

Veronika Vaclavik

List of Publications by Year in descending order

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#	ARTICLE	IF	CITATIONS
1	New <i>COL6A6</i> Variant Causes Autosomal Dominant Retinitis Pigmentosa in a Four-Generation Family. , 2022, 63, 23.		1
2	Contribution of Whole-Genome Sequencing and Transcript Analysis to Decipher Retinal Diseases Associated with MFSD8 Variants. International Journal of Molecular Sciences, 2022, 23, 4294.	4.1	3
3	Malattia Leventinese: EFEMP1 R345W Variant Is a Hot Spot Mutation, Not a Founder Mutation. Ophthalmology Retina, 2020, 4, 1023.	2.4	1
4	Statins in ophthalmology. Survey of Ophthalmology, 2019, 64, 401-432.	4.0	29
5	Variability in clinical phenotypes of <i>PRPF8</i> -linked autosomal dominant retinitis pigmentosa correlates with differential PRPF8/SNRNP200 interactions. Ophthalmic Genetics, 2018, 39, 80-86.	1.2	8
6	CRX-linked macular dystrophy with intrafamilial variable expressivity. Ophthalmic Genetics, 2018, 39, 637-641.	1.2	5