

Stuart MacGregor

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

263
papers

17,754
citations

63
h-index

129
g-index

317
ext. papers

21,159
ext. citations

9.7
avg, IF

6.42
L-index

#	Paper	IF	Citations
263	Is Genetic Risk for Sleep Apnea Causally Linked With Glaucoma Susceptibility? <i>2022</i> , 63, 25		0
262	Genetic Risk of Cardiovascular Disease Is Associated with Macular Ganglion Cell Inner Plexiform Layer Thinning in an Early Glaucoma Cohort. <i>Ophthalmology Science</i> , 2022 , 2, 100108		
261	Genetic overlap analysis of endometriosis and asthma identifies shared loci implicating sex hormones and thyroid signalling pathways.. <i>Human Reproduction</i> , 2022 , 37, 366-383	5.7	2
260	Evaluating a causal relationship between Complement Factor I protein level and advanced age-related macular degeneration using Mendelian Randomisation. <i>Ophthalmology Science</i> , 2022 , 100146		0
259	The APOE E4 allele is associated with faster rates of neuroretinal thinning in a prospective cohort study of suspect and early glaucoma. <i>Ophthalmology Science</i> , 2022 , 100159		0
258	A Polygenic Risk Score Predicts Intraocular Pressure Readings Outside Office Hours and Early Morning Spikes as Measured by Home Tonometry. <i>Ophthalmology Glaucoma</i> , 2021 , 4, 411-420	2.2	3
257	Germline variants are associated with increased primary melanoma tumor thickness at diagnosis. <i>Human Molecular Genetics</i> , 2021 , 29, 3578-3587	5.6	1
256	Attitudes towards polygenic risk testing in individuals with glaucoma. <i>Ophthalmology Glaucoma</i> , 2021 ,	2.2	2
255	Genetically determined risk of keratinocyte carcinoma and risk of other cancers. <i>International Journal of Epidemiology</i> , 2021 , 50, 1316-1324	7.8	
254	A multi-ethnic genome-wide association study implicates collagen matrix integrity and cell differentiation pathways in keratoconus. <i>Communications Biology</i> , 2021 , 4, 266	6.7	10
253	Normal-tension glaucoma is associated with cognitive impairment. <i>British Journal of Ophthalmology</i> , 2021 ,	5.5	4
252	Time spent outdoors in childhood is associated with reduced risk of myopia as an adult. <i>Scientific Reports</i> , 2021 , 11, 6337	4.9	9
251	IMI 2021 Yearly Digest 2021 , 62, 7		6
250	Symptom-level modelling unravels the shared genetic architecture of anxiety and depression. <i>Nature Human Behaviour</i> , 2021 , 5, 1432-1442	12.8	7
249	Genetic variation affects morphological retinal phenotypes extracted from UK Biobank optical coherence tomography images. <i>PLoS Genetics</i> , 2021 , 17, e1009497	6	5
248	Associations of sleep apnoea with glaucoma and age-related macular degeneration: an analysis in the United Kingdom Biobank and the Canadian Longitudinal Study on Aging. <i>BMC Medicine</i> , 2021 , 19, 104	11.4	3
247	Polyunsaturated Fatty Acid Levels and the Risk of Keratinocyte Cancer: A Mendelian Randomization Analysis. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2021 , 30, 1591-1598	4	3

246	Polygenic Risk Scores Stratify Keratinocyte Cancer Risk among Solid Organ Transplant Recipients with Chronic Immunosuppression in a High Ultraviolet Radiation Environment. <i>Journal of Investigative Dermatology</i> , 2021 , 141, 2866-2875.e2	4.3	1
245	Identification of a Locus Near Associated With Progression-Free Survival in Ovarian Cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2021 , 30, 1669-1680	4	2
244	Multitrait genetic association analysis identifies 50 new risk loci for gastro-oesophageal reflux, seven new loci for Barrett's oesophagus and provides insights into clinical heterogeneity in reflux diagnosis. <i>Gut</i> , 2021 ,	19.2	2
243	Evaluation of Shared Genetic Susceptibility to High and Low Myopia and Hyperopia. <i>JAMA Ophthalmology</i> , 2021 , 139, 601-609	3.9	4
242	Polygenic Risk Scores Allow Risk Stratification for Keratinocyte Cancer in Organ-Transplant Recipients. <i>Journal of Investigative Dermatology</i> , 2021 , 141, 325-333.e6	4.3	4
241	Cardiovascular Disease Predicts Structural and Functional Progression in Early Glaucoma. <i>Ophthalmology</i> , 2021 , 128, 58-69	7.3	10
240	Evaluating the role of alcohol consumption in breast and ovarian cancer susceptibility using population-based cohort studies and two-sample Mendelian randomization analyses. <i>International Journal of Cancer</i> , 2021 , 148, 1338-1350	7.5	4
239	The effects of eight serum lipid biomarkers on age-related macular degeneration risk: a Mendelian randomization study. <i>International Journal of Epidemiology</i> , 2021 , 50, 325-336	7.8	4
238	Germline variation in the insulin-like growth factor pathway and risk of Barrett's esophagus and esophageal adenocarcinoma. <i>Carcinogenesis</i> , 2021 , 42, 369-377	4.6	4
237	Genetic analysis of endometriosis and depression identifies shared loci and implicates causal links with gastric mucosa abnormality. <i>Human Genetics</i> , 2021 , 140, 529-552	6.3	8
236	A comprehensive re-assessment of the association between vitamin D and cancer susceptibility using Mendelian randomization. <i>Nature Communications</i> , 2021 , 12, 246	17.4	12
235	Genome-wide meta-analysis identifies 127 open-angle glaucoma loci with consistent effect across ancestries. <i>Nature Communications</i> , 2021 , 12, 1258	17.4	47
234	Methotrexate-related central neurotoxicity: clinical characteristics, risk factors and genome-wide association study in children treated for acute lymphoblastic leukemia. <i>Haematologica</i> , 2021 ,	6.6	2
233	Predicting the Future of Genetic Risk Profiling of Glaucoma: A Narrative Review. <i>JAMA Ophthalmology</i> , 2021 , 139, 224-231	3.9	2
232	Genetic analyses of gynecological disease identify genetic relationships between uterine fibroids and endometrial cancer, and a novel endometrial cancer genetic risk region at the WNT4 1p36.12 locus. <i>Human Genetics</i> , 2021 , 140, 1353-1365	6.3	5
231	Automated AI labeling of optic nerve head enables insights into cross-ancestry glaucoma risk and genetic discovery in >280,000 images from UKB and CLSA. <i>American Journal of Human Genetics</i> , 2021 , 108, 1204-1216	11	6
230	Association of Monogenic and Polygenic Risk With the Prevalence of Open-Angle Glaucoma. <i>JAMA Ophthalmology</i> , 2021 , 139, 1023-1028	3.9	4
229	Assessing the genetic relationship between gastro-esophageal reflux disease and risk of COVID-19 infection. <i>Human Molecular Genetics</i> , 2021 ,	5.6	1

