

Li Jia Chen

List of Publications by Year in descending order

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131
papers

6,052
citations

94433

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134
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#	ARTICLE	IF	CITATIONS
1	Association of <i>SIX1-SIX6</i> polymorphisms with peripapillary retinal nerve fibre layer thickness in children. <i>British Journal of Ophthalmology</i> , 2023, 107, 1216-1222.	3.9	0
2	Differential compensatory role of internal astigmatism in school children and adults: The Hong Kong Children Eye Study. <i>Eye</i> , 2023, 37, 1107-1113.	2.1	2
3	Increase in Bruchâ€™s membrane opening minimum rim width with age in healthy children: the Hong Kong Children Eye Study. <i>British Journal of Ophthalmology</i> , 2023, 107, 1344-1349.	3.9	1
4	Genetic associations of central serous chorioretinopathy: a systematic review and meta-analysis. <i>British Journal of Ophthalmology</i> , 2022, 106, 1542-1548.	3.9	12
5	Myopia incidence and lifestyle changes among school children during the COVID-19 pandemic: a population-based prospective study. <i>British Journal of Ophthalmology</i> , 2022, 106, 1772-1778.	3.9	84
6	Three-Year Clinical Trial of Low-Concentration Atropine for Myopia Progression (LAMP) Study: Continued Versus Washout. <i>Ophthalmology</i> , 2022, 129, 308-321.	5.2	79
7	Delayed Diagnosis of Amblyopia in Children of Lower Socioeconomic Families: The Hong Kong Children Eye Study. <i>Ophthalmic Epidemiology</i> , 2022, 29, 621-628.	1.7	4
8	Global retinoblastoma survival and globe preservation: a systematic review and meta-analysis of associations with socioeconomic and health-care factors. <i>The Lancet Global Health</i> , 2022, 10, e380-e389.	6.3	25
9	The Association of Choroidal Thickening by Atropine With Treatment Effects for Myopia: Two-Year Clinical Trial of the Low-concentration Atropine for Myopia Progression (LAMP) Study. <i>American Journal of Ophthalmology</i> , 2022, 237, 130-138.	3.3	39
10	Thicker Retinal Nerve Fiber Layer with Age among Schoolchildren: The Hong Kong Children Eye Study. <i>Diagnostics</i> , 2022, 12, 500.	2.6	8
11	Myopia Genetics and Heredity. <i>Children</i> , 2022, 9, 382.	1.5	20
12	Vitamin D and Ocular Diseases: A Systematic Review. <i>International Journal of Molecular Sciences</i> , 2022, 23, 4226.	4.1	26
13	Prevalence and predictors of myopic macular degeneration among Asian adults: pooled analysis from the Asian Eye Epidemiology Consortium. <i>British Journal of Ophthalmology</i> , 2021, 105, 1140-1148.	3.9	19
14	Analysis of choriocapillaris perfusion and choroidal layer changes in patients with chronic central serous chorioretinopathy randomised to micropulse laser or photodynamic therapy. <i>British Journal of Ophthalmology</i> , 2021, 105, 555-560.	3.9	34
15	Ellipsoid zone optical intensity reduction as an early biomarker for retinitis pigmentosa. <i>Acta Ophthalmologica</i> , 2021, 99, e215-e221.	1.1	11
16	Exposure to Secondhand Smoke in Children is Associated with a Thinner Retinal Nerve Fiber Layer: The Hong Kong Children Eye Study. <i>American Journal of Ophthalmology</i> , 2021, 223, 91-99.	3.3	14
17	Genome-wide meta-analysis identifies 127 open-angle glaucoma loci with consistent effect across ancestries. <i>Nature Communications</i> , 2021, 12, 1258.	12.8	196
18	Association of polymorphisms in <i>ZFX1B</i> , <i>KCNQ5</i> and <i>GJD2</i> with myopia progression and polygenic risk prediction in children. <i>British Journal of Ophthalmology</i> , 2021, 105, 1751-1757.	3.9	5

