Ingele Casteels

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/3513447/publications.pdf

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32	383	759233 12	839539 18
papers	citations	h-index	g-index

32	32	32	810
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Mutations in Splicing Factor Genes Are a Major Cause of Autosomal Dominant Retinitis Pigmentosa in Belgian Families. PLoS ONE, 2017, 12, e0170038.	2.5	47
2	Abnormal differentiation of B cells and megakaryocytes in patients with Roifman syndrome. Journal of Allergy and Clinical Immunology, 2018, 142, 630-646.	2.9	36
3	Current knowledge on the human eye microbiome: a systematic review of available amplicon and metagenomic sequencing data. Acta Ophthalmologica, 2021, 99, 16-25.	1.1	36
4	Educational paper. European Journal of Pediatrics, 2012, 171, 887-893.	2.7	30
5	Where are the missing gene defects in inherited retinal disorders? Intronic and synonymous variants contribute at least to 4% of CACNA1F â€mediated inherited retinal disorders. Human Mutation, 2019, 40, 765-787.	2.5	24
6	The majority of autosomal recessive nanophthalmos and posterior microphthalmia can be attributed to biallelic sequence and structural variants in MFRP and PRSS56. Scientific Reports, 2020, 10, 1289.	3.3	24
7	Nonâ€organic visual loss in children: prospective and retrospective analysis of associated psychosocial problems and stress factors. Acta Ophthalmologica, 2016, 94, e312-6.	1.1	22
8	Clinical Practice: A Proposed Standardized Ophthalmological Assessment for Patients with Cystinosis. Ophthalmology and Therapy, 2017, 6, 93-104.	2.3	20
9	Insights into multiple sclerosisâ€associated uveitis: a scoping review. Acta Ophthalmologica, 2021, 99, 592-603.	1.1	18
10	Nature of the Visual Loss in Observers With Leber's Congenital Amaurosis Caused by Specific Mutations in RPE65. Investigative Ophthalmology and Visual Science, 2014, 55, 6817-6828.	3.3	15
11	Mutation spectrum and clinical investigation of achromatopsia patients with mutations in the GNAT2 gene. Human Mutation, 2019, 40, 1145-1155.	2.5	15
12	Optical Coherence Tomography Angiography of Retinal Microvascular Changes Overlying Choroidal Nodules in Neurofibromatosis Type 1. Case Reports in Ophthalmology, 2017, 8, 214-220.	0.7	14
13	Cystinosis: a new perspective. Acta Clinica Belgica, 2016, 71, 131-137.	1.2	12
14	Effect of Storage Conditions on Stability of Ophthalmological Compounded Cysteamine Eye Drops. JIMD Reports, 2017, 42, 47-51.	1.5	11
15	Association between near viewing and acute acquired esotropia in children during tablet and smartphone use. Strabismus, 2022, 30, 59-64.	0.7	10
16	The Effect of Topical Anesthetics on 16S Ribosomal Ribonucleic Acid Amplicon Sequencing Results in Ocular Surface Microbiome Research. Translational Vision Science and Technology, 2022, 11, 2.	2.2	9
17	Hematopoietic Stem Cell Transplantation Cures Chronic Aichi Virus Infection in a Patient with X-linked Agammaglobulinemia. Journal of Clinical Immunology, 2021, 41, 1403-1405.	3.8	8
18	Use of the Masquerade Flap in Ablepharonâ€“Macrostomia Syndrome: A Case Report. Cornea, 2018, 37, 929-932.	1.7	7

#	Article	IF	Citations
19	A Retrospective Case Series of Uveal Effusion Syndrome. Journal of Glaucoma, 2020, 29, 995-998.	1.6	6
20	Clinically Relevant Response to Cisplatin-5-Fluorouracyl in Intestinal-Type Sinonasal Adenocarcinoma with Loss of Vision: A Case Report. Case Reports in Oncology, 2019, 12, 277-281.	0.7	4
21	Two siblings with Heimler syndrome caused by PEX1 variants: follow-up of ophthalmologic findings. Ophthalmic Genetics, 2021, 42, 480-485.	1.2	4
22	Bilateral congenital cataract with suspected lens-induced granulomatous uveitis. Journal of AAPOS, 2014, 18, 492-494.	0.3	3
23	Ophthalmological examination in neurofibromatosis type 1: a longâ€term retrospective analysis. Acta Ophthalmologica, 2018, 96, e1044-e1046.	1.1	3
24	How to approach complications of acute rhinosinusitis in children?. International Journal of Pediatric Otorhinolaryngology, 2020, 136, 110155.	1.0	3
25	Exudative Type 3 Retinal Arteriovenous Malformation in a Pediatric Patient. Case Reports in Ophthalmology, 2019, 9, 504-509.	0.7	2
26	Nephro-urologic, gastrointestinal, and ophthalmic findings. , 2005, , 105-122.		0
27	A Conjunctival Vascular Malformation as a Rare Presenting Sign of Wyburnâ€“Mason Syndrome. Journal of Pediatric Neurology, 2018, 16, 239-242.	0.2	0
28	Subconjunctival Hemorrhage in a Child with the Blue Rubber Bleb Nevus Syndrome on Treatment with Oral Propranolol. Case Reports in Ophthalmology, 2021, 12, 451-456.	0.7	0
29	Extraocular muscle hypoplasia associated with Axenfeld-Rieger syndrome. Strabismus, 2021, 29, 216-220.	0.7	0
30	Primary vitreous cysts. GMS Ophthalmology Cases, 2020, 10, Doc18.	0.1	0
31	Acute bilateral serous retinal detachments with spontaneous resolution in a 6-year-old boy. GMS Ophthalmology Cases, 2020, 10, Doc37.	0.1	0
32	High myopia and vitreal veils in a patient with Porettiâ€“ Boltshauser syndrome due to a novel homozygous LAMA1 mutation. Ophthalmic Genetics, 2022, , 1-5.	1.2	0