228	Characteristics of p.Gln368Ter Myocilin Variant and Influence of Polygenic Risk on Glaucoma Penetrance in the UK Biobank. <i>Ophthalmology</i> , 2021 , 128, 1300-1311	7.3	5
227	Coffee consumption and risk of breast cancer: A Mendelian randomization study. <i>PLoS ONE</i> , 2021 , 16, e0236904	3.7	0
226	Gene Discovery Using Twins. <i>Twin Research and Human Genetics</i> , 2020 , 23, 90-93	2.2	
225	Association of Myopia and Intraocular Pressure With Retinal Detachment in European Descent Participants of the UK Biobank Cohort: A Mendelian Randomization Study. <i>JAMA Ophthalmology</i> , 2020 , 138, 671-678	3.9	6
224	Genome-wide association meta-analysis of corneal curvature identifies novel loci and shared genetic influences across axial length and refractive error. <i>Communications Biology</i> , 2020 , 3, 133	6.7	9
223	Meta-analysis of 542,934 subjects of European ancestry identifies new genes and mechanisms predisposing to refractive error and myopia. <i>Nature Genetics</i> , 2020 , 52, 401-407	36.3	68
222	Assessment of polygenic architecture and risk prediction based on common variants across fourteen cancers. <i>Nature Communications</i> , 2020 , 11, 3353	17.4	32
221	Body mass index and height and risk of cutaneous melanoma: Mendelian randomization analyses. <i>International Journal of Epidemiology</i> , 2020 , 49, 1236-1245	7.8	9
220	Multitrait analysis of glaucoma identifies new risk loci and enables polygenic prediction of disease susceptibility and progression. <i>Nature Genetics</i> , 2020 , 52, 160-166	36.3	78
219	Investigating the genetic and causal relationship between initiation or use of alcohol, caffeine, cannabis and nicotine. <i>Drug and Alcohol Dependence</i> , 2020 , 210, 107966	4.9	5
218	Genome-wide meta-analysis identifies novel loci associated with age-related macular degeneration. <i>Journal of Human Genetics</i> , 2020 , 65, 657-665	4.3	21
217	Genome-wide association meta-analyses combining multiple risk phenotypes provide insights into the genetic architecture of cutaneous melanoma susceptibility. <i>Nature Genetics</i> , 2020 , 52, 494-504	36.3	39
216	Genome-Wide Association Meta-Analysis of Single-Nucleotide Polymorphisms and Symptomatic Venous Thromboembolism during Therapy for Acute Lymphoblastic Leukemia and Lymphoma in Caucasian Children. <i>Cancers</i> , 2020 , 12,	6.6	2
215	The Genetics of Myopia 2020 , 95-132		3
214	An Intraocular Pressure Polygenic Risk Score Stratifies Multiple Primary Open-Angle Glaucoma Parameters Including Treatment Intensity. <i>Ophthalmology</i> , 2020 , 127, 901-907	7.3	12
213	Using Mendelian randomization to evaluate the causal relationship between serum C-reactive protein levels and age-related macular degeneration. <i>European Journal of Epidemiology</i> , 2020 , 35, 139-146	12.1	19
212	Association of Genetic Variation With Keratoconus. <i>JAMA Ophthalmology</i> , 2020 , 138, 174-181	3.9	24
211	Does polygenic risk influence associations between sun exposure and melanoma? A prospective cohort analysis. <i>British Journal of Dermatology</i> , 2020 , 183, 303-310	4	4