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19	Fundus Autofluorescence and Optical Coherence Tomography Characteristics in Different Stages of Central Serous Chorioretinopathy. <i>Journal of Ophthalmology</i> , 2021, 2021, 1-9.	1.3	2
20	The association between attention-deficit/hyperactivity disorder and retinal nerve fiber/ganglion cell layer thickness measured by optical coherence tomography: a systematic review and meta-analysis. <i>International Ophthalmology</i> , 2021, 41, 3211-3221.	1.4	8
21	Optical Coherence Tomography Angiography Compared with Multimodal Imaging for Diagnosing Neovascular Central Serous Chorioretinopathy. <i>American Journal of Ophthalmology</i> , 2021, 232, 70-82.	3.3	10
22	Evaluation of Shared Genetic Susceptibility to High and Low Myopia and Hyperopia. <i>JAMA Ophthalmology</i> , 2021, 139, 601.	2.5	22
23	Optical coherence tomography biomarkers of photoreceptor degeneration in retinitis pigmentosa. <i>International Ophthalmology</i> , 2021, 41, 3949-3959.	1.4	5
24	A Multitask Deep-Learning System to Classify Diabetic Macular Edema for Different Optical Coherence Tomography Devices: A Multicenter Analysis. <i>Diabetes Care</i> , 2021, 44, 2078-2088.	8.6	27
25	Prevalence of strabismus and its risk factors among school aged children: The Hong Kong Children Eye Study. <i>Scientific Reports</i> , 2021, 11, 13820.	3.3	15
26	Comparison of choroidal thickness measurements between spectral domain optical coherence tomography and swept source optical coherence tomography in children. <i>Scientific Reports</i> , 2021, 11, 13749.	3.3	4
27	Age Effect on Treatment Responses to 0.05%, 0.025%, and 0.01% Atropine. <i>Ophthalmology</i> , 2021, 128, 1180-1187.	5.2	50
28	Association of Corneal Biomechanics Properties with Myopia in a Child and a Parent Cohort: Hong Kong Children Eye Study. <i>Diagnostics</i> , 2021, 11, 2357.	2.6	4
29	Association of the ZC3H11B, ZFH1B and SNTB1 genes with myopia of different severities. <i>British Journal of Ophthalmology</i> , 2020, 104, 1472-1476.	3.9	14
30	Two-Year Clinical Trial of the Low-Concentration Atropine for Myopia Progression (LAMP) Study. <i>Ophthalmology</i> , 2020, 127, 910-919.	5.2	164
31	Association of WNT7B and RSPO1 with Axial Length in School Children. , 2020, 61, 11.		6
32	Genetic associations of myopia severities and endophenotypes in children. <i>British Journal of Ophthalmology</i> , 2020, 105, bjophthalmol-2020-316728.	3.9	9
33	rad21 Is Involved in Corneal Stroma Development by Regulating Neural Crest Migration. <i>International Journal of Molecular Sciences</i> , 2020, 21, 7807.	4.1	3
34	Genetic Association of Age-Related Macular Degeneration and Polypoidal Choroidal Vasculopathy. <i>Asia-Pacific Journal of Ophthalmology</i> , 2020, 9, 104-109.	2.5	16
35	Independent Influence of Parental Myopia on Childhood Myopia in a Dose-Related Manner in 2,055 Trios: The Hong Kong Children Eye Study. <i>American Journal of Ophthalmology</i> , 2020, 218, 199-207.	3.3	25
36	Differential Effects on Ocular Biometrics by 0.05%, 0.025%, and 0.01% Atropine. <i>Ophthalmology</i> , 2020, 127, 1603-1611.	5.2	46

