602.	1.2	6

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19	Macular hole in Stargardt disease: Clinical and ultra-structural observation. Ophthalmic Genetics, 2017, 38, 486-489.	1.2	5
20	CFH Y402H polymorphism in Italian patients with age-related macular degeneration, retinitis pigmentosa, and Stargardt disease. Ophthalmic Genetics, 2018, 39, 699-705.	1.2	5
21	Italian real-life experience on the use of ocriplasmin. BMJ Open Ophthalmology, 2018, 3, e000110.	1.6	5
22	EDI OCT evaluation of choroidal thickness in Stargardt disease. PLoS ONE, 2018, 13, e0190780.	2.5	5
23	Optical coherence tomography (OCT) features of cystoid spaces in choroideremia (CHM). Graefe's Archive for Clinical and Experimental Ophthalmology, 2019, 257, 2655-2663.	1.9	5
24	En face OCT in choroideremia. Ophthalmic Genetics, 2019, 40, 514-520.	1.2	5
25	CHOROIDAL VASCULARITY INDEX IN YOUNG CHOROIDEREMIA PATIENTS. Retina, 2021, 41, 1018-1025.	1.7	5
26	OCTA Imaging of Choroidal Neovascularization Treated Using Photodynamic Therapy in a Young Patient With Best Macular Dystrophy. Ophthalmic Surgery Lasers and Imaging Retina, 2018, 49, 969-973.	0.7	5
27	Multimodal imaging of benign yellow dot maculopathy. Ophthalmic Genetics, 2019, 40, 135-140.	1.2	4
28	ULTRASOUND IN VITRECTOMY. Retina, 2020, 40, 24-32.	1.7	4
29	Optical Coherence Tomography Angiography (OCT-A) in Choroideremia (CHM) carriers. Ophthalmic Genetics, 2020, 41, 146-151.	1.2	4
30	Adaptive Optics Imaging in Patients Affected by Pseudoxanthoma Elasticum. American Journal of Ophthalmology, 2021, 224, 84-95.	3.3	4
31	Peculiar Clinical Findings in Young Choroideremia Patients: A Retrospective Case Review. Ophthalmologica, 2019, 242, 195-207.	1.9	3
32	Use of anti-vascular endothelial growth factor drugs for eye disease in Tuscany: Development and test of indicators of treatment intensity. European Journal of Ophthalmology, 2020, 30, 1440-1447.	1.3	3
33	Long-Term Follow-Up of an Atypical Case of Idiopathic Macular Telangiectasia. Ophthalmic Surgery Lasers and Imaging Retina, 2018, 49, e278-e283.	0.7	3
34	Choroidal Caverns in Stargardt Disease. , 2022, 63, 25.		3
35	Near-infrared autofluorescence in young choroideremia patients. Ophthalmic Genetics, 2019, 40, 421-427.	1.2	2
36	The Veterans Affairs Low-Vision Visual Functioning Questionnaire-48 (VA LV VFQ-48): Performance of the Italian version. European Journal of Ophthalmology, 2020, 30, 1014-1018.	1.3	2

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37	Optical coherence tomography angiography cyclic remodeling of CNV in patients affected by Best macular dystrophy. Ophthalmic Genetics, 2020, 41, 440-447.	1.2	2
38	Outer nuclear layer relevance in visual function correlated to quantitative enface OCT parameters in Stargardt disease. European Journal of Ophthalmology, 2021, 31, 112067212199057.	1.3	2
39	Clinical and molecular findings in patients with pattern dystrophy. Ophthalmic Genetics, 2021, 42, 577-587.	1.2	2
40	Lamellar Hole-associated Epiretinal Proliferation in choroideremia: a case report. International Journal of Retina and Vitreous, 2021, 7, 63.	1.9	2
41	Clinical and genetic findings in Italian patients with sector retinitis pigmentosa. Molecular Vision, 2021, 27, 78-94.	1.1	2
42	Intraretinal hyperreflective foci in PXE-related retinopathy with acquired vitelliform lesions: a long-term follow-up. Ophthalmic Genetics, 2019, 40, 385-387.	1.2	1
43	Circulating endothelial and progenitor cells in age-related macular degeneration. European Journal of Ophthalmology, 2020, 30, 956-965.	1.3	1
44	Acquired retinoschisis and vitreous hemorrhage as unusual findings in choroideremia: Case report. European Journal of Ophthalmology, 2020, 31, 112067212094657.	1.3	1
45	A Case Report of Pseudoxanthoma elasticum with Rare Sequence Variants in Genes Related to Inherited Retinal Diseases. Diagnostics, 2021, 11, 1800.	2.6	1
46	A Nine-Year Follow-Up of Macular Complications in Retinitis Pigmentosa and Diabetes Mellitus. Ophthalmic Surgery Lasers and Imaging Retina, 2017, 48, 340-344.	0.7	1
47	Outcome and genetic analysis of patients affected by retinal capillary hemangioblastoma in von Hippel Lindau syndrome. Molecular Vision, 2021, 27, 542-554.	1.1	1
48	Peripheral retinal neovascularization in a patient with pilocytic astrocytoma. European Journal of Ophthalmology, 2020, 30, NP82-NP85.	1.3	0
49	Long-term follow-up and â€œdouble layer signâ€•in patients affected by circumscribed choroidal hemangioma. Photodiagnosis and Photodynamic Therapy, 2020, 31, 101960.	2.6	0
50	From Clinical Diagnosis to the Discovery of Multigene Rare Sequence Variants in Pseudoxanthoma elasticum: A Case Report. Frontiers in Medicine, 2021, 8, 726856.	2.6	0
51	Choroidal Vascularity Index in CHM Carriers. Frontiers in Ophthalmology, 2021, 1, .	0.5	0