210	Overlapping genetic architecture between Parkinson disease and melanoma. <i>Acta Neuropathologica</i> , 2020 , 139, 347-364	14.3	12
209	Multiplex melanoma families are enriched for polygenic risk. <i>Human Molecular Genetics</i> , 2020 , 29, 2976-2985	5.8	3
208	Is there a causal relationship between vitamin D and melanoma risk? A Mendelian randomization study. <i>British Journal of Dermatology</i> , 2020 , 182, 97-103	4	8
207	Potential influence of socioeconomic status on genetic correlations between alcohol consumption measures and mental health. <i>Psychological Medicine</i> , 2020 , 50, 484-498	6.9	29
206	Genetic heterogeneity in self-reported depressive symptoms identified through genetic analyses of the PHQ-9. <i>Psychological Medicine</i> , 2020 , 50, 2385-2396	6.9	19
205	Rationale and protocol for the 7- and 8-year longitudinal assessments of eye health in a cohort of young adults in the Raine Study. <i>BMJ Open</i> , 2020 , 10, e033440	3	4
204	Genome-wide association analysis of 95 549 individuals identifies novel loci and genes influencing optic disc morphology. <i>Human Molecular Genetics</i> , 2019 , 28, 3680-3690	5.6	11
203	Gastroesophageal reflux GWAS identifies risk loci that also associate with subsequent severe esophageal diseases. <i>Nature Communications</i> , 2019 , 10, 4219	17.4	15
202	Genetic Correlations Between Diabetes and Glaucoma: An Analysis of Continuous and Dichotomous Phenotypes. <i>American Journal of Ophthalmology</i> , 2019 , 206, 245-255	4.9	6
201	Combined analysis of keratinocyte cancers identifies novel genome-wide loci. <i>Human Molecular Genetics</i> , 2019 , 28, 3148-3160	5.6	20
200	Exome-Derived Adiponectin-Associated Variants Implicate Obesity and Lipid Biology. <i>American Journal of Human Genetics</i> , 2019 , 105, 15-28	11	12
199	Assessment of melanoma candidate genes in a meta-analysis of 16534 melanoma cases. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2019 , 33, e369-e370	4.6	
198	New insight into human sweet taste: a genome-wide association study of the perception and intake of sweet substances. <i>American Journal of Clinical Nutrition</i> , 2019 , 109, 1724-1737	7	29
197	Implementing MR-PRESSO and GCTA-GSMR for pleiotropy assessment in Mendelian randomization studies from a practitioner's perspective. <i>Genetic Epidemiology</i> , 2019 , 43, 609-616	2.6	18
196	Risk factors for symptomatic venous thromboembolism during therapy for childhood acute lymphoblastic leukemia. <i>Thrombosis Research</i> , 2019 , 178, 132-138	8.2	8
195	Mendelian Randomization Study for Genetically Predicted Polyunsaturated Fatty Acids Levels on Overall Cancer Risk and Mortality. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2019 , 28, 1015-1023	4	10
194	Evidence of causal effect of major depression on alcohol dependence: findings from the psychiatric genomics consortium. <i>Psychological Medicine</i> , 2019 , 49, 1218-1226	6.9	33
193	No Association Between Vitamin D Status and Risk of Barrett's Esophagus or Esophageal Adenocarcinoma: A Mendelian Randomization Study. <i>Clinical Gastroenterology and Hepatology</i> , 2019 , 17, 2227-2235.e1	6.9	8

192	Effect of increased body mass index on risk of diagnosis or death from cancer. <i>British Journal of Cancer</i> , 2019 , 120, 565-570	8.7	13
191	Inherited Contributions to Melanoma Risk 2019 , 225-248		
190	Association between coffee consumption and overall risk of being diagnosed with or dying from cancer among >300 000 UK Biobank participants in a large-scale Mendelian randomization study. <i>International Journal of Epidemiology</i> , 2019 , 48, 1447-1456	7.8	15
189	Determining Possible Shared Genetic Architecture Between Myopia and Primary Open-Angle Glaucoma 2019 , 60, 3142-3149		5
188	Protein-coding variants implicate novel genes related to lipid homeostasis contributing to body-fat distribution. <i>Nature Genetics</i> , 2019 , 51, 452-469	36.3	44
187	Multi-trait genome-wide association study identifies new loci associated with optic disc parameters. <i>Communications Biology</i> , 2019 , 2, 435	6.7	10
186	Myocilin Gene Gln368Ter Variant Penetrance and Association With Glaucoma in Population-Based and Registry-Based Studies. <i>JAMA Ophthalmology</i> , 2019 , 137, 28-35	3.9	14
185	Polyunsaturated fatty acids and risk of melanoma: A Mendelian randomisation analysis. <i>International Journal of Cancer</i> , 2018 , 143, 508-514	7.5	11
184	Genetic overlap between endometriosis and endometrial cancer: evidence from cross-disease genetic correlation and GWAS meta-analyses. <i>Cancer Medicine</i> , 2018 , 7, 1978-1987	4.8	40
183	Genome-wide association study identifies seven novel susceptibility loci for primary open-angle glaucoma. <i>Human Molecular Genetics</i> , 2018 , 27, 1486-1496	5.6	72
182	Analysis combining correlated glaucoma traits identifies five new risk loci for open-angle glaucoma. <i>Scientific Reports</i> , 2018 , 8, 3124	4.9	25
181	Genome-wide association study of paclitaxel and carboplatin disposition in women with epithelial ovarian cancer. <i>Scientific Reports</i> , 2018 , 8, 1508	4.9	3
180	Assessment of moderate coffee consumption and risk of epithelial ovarian cancer: a Mendelian randomization study. <i>International Journal of Epidemiology</i> , 2018 , 47, 450-459	7.8	8
179	Height and overall cancer risk and mortality: evidence from a Mendelian randomisation study on 310,000 UK Biobank participants. <i>British Journal of Cancer</i> , 2018 , 118, 1262-1267	8.7	27
178	Interactions Between Genetic Variants and Environmental Factors Affect Risk of Esophageal Adenocarcinoma and Barrett's Esophagus. <i>Clinical Gastroenterology and Hepatology</i> , 2018 , 16, 1598-1606	6.9	14
177	Association Between Population Density and Genetic Risk for Schizophrenia. <i>JAMA Psychiatry</i> , 2018 , 75, 901-910	14.5	35
176	Accuracy of Inferred APOE Genotypes for a Range of Genotyping Arrays and Imputation Reference Panels. <i>Journal of Alzheimer's Disease</i> , 2018 , 64, 49-54	4.3	5
175	Combining common genetic variants and non-genetic risk factors to predict risk of cutaneous melanoma. <i>Human Molecular Genetics</i> , 2018 , 27, 4145-4156	5.6	21

