Evaluation of TGFBI corneal dystrophy and molecular d

Eye

33, 874-881

DOI: 10.1038/s41433-019-0346-x

Citation Report

#	Article	IF	Citations
1	Aberrant DNA methylation of miRNAs in Fuchs endothelial corneal dystrophy. Scientific Reports, 2019, 9, 16385.	3.3	16
2	CRISPR Diagnosis and Therapeutics with Single Base Pair Precision. Trends in Molecular Medicine, 2020, 26, 337-350.	6.7	30
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10	Further evaluation of differential expression of keratoconus candidate genes in human corneas. PeerJ, 2020, 8, e9793.	2.0	16
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12	Recurrence and Visual Outcomes of Phototherapeutic Keratectomy in Lattice Corneal Dystrophy: A Cohort Study. Journal of Refractive Surgery, 2022, 38, 43-49.	2.3	2
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17	Autophagy in the normal and diseased cornea. Experimental Eye Research, 2022, 225, 109274.	2.6	5
18	Clinical and Histopathologic Characteristics and Template of the TGFBI p.(His626Arg) Missense Variant Lattice Corneal Dystrophy. Cornea, 2023, 42, 1124-1132.	1.7	1

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19	Variant Landscape of 15 Genes Involved in Corneal Dystrophies: Report of 30 Families and Comprehensive Analysis of the Literature. International Journal of Molecular Sciences, 2023, 24, 5012.	4.1	0
20	Release of frustration drives corneal amyloid disaggregation by brain chaperone. Communications Biology, 2023, 6, .	4.4	0
21	Novel Manifestation of Corneal Dystrophy After Keratorefractive Surgery. Cornea, 2024, 43, 404-408.	1.7	0