Huann-Sheng Chen

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Data-driven choice of a model selection method in joinpoint regression. Journal of Applied Statistics, 2023, 50, 1992-2013.	0.6	5
2	Twenty years since Joinpoint 1.0: Two major enhancements, their justification, and impact. Statistics in Medicine, 2022, 41, 3102-3130.	0.8	13
3	On the application, reporting, and sharing of in silico simulations for genetic studies. Genetic Epidemiology, 2021, 45, 131-141.	0.6	4
4	Updated Methodology for Projecting U.S and State-Level Cancer Counts for the Current Calendar Year: Part I: Spatio-temporal Modeling for Cancer Incidence. Cancer Epidemiology Biomarkers and Prevention, 2021, 30, 1620-1626.	1.1	6
5	Updated Methodology for Projecting U.S and State-Level Cancer Counts for the Current Calendar Year: Part II: Evaluation of Incidence and Mortality Projection Methods. Cancer Epidemiology Biomarkers and Prevention, 2021, 30, 1993-2000.	1.1	9
6	Regional differences in tobacco smoking and lung cancer in Portugal in 2018: a population-based analysis using nationwide incidence and mortality data. BMJ Open, 2020, 10, e038937.	0.8	4
7	The Joinpoint-Jump and Joinpoint-Comparability Ratio Model for Trend Analysis with Applications to Coding Changes in Health Statistics. Journal of Official Statistics, 2020, 36, 49-62.	0.1	12
8	CP*Trends: An Online Tool for Comparing Cohort and Period Trends Across Cancer Sites. American Journal of Epidemiology, 2019, 188, 1361-1370.	1.6	3
9	Genetic Simulation Resources and the GSR Certification Program. Bioinformatics, 2019, 35, 709-710.	1.8	6
10	Early estimates of cancer incidence for 2015: Expanding to include estimates for white and black races. Cancer, 2018, 124, 2192-2204.	2.0	9
11	Early estimates of SEER cancer incidence, 2014. Cancer, 2017, 123, 2524-2534.	2.0	39
12	Improved confidence interval for average annual percent change in trend analysis. Statistics in Medicine, 2017, 36, 3059-3074.	0.8	59
13	Up For A Challenge (U4C): Stimulating innovation in breast cancer genetic epidemiology. PLoS Genetics, 2017, 13, e1006945.	1.5	3
14	Bottom-up GGM algorithm for constructing multilayered hierarchical gene regulatory networks that govern biological pathways or processes. BMC Bioinformatics, 2016, 17, 132.	1.2	19
15	Preliminary estimates of SEER cancer incidence for 2013. Cancer, 2016, 122, 1579-1587.	2.0	10
16	Genetic Data Simulators and their Applications: An Overview. Genetic Epidemiology, 2015, 39, 2-10.	0.6	26
17	Early estimates of <scp>SEER</scp> cancer incidence for 2012: Approaches, opportunities, and cautions for obtaining preliminary estimates of cancer incidence. Cancer, 2015, 121, 2053-2062.	2.0	13
18	Genetic Simulation Tools for Postâ€Genome Wide Association Studies of Complex Diseases. Genetic Epidemiology, 2015, 39, 11-19.	0.6	22

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19	Clustering of trend data using joinpoint regression models. Statistics in Medicine, 2014, 33, 4087-4103.	0.8	30
20	Developments and challenges in statistical methods in cancer surveillance. Statistics and Its Interface, 2014, 7, 135-151.	0.2	5
21	A combined p-value test for multiple hypothesis testing. Journal of Statistical Planning and Inference, 2013, 143, 764-770.	0.4	17
22	Genetic Simulation Resources: a website for the registration and discovery of genetic data simulators. Bioinformatics, 2013, 29, 1101-1102.	1.8	29
23	A Powerful Method for Combining <i>P</i> â€Values in Genomic Studies. Genetic Epidemiology, 2013, 37, 814-819.	0.6	5
24	Evaluation of Gene Association Methods for Coexpression Network Construction and Biological Knowledge Discovery. PLoS ONE, 2012, 7, e50411.	1.1	100
25	Predicting US―and stateâ€level cancer counts for the current calendar year. Cancer, 2012, 118, 1091-1099.	2.0	44
26	Predicting US―and stateâ€level cancer counts for the current calendar year. Cancer, 2012, 118, 1100-1109.	2.0	48
27	Entropy-based information gain approaches to detect and to characterize gene-gene and gene-environment interactions/correlations of complex diseases. Genetic Epidemiology, 2011, 35, 706-721.	0.6	54
28	TF-finder: A software package for identifying transcription factors involved in biological processes using microarray data and existing knowledge base. BMC Bioinformatics, 2010, 11, 425.	1.2	13
29	A combinatorial approach for detecting gene-gene interaction using multiple traits of Genetic Analysis Workshop 16 rheumatoid arthritis data. BMC Proceedings, 2009, 3, S43.	1.8	4
30	Association Between Two Unlinked Loci at 8q24 and Prostate Cancer Risk Among European Americans. Journal of the National Cancer Institute, 2007, 99, 1525-1533.	3.0	126
31	Genome-wide association tests by using block information in family data. BMC Proceedings, 2007, 1, S149.	1.8	1
32	A new association test using haplotype similarity. Genetic Epidemiology, 2007, 31, 577-593.	0.6	21
33	Multiple testing in the genomics era: Findings from Genetic Analysis Workshop 15, Group 15. Genetic Epidemiology, 2007, 31, S124-S131.	0.6	14
34	Haplotype sharing transmission/disequilibrium tests that allow for genotyping errors. Genetic Epidemiology, 2005, 28, 341-351.	0.6	11
35	Metabolic Profiling of the Sink-to-Source Transition in Developing Leaves of Quaking Aspen. Plant Physiology, 2004, 136, 3364-3375.	2.3	81
36	Joint analysis of two microarray gene-expression data sets to select lung adenocarcinoma marker genes. BMC Bioinformatics, 2004, 5, 81.	1.2	228

#	Article	IF	CITATIONS
37	Reply to Knapp and Becker. American Journal of Human Genetics, 2004, 74, 591-593.	2.6	8
38	Transmission/Disequilibrium Test Based on Haplotype Sharing for Tightly Linked Markers. American Journal of Human Genetics, 2003, 73, 566-579.	2.6	73
39	EfficientL1estimation and related inferences in linear regression with unknown form of heteroscedasticity. Journal of Nonparametric Statistics, 2002, 14, 607-622.	0.4	0
40	Estimation of the exponential mean under type I censorded sampling. Journal of Statistical Planning and Inference, 1992, 33, 187-196.	0.4	3