174	Genome-wide association study of intraocular pressure uncovers new pathways to glaucoma. <i>Nature Genetics</i> , 2018 , 50, 1067-1071	36.3	86
173	Family-Based Genome-Wide Association Study of South Indian Pedigrees Supports WNT7B as a Central Corneal Thickness Locus 2018 , 59, 2495-2502		7
172	Genome-Wide Association Study Identifies a Susceptibility Locus for Comitant Esotropia and Suggests a Parent-of-Origin Effect 2018 , 59, 4054-4064		10
171	GWAS of lifetime cannabis use reveals new risk loci, genetic overlap with psychiatric traits, and a causal influence of schizophrenia. <i>Nature Neuroscience</i> , 2018 , 21, 1161-1170	25.5	270
170	Vitamin D and overall cancer risk and cancer mortality: a Mendelian randomization study. <i>Human Molecular Genetics</i> , 2018 , 27, 4315-4322	5.6	32
169	Assessing the Incremental Contribution of Common Genomic Variants to Melanoma Risk Prediction in Two Population-Based Studies. <i>Journal of Investigative Dermatology</i> , 2018 , 138, 2617-2624	4.3	36
168	Testosterone Pathway Genetic Polymorphisms in Relation to Primary Open-Angle Glaucoma: An Analysis in Two Large Datasets 2018 , 59, 629-636		9
167	Genomic locus modulating corneal thickness in the mouse identifies POU6F2 as a potential risk of developing glaucoma. <i>PLoS Genetics</i> , 2018 , 14, e1007145	6	19
166	Inherited Contributions to Melanoma Risk 2018 , 1-23		
165	Novel pleiotropic risk loci for melanoma and nevus density implicate multiple biological pathways. <i>Nature Communications</i> , 2018 , 9, 4774	17.4	47
164	Understanding the role of bitter taste perception in coffee, tea and alcohol consumption through Mendelian randomization. <i>Scientific Reports</i> , 2018 , 8, 16414	4.9	22
163	Cell-type-specific eQTL of primary melanocytes facilitates identification of melanoma susceptibility genes. <i>Genome Research</i> , 2018 , 28, 1621-1635	9.7	33
162	Genome-wide association meta-analysis highlights light-induced signaling as a driver for refractive error. <i>Nature Genetics</i> , 2018 , 50, 834-848	36.3	135
161	Cross-ancestry genome-wide association analysis of corneal thickness strengthens link between complex and Mendelian eye diseases. <i>Nature Communications</i> , 2018 , 9, 1864	17.4	37
160	New insights into the genetics of primary open-angle glaucoma based on meta-analyses of intraocular pressure and optic disc characteristics. <i>Human Molecular Genetics</i> , 2017 , 26, 438-453	5.6	80
159	A Novel Approach for Pathway Analysis of GWAS Data Highlights Role of BMP Signaling and Muscle Cell Differentiation in Colorectal Cancer Susceptibility. <i>Twin Research and Human Genetics</i> , 2017 , 20, 1-9	2.2	24
158	Haplotype reference consortium panel: Practical implications of imputations with large reference panels. <i>Human Mutation</i> , 2017 , 38, 1025-1032	4.7	20
157	Genome-Wide Association Shows that Pigmentation Genes Play a Role in Skin Aging. <i>Journal of Investigative Dermatology</i> , 2017 , 137, 1887-1894	4.3	30

156	Genetically low vitamin D concentrations and myopic refractive error: a Mendelian randomization study. <i>International Journal of Epidemiology</i> , 2017 , 46, 1882-1890	7.8	34
155	Meta-analysis identifies five novel loci associated with endometriosis highlighting key genes involved in hormone metabolism. <i>Nature Communications</i> , 2017 , 8, 15539	17.4	151
154	Germline variation in inflammation-related pathways and risk of Barrett's oesophagus and oesophageal adenocarcinoma. <i>Gut</i> , 2017 , 66, 1739-1747	19.2	24
153	A common intronic variant of PARP1 confers melanoma risk and mediates melanocyte growth via regulation of MITF. <i>Nature Genetics</i> , 2017 , 49, 1326-1335	36.3	36
152	Contribution of Mutations in Known Mendelian Glaucoma Genes to Advanced Early-Onset Primary Open-Angle Glaucoma 2017 , 58, 1537-1544		11
151	Whole exome sequencing implicates eye development, the unfolded protein response and plasma membrane homeostasis in primary open-angle glaucoma. <i>PLoS ONE</i> , 2017 , 12, e0172427	3.7	8
150	Analyses of germline variants associated with ovarian cancer survival identify functional candidates at the 1q22 and 19p12 outcome loci. <i>Oncotarget</i> , 2017 , 8, 64670-64684	3.3	5
149	Genome-wide association studies in oesophageal adenocarcinoma and Barrett's oesophagus: a large-scale meta-analysis. <i>Lancet Oncology</i> , 2016 , 17, 1363-1373	21.7	94
148	Assessment of polygenic effects links primary open-angle glaucoma and age-related macular degeneration. <i>Scientific Reports</i> , 2016 , 6, 26885	4.9	17
147	Pooled genome wide association detects association upstream of FCRL3 with Graves' disease. <i>BMC Genomics</i> , 2016 , 17, 939	4.5	7
146	Meta-analysis of gene-environment-wide association scans accounting for education level identifies additional loci for refractive error. <i>Nature Communications</i> , 2016 , 7, 11008	17.4	79
145	Childhood gene-environment interactions and age-dependent effects of genetic variants associated with refractive error and myopia: The CREAM Consortium. <i>Scientific Reports</i> , 2016 , 6, 25853	4.9	57
144	Chronic gastroesophageal reflux disease shares genetic background with esophageal adenocarcinoma and Barrett's esophagus. <i>Human Molecular Genetics</i> , 2016 , 25, 828-35	5.6	26
143	Genome-wide association analysis identifies TXNRD2, ATXN2 and FOXC1 as susceptibility loci for primary open-angle glaucoma. <i>Nature Genetics</i> , 2016 , 48, 189-94	36.3	159
142	Germline polymorphisms in an enhancer of PSIP1 are associated with progression-free survival in epithelial ovarian cancer. <i>Oncotarget</i> , 2016 , 7, 6353-68	3.3	19
141	Association of Polymorphisms in MACRO Domain Containing 2 With Thyroid-Associated Orbitopathy 2016 , 57, 3129-37		6
140	Author Response: Stronger Association of CDKN2B-AS1 Variants in Female Normal-Tension Glaucoma Patients in a Japanese Population 2016 , 57, 6418		
139	Genetic Association at the 9p21 Glaucoma Locus Contributes to Sex Bias in Normal-Tension Glaucoma 2016 , 57, 3416-21		23