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37	Association of the <i>CAV1</i> & <i>CAV2</i> locus with normal-tension glaucoma in Chinese and Japanese. <i>Clinical and Experimental Ophthalmology</i> , 2020, 48, 658-665.	2.6	10
38	Identification of TIE2 as a susceptibility gene for neovascular age-related macular degeneration and polypoidal choroidal vasculopathy. <i>British Journal of Ophthalmology</i> , 2020, 105, bjophthalmol-2019-315746.	3.9	6
39	High prevalence of myopia in children and their parents in Hong Kong Chinese Population: the Hong Kong Children Eye Study. <i>Acta Ophthalmologica</i> , 2020, 98, e639.	1.1	83
40	Reduced photoreceptor outer segment layer thickness in mild commotio retinae without ellipsoid zone disruption. <i>Graefe's Archive for Clinical and Experimental Ophthalmology</i> , 2020, 258, 1437-1442.	1.9	1
41	Topical immunosuppressants for blepharitis in adults. <i>The Cochrane Library</i> , 2020, , .	2.8	0
42	Retrospective analysis of the possibility of predicting the COVID-19 outbreak from Internet searches and social media data, China, 2020. <i>Eurosurveillance</i> , 2020, 25, .	7.0	262
43	Clinical features and treatment outcomes of endogenous <i>Klebsiella</i> endophthalmitis: a 12-year review. <i>International Journal of Ophthalmology</i> , 2020, 13, 1933-1940.	1.1	4
44	Quantitative retinal microvasculature in children using swept-source optical coherence tomography: the Hong Kong Children Eye Study. <i>British Journal of Ophthalmology</i> , 2019, 103, 672-679.	3.9	51
45	Vitamin D and its pathway genes in myopia: systematic review and meta-analysis. <i>British Journal of Ophthalmology</i> , 2019, 103, 8-17.	3.9	27
46	Low-Concentration Atropine for Myopia Progression (LAMP) Study. <i>Ophthalmology</i> , 2019, 126, 113-124.	5.2	371
47	Identification and characterization of a novel promoter variant in placental growth factor for neovascular age-related macular degeneration. <i>Experimental Eye Research</i> , 2019, 187, 107748.	2.6	2
48	Experience of using adalimumab in treating sight-threatening paediatric or adolescent Behçet's disease-related uveitis. <i>Journal of Ophthalmic Inflammation and Infection</i> , 2019, 9, 14.	2.2	16
49	Association of Polymorphisms at the <i>SIX1-SIX6</i> Locus With Primary Open-Angle Glaucoma. , 2019, 60, 2914.		13
50	Association of Secondhand Smoking Exposure With Choroidal Thinning in Children Aged 6 to 8 Years. <i>JAMA Ophthalmology</i> , 2019, 137, 1406.	2.5	31
51	Coding Region Mutation Screening in Optineurin in Chinese Normal-Tension Glaucoma Patients. <i>Disease Markers</i> , 2019, 2019, 1-5.	1.3	9
52	Comorbidity of dementia and age-related macular degeneration calls for clinical awareness: a meta-analysis. <i>British Journal of Ophthalmology</i> , 2019, 103, bjophthalmol-2018-313277.	3.9	33
53	Latest Developments in Normal-Pressure Glaucoma: Diagnosis, Epidemiology, Genetics, Etiology, Causes and Mechanisms to Management. <i>Asia-Pacific Journal of Ophthalmology</i> , 2019, 8, 457-468.	2.5	40
54	Evaluation of the association of C5 with neovascular age-related macular degeneration and polypoidal choroidal vasculopathy. <i>Eye and Vision (London, England)</i> , 2019, 6, 34.	3.0	8

