## Yi-Hsiang Hsu

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	<i>FTO</i> Obesity Variant Circuitry and Adipocyte Browning in Humans. New England Journal of Medicine, 2015, 373, 895-907.	27.0	1,105
2	Genome-wide meta-analysis identifies 56 bone mineral density loci and reveals 14 loci associated with risk of fracture. Nature Genetics, 2012, 44, 491-501.	21.4	1,100
3	Twenty bone-mineral-density loci identified by large-scale meta-analysis of genome-wide association studies. Nature Genetics, 2009, 41, 1199-1206.	21.4	660
4	An atlas of genetic influences on osteoporosis in humans and mice. Nature Genetics, 2019, 51, 258-266.	21.4	557
5	Wholeâ€genome sequencing identifies EN1 as a determinant of bone density and fracture. Nature, 2015, 526, 112-117.	27.8	483
6	Relation of body composition, fat mass, and serum lipids to osteoporotic fractures and bone mineral density in Chinese men and women. American Journal of Clinical Nutrition, 2006, 83, 146-154.	4.7	441
7	Incidence of Transitional Cell Carcinoma and Arsenic in Drinking Water: A Follow-up Study of 8,102 Residents in an Arseniasis-endemic Area in Northeastern Taiwan. American Journal of Epidemiology, 2001, 153, 411-418.	3.4	371
8	Collaborative Meta-analysis: Associations of 150 Candidate Genes With Osteoporosis and Osteoporotic Fracture. Annals of Internal Medicine, 2009, 151, 528.	3.9	250
9	Dose-Response Relationship Between Prevalence of Cerebrovascular Disease and Ingested Inorganic Arsenic. Stroke, 1997, 28, 1717-1723.	2.0	234
10	An Integration of Genome-Wide Association Study and Gene Expression Profiling to Prioritize the Discovery of Novel Susceptibility Loci for Osteoporosis-Related Traits. PLoS Genetics, 2010, 6, e1000977.	3.5	191
11	Association of JAG1 with Bone Mineral Density and Osteoporotic Fractures: A Genome-wide Association Study and Follow-up Replication Studies. American Journal of Human Genetics, 2010, 86, 229-239.	6.2	188
12	Arsenic methylation capacity, body retention, and null genotypes of glutathione S-transferase M1 and T1 among current arsenic-exposed residents in Taiwan. Mutation Research - Reviews in Mutation Research, 1997, 386, 197-207.	5.5	181
13	Biomarkers of exposure, effect, and susceptibility of arsenic-induced health hazards in Taiwan. Toxicology and Applied Pharmacology, 2005, 206, 198-206.	2.8	170
14	Large meta-analysis of genome-wide association studies identifies five loci for lean body mass. Nature Communications, 2017, 8, 80.	12.8	147
15	Leveraging Cross-Species Transcription Factor Binding Site Patterns: From Diabetes Risk Loci to Disease Mechanisms. Cell, 2014, 156, 343-358.	28.9	113
16	Low-Frequency Synonymous Coding Variation in CYP2R1 Has Large Effects on Vitamin D Levels and Risk of Multiple Sclerosis. American Journal of Human Genetics, 2017, 101, 227-238.	6.2	112
17	Pleiotropic genes for metabolic syndrome and inflammation. Molecular Genetics and Metabolism, 2014, 112, 317-338.	1.1	107
18	Variation in genes involved in the RANKL/RANK/OPG bone remodeling pathway are associated with bone mineral density at different skeletal sites in men. Human Genetics, 2006, 118, 568-577.	3.8	103

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19	Association of Polygenic Score for Schizophrenia and HLA Antigen and Inflammation Genes With Response to Lithium in Bipolar Affective Disorder. JAMA Psychiatry, 2018, 75, 65-74.	11.0	102
20	Genome-Wide Association Studies of Skeletal Phenotypes: What We Have Learned and Where We Are Headed. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E1958-E1977.	3.6	99