138	Polymorphisms in genes in the androgen pathway and risk of Barrett's esophagus and esophageal adenocarcinoma. <i>International Journal of Cancer</i> , 2016 , 138, 1146-52	7.5	10
137	Rare variants in optic disc area gene enriched in primary open-angle glaucoma. <i>Molecular Genetics & Genomic Medicine</i> , 2016 , 4, 624-633	2.3	5
136	GWAS study using DNA pooling strategy identifies association of variant rs4910623 in OR52B4 gene with anti-VEGF treatment response in age-related macular degeneration. <i>Scientific Reports</i> , 2016 , 6, 37924	4.9	18
135	Assessing the genetic architecture of epithelial ovarian cancer histological subtypes. <i>Human Genetics</i> , 2016 , 135, 741-56	6.3	18
134	When do myopia genes have their effect? Comparison of genetic risks between children and adults. <i>Genetic Epidemiology</i> , 2016 , 40, 756-766	2.6	26
133	Association of vitamin D levels and risk of ovarian cancer: a Mendelian randomization study. <i>International Journal of Epidemiology</i> , 2016 , 45, 1619-1630	7.8	77
132	Sweet Taste Perception is Associated with Body Mass Index at the Phenotypic and Genotypic Level. <i>Twin Research and Human Genetics</i> , 2016 , 19, 465-71	2.2	8
131	Assessing the Genetic Predisposition of Education on Myopia: A Mendelian Randomization Study. <i>Genetic Epidemiology</i> , 2016 , 40, 66-72	2.6	39
130	Genome-wide meta-analysis identifies five new susceptibility loci for cutaneous malignant melanoma. <i>Nature Genetics</i> , 2015 , 47, 987-995	36.3	162
129	Genome-wide Analysis Identifies Novel Loci Associated with Ovarian Cancer Outcomes: Findings from the Ovarian Cancer Association Consortium. <i>Clinical Cancer Research</i> , 2015 , 21, 5264-76	12.9	24
128	Retinal microvessels reflect familial vulnerability to psychotic symptoms: A comparison of twins discordant for psychotic symptoms and controls. <i>Schizophrenia Research</i> , 2015 , 164, 47-52	3.6	33
127	Genome-wide association study for refractive astigmatism reveals genetic co-determination with spherical equivalent refractive error: the CREAM consortium. <i>Human Genetics</i> , 2015 , 134, 131-46	6.3	20
126	LocusTrack: Integrated visualization of GWAS results and genomic annotation. <i>Source Code for Biology and Medicine</i> , 2015 , 10, 1	1.9	24
125	Genome-wide enrichment analysis between endometriosis and obesity-related traits reveals novel susceptibility loci. <i>Human Molecular Genetics</i> , 2015 , 24, 1185-99	5.6	57
124	VEGAS2: Software for More Flexible Gene-Based Testing. <i>Twin Research and Human Genetics</i> , 2015 , 18, 86-91	2.2	188
123	Identification of myopia-associated WNT7B polymorphisms provides insights into the mechanism underlying the development of myopia. <i>Nature Communications</i> , 2015 , 6, 6689	17.4	56
122	Shared genetics underlying epidemiological association between endometriosis and ovarian cancer. <i>Human Molecular Genetics</i> , 2015 , 24, 5955-64	5.6	48
121	Survival outcomes in patients with multiple primary melanomas. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2015 , 29, 2120-7	4.6	18

120	Pleiotropic analysis of cancer risk loci on esophageal adenocarcinoma risk. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2015 , 24, 1801-3	4	7
119	WNT10A exonic variant increases the risk of keratoconus by decreasing corneal thickness. <i>Human Molecular Genetics</i> , 2015 , 24, 5060-8	5.6	50
118	PARP1 polymorphisms play opposing roles in melanoma occurrence and survival. <i>International Journal of Cancer</i> , 2015 , 136, 2488-9	7.5	6
117	Polymorphisms in Genes of Relevance for Oestrogen and Oxytocin Pathways and Risk of Barrett's Oesophagus and Oesophageal Adenocarcinoma: A Pooled Analysis from the BEACON Consortium. <i>PLoS ONE</i> , 2015 , 10, e0138738	3.7	9
116	Accurate Imputation-Based Screening of Gln368Ter Myocilin Variant in Primary Open-Angle Glaucoma 2015 , 56, 5087-93		15
115	Genetic burden associated with varying degrees of disease severity in endometriosis. <i>Molecular Human Reproduction</i> , 2015 , 21, 594-602	4.4	22
114	Genetic and environmental factors in conjunctival UV autofluorescence. <i>JAMA Ophthalmology</i> , 2015 , 133, 406-12	3.9	16
113	A common variant near TGFBR3 is associated with primary open angle glaucoma. <i>Human Molecular Genetics</i> , 2015 , 24, 3880-92	5.6	84
112	Association between endometriosis and the interleukin 1A (IL1A) locus. <i>Human Reproduction</i> , 2015 , 30, 239-48	5.7	49
111	ARHGEF12 influences the risk of glaucoma by increasing intraocular pressure. <i>Human Molecular Genetics</i> , 2015 , 24, 2689-99	5.6	62
110	Meta-analysis of Genome-Wide Association Studies Identifies Novel Loci Associated With Optic Disc Morphology. <i>Genetic Epidemiology</i> , 2015 , 39, 207-16	2.6	58
109	Genome-wide analysis of multi-ancestry cohorts identifies new loci influencing intraocular pressure and susceptibility to glaucoma. <i>Nature Genetics</i> , 2014 , 46, 1126-1130	36.3	171
108	Common variants near ABCA1, AFAP1 and GMDS confer risk of primary open-angle glaucoma. <i>Nature Genetics</i> , 2014 , 46, 1120-1125	36.3	141
107	Most common 'sporadic' cancers have a significant germline genetic component. <i>Human Molecular Genetics</i> , 2014 , 23, 6112-8	5.6	74
106	Identification of a melanoma susceptibility locus and somatic mutation in TET2. <i>Carcinogenesis</i> , 2014 , 35, 2097-101	4.6	38
105	Meta-analysis of genome-wide association studies identifies novel loci that influence cupping and the glaucomatous process. <i>Nature Communications</i> , 2014 , 5, 4883	17.4	71
104	ABCA transporter gene expression and poor outcome in epithelial ovarian cancer. <i>Journal of the National Cancer Institute</i> , 2014 , 106,	9.7	79
103	Obesity and risk of esophageal adenocarcinoma and Barrett's esophagus: a Mendelian randomization study. <i>Journal of the National Cancer Institute</i> , 2014 , 106,	9.7	105

