

Matthew H Law

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.

83
papers

2,363
citations

25
h-index

47
g-index

103
ext. papers

3,196
ext. citations

9.1
avg, IF

4.56
L-index

| # | Paper | IF | Citations |
|----|---|------|-----------|
| 83 | Is Genetic Risk for Sleep Apnea Causally Linked With Glaucoma Susceptibility? <i>2022</i> , 63, 25 | | 0 |
| 82 | Germline variants are associated with increased primary melanoma tumor thickness at diagnosis. <i>Human Molecular Genetics</i> , 2021 , 29, 3578-3587 | 5.6 | 1 |
| 81 | Genetically determined risk of keratinocyte carcinoma and risk of other cancers. <i>International Journal of Epidemiology</i> , 2021 , 50, 1316-1324 | 7.8 | |
| 80 | Genomic Risk Score for Melanoma in a Prospective Study of Older Individuals. <i>Journal of the National Cancer Institute</i> , 2021 , 113, 1379-1385 | 9.7 | 4 |
| 79 | 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 |
| 78 | 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 |
| 77 | 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 |
| 76 | 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 |
| 75 | Risk factors for melanoma by anatomical site: an evaluation of aetiological heterogeneity. <i>British Journal of Dermatology</i> , 2021 , 184, 1085-1093 | 4 | 3 |
| 74 | 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 |
| 73 | Impact of personal genomic risk information on melanoma prevention behaviors and psychological outcomes: a randomized controlled trial. <i>Genetics in Medicine</i> , 2021 , 23, 2394-2403 | 8.1 | 2 |
| 72 | Cell-type-specific meQTLs extend melanoma GWAS annotation beyond eQTLs and inform melanocyte gene-regulatory mechanisms. <i>American Journal of Human Genetics</i> , 2021 , 108, 1631-1646 | 11 | 2 |
| 71 | A UVB-responsive common variant at chromosome band 7p21.1 confers tanning response and melanoma risk via regulation of the aryl hydrocarbon receptor, AHR. <i>American Journal of Human Genetics</i> , 2021 , 108, 1611-1630 | 11 | 0 |
| 70 | Independent evaluation of melanoma polygenic risk scores in UK and Australian prospective cohorts.. <i>British Journal of Dermatology</i> , 2021 , | 4 | 1 |
| 69 | Massively parallel reporter assays of melanoma risk variants identify MX2 as a gene promoting melanoma. <i>Nature Communications</i> , 2020 , 11, 2718 | 17.4 | 24 |
| 68 | The Melanoma Genomics Managing Your Risk Study randomised controlled trial: statistical analysis plan. <i>Trials</i> , 2020 , 21, 594 | 2.8 | 1 |
| 67 | Assessment of polygenic architecture and risk prediction based on common variants across fourteen cancers. <i>Nature Communications</i> , 2020 , 11, 3353 | 17.4 | 32 |

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| 66 | The emerging field of polygenic risk scores and perspective for use in clinical care. <i>Human Molecular Genetics</i> , 2020 , 29, R165-R176 | 5.6 | 16 |
| 65 | 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 |
| 64 | 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 |
| 63 | 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 |
| 62 | A risk prediction model for the development of subsequent primary melanoma in a population-based cohort. <i>British Journal of Dermatology</i> , 2020 , 182, 1148-1157 | 4 | 12 |
| 61 | 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 |
| 60 | Overlapping genetic architecture between Parkinson disease and melanoma. <i>Acta Neuropathologica</i> , 2020 , 139, 347-364 | 14.3 | 12 |
| 59 | Making sense of different measures of skin ageing. <i>British Journal of Dermatology</i> , 2020 , 182, 1323-1324 | | |
| 58 | Multiplex melanoma families are enriched for polygenic risk. <i>Human Molecular Genetics</i> , 2020 , 29, 2976-2985 | 3 | 3 |
| 57 | 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 |
| 56 | Neural crest-derived tumor neuroblastoma and melanoma share 1p13.2 as susceptibility locus that shows a long-range interaction with the SLC16A1 gene. <i>Carcinogenesis</i> , 2020 , 41, 284-295 | 4.6 | 11 |
| 55 | Gastroesophageal reflux GWAS identifies risk loci that also associate with subsequent severe esophageal diseases. <i>Nature Communications</i> , 2019 , 10, 4219 | 17.4 | 15 |
| 54 | Combined analysis of keratinocyte cancers identifies novel genome-wide loci. <i>Human Molecular Genetics</i> , 2019 , 28, 3148-3160 | 5.6 | 20 |
| 53 | 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 | |
| 52 | 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 |
| 51 | 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 |
| 50 | Inherited Contributions to Melanoma Risk 2019 , 225-248 | | |
| 49 | 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 |