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55	A Cohesin Subunit Variant Identified from a Peripheral Sclerocornea Pedigree. <i>Disease Markers</i> , 2019, 2019, 1-8.	1.3	6
56	Association of the SIX6 locus with primary open angle glaucoma in southern Chinese and Japanese. <i>Experimental Eye Research</i> , 2019, 180, 129-136.	2.6	12
57	Spectral-Domain OCT Measurements in Alzheimer's Disease. <i>Ophthalmology</i> , 2019, 126, 497-510.	5.2	236
58	Association of antenatal steroid and risk of retinopathy of prematurity: a systematic review and meta-analysis. <i>British Journal of Ophthalmology</i> , 2018, 102, 1336-1341.	3.9	19
59	Association of the <i>PAX6</i> gene with extreme myopia rather than lower grade myopias. <i>British Journal of Ophthalmology</i> , 2018, 102, 570-574.	3.9	19
60	Analysis of multiple genetic loci reveals MPDZ-NF1B rs1324183 as a putative genetic marker for keratoconus. <i>British Journal of Ophthalmology</i> , 2018, 102, 1736-1741.	3.9	13
61	Topical Olopatadine in the Treatment of Allergic Conjunctivitis: A Systematic Review and Meta-analysis. <i>Ocular Immunology and Inflammation</i> , 2017, 25, 668-682.	1.8	25
62	Genome-Wide Association Study of Age-Related Eye Diseases in Chinese Population. <i>Essentials in Ophthalmology</i> , 2017, , 209-229.	0.1	0
63	Infectious keratitis and orthokeratology lens use: a systematic review. <i>Infection</i> , 2017, 45, 727-735.	4.7	60
64	Shared genetic variants for polypoidal choroidal vasculopathy and typical neovascular age-related macular degeneration in East Asians. <i>Journal of Human Genetics</i> , 2017, 62, 1049-1055.	2.3	35
65	Genetic associations for keratoconus: a systematic review and meta-analysis. <i>Scientific Reports</i> , 2017, 7, 4620.	3.3	54
66	Protective effects of an HTRA1 insertion-deletion variant against age-related macular degeneration in the Chinese populations. <i>Laboratory Investigation</i> , 2017, 97, 43-52.	3.7	8
67	HDL-cholesterol levels and risk of age-related macular degeneration: a multiethnic genetic study using Mendelian randomization. <i>International Journal of Epidemiology</i> , 2017, 46, 1891-1902.	1.9	73
68	Novel Mutations in PRPF31 Causing Retinitis Pigmentosa Identified Using Whole-Exome Sequencing. , 2017, 58, 6342.		23
69	Genetic Association of the <i>PARL-ABCC5-HTR3D-HTR3C</i> Locus With Primary Angle-Closure Glaucoma in Chinese. , 2017, 58, 4384.		10
70	Identification of <i>ANGPT2</i> as a New Gene for Neovascular Age-Related Macular Degeneration and Polypoidal Choroidal Vasculopathy in the Chinese and Japanese Populations. , 2017, 58, 1076.		29
71	Corneal blindness and current major treatment concern-graft scarcity. <i>International Journal of Ophthalmology</i> , 2017, 10, 1154-1162.	1.1	29
72	Molecular and Clinical Genetics of Retinoblastoma. <i>Essentials in Ophthalmology</i> , 2017, , 243-258.	0.1	0

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73	Association of <i>ABCG1</i> With Neovascular Age-Related Macular Degeneration and Polypoidal Choroidal Vasculopathy in Chinese and Japanese. , 2016, 57, 5758.		11
74	Identification of <i>PGF</i> as a New Gene for Neovascular Age-Related Macular Degeneration in a Chinese Population. , 2016, 57, 1714.		19
75	Association of Gestational Hypertensive Disorders with Retinopathy of prematurity: A Systematic Review and Meta-analysis. Scientific Reports, 2016, 6, 30732.	3.3	22
76	Antagonists of growth hormone-releasing hormone receptor induce apoptosis specifically in retinoblastoma cells. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 14396-14401.	7.1	30
77	Ethnic specific association of the <i>CAV1/CAV2</i> locus with primary open-angle glaucoma. Scientific Reports, 2016, 6, 27837.	3.3	29
78	Myopia Genetics—The Asia-Pacific Perspective. Asia-Pacific Journal of Ophthalmology, 2016, 5, 236-244.	2.5	22
79	Association of toll-like receptor 3 polymorphism rs3775291 with age-related macular degeneration: a systematic review and meta-analysis. Scientific Reports, 2016, 6, 19718.	3.3	18
80	Genome-wide association study identifies five new susceptibility loci for primary angle closure glaucoma. Nature Genetics, 2016, 48, 556-562.	21.4	147
81	Genetic risk of extranodal natural killer T-cell lymphoma: a genome-wide association study. Lancet Oncology, The, 2016, 17, 1240-1247.	10.7	84
82	Meta-analysis of gene-environment-wide association scans accounting for education level identifies additional loci for refractive error. Nature Communications, 2016, 7, 11008.	12.8	104
83	Refractive Errors and Concomitant Strabismus: A Systematic Review and Meta-analysis. Scientific Reports, 2016, 6, 35177.	3.3	32
84	HTRA1 promoter variant differentiates polypoidal choroidal vasculopathy from exudative age-related macular degeneration. Scientific Reports, 2016, 6, 28639.	3.3	24
85	Genetic Associations of Primary Angle-Closure Disease. Ophthalmology, 2016, 123, 1211-1221.	5.2	32
86	Genetic Associations of Interleukin-related Genes with Graves' Ophthalmopathy: a Systematic Review and Meta-analysis. Scientific Reports, 2015, 5, 16672.	3.3	21
87	SPP2 Mutations Cause Autosomal Dominant Retinitis Pigmentosa. Scientific Reports, 2015, 5, 14867.	3.3	24
88	Ethnic differences in the association of <i>SERPING1</i> with age-related macular degeneration and polypoidal choroidal vasculopathy. Scientific Reports, 2015, 5, 9424.	3.3	27
89	Association between hyperglycemia and retinopathy of prematurity: a systemic review and meta-analysis. Scientific Reports, 2015, 5, 9091.	3.3	46
90	A common variant near <i>TGFBR3</i> is associated with primary open angle glaucoma. Human Molecular Genetics, 2015, 24, 3880-3892.	2.9	105

