

# Jane Ashworth

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4121598/publications.pdf>

Version: 2024-02-01

26  
papers

578  
citations

840119

11  
h-index

642321

23  
g-index

26  
all docs

26  
docs citations

26  
times ranked

872  
citing authors

#	ARTICLE	IF	CITATIONS
1	Progression of eye disease over 15 years in a patient with mucopolysaccharidosis type VI on enzyme replacement therapy. <i>BMJ Case Reports</i> , 2021, 14, e238544.	0.2	4
2	Management of Corneal Clouding in Patients with Mucopolysaccharidosis. <i>Journal of Clinical Medicine</i> , 2021, 10, 3263.	1.0	5
3	Areas of agreement in the management of childhood non-infectious chronic anterior uveitis in the UK. <i>British Journal of Ophthalmology</i> , 2020, 104, 11-16.	2.1	37
4	Favourable outcome in paediatric endogenous endophthalmitis secondary to <i>Neisseria meningitidis</i> following pars plana vitrectomy. <i>BMJ Case Reports</i> , 2020, 13, e233133.	0.2	2
5	Variability in the ocular phenotype in mucopolysaccharidosis. <i>British Journal of Ophthalmology</i> , 2019, 103, 504-510.	2.1	16
6	Incidence, management and outcome of raised intraocular pressure in childhood-onset uveitis at a tertiary referral centre. <i>British Journal of Ophthalmology</i> , 2019, 103, 748-752.	2.1	5
7	Diagnosis and management of individuals with Fetal Valproate Spectrum Disorder; a consensus statement from the European Reference Network for Congenital Malformations and Intellectual Disability. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 180.	1.2	33
8	An update on the modern management of paediatric uveitis. <i>British Journal of Ophthalmology</i> , 2019, 103, bjophthalmol-2019-314212.	2.1	11
9	<p>Determining the needs of ophthalmic trainees entering into specialist training and how they can be met</p>. <i>Advances in Medical Education and Practice</i> , 2019, Volume 10, 201-206.	0.7	1
10	Recommendations for the management of MPS VI: systematic evidence- and consensus-based guidance. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 118.	1.2	30
11	Recommendations for the management of MPS IVA: systematic evidence- and consensus-based guidance. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 137.	1.2	62
12	Traboulsi syndrome due to ASPH mutation: an under-recognised cause of ectopia lentis. <i>Clinical Dysmorphology</i> , 2019, 28, 184-189.	0.1	10
13	Nonaccidental injury presenting as unilateral retinal detachment in two infants. <i>Journal of AAPOS</i> , 2018, 22, 231-233.	0.2	2
14	The effect of haemopoietic stem cell transplantation on the ocular phenotype in mucopolysaccharidosis type I (Hurler). <i>Acta Ophthalmologica</i> , 2018, 96, 494-498.	0.6	5
15	Varicella-zoster virus necrotising retinitis, retinal vasculitis and panuveitis following uncomplicated chickenpox in an immunocompetent child. <i>BMJ Case Reports</i> , 2018, 2018, bcr-2017-223823.	0.2	3
16	Outcomes of keratoplasty in the mucopolysaccharidoses: an international perspective. <i>British Journal of Ophthalmology</i> , 2017, 101, 909-912.	2.1	22
17	Objective Quantification of Changes in Corneal Clouding Over Time in Patients With Mucopolysaccharidosis. , 2017, 58, 954.		17
18	Use of new imaging in detecting and monitoring ocular manifestations of the mucopolysaccharidoses. <i>Acta Ophthalmologica</i> , 2016, 94, e676-e682.	0.6	11

#	ARTICLE	IF	CITATIONS
19	The Novel Evidenced Assessment of Tortuosity system: interobserver reliability and agreement with clinical assessment. <i>Acta Ophthalmologica</i> , 2016, 94, e421-6.	0.6	3
20	Automated Measurement of Visual Acuity in Pediatric Ophthalmic Patients Using Principles of Game Design and Tablet Computers. <i>American Journal of Ophthalmology</i> , 2016, 170, 223-227.	1.7	12
21	Outcomes of Long-Term Treatment with Laronidase in Patients with Mucopolysaccharidosis Type I. <i>Journal of Pediatrics</i> , 2016, 178, 219-226.e1.	0.9	44
22	Next-generation Sequencing in the Diagnosis of Metabolic Disease Marked by Pediatric Cataract. <i>Ophthalmology</i> , 2016, 123, 217-220.	2.5	44
23	Multifocal chorioretinitis with progressive subretinal fibrosis in a young child. <i>BMJ Case Reports</i> , 2015, 2015, bcr2015212526.	0.2	6
24	Assessment and diagnosis of suspected glaucoma in patients with mucopolysaccharidosis. <i>Acta Ophthalmologica</i> , 2015, 93, e1111-7.	0.6	22
25	Personalized Diagnosis and Management of Congenital Cataract by Next-Generation Sequencing. <i>Ophthalmology</i> , 2014, 121, 2124-2137.e2.	2.5	153
26	Oculoplastic reconstruction following Mohs surgery. <i>Eye</i> , 1998, 12, 214-218.	1.1	18