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| 48 | Polyunsaturated fatty acids and risk of melanoma: A Mendelian randomisation analysis. <i>International Journal of Cancer</i> , 2018 , 143, 508-514 | 7.5 | 11 |
| 47 | Analysis combining correlated glaucoma traits identifies five new risk loci for open-angle glaucoma. <i>Scientific Reports</i> , 2018 , 8, 3124 | 4.9 | 25 |
| 46 | 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 |
| 45 | 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 |
| 44 | Genome-wide association study of intraocular pressure uncovers new pathways to glaucoma. <i>Nature Genetics</i> , 2018 , 50, 1067-1071 | 36.3 | 86 |
| 43 | 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 |
| 42 | 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 |
| 41 | Inherited Contributions to Melanoma Risk 2018 , 1-23 | | |
| 40 | Novel pleiotropic risk loci for melanoma and nevus density implicate multiple biological pathways. <i>Nature Communications</i> , 2018 , 9, 4774 | 17.4 | 47 |
| 39 | Cell-type-specific eQTL of primary melanocytes facilitates identification of melanoma susceptibility genes. <i>Genome Research</i> , 2018 , 28, 1621-1635 | 9.7 | 33 |
| 38 | The melanoma genomics managing your risk study: A protocol for a randomized controlled trial evaluating the impact of personal genomic risk information on skin cancer prevention behaviors. <i>Contemporary Clinical Trials</i> , 2018 , 70, 106-116 | 2.3 | 13 |
| 37 | Association Between Telomere Length and Risk of Cancer and Non-Neoplastic Diseases: A Mendelian Randomization Study. <i>JAMA Oncology</i> , 2017 , 3, 636-651 | 13.4 | 236 |
| 36 | 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 |
| 35 | 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 |
| 34 | A Pilot Randomized Controlled Trial of the Feasibility, Acceptability, and Impact of Giving Information on Personalized Genomic Risk of Melanoma to the Public. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2017 , 26, 212-221 | 4 | 33 |
| 33 | Rare variants analysis of cutaneous malignant melanoma genes in Parkinson's disease. <i>Neurobiology of Aging</i> , 2016 , 48, 222.e1-222.e7 | 5.6 | 12 |
| 32 | A pilot randomised controlled trial examining the feasibility, acceptability and impact of giving information on personalised genomic risk of melanoma to the public, for motivating preventive behaviours. <i>Journal of Clinical Oncology</i> , 2016 , 34, 1556-1556 | 2.2 | |
| 31 | Variants of EVER1 and EVER2 (TMC6 and TMC8) and human papillomavirus status in patients with mucosal squamous cell carcinoma of the head and neck. <i>Cancer Causes and Control</i> , 2016 , 27, 809-15 | 2.8 | 6 |

