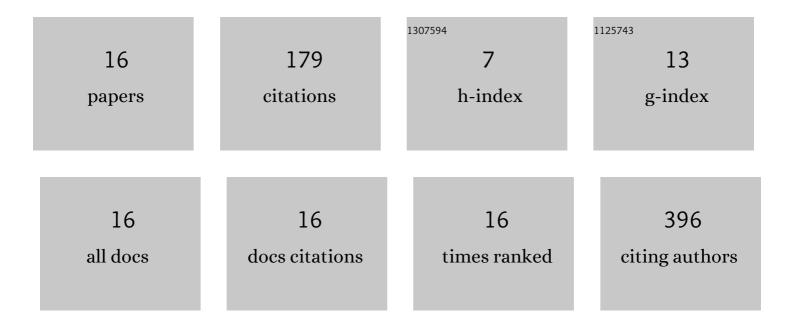
Hsing-Fang Lu

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2014841/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Genome-wide association study identifies 14 previously unreported susceptibility loci for adolescent idiopathic scoliosis in Japanese. Nature Communications, 2019, 10, 3685.	12.8	47
2	Gene expression profiling combined with functional analysis identify integrin beta1 (ITGB1) as a potential prognosis biomarker in triple negative breast cancer. Pharmacological Research, 2016, 104, 31-37.	7.1	39
3	Interaction between HLA-B60 and HLA-B27 as a Better Predictor of Ankylosing Spondylitis in a Taiwanese Population. PLoS ONE, 2015, 10, e0137189.	2.5	23
4	rs2841277 (<i>PLD4</i>) is associated with susceptibility and rs4672495 is associated with disease activity in rheumatoid arthritis. Oncotarget, 2017, 8, 64180-64190.	1.8	11
5	Study of the Association between <i>ITPKC</i> Genetic Polymorphisms and Calcium Nephrolithiasis. BioMed Research International, 2014, 2014, 1-6.	1.9	10
6	Meta-Analysis of Genome-Wide Association Studies Identifies Three Loci Associated With Stiffness Index of the Calcaneus. Journal of Bone and Mineral Research, 2019, 34, 1275-1283.	2.8	8
7	Pharmacogenomics Study for Raloxifene in Postmenopausal Female with Osteoporosis. Disease Markers, 2020, 2020, 1-8.	1.3	8
8	Correlation of HAMP gene polymorphisms and expression with the susceptibility and length of hospital stays in Taiwanese children with Kawasaki disease. Oncotarget, 2017, 8, 51859-51868.	1.8	8
9	rs657075 (CSF2) Is Associated with the Disease Phenotype (BAS-G) of Ankylosing Spondylitis. International Journal of Molecular Sciences, 2017, 18, 83.	4.1	6
10	Functional correlations between CXCL10/IP10 gene polymorphisms and risk of Kawasaki disease. Pediatric Allergy and Immunology, 2021, 32, 363-370.	2.6	5
11	Polygenic Risk Score of Adolescent Idiopathic Scoliosis for Potential Clinical Use. Journal of Bone and Mineral Research, 2020, 36, 1481-1491.	2.8	5
12	Association Study between the FTCDNL1 (FONG) and Susceptibility to Osteoporosis. PLoS ONE, 2015, 10, e0140549.	2.5	4
13	Associations of genetic variants of endothelin with cardiovascular complications in patients with renal failure. BMC Nephrology, 2017, 18, 291.	1.8	2
14	Genomic interrogation of familial short stature contributes to the discovery of the pathophysiological mechanisms and pharmaceutical drug repositioning. Journal of Biomedical Science, 2019, 26, 91.	7.0	2
15	Integrative genomic analysis for the functional roles of <i>ITPKC</i> in bone mineral density. Bioscience Reports, 2018, 38, .	2.4	1
16	Association of endothelin genetic variants and hospitalized infection complications in end-stage renal disease (ESRD) patients. BMC Nephrology, 2019, 20, 203.	1.8	0