

Kirk A J Stephenson

List of Publications by Year in descending order

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29
papers

221
citations

1307594

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docs citations

29
times ranked

376
citing authors

#	ARTICLE	IF	CITATIONS
1	Panel-Based Population Next-Generation Sequencing for Inherited Retinal Degenerations. <i>Scientific Reports</i> , 2016, 6, 33248.	3.3	49
2	Target 5000: Target Capture Sequencing for Inherited Retinal Degenerations. <i>Genes</i> , 2017, 8, 304.	2.4	46
3	Findings from a Genotyping Study of over 1000 People with Inherited Retinal Disorders in Ireland. <i>Genes</i> , 2020, 11, 105.	2.4	38
4	Final anatomic and visual outcomes appear independent of duration of silicone oil intraocular tamponade in complex retinal detachment surgery. <i>International Journal of Ophthalmology</i> , 2018, 11, 83-88.	1.1	14
5	Target 5000: a standardized all-Ireland pathway for the diagnosis and management of inherited retinal degenerations. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 200.	2.7	10
6	Paediatric retinal detachment: aetiology, characteristics and outcomes. <i>International Journal of Ophthalmology</i> , 2018, 11, 262-266.	1.1	9
7	A FBN1 variant manifesting as non-syndromic ectopia lentis with retinal detachment: clinical and genetic characteristics. <i>Eye</i> , 2020, 34, 690-694.	2.1	9
8	Clinical and Genetic Re-Evaluation of Inherited Retinal Degeneration Pedigrees following Initial Negative Findings on Panel-Based Next Generation Sequencing. <i>International Journal of Molecular Sciences</i> , 2022, 23, 995.	4.1	8
9	A Novel FLVCR1 Variant Implicated in Retinitis Pigmentosa. <i>Advances in Experimental Medicine and Biology</i> , 2019, 1185, 203-207.	1.6	6
10	The Natural History of Leber's Hereditary Optic Neuropathy in an Irish Population and Assessment for Prognostic Biomarkers. <i>Neuro-Ophthalmology</i> , 2022, 46, 159-170.	1.0	6
11	Multimodal imaging in a pedigree of X-linked Retinoschisis with a novel RS1 variant. <i>BMC Medical Genetics</i> , 2018, 19, 195.	2.1	5
12	Management of Keratoconus in Down Syndrome and Other Intellectual Disability. <i>Cornea</i> , 2021, Publish Ahead of Print, 456-461.	1.7	5
13	Electrophysiology-Guided Genetic Characterisation Maximises Molecular Diagnosis in an Irish Paediatric Inherited Retinal Degeneration Population. <i>Genes</i> , 2022, 13, 615.	2.4	4
14	The Use of Oral Midazolam to Facilitate the Ophthalmic Examination of Children with Autism and Developmental Disorders. <i>Journal of Autism and Developmental Disorders</i> , 2021, 51, 1678-1682.	2.7	3
15	Management of significant secondary genetic findings in an ophthalmic genetics clinic. <i>Eye</i> , 2022, 36, 896-898.	2.1	3
16	Acute Exudative Polymorphous Vitelliform Maculopathy Syndrome; natural history and evolution of fundal and OCT images over time. <i>BMJ Case Reports</i> , 2018, 2018, bcr-2018-224241.	0.5	3
17	Management of cataract in a patient with anterior megalophthalmos. <i>BMJ Case Reports</i> , 2021, 14, e241659.	0.5	1
18	Retinal imaging via the implantable miniature telescope. <i>BMJ Case Reports</i> , 2021, 14, e243242.	0.5	1

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19	Postpartum haemorrhage associated choroidopathy. BMJ Case Reports, 2022, 15, e249226.	0.5	1
20	An ocular motility conundrum. BMJ Case Reports, 2014, 2014, bcr2014206862-bcr2014206862.	0.5	0
21	Monocular syphilitic uveitis. BMJ Case Reports, 2021, 14, e241403.	0.5	0
22	Behçet's disease presenting as bilateral occlusive retinal vasculitis in a young woman. BMJ Case Reports, 2021, 14, e241794.	0.5	0
23	Monitoring of non-progressive retinoschisis detachment with posterior outer leaf break. BMJ Case Reports, 2021, 14, e241848.	0.5	0
24	Visual field loss in an elderly vasculopath: clinical significance of multimodal imaging. BMJ Case Reports, 2021, 14, e245115.	0.5	0
25	Persistent photopsia: multiple evanescent white dot syndrome in a sexagenarian. BMJ Case Reports, 2021, 14, e246140.	0.5	0
26	Pigmentary retinopathy masked by asymmetric acquired phenomena. BMJ Case Reports, 2021, 14, e246982.	0.5	0
27	Colonic adenocarcinoma presenting as monocular metamorphopsia. BMJ Case Reports, 2022, 15, e245828.	0.5	0
28	Coats-like exudative vitreoretinopathy (CLEVER) in CEP290 inherited retinal degeneration. BMJ Case Reports, 2022, 15, e247229.	0.5	0
29	Insidious ocular surface lesion in an 81-year-old woman. BMJ Case Reports, 2022, 15, e248725.	0.5	0