Henry J Lin

List of Publications by Year
in descending order

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41 papers	1,609 citations	430874 18 h-index	345221 36 g-index
45 all docs	45 docs citations	45 times ranked	3452 citing authors

#	Article	IF	Citations
1	Glucose-6-phosphate dehydrogenase deficiency presenting with rhabdomyolysis in a patient with coronavirus disease 2019 pneumonia: aÂcase report. Journal of Medical Case Reports, 2022, 16, 106.	0.8	5
2	Monogenic and Polygenic Contributions to QTc Prolongation in the Population. Circulation, 2022, 145, 1524-1533.	1.6	14
3	A multiancestry genome-wide association study of unexplained chronic ALT elevation as a proxy for nonalcoholic fatty liver disease with histological and radiological validation. Nature Genetics, 2022, 54, 761-771.	21.4	68
4	A multi-ethnic polygenic risk score is associated with hypertension prevalence and progression throughout adulthood. Nature Communications, 2022, 13, .	12.8	27
5	Rare Coding Variants Associated With Electrocardiographic Intervals Identify Monogenic Arrhythmia Susceptibility Genes: A Multi-Ancestry Analysis. Circulation Genomic and Precision Medicine, 2021, 14, e003300.	3.6	7
6	Identification of Functional Genetic Determinants of Cardiac Troponin T and I in a Multiethnic Population and Causal Associations With Atrial Fibrillation. Circulation Genomic and Precision Medicine, 2021, 14, CIRCGEN121003460.	3.6	5
7	An Outbreak of Polygenic Scores for Coronary Artery Disease. Journal of the American College of Cardiology, 2020, 75, 2781-2784.	2.8	11
8	Multi-ancestry GWAS of the electrocardiographic PR interval identifies 202 loci underlying cardiac conduction. Nature Communications, 2020, 11, 2542.	12.8	59
9	GWAS of QRS duration identifies new loci specific to Hispanic/Latino populations. PLoS ONE, 2019, 14, e0217796.	2.5	8
10	Effects of Calcium, Magnesium, and Potassium Concentrations on Ventricular Repolarization in Unselected Individuals. Journal of the American College of Cardiology, 2019, 73, 3118-3131.	2.8	27
11	Home use of a compact, 12â€lead ECG recording system for newborns. Journal of Electrocardiology, 2019, 53, 89-94.	0.9	7
12	Lego bricks and the octet rule: Molecular models for biochemical pathways with plastic, interlocking toy bricks. Biochemistry and Molecular Biology Education, 2018, 46, 54-57.	1.2	10
13	Community Partnership in Precision Medicine: Themes from a Community Engagement Conference. Ethnicity and Disease, 2018, 28, 503-510.	2.3	6
14	ExomeChip-Wide Analysis of 95 626 Individuals Identifies 10 Novel Loci Associated With QT and JT Intervals. Circulation Genomic and Precision Medicine, 2018, 11, e001758.	3.6	27
15	Common and Rare Coding Genetic Variation Underlying the Electrocardiographic PR Interval. Circulation Genomic and Precision Medicine, 2018, 11, e002037.	3.6	19
16	PR interval genome-wide association meta-analysis identifies 50 loci associated with atrial and atrioventricular electrical activity. Nature Communications, 2018, 9, 2904.	12.8	71
17	Multi-ethnic genome-wide association study for atrial fibrillation. Nature Genetics, 2018, 50, 1225-1233.	21.4	552
18	Genetic loci associated with heart rate variability and their effects on cardiac disease risk. Nature Communications, 2017, 8, 15805.	12.8	95

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19	Genome-wide association study of iron traits and relation to diabetes in the Hispanic Community Health Study/Study of Latinos (HCHS/SOL): potential genomic intersection of iron and glucose regulation?. Human Molecular Genetics, 2017, 26, 1966-1978.	2.9	31
20	Genetic Risk Prediction of Atrial Fibrillation. Circulation, 2017, 135, 1311-1320.	1.6	87
21	Evaluating p97 Inhibitor Analogues for Potency against p97â€“p37 and p97â€“Npl4â€“Ufd1 Complexes. ChemMedChem, 2016, 11, 953-957.	3.2	13
22	Altered cofactor regulation with disease-associated p97/VCP mutations. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, E1705-14.	7.1	87
23	A tortuous proximal urethra in urorectal septum malformation sequence?. American Journal of Medical Genetics, Part A, 2014, 164, 1298-1303.	1.2	2
24	Intestinal tumor suppression in Apc Min/+ mice by prostaglandin D 2 receptor PTGDR. Cancer Medicine, 2014, 3, 1041-1051.	2.8	26
25	High Dietary Niacin May Increase Prostaglandin Formation but Does Not Increase Tumor Formation inApc^{Min/+}Mice. Nutrition and Cancer, 2011, 63, 950-959.	2.0	2
26	Asymptomatic Maternal Combined Homocystinuria and Methylmalonic Aciduria (cblC) Detected through Low Carnitine Levels on Newborn Screening. Journal of Pediatrics, 2009, 155, 924-927.	1.8	29
27	Use of HLA marker associations and HLA haplotype linkage to estimate disease risks in families with gluten-sensitive enteropathy. Clinical Genetics, 2008, 28, 185-198.	2.0	10
28	Hematopoietic Prostaglandin D Synthase Suppresses Intestinal Adenomas in ApcMin/+ Mice. Cancer Research, 2007, 67, 881-889.	0.9	55
29	Naturally occurring Phe151Leu substitution near a conserved folding module lowers stability of glutathione transferase P1â€“1. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2003, 1649, 16-23.	2.3	13
30	Glutathione transferase GSTT1, broccoli, and prevalence of colorectal adenomas. Pharmacogenetics and Genomics, 2002, 12, 175-179.	5.7	27
31	Neonatal Marfan Syndrome. Fetal and Pediatric Pathology, 2001, 20, 235-239.	0.3	0
32	NEONATAL MARFAN SYNDROME. Fetal and Pediatric Pathology, 2001, 20, 235-239.	0.3	0
33	Mosaic tetrasomy 8q: Inverted duplication of 8q23.3qter in an analphoid marker. American Journal of Medical Genetics Part A, 2000, 92, 69-76.	2.4	27
34	Intrachromosomal triplication of 2q11.2-q21 in a severely malformed infant: Case report and review of triplications and their possible mechanism. American Journal of Medical Genetics Part A, 1999, 82, 312-317.	2.4	32
35	Anomalous inferior and superior venae cavae with oculoauriculovertebral defect: Review of Goldenhar complex and malformations of left-right asymmetry. , 1998, 75, 88-94.		32
36	Omphalocele with absent radial ray (ORR): A case with diploid-triploid mixoploidy. American Journal of Medical Genetics Part A, 1998, 75, 235-239.	2.4	12

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37	Infantile Marfanâ€™s Syndrome. Circulation, 1998, 97, 1103-1104.	1.6	0
38	Occipital encephalocele and MURCS association: Case report and review of central nervous system anomalies in MURCS patients. , 1996, 61, 59-62.		22
39	Exstrophy of the cloaca in a 47,XXX child: Review of genitourinary malformations in tripleâ€X patients. American Journal of Medical Genetics Part A, 1993, 45, 761-763.	2.4	50
40	DOOR syndrome (deafness, onycho-osteodystrophy, and mental retardation): A new patient and delineation of neurologic variability among recessive cases. American Journal of Medical Genetics Part A, 1993, 47, 534-539.	2.4	14
41	Disease risk estimates from marker association data: Application to individuals at risk for hemochromatosis. Clinical Genetics, 1985, 27, 127-133.	2.0	3