102	Associations between depression and anxiety symptoms and retinal vessel caliber in adolescents and young adults. <i>Psychosomatic Medicine</i> , 2014 , 76, 732-8	3.7	19
101	The effect on melanoma risk of genes previously associated with telomere length. <i>Journal of the National Cancer Institute</i> , 2014 , 106,	9.7	97
100	A genome-wide association study of intra-ocular pressure suggests a novel association in the gene FAM125B in the TwinsUK cohort. <i>Human Molecular Genetics</i> , 2014 , 23, 3343-8	5.6	32
99	Genome-wide association study for identification of candidate SNPs associated with carboplatin and paclitaxel clearance in ovarian cancer patients.. <i>Journal of Clinical Oncology</i> , 2014 , 32, 5563-5563	2.2	
98	ABCB1 (MDR1) polymorphisms and ovarian cancer progression and survival: a comprehensive analysis from the Ovarian Cancer Association Consortium and The Cancer Genome Atlas. <i>Gynecologic Oncology</i> , 2013 , 131, 8-14	4.9	39
97	A genome wide survey supports the involvement of large copy number variants in schizophrenia with and without intellectual disability. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2013 , 162B, 847-54	3.5	13
96	Germline genetic contributions to risk for esophageal adenocarcinoma, Barrett's esophagus, and gastroesophageal reflux. <i>Journal of the National Cancer Institute</i> , 2013 , 105, 1711-8	9.7	75
95	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, 264-77	11	116
94	Genome-wide meta-analyses of multiancestry cohorts identify multiple new susceptibility loci for refractive error and myopia. <i>Nature Genetics</i> , 2013 , 45, 314-8	36.3	314
93	A variant in FTO shows association with melanoma risk not due to BMI. <i>Nature Genetics</i> , 2013 , 45, 428-32, 432e1	36.3	95
92	Association between putative functional variants in the PSMB9 gene and risk of melanoma--re-analysis of published melanoma genome-wide association studies. <i>Pigment Cell and Melanoma Research</i> , 2013 , 26, 392-401	4.5	4
91	A genome-wide association study identifies new susceptibility loci for esophageal adenocarcinoma and Barrett's esophagus. <i>Nature Genetics</i> , 2013 , 45, 1487-93	36.3	151
90	Association between functional polymorphisms in genes involved in the MAPK signaling pathways and cutaneous melanoma risk. <i>Carcinogenesis</i> , 2013 , 34, 885-92	4.6	6
89	Genome-wide association analyses identify multiple loci associated with central corneal thickness and keratoconus. <i>Nature Genetics</i> , 2013 , 45, 155-63	36.3	222
88	Evaluating the association between keratoconus and the corneal thickness genes in an independent Australian population 2013 , 54, 8224-8		48
87	Identification of a candidate gene for astigmatism 2013 , 54, 1260-7		23
86	Copy number variation at chromosome 5q21.2 is associated with intraocular pressure 2013 , 54, 3607-12		10
85	Genetic loci for retinal arteriolar microcirculation. <i>PLoS ONE</i> , 2013 , 8, e65804	3.7	19

84	Interrogation of the platelet-derived growth factor receptor alpha locus and corneal astigmatism in Australians of Northern European ancestry: results of a genome-wide association study. <i>Molecular Vision</i> , 2013 , 19, 1238-46	2.3	7
83	Genome-wide association meta-analysis identifies new endometriosis risk loci. <i>Nature Genetics</i> , 2012 , 44, 1355-9	36.3	214
82	Promoter polymorphisms in two overlapping 6p25 genes implicate mitochondrial proteins in cognitive deficit in schizophrenia. <i>Molecular Psychiatry</i> , 2012 , 17, 1328-39	15.1	14
81	A genome-wide analysis of 'Bounty' descendants implicates several novel variants in migraine susceptibility. <i>Neurogenetics</i> , 2012 , 13, 261-6	3	28
80	Association between in vivo alcohol metabolism and genetic variation in pathways that metabolize the carbon skeleton of ethanol and NADH reoxidation in the alcohol challenge twin study. <i>Alcoholism: Clinical and Experimental Research</i> , 2012 , 36, 2074-85	3.7	9
79	Genetic variants near PDGFRA are associated with corneal curvature in Australians 2012 , 53, 7131-6		31
78	Melanoma genetics: recent findings take us beyond well-traveled pathways. <i>Journal of Investigative Dermatology</i> , 2012 , 132, 1763-74	4.3	65
77	Genome-wide association study of major depressive disorder: new results, meta-analysis, and lessons learned. <i>Molecular Psychiatry</i> , 2012 , 17, 36-48	15.1	335
76	Common genetic determinants of intraocular pressure and primary open-angle glaucoma. <i>PLoS Genetics</i> , 2012 , 8, e1002611	6	131
75	Meta-analysis combining new and existing data sets confirms that the TERT-CLPTM1L locus influences melanoma risk. <i>Journal of Investigative Dermatology</i> , 2012 , 132, 485-7	4.3	38
74	Common variants at the MHC locus and at chromosome 16q24.1 predispose to Barrett's esophagus. <i>Nature Genetics</i> , 2012 , 44, 1131-6	36.3	139
73	Genome-wide association study for ovarian cancer susceptibility using pooled DNA. <i>Twin Research and Human Genetics</i> , 2012 , 15, 615-623	2.2	8
72	Genome-wide association study identifies three new melanoma susceptibility loci. <i>Nature Genetics</i> , 2011 , 43, 1108-13	36.3	203
71	Association of polymorphisms in the hepatocyte growth factor gene promoter with keratoconus 2011 , 52, 8514-9		101
70	Functional polymorphisms in the TERT promoter are associated with risk of serous epithelial ovarian and breast cancers. <i>PLoS ONE</i> , 2011 , 6, e24987	3.7	41
69	Genome-wide association study identifies a locus at 7p15.2 associated with endometriosis. <i>Nature Genetics</i> , 2011 , 43, 51-4	36.3	227
68	Genome-wide association study identifies susceptibility loci for open angle glaucoma at TMCO1 and CDKN2B-AS1. <i>Nature Genetics</i> , 2011 , 43, 574-8	36.3	329
67	GWAS findings for human iris patterns: associations with variants in genes that influence normal neuronal pattern development. <i>American Journal of Human Genetics</i> , 2011 , 89, 334-43	11	47