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91	New loci and coding variants confer risk for age-related macular degeneration in East Asians. <i>Nature Communications</i> , 2015, 6, 6063.	12.8	147
92	Association of Genetic Variants with Polypoidal Choroidal Vasculopathy. <i>Ophthalmology</i> , 2015, 122, 1854-1865.	5.2	61
93	Association of PEDF polymorphisms with age-related macular degeneration and polypoidal choroidal vasculopathy: a systematic review and meta-analysis. <i>Scientific Reports</i> , 2015, 5, 9497.	3.3	13
94	Efficacy and Safety of Topical 0.05% Cyclosporine Eye Drops in the Treatment of Dry Eye Syndrome: A Systematic Review and Meta-analysis. <i>Ocular Surface</i> , 2015, 13, 213-225.	4.4	102
95	Whole-exome sequencing implicates UBE3D in age-related macular degeneration in East Asian populations. <i>Nature Communications</i> , 2015, 6, 6687.	12.8	40
96	ABCC5, a Gene That Influences the Anterior Chamber Depth, Is Associated with Primary Angle Closure Glaucoma. <i>PLoS Genetics</i> , 2014, 10, e1004089.	3.5	68
97	PAX6 Gene Associated with High Myopia. <i>Optometry and Vision Science</i> , 2014, 91, 419-429.	1.2	35
98	PRPF4 mutations cause autosomal dominant retinitis pigmentosa. <i>Human Molecular Genetics</i> , 2014, 23, 2926-2939.	2.9	98
99	Genes in the High-Density Lipoprotein Metabolic Pathway in Age-related Macular Degeneration and Polypoidal Choroidal Vasculopathy. <i>Ophthalmology</i> , 2014, 121, 911-916.	5.2	56
100	Common variants near ABCA1 and in PMM2 are associated with primary open-angle glaucoma. <i>Nature Genetics</i> , 2014, 46, 1115-1119.	21.4	160
101	Gender specific association of a complement component 3 polymorphism with polypoidal choroidal vasculopathy. <i>Scientific Reports</i> , 2014, 4, 7018.	3.3	27
102	Diabetes Mellitus and Risk of Age-Related Macular Degeneration: A Systematic Review and Meta-Analysis. <i>PLoS ONE</i> , 2014, 9, e108196.	2.5	70
103	Association of Common Variants in TCF4 and PTPRG with Fuchs' Corneal Dystrophy: A Systematic Review and Meta-Analysis. <i>PLoS ONE</i> , 2014, 9, e109142.	2.5	13
104	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.	6.2	139
105	Associations of the C2-CFB-RDBP-SKIV2L Locus with Age-related Macular Degeneration and Polypoidal Choroidal Vasculopathy. <i>Ophthalmology</i> , 2013, 120, 837-843.	5.2	38
106	Topical Cyclosporine in the Treatment of Allergic Conjunctivitis. <i>Ophthalmology</i> , 2013, 120, 2197-2203.	5.2	58
107	Genome-wide meta-analyses of multiethnicity cohorts identify multiple new susceptibility loci for refractive error and myopia. <i>Nature Genetics</i> , 2013, 45, 314-318.	21.4	398
108	Age-Related Macular Degeneration. <i>Asia-Pacific Journal of Ophthalmology</i> , 2013, 2, 211-212.	2.5	1