21	Genetic determinants of heel bone properties: genome-wide association meta-analysis and replication in the GEFOS/GENOMOS consortium. Human Molecular Genetics, 2014, 23, 3054-3068.	2.9	90
22	Impact of seafood and fruit consumption on bone mineral density. Maturitas, 2007, 56, 1-11.	2.4	87
23	Bivariate genome-wide association meta-analysis of pediatric musculoskeletal traits reveals pleiotropic effects at the SREBF1/TOM1L2 locus. Nature Communications, 2017, 8, 121.	12.8	82
24	<i>METTL21C</i> Is a Potential Pleiotropic Gene for Osteoporosis and Sarcopenia Acting Through the Modulation of the NF-κB Signaling Pathway. Journal of Bone and Mineral Research, 2014, 29, 1531-1540.	2.8	80
25	Meta-analysis of epigenome-wide association studies of cognitive abilities. Molecular Psychiatry, 2018, 23, 2133-2144.	7.9	68
26	Bivariate genome-wide association analyses of the broad depression phenotype combined with major depressive disorder, bipolar disorder or schizophrenia reveal eight novel genetic loci for depression. Molecular Psychiatry, 2020, 25, 1420-1429.	7.9	68
27	Identification of homogeneous genetic architecture of multiple genetically correlated traits by block clustering of genome-wide associations. Journal of Bone and Mineral Research, 2011, 26, 1261-1271.	2.8	56
28	Mouse BMD quantitative trait loci show improved concordance with human genome-wide association loci when recalculated on a new, common mouse genetic map. Journal of Bone and Mineral Research, 2010, 25, 1808-1820.	2.8	53
29	Distinct DNA methylation profiles in bone and blood of osteoporotic and healthy postmenopausal women. Epigenetics, 2017, 12, 674-687.	2.7	53
30	Bicc1 is a genetic determinant of osteoblastogenesis and bone mineral density. Journal of Clinical Investigation, 2014, 124, 2736-2749.	8.2	51
31	Genome-wide pleiotropy of osteoporosis-related phenotypes: The framingham study. Journal of Bone and Mineral Research, 2010, 25, 1555-1563.	2.8	50
32	Epigenome-wide Association of DNA Methylation in Whole Blood With Bone Mineral Density. Journal of Bone and Mineral Research, 2017, 32, 1644-1650.	2.8	49
33	Association of the methylenetetrahydrofolate reductase C677T polymorphism and fracture risk in Chinese postmenopausal women. Bone, 2007, 40, 737-742.	2.9	47
34	Assessment of gene-by-sex interaction effect on bone mineral density. Journal of Bone and Mineral Research, 2012, 27, 2051-2064.	2.8	47
35	Association of polygenic score for major depression with response to lithium in patients with bipolar disorder. Molecular Psychiatry, 2021, 26, 2457-2470.	7.9	44
36	Novel Genetic Variants Associated With Increased Vertebral Volumetric BMD, Reduced Vertebral Fracture Risk, and Increased Expression of <i>SLC1A3</i> and <i>EPHB2</i> . Journal of Bone and Mineral Research, 2016, 31, 2085-2097.	2.8	42

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37	Spontaneous and induced sister chromatid exchanges and delayed cell proliferation in peripheral lymphocytes of Bowen's disease patients and matched controls of arseniasis-hyperendemic villages in Taiwan. Mutation Research - Reviews in Mutation Research, 1997, 386, 241-251.	5.5	40
38	Association of the Polygenic Scores for Personality Traits and Response to Selective Serotonin Reuptake Inhibitors in Patients with Major Depressive Disorder. Frontiers in Psychiatry, 2018, 9, 65.	2.6	38
39	Disentangling the genetics of lean mass. American Journal of Clinical Nutrition, 2019, 109, 276-287.	4.7	38
40	Phenomics-Based Quantification of CRISPR-Induced Mosaicism in Zebrafish. Cell Systems, 2020, 10, 275-286.e5.	6.2	38
