

Eyvind RÃ¸dahl

List of Publications by Year in descending order

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Version: 2024-02-01

21
papers

430
citations

759233

12
h-index

794594

19
g-index

22
all docs

22
docs citations

22
times ranked

637
citing authors

#	ARTICLE	IF	CITATIONS
1	Primary Sjögren's syndrome and the eye. <i>Survey of Ophthalmology</i> , 2020, 65, 119-132.	4.0	79
2	A Second Decorin Frame Shift Mutation in a Family With Congenital Stromal Corneal Dystrophy. <i>American Journal of Ophthalmology</i> , 2006, 142, 520-521.	3.3	59
3	The intronic <i>ABCA4</i> c.5461T>C variant, frequently seen in patients with Stargardt disease, causes splice defects and reduced <i>ABCA4</i> protein level. <i>Acta Ophthalmologica</i> , 2017, 95, 240-246.	1.1	35
4	Mutations in <i>MAPKBP1</i> Cause Juvenile or Late-Onset Cilia-Independent Nephronophthisis. <i>American Journal of Human Genetics</i> , 2017, 100, 323-333.	6.2	29
5	Variants of Anterior Segment Dysgenesis and Cerebral Involvement in a Large Family With a Novel <i>COL4A1</i> Mutation. <i>American Journal of Ophthalmology</i> , 2013, 155, 946-953.e2.	3.3	27
6	Temporal hollowing and other adverse effects after lateral orbital wall decompression. <i>Acta Ophthalmologica</i> , 2016, 94, 793-797.	1.1	24
7	Role of Decorin Core Protein in Collagen Organisation in Congenital Stromal Corneal Dystrophy (CSCD). <i>PLoS ONE</i> , 2016, 11, e0147948.	2.5	23
8	Absence of the proteoglycan decorin reduces glucose tolerance in overfed male mice. <i>Scientific Reports</i> , 2019, 9, 4614.	3.3	21
9	A tyrosine kinase-activating variant Asn666Ser in <i>PDGFRB</i> causes a progeria-like condition in the severe end of Penttinen syndrome. <i>European Journal of Human Genetics</i> , 2019, 27, 574-581.	2.8	20
10	Recurrent, Activating Variants in the Receptor Tyrosine Kinase <i>DDR2</i> Cause Warburg-Cinotti Syndrome. <i>American Journal of Human Genetics</i> , 2018, 103, 976-983.	6.2	17
11	Chromosomal imbalances in some benign orbital tumours. <i>Acta Ophthalmologica</i> , 2005, 83, 385-391.	0.3	15
12	Development of Congenital Stromal Corneal Dystrophy Is Dependent on Export and Extracellular Deposition of Truncated Decorin. , 2015, 56, 2909.		15
13	High Myopia-Excavated Optic Disc Anomaly Associated With a Frameshift Mutation in the <i>MYC</i> -Binding Protein 2 Gene (<i>MYCBP2</i>). <i>American Journal of Ophthalmology</i> , 2015, 159, 973-979.e2.	3.3	13
14	Iris Malformation and Anterior Segment Dysgenesis in Mice and Humans With a Mutation in PI 3-Kinase. , 2017, 58, 3100.		11
15	Vision-related quality of life in patients with occipital stroke. <i>Acta Neurologica Scandinavica</i> , 2020, 141, 509-518.	2.1	9
16	Hyaluronic acid is superior to autologous fat for treatment of temporal hollowing after lateral orbital wall decompression: A prospective interventional trial. <i>Journal of Plastic, Reconstructive and Aesthetic Surgery</i> , 2019, 72, 973-981.	1.0	7
17	Clinical features and molecular genetics of patients with <i>ABCA4</i> -retinal dystrophies. <i>Acta Ophthalmologica</i> , 2021, 99, e733-e746.	1.1	6
18	Temperature-dependent autoactivation associated with clinical variability of <i>PDGFRB</i> Asn666 substitutions. <i>Human Molecular Genetics</i> , 2021, 30, 72-77.	2.9	6

#	ARTICLE	IF	CITATIONS
19	K ⁺ regulates relocation of Pellino ² to the site of NLRP3 inflammasome activation in macrophages. FEBS Letters, 2021, 595, 2437-2446.	2.8	6
20	Novel inflammatory biomarkers in thyroid eye disease. European Journal of Endocrinology, 2022, 187, 293-300.	3.7	5
21	Pellino ² in nonimmune cells: novel interaction partners and intracellular localization. FEBS Letters, 2021, 595, 2909-2921.	2.8	3