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109	Screening and Referral of Diabetic Retinopathy. <i>Asia-Pacific Journal of Ophthalmology</i> , 2013, 2, 310-316.	2.5	1
110	Genome-wide association study identifies ZFHX1B as a susceptibility locus for severe myopia. <i>Human Molecular Genetics</i> , 2013, 22, 5288-5294.	2.9	59
111	Targeted Sequencing of 179 Genes Associated with Hereditary Retinal Dystrophies and 10 Candidate Genes Identifies Novel and Known Mutations in Patients with Various Retinal Diseases. , 2013, 54, 2186.		63
112	Association of Genetic Variants on 8p21 and 4q12 with Age-Related Macular Degeneration in Asian Populations. , 2012, 53, 6576.		22
113	Genome-wide association analyses identify three new susceptibility loci for primary angle closure glaucoma. <i>Nature Genetics</i> , 2012, 44, 1142-1146.	21.4	196
114	Differentiation of Exudative Age-Related Macular Degeneration and Polypoidal Choroidal Vasculopathy in the <i>ARMS2</i> / <i>HTRA1</i> Locus. , 2012, 53, 3175.		43
115	SNP rs1533428 at 2p16.3 as a marker for late-onset primary open-angle glaucoma. <i>Molecular Vision</i> , 2012, 18, 1629-39.	1.1	28
116	Evaluation of NTF4 as a causative gene for primary open-angle glaucoma. <i>Molecular Vision</i> , 2012, 18, 1763-72.	1.1	18
117	Adjunctive Effect of Acupuncture to Refractive Correction on Anisometropic Amblyopia. <i>Ophthalmology</i> , 2011, 118, 1501-1511.	5.2	35
118	Interactive Expressions of HtrA1 and VEGF in Human Vitreous Humors and Fetal RPE Cells. , 2011, 52, 3706.		18
119	Long-Term In Vivo Imaging and Measurement of Dendritic Shrinkage of Retinal Ganglion Cells. , 2011, 52, 1539.		104
120	Acupuncture and Amblyopia—Reply. <i>JAMA Ophthalmology</i> , 2011, 129, 962.	2.4	1
121	Unfair Comparison of In-Office Acupuncture vs At-Home Patching for Amblyopia—Reply. <i>JAMA Ophthalmology</i> , 2011, 129, 963.	2.4	1
122	Association of NR2E3 but Not NRL Mutations with Retinitis Pigmentosa in the Chinese Population. , 2010, 51, 2229.		17
123	Randomized Controlled Trial of Patching vs Acupuncture for Anisometropic Amblyopia in Children Aged 7 to 12 Years. <i>JAMA Ophthalmology</i> , 2010, 128, 1510.	2.4	43
124	Compound Heterozygosity of Two Novel Truncation Mutations in <i>RP1</i> Causing Autosomal Recessive Retinitis Pigmentosa. , 2010, 51, 2236.		54
125	Development of novel drugs for ocular diseases: possibilities for individualized therapy. <i>Personalized Medicine</i> , 2010, 7, 371-386.	1.5	5
126	Evaluation of SPARC as a candidate gene of juvenile-onset primary open-angle glaucoma by mutation and copy number analyses. <i>Molecular Vision</i> , 2010, 16, 2016-25.	1.1	7

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127	Autosomal-Dominant Retinitis Pigmentosa Caused by a Mutation in SNRNP200, a Gene Required for Unwinding of U4/U6 snRNAs. <i>American Journal of Human Genetics</i> , 2009, 85, 617-627.	6.2	141
128	Multiple Gene Polymorphisms in the Complement Factor H Gene Are Associated with Exudative Age-Related Macular Degeneration in Chinese. , 2008, 49, 3312.		82
129	HTRA1 Variants in Exudative Age-Related Macular Degeneration and Interactions with Smoking and CFH. , 2008, 49, 2357.		81
130	Association of complement factor H polymorphisms with exudative age-related macular degeneration. <i>Molecular Vision</i> , 2006, 12, 1536-42.	1.1	74
131	Intracameral injection of lidocaine and carbachol. <i>Journal of Cataract and Refractive Surgery</i> , 2005, 31, 1855.	1.5	0