66	Genome-wide association study identifies novel loci predisposing to cutaneous melanoma. <i>Human Molecular Genetics</i> , 2011 , 20, 5012-23	5.6	164
65	A 3p26-3p25 genetic linkage finding for DSM-IV major depression in heavy smoking families. <i>American Journal of Psychiatry</i> , 2011 , 168, 848-52	11.9	33
64	Platinum sensitivity-related germline polymorphism discovered via a cell-based approach and analysis of its association with outcome in ovarian cancer patients. <i>Clinical Cancer Research</i> , 2011 , 17, 5490-500	12.9	55
63	The role of KRAS rs61764370 in invasive epithelial ovarian cancer: implications for clinical testing. <i>Clinical Cancer Research</i> , 2011 , 17, 3742-50	12.9	45
62	A novel recurrent mutation in MITF predisposes to familial and sporadic melanoma. <i>Nature</i> , 2011 , 480, 99-103	50.4	335
61	Genome-wide association study identifies a new melanoma susceptibility locus at 1q21.3. <i>Nature Genetics</i> , 2011 , 43, 1114-8	36.3	126
60	Legacy of mutiny on the Bounty: founder effect and admixture on Norfolk Island. <i>European Journal of Human Genetics</i> , 2010 , 18, 67-72	5.3	23
59	European and Polynesian admixture in the Norfolk Island population. <i>Heredity</i> , 2010 , 105, 229-34	3.6	18
58	Common variants near CAV1 and CAV2 are associated with primary open-angle glaucoma. <i>Nature Genetics</i> , 2010 , 42, 906-9	36.3	303
57	A genome-wide association study for myopia and refractive error identifies a susceptibility locus at 15q25. <i>Nature Genetics</i> , 2010 , 42, 902-5	36.3	179
56	A genomewide association study of nicotine and alcohol dependence in Australian and Dutch populations. <i>Twin Research and Human Genetics</i> , 2010 , 13, 10-29	2.2	90
55	Common genetic variants near the Brittle Cornea Syndrome locus ZNF469 influence the blinding disease risk factor central corneal thickness. <i>PLoS Genetics</i> , 2010 , 6, e1000947	6	106
54	Genome-wide association identifies ATOH7 as a major gene determining human optic disc size. <i>Human Molecular Genetics</i> , 2010 , 19, 2716-24	5.6	118
53	Evaluation of candidate stromal epithelial cross-talk genes identifies association between risk of serous ovarian cancer and TERT, a cancer susceptibility "hot-spot". <i>PLoS Genetics</i> , 2010 , 6, e1001016	6	42
52	Four novel Loci (19q13, 6q24, 12q24, and 5q14) influence the microcirculation in vivo. <i>PLoS Genetics</i> , 2010 , 6, e1001184	6	111
51	A Genomewide Association Study of Nicotine and Alcohol Dependence in Australian and Dutch Populations. <i>Twin Research and Human Genetics</i> , 2010 , 13, 11-29	2.2	2
50	A versatile gene-based test for genome-wide association studies. <i>American Journal of Human Genetics</i> , 2010 , 87, 139-45	11	648
49	Optimal selection of markers from DNA pooling experiments. <i>Behavior Genetics</i> , 2010 , 40, 46-7; discussion 48	3.2	1

48	Hormonal responses differ when playing violent video games against an ingroup and outgroup. <i>Evolution and Human Behavior</i> , 2010 , 31, 201-209	4	53
47	A simple and fast two-locus quality control test to detect false positives due to batch effects in genome-wide association studies. <i>Genetic Epidemiology</i> , 2010 , 34, 854-62	2.6	30
46	Evaluation of association of HNF1B variants with diverse cancers: collaborative analysis of data from 19 genome-wide association studies. <i>PLoS ONE</i> , 2010 , 5, e10858	3.7	24
45	Associations of ADH and ALDH2 gene variation with self report alcohol reactions, consumption and dependence: an integrated analysis. <i>Human Molecular Genetics</i> , 2009 , 18, 580-93	5.6	157
44	Rapid inexpensive genome-wide association using pooled whole blood. <i>Genome Research</i> , 2009 , 19, 2075-80	9.7	35
43	Bipolar disorder in the Bulgarian Gypsies: genetic heterogeneity in a young founder population. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2009 , 150B, 191-201	3.5	9
42	P2RX7: A bipolar and unipolar disorder candidate susceptibility gene?. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2009 , 150B, 1063-9	3.5	47
41	Neuregulin 1 and age of onset in the major psychoses. <i>Journal of Neural Transmission</i> , 2009 , 116, 479-86	4.3	5
40	Common polygenic variation contributes to risk of schizophrenia and bipolar disorder. <i>Nature</i> , 2009 , 460, 748-52	50.4	3568
39	Common variants in the trichohyalin gene are associated with straight hair in Europeans. <i>American Journal of Human Genetics</i> , 2009 , 85, 750-5	11	200
38	Rare chromosomal deletions and duplications increase risk of schizophrenia. <i>Nature</i> , 2008 , 455, 237-41	50.4	1251
37	Common sequence variants on 20q11.22 confer melanoma susceptibility. <i>Nature Genetics</i> , 2008 , 40, 838-43	3.4	188
36	The role of GABRA2 in alcohol dependence, smoking, and illicit drug use in an Australian population sample. <i>Alcoholism: Clinical and Experimental Research</i> , 2008 , 32, 1721-31	3.7	44
35	Vitamin D receptor gene polymorphisms have negligible effect on human height. <i>Twin Research and Human Genetics</i> , 2008 , 11, 488-94	2.2	11
34	Effects of GABRA2 variation on physiological, psychomotor and subjective responses in the alcohol challenge twin study. <i>Twin Research and Human Genetics</i> , 2008 , 11, 174-82	2.2	16
33	Linkage analysis in a large family from Pakistan with depression and a high incidence of consanguineous marriages. <i>Human Heredity</i> , 2008 , 66, 190-8	1.1	2
32	Highly cost-efficient genome-wide association studies using DNA pools and dense SNP arrays. <i>Nucleic Acids Research</i> , 2008 , 36, e35	20.1	87
31	Optimal two-stage testing for family-based genome-wide association studies. <i>American Journal of Human Genetics</i> , 2008 , 82, 797-9; author reply 799-800	11	6