41	Metabolomics Insights into Osteoporosis Through Association With Bone Mineral Density. Journal of Bone and Mineral Research, 2020, 36, 729-738.	2.8	37
42	Large-Scale Genome-Wide Linkage Analysis for Loci Linked to BMD at Different Skeletal Sites in Extreme Selected Sibships. Journal of Bone and Mineral Research, 2006, 22, 184-194.	2.8	36
43	A genome-wide copy number association study of osteoporotic fractures points to the 6p25.1 locus. Journal of Medical Genetics, 2014, 51, 122-131.	3.2	36
44	Genome-wide association study for radiographic vertebral fractures: A potential role for the 16q24 BMD locus. Bone, 2014, 59, 20-27.	2.9	32
45	A combined reference panel from the 1000 Genomes and UK10K projects improved rare variant imputation in European and Chinese samples. Scientific Reports, 2016, 6, 39313.	3.3	32
46	Impact of Common Variation in Bone-Related Genes on Type 2 Diabetes and Related Traits. Diabetes, 2012, 61, 2176-2186.	0.6	31
47	A regulatory variant at 3q21.1 confers an increased pleiotropic risk for hyperglycemia and altered bone mineral density. Cell Metabolism, 2021, 33, 615-628.e13.	16.2	28
48	Meta-Analysis of Genomewide Association Studies Reveals Genetic Variants for Hip Bone Geometry. Journal of Bone and Mineral Research, 2019, 34, 1284-1296.	2.8	27
49	The association of obesity and coronary artery disease genes with response to SSRIs treatment in major depression. Journal of Neural Transmission, 2019, 126, 35-45.	2.8	27
50	Environmental Mold and Mycotoxin Exposures Elicit Specific Cytokine and Chemokine Responses. PLoS ONE, 2015, 10, e0126926.	2.5	26
51	Combining schizophrenia and depression polygenic risk scores improves the genetic prediction of lithium response in bipolar disorder patients. Translational Psychiatry, 2021, 11, 606.	4.8	25
52	Identification of a novel locus on chromosome 2q13, which predisposes to clinical vertebral fractures independently of bone density. Annals of the Rheumatic Diseases, 2018, 77, 378-385.	0.9	21
53	Genome-wide association meta-analyses to identify common genetic variants associated with hallux valgus in Caucasian and African Americans. Journal of Medical Genetics, 2015, 52, 762-769.	3.2	18
54	Genetic basis of falling risk susceptibility in the UK Biobank Study. Communications Biology, 2020, 3, 543.	4.4	17

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55	Genome-wide association study for radiographic vertebral fractures: a potential role for the 16q24 BMD locus. Bone, 2014, 59, 20-7.	2.9	17
56	Genetic variation in TRPS1 may regulate hip geometry as well as bone mineral density. Bone, 2012, 50, 1188-1195.	2.9	16
57	CYP19A1 polymorphisms are associated with bone mineral density in Chinese men. Human Genetics, 2007, 121, 491-500.	3.8	14
58	Percent fat mass is inversely associated with bone mass and hip geometry in rural Chinese adolescents. Journal of Bone and Mineral Research, 2010, 25, 1544-1554.	2.8	14
59	Meta-analysis of genome-wide association studies identifies two loci associated with circulating osteoprotegerin levels. Human Molecular Genetics, 2014, 23, 6684-6693.	2.9	14
60	Using polygenic scores and clinical data for bipolar disorder patient stratification and lithium response prediction: machine learning approach. British Journal of Psychiatry, 2022, 220, 219-228.	2.8	11
61	Individual and Joint Associations of Methylenetetrahydrofolate Reductase C677T Genotype and Plasma Homocysteine With Dyslipidemia in a Chinese Population With Hypertension. Clinical and Applied Thrombosis/Hemostasis, 2017, 23, 287-293.	1.7	10