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| 30 | Vitamin D Pathway Gene Polymorphisms and Keratinocyte Cancers: A Nested Case-Control Study and Meta-Analysis. <i>Anticancer Research</i> , 2016 , 36, 2145-52 | 2.3 | 7 |
| 29 | Genome-wide meta-analysis identifies five new susceptibility loci for cutaneous malignant melanoma. <i>Nature Genetics</i> , 2015 , 47, 987-995 | 36.3 | 162 |
| 28 | Cardiometabolic effects of genetic upregulation of the interleukin 1 receptor antagonist: a Mendelian randomisation analysis. <i>Lancet Diabetes and Endocrinology</i> , 2015 , 3, 243-53 | 18.1 | 81 |
| 27 | 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 |
| 26 | PARP1 polymorphisms play opposing roles in melanoma occurrence and survival. <i>International Journal of Cancer</i> , 2015 , 136, 2488-9 | 7.5 | 6 |
| 25 | Accurate Imputation-Based Screening of Gln368Ter Myocilin Variant in Primary Open-Angle Glaucoma 2015 , 56, 5087-93 | | 15 |
| 24 | 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 |
| 23 | 12-hydroxyeicosatetraenoic acid is associated with variability in aspirin-induced platelet inhibition. <i>Journal of Inflammation</i> , 2014 , 11, 33 | 6.7 | 15 |
| 22 | Identification of a melanoma susceptibility locus and somatic mutation in TET2. <i>Carcinogenesis</i> , 2014 , 35, 2097-101 | 4.6 | 38 |
| 21 | 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 |
| 20 | 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 |
| 19 | 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 |
| 18 | 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 |
| 17 | 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 |
| 16 | Melanoma genetics: recent findings take us beyond well-traveled pathways. <i>Journal of Investigative Dermatology</i> , 2012 , 132, 1763-74 | 4.3 | 65 |
| 15 | A study of circulating gliadin antibodies in schizophrenia among a Chinese population. <i>Schizophrenia Bulletin</i> , 2012 , 38, 514-8 | 1.3 | 29 |
| 14 | 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 |
| 13 | The functional significance of the TGM2 gene in schizophrenia: a correlation of SNPs and circulating IL-2 levels. <i>Journal of Neuroimmunology</i> , 2011 , 232, 5-7 | 3.5 | 4 |

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| 12 | No association observed between schizophrenia and non-HLA coeliac disease genes: integration with the initial MYO9B association with coeliac disease. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2011 , 156B, 709-19 | 3.5 | 3 |
| 11 | A novel recurrent mutation in MITF predisposes to familial and sporadic melanoma. <i>Nature</i> , 2011 , 480, 99-103 | 50.4 | 335 |
| 10 | Genetic association of the AKT1 gene with schizophrenia in a British population. <i>Psychiatric Genetics</i> , 2010 , 20, 118-22 | 2.9 | 20 |
| 9 | The TGM2 gene is associated with schizophrenia in a British population. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2009 , 150B, 335-40 | 3.5 | 11 |
| 8 | No association between the PPARG gene and schizophrenia in a British population. <i>Prostaglandins Leukotrienes and Essential Fatty Acids</i> , 2009 , 81, 273-7 | 2.8 | 4 |
| 7 | 273 IA study of gluten antibody levels in serum among patients with schizophrenia. <i>Schizophrenia Research</i> , 2008 , 98, 145 | 3.6 | 4 |
| 6 | Chemical cleavage of mismatch (CCM) to locate base mismatches in heteroduplex DNA. <i>Nature Protocols</i> , 2006 , 1, 2297-304 | 18.8 | 11 |
| 5 | The role of phospholipases A2 in schizophrenia. <i>Molecular Psychiatry</i> , 2006 , 11, 547-56 | 15.1 | 54 |
| 4 | Global Biobank Meta-analysis Initiative: powering genetic discovery across human diseases | | 6 |
| 3 | Massively parallel reporter assays combined with cell-type specific eQTL informed multiple melanoma loci and identified a pleiotropic function of HIV-1 restriction gene, MX2, in melanoma promotion | | 2 |
| 2 | Genome-wide analyses in 1,987,836 participants identify 39 genetic loci associated with sleep apnoea | | 4 |
| 1 | Assessment of Polygenic Architecture and Risk Prediction based on Common Variants Across Fourteen Cancers | | 1 |