Eyvind RÃ¸dahl

List of Publications by Year
in descending order

21	430	759233 12	794594 19
papers	citations	h-index	g-index

22	22	22	637
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Novel inflammatory biomarkers in thyroid eye disease. European Journal of Endocrinology, 2022, 187, 293-300.	3.7	5
2	Clinical features and molecular genetics of patients with ABCA4â€retinal dystrophies. Acta Ophthalmologica, 2021, 99, e733-e746.	1.1	6
3	Temperature-dependent autoactivation associated with clinical variability of PDGFRB Asn666 substitutions. Human Molecular Genetics, 2021, 30, 72-77.	2.9	6
4	K⁺ regulates relocation of Pellinoâ€2 to the site of NLRP3 inflammasome activation in macrophages. FEBS Letters, 2021, 595, 2437-2446.	2.8	6
5	Pellinoâ€2 in nonimmune cells: novel interaction partners and intracellular localization. FEBS Letters, 2021, 595, 2909-2921.	2.8	3
6	Primary SjÃ¶grenâ€™s syndrome and the eye. Survey of Ophthalmology, 2020, 65, 119-132.	4.0	79
7	Visionâ€related quality of life in patients with occipital stroke. Acta Neurologica Scandinavica, 2020, 141, 509-518.	2.1	9
8	Absence of the proteoglycan decorin reduces glucose tolerance in overfed male mice. Scientific Reports, 2019, 9, 4614.	3.3	21
9	A tyrosine kinase-activating variant Asn666Ser in PDGFRB causes a progeria-like condition in the severe end of Penttinen syndrome. European Journal of Human Genetics, 2019, 27, 574-581.	2.8	20
10	Hyaluronic acid is superior to autologous fat for treatment of temporal hollowing after lateral orbital wall decompression: A prospective interventional trial. Journal of Plastic, Reconstructive and Aesthetic Surgery, 2019, 72, 973-981.	1.0	7
11	Recurrent, Activating Variants in the Receptor Tyrosine Kinase DDR2 Cause Warburg-Cinotti Syndrome. American Journal of Human Genetics, 2018, 103, 976-983.	6.2	17
12	Mutations in MAPKBP1 Cause Juvenile or Late-Onset Cilia-Independent Nephronophthisis. American Journal of Human Genetics, 2017, 100, 323-333.	6.2	29
13	The intronic <scp>ABCA</scp>4 c.5461â€10T>C variant, frequently seen in patients with Stargardt disease, causes splice defects and reduced <scp>ABCA</scp>4 protein level. Acta Ophthalmologica, 2017, 95, 240-246.	1.1	35
14	Iris Malformation and Anterior Segment Dysgenesis in Mice and Humans With a Mutation in PI 3-Kinase. , 2017, 58, 3100.		11
15	Role of Decorin Core Protein in Collagen Organisation in Congenital Stromal Corneal Dystrophy (CSCD). PLoS ONE, 2016, 11, e0147948.	2.5	23
16	Temporal hollowing and other adverse effects after lateral orbital wall decompression. Acta Ophthalmologica, 2016, 94, 793-797.	1.1	24
17	Development of Congenital Stromal Corneal Dystrophy Is Dependent on Export and Extracellular Deposition of Truncated Decorin. , 2015, 56, 2909.		15
18	High Myopiaâ€“Excavated Optic Disc Anomaly Associated With a Frameshift Mutation in the MYC-Binding Protein 2 Gene (MYCBP2). American Journal of Ophthalmology, 2015, 159, 973-979.e2.	3.3	13

#	ARTICLE	IF	CITATIONS
19	Variants of Anterior Segment Dysgenesis and Cerebral Involvement in a Large Family With a Novel COL4A1 Mutation. American Journal of Ophthalmology, 2013, 155, 946-953.e2.	3.3	27
20	A Second Decorin Frame Shift Mutation in a Family With Congenital Stromal Corneal Dystrophy. American Journal of Ophthalmology, 2006, 142, 520-521.	3.3	59
21	Chromosomal imbalances in some benign orbital tumours. Acta Ophthalmologica, 2005, 83, 385-391.	0.3	15