62	HLA-DRB1 and HLA-DQB1 genetic diversity modulates response to lithium in bipolar affective disorders. Scientific Reports, 2021, 11, 17823.	3.3	10
63	A Meta-Analysis of the Transferability of Bone Mineral Density Genetic Loci Associations From European to African Ancestry Populations. Journal of Bone and Mineral Research, 2020, 36, 469-479.	2.8	9
64	DNA methylation-based subclassification of psoriasis in the Chinese Han population. Frontiers of Medicine, 2018, 12, 717-725.	3.4	8
65	Trans-Ethnic Polygenic Analysis Supports Genetic Overlaps of Lumbar Disc Degeneration With Height, Body Mass Index, and Bone Mineral Density. Frontiers in Genetics, 2018, 9, 267.	2.3	8
66	Associations between prenatal exposure to perfluoroalkyl substances, hypomethylation of MEST imprinted gene and birth outcomes. Environmental Pollution, 2022, 304, 119183.	7.5	8
67	Targeted sequencing of genome wide significant loci associated with bone mineral density (BMD) reveals significant novel and rare variants: the Cohorts for Heart and Aging Research in Genomic Epidemiology (CHARGE) targeted sequencing study. Human Molecular Genetics, 2016, 25, ddw289.	2.9	7
68	Evaluation of power of the Illumina HumanOmni5M-4v1 BeadChip to detect risk variants for human complex diseases. European Journal of Human Genetics, 2016, 24, 1029-1034.	2.8	7
69	Associations of methylenetetrahydrofolate reductase C677T genotype with blood pressure levels in Chinese population with essential hypertension. Clinical and Experimental Hypertension, 2018, 40, 207-212.	1.3	7
70	Elevation in Total Homocysteine Levels in Chinese Patients With Essential Hypertension Treated With Antihypertensive Benazepril. Clinical and Applied Thrombosis/Hemostasis, 2016, 22, 191-198.	1.7	5
71	Genetic Determinants and Pharmacogenetics of Osteoporosis and Osteoporotic Fracture. Contemporary Endocrinology, 2020, , 485-506.	0.1	4
72	Association Study between the FTCDNL1 (FONG) and Susceptibility to Osteoporosis. PLoS ONE, 2015, 10, e0140549.	2.5	4

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73	Understanding the complex genetic architecture connecting rheumatoid arthritis, osteoporosis and inflammation: discovering causal pathways. Human Molecular Genetics, 2022, , .	2.9	3
74	Musculoskeletal genetics and -omics: Meeting report from the 32nd Annual Meeting of the American Society for Bone and Mineral Research. IBMS BoneKEy, 2011, 8, 112-122.	0.0	1
75	Interactive Effect of the KCNJ11 Ile337Val Polymorphism and Cigarette Smoking on the Antihypertensive Response to Irbesartan in Chinese Hypertensive Patients. American Journal of Hypertension, 2016, 29, 553-559.	2.0	1
76	Genetic determinants of bone mass and osteoporotic fracture. , 2020, , 1615-1630.		1
77	Effect of simvastatin on plasma homocysteine levels and its modification by MTHFR C677T polymorphism in Chinese patients with primary hyperlipidemia. Cardiovascular Therapeutics, 2012, 31, n/a-n/a.	2.5	1
78	Response to "fat mass is a positive predictor of bone mass in adolescents― Journal of Bone and Mineral Research, 2011, 26, 674-675.	2.8	0
79	ISDN2014_0385: REMOVED: A bivariate Genome Wide Association Study (GWAS) of depressive symptoms and lipid levels has identified pleiotropic gene loci. International Journal of Developmental Neuroscience, 2015, 47, 113-114.	1.6	0
80	METTL21C: From GWAS to in vitro function in skeletal muscle cells. FASEB Journal, 2013, 27, 942.5.	0.5	0
81	Whole genome shotgun metagenomic sequencing to identify differential abundant microbiome features between dementia and mild cognitive impairment (MCI) in AD subjects. Alzheimer's and Dementia, 2021, 17, .	0.8	0