

Genome-wide meta-analyses of multiancestry cohorts identify new loci for refractive error and myopia

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Mutations in LRPAP1 Are Associated with Severe Myopia in Humans. <i>American Journal of Human Genetics</i> , 2013, 93, 313-320.	2.6	104
2	Genetic susceptibility and mechanisms for refractive error. <i>Clinical Genetics</i> , 2013, 84, 102-108.	1.0	42
3	Whole genome expression profiling of normal human fetal and adult ocular tissues. <i>Experimental Eye Research</i> , 2013, 116, 265-278.	1.2	19
4	Education influences the role of genetics in myopia. <i>European Journal of Epidemiology</i> , 2013, 28, 973-980.	2.5	102
5	Advances in the genomics of common eye diseases. <i>Human Molecular Genetics</i> , 2013, 22, R59-R65.	1.4	46
6	The Rotterdam Study: 2014 objectives and design update. <i>European Journal of Epidemiology</i> , 2013, 28, 889-926.	2.5	282
7	Nine Loci for Ocular Axial Length Identified through Genome-wide Association Studies, Including Shared Loci with Refractive Error. <i>American Journal of Human Genetics</i> , 2013, 93, 264-277.	2.6	139
9	Exome sequencing reveals CCDC111 mutation associated with high myopia. <i>Human Genetics</i> , 2013, 132, 913-921.	1.8	74
10	Birth Order and Myopia: What are the Messages to Readers?. <i>Ophthalmic Epidemiology</i> , 2013, 20, 333-334.	0.8	5
11	Focusing In on the Complex Genetics of Myopia. <i>PLoS Genetics</i> , 2013, 9, e1003442.	1.5	58
12	Advances in the genetics of eye diseases. <i>Current Opinion in Pediatrics</i> , 2013, 25, 645-652.	1.0	6
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15	Genome-wide association study identifies genetic risk underlying primary rhegmatogenous retinal detachment. <i>Human Molecular Genetics</i> , 2013, 22, 3174-3185.	1.4	34
16	Age of myopia onset in a British population-based twin cohort. <i>Ophthalmic and Physiological Optics</i> , 2013, 33, 339-345.	1.0	33
17	The Syndrome of Microcornea, Myopic Chorioretinal Atrophy, and Telecanthus (MMCAT) Is Caused by Mutations in <i>ADAMTS18</i> . <i>Human Mutation</i> , 2013, 34, 1195-1199.	1.1	56
19	Myopia Stabilization and Associated Factors Among Participants in the Correction of Myopia Evaluation Trial (COMET)., 2013, 54, 7871.		148
20	A Functional Polymorphism at the FGF10 Gene Is Associated With Extreme Myopia. , 2013, 54, 3265.		25

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22	Association study of IGF1 polymorphisms with susceptibility to high myopia in a Japanese population. <i>Clinical Ophthalmology</i> , 2013, 7, 2057.	0.9	10
23	The power of regional heritability analysis for rare and common variant detection: simulations and application to eye biometrical traits. <i>Frontiers in Genetics</i> , 2013, 4, 232.	1.1	36
24	Genome-Wide Meta-Analysis of Myopia and Hyperopia Provides Evidence for Replication of 11 Loci. <i>PLoS ONE</i> , 2014, 9, e107110.	1.1	40
25	Gene Profiling of Postnatal Mfrprd6 Mutant Eyes Reveals Differential Accumulation of Prss56, Visual Cycle and Phototransduction mRNAs. <i>PLoS ONE</i> , 2014, 9, e110299.	1.1	26
26	Genetic variation of the RASGRF1 regulatory region affects human hippocampus-dependent memory. <i>Frontiers in Human Neuroscience</i> , 2014, 8, 260.	1.0	22
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33	Visual activity and its association with myopia stabilisation. <i>Ophthalmic and Physiological Optics</i> , 2014, 34, 353-361.	1.0	31
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