30	Most pooling variation in array-based DNA pooling is attributable to array error rather than pool construction error. <i>European Journal of Human Genetics</i> , 2007 , 15, 501-4	5.3	45
29	Genome partitioning of genetic variation for height from 11,214 sibling pairs. <i>American Journal of Human Genetics</i> , 2007 , 81, 1104-10	11	110
28	Analysis of pooled DNA samples on high density arrays without prior knowledge of differential hybridization rates. <i>Nucleic Acids Research</i> , 2006 , 34, e55	20.1	48
27	Convergent evidence that oligodendrocyte lineage transcription factor 2 (OLIG2) and interacting genes influence susceptibility to schizophrenia. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2006 , 103, 12469-74	11.5	101
26	False Disease Region Identification From Identity-By-Descent Haplotype Sharing in the Presence of Phenocopies. <i>Twin Research and Human Genetics</i> , 2006 , 9, 9-16	2.2	1
25	Genetic variation of brain-derived neurotrophic factor (BDNF) in bipolar disorder: case-control study of over 3000 individuals from the UK. <i>British Journal of Psychiatry</i> , 2006 , 188, 21-5	5.4	119
24	A single nucleotide polymorphism in CHAT influences response to acetylcholinesterase inhibitors in Alzheimer's disease. <i>Pharmacogenetics and Genomics</i> , 2006 , 16, 75-7	1.9	31
23	Variation at the DAOA/G30 locus influences susceptibility to major mood episodes but not psychosis in schizophrenia and bipolar disorder. <i>Archives of General Psychiatry</i> , 2006 , 63, 366-73		124
22	GAIA: an easy-to-use web-based application for interaction analysis of case-control data. <i>BMC Medical Genetics</i> , 2006 , 7, 34	2.1	22
21	Use of phenotypic covariates in association analysis by sequential addition of cases. <i>European Journal of Human Genetics</i> , 2006 , 14, 529-34	5.3	18
20	Bias, precision and heritability of self-reported and clinically measured height in Australian twins. <i>Human Genetics</i> , 2006 , 120, 571-80	6.3	117
19	False disease region identification from identity-by-descent haplotype sharing in the presence of phenocopies. <i>Twin Research and Human Genetics</i> , 2006 , 9, 9-16	2.2	1
18	Bipolar disorder and polymorphisms in the dysbindin gene (DTNBP1). <i>Biological Psychiatry</i> , 2005 , 57, 696-701	7.9	110
17	Operation of the schizophrenia susceptibility gene, neuregulin 1, across traditional diagnostic boundaries to increase risk for bipolar disorder. <i>Archives of General Psychiatry</i> , 2005 , 62, 642-8		209
16	Covariate linkage analysis of GAW14 simulated data incorporating subclinical phenotype, sex, population, parent-of-origin, and interaction. <i>BMC Genetics</i> , 2005 , 6 Suppl 1, S45	2.6	8
15	Quantitative trait locus analysis of longitudinal quantitative trait data in complex pedigrees. <i>Genetics</i> , 2005 , 171, 1365-76	4	23
14	No association between schizophrenia and polymorphisms in COMT in two large samples. <i>American Journal of Psychiatry</i> , 2005 , 162, 1736-8	11.9	70
13	A genome scan and follow-up study identify a bipolar disorder susceptibility locus on chromosome 1q42. <i>Molecular Psychiatry</i> , 2004 , 9, 1083-90	15.1	90

12	Longitudinal data analysis in pedigree studies. <i>Genetic Epidemiology</i> , 2003 , 25 Suppl 1, S18-28	2.6	29
11	Longitudinal variance-components analysis of the Framingham Heart Study data. <i>BMC Genetics</i> , 2003 , 4 Suppl 1, S22	2.6	6
10	Genome scan meta-analysis of schizophrenia and bipolar disorder, part III: Bipolar disorder. <i>American Journal of Human Genetics</i> , 2003 , 73, 49-62	11	353
9	Genetics of schizophrenia and bipolar affective disorder: strategies to identify candidate genes. <i>Cold Spring Harbor Symposia on Quantitative Biology</i> , 2003 , 68, 383-94	3.9	5
8	Is schizophrenia linked to chromosome 1q?. <i>Science</i> , 2002 , 298, 2277; author reply 2277	33.3	7
7	Global Biobank Meta-analysis Initiative: powering genetic discovery across human diseases		6
6	Investigating Genetic Heterogeneity in Major Depression Through Item-level Genetic Analyses of the PHQ-9		2
5	A large cross-ancestry meta-analysis of genome-wide association studies identifies 69 novel risk loci for primary open-angle glaucoma and includes a genetic link with Alzheimer's disease		4
4	Symptom-level genetic modelling identifies novel risk loci and unravels the shared genetic architecture of anxiety and depression		1
3	Genome-wide analyses in 1,987,836 participants identify 39 genetic loci associated with sleep apnoea		4
2	Evidence of causal effect of major depression on alcohol dependence: Findings from the Psychiatric Genomics Consortium		4
1	Assessment of Polygenic Architecture and Risk Prediction based on Common Variants Across Fourteen Cancers		1