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	Association between near viewing and acute acquired esotropia in children during tablet and smartphone use. Strabismus, 2022, 30, 59-64.	0.7	10
2	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
3	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
4	Insights into multiple sclerosisâ€associated uveitis: a scoping review. Acta Ophthalmologica, 2021, 99, 592-603.	1.1	18
5	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
6	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
7	Two siblings with Heimler syndrome caused by PEX1 variants: follow-up of ophthalmologic findings. Ophthalmic Genetics, 2021, 42, 480-485.	1.2	4
8	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
9	Extraocular muscle hypoplasia associated with Axenfeld-Rieger syndrome. Strabismus, 2021, 29, 216-220.	0.7	0
10	A Retrospective Case Series of Uveal Effusion Syndrome. Journal of Glaucoma, 2020, 29, 995-998.	1.6	6
11	How to approach complications of acute rhinosinusitis in children?. International Journal of Pediatric Otorhinolaryngology, 2020, 136, 110155.	1.0	3
12	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
13	Primary vitreous cysts. GMS Ophthalmology Cases, 2020, 10, Doc18.	0.1	0
14	Acute bilateral serous retinal detachments with spontaneous resolution in a 6-year-old boy. GMS Ophthalmology Cases, 2020, 10, Doc37.	0.1	0
15	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
16	Mutation spectrum and clinical investigation of achromatopsia patients with mutations in the GNAT2 gene. Human Mutation, 2019, 40, 1145-1155.	2.5	15
17	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
18	Exudative Type 3 Retinal Arteriovenous Malformation in a Pediatric Patient. Case Reports in Ophthalmology, 2019, 9, 504-509.	0.7	2

#	Article	IF	Citations
19	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
20	A Conjunctival Vascular Malformation as a Rare Presenting Sign of Wyburnâ€“Mason Syndrome. Journal of Pediatric Neurology, 2018, 16, 239-242.	0.2	0
21	Ophthalmological examination in neurofibromatosis type 1: a longâ€term retrospective analysis. Acta Ophthalmologica, 2018, 96, e1044-e1046.	1.1	3
22	Use of the Masquerade Flap in Ablepharonâ€“Macrostomia Syndrome: A Case Report. Cornea, 2018, 37, 929-932.	1.7	7
23	Clinical Practice: A Proposed Standardized Ophthalmological Assessment for Patients with Cystinosis. Ophthalmology and Therapy, 2017, 6, 93-104.	2.3	20
24	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
25	Effect of Storage Conditions on Stability of Ophthalmological Compounded Cysteamine Eye Drops. JIMD Reports, 2017, 42, 47-51.	1.5	11
26	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
27	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
28	Cystinosis: a new perspective. Acta Clinica Belgica, 2016, 71, 131-137.	1.2	12
29	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
30	Bilateral congenital cataract with suspected lens-induced granulomatous uveitis. Journal of AAPOS, 2014, 18, 492-494.	0.3	3
31	Educational paper. European Journal of Pediatrics, 2012, 171, 887-893.	2.7	30
32	Nephro-urologic, gastrointestinal, and ophthalmic findings. , 2005, , 105-122.		0