## Shelley B Bull

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	New Susceptibility Loci Associated with Kidney Disease in Type 1 Diabetes. PLoS Genetics, 2012, 8, e1002921.	3.5	216
2	Genome-Wide Association Study of Diabetic Kidney Disease Highlights Biology Involved in Glomerular Basement Membrane Collagen. Journal of the American Society of Nephrology: JASN, 2019, 30, 2000-2016.	6.1	135
3	Comparative Expression of the Mitotic Regulators SAK and PLK in Colorectal Cancer. Annals of Surgical Oncology, 2001, 8, 729-740.	1.5	105
4	A Genome-Wide Association Study Identifies a Novel Major Locus for Glycemic Control in Type 1 Diabetes, as Measured by Both A1C and Glucose. Diabetes, 2010, 59, 539-549.	0.6	103
5	The Combination of p53 Mutation and neu/erbB-2 Amplification Is Associated With Poor Survival in Node-Negative Breast Cancer. Journal of Clinical Oncology, 2004, 22, 86-96.	1.6	90
6	Multiple Superoxide Dismutase 1/Splicing Factor Serine Alanine 15 Variants Are Associated With the Development and Progression of Diabetic Nephropathy. Diabetes, 2008, 57, 218-228.	0.6	89
7	Multiple Variants in Vascular Endothelial Growth Factor (VEGFA) Are Risk Factors for Time to Severe Retinopathy in Type 1 Diabetes. Diabetes, 2007, 56, 2161-2168.	0.6	88
8	MDR1 Gene Expression and Outcome in Osteosarcoma: A Prospective, Multicenter Study. Journal of Clinical Oncology, 2000, 18, 2685-2694.	1.6	80
9	Reduction of selection bias in genomewide studies by resampling. Genetic Epidemiology, 2005, 28, 352-367.	1.3	67
10	Distinguishing luminal breast cancer subtypes by Ki67, progesterone receptor or TP53 status provides prognostic information. Modern Pathology, 2014, 27, 554-561.	5.5	66
11	The association of previously reported polymorphisms for microvascular complications in a meta-analysis of diabetic retinopathy. Human Genetics, 2015, 134, 247-257.	3.8	60
12	BR-squared: a practical solution to the winner's curse in genome-wide scans. Human Genetics, 2011, 129, 545-552.	3.8	56
13	Preferential allelic expression can lead to reduced expression ofBRCA1 in sporadic breast cancers. , 1998, 77, 1-6.		46
14	Confidence intervals for multinomial logistic regression in sparse data. Statistics in Medicine, 2007, 26, 903-918.	1.6	44
15	Prognostic Effect of Basal-Like Breast Cancers Is Time Dependent: Evidence from Tissue Microarray Studies on a Lymph Node–Negative Cohort. Clinical Cancer Research, 2008, 14, 4168-4174.	7.0	37
16	The Efficiency of Multinomial Logistic Regression Compared with Multiple Group Discriminant Analysis. Journal of the American Statistical Association, 1987, 82, 1118-1122.	3.1	36
17	Re-Ranking Sequencing Variants in the Post-GWAS Era for Accurate Causal Variant Identification. PLoS Genetics, 2013, 9, e1003609.	3.5	36
18	A flexible genomeâ€wide bootstrap method that accounts for rankingand thresholdâ€selection bias in GWAS interpretation and replication study design. Statistics in Medicine, 2011, 30, 1898-1912.	1.6	32

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19	Elevated expression of podocalyxin is associated with lymphatic invasion, basal-like phenotype, and clinical outcome in axillary lymph node-negative breast cancer. Breast Cancer Research and Treatment, 2013, 137, 709-719.	2.5	30
20	Incorporation of covariates into genome scanning using sib-pair analysis in bipolar affective disorder. Genetic Epidemiology, 1997, 14, 635-640.	1.3	28
21	Menacalc, a quantitative method of metastasis assessment, as a prognostic marker for axillary node-negative breast cancer. BMC Cancer, 2015, 15, 483.	2.6	27
22	Validation of Intratumoral T-bet+ Lymphoid Cells as Predictors of Disease-Free Survival in Breast Cancer. Cancer Immunology Research, 2016, 4, 41-48.	3.4	25
23	Insights From Mixture Cure Modeling of Molecular Markers for Prognosis in Breast Cancer. Journal of Clinical Oncology, 2013, 31, 2047-2054.	1.6	23
24	Molecular and Epidemiologic Study of Multiresistant Serratia marcescens Infections in a Spinal Cord Injury Rehabilitation Unit. Infection Control and Hospital Epidemiology, 1988, 9, 20-27.	1.8	17
25	Type of TP53 mutation and ERBB2 amplification affects survival in node-negative breast cancer. Breast Cancer Research and Treatment, 2007, 105, 255-265.	2.5	17
26	JACKKNIFE BIAS REDUCTION FOR POLYCHOTOMOUS LOGISTIC REGRESSION. Statistics in Medicine, 1997, 16, 545-560.	1.6	16
27	A hierarchical clustering method for estimating copy number variation. Biostatistics, 2007, 8, 632-653.	1.5	16
28	Bivariate genetic association analysis of systolic and diastolic blood pressure by copula models. BMC Proceedings, 2014, 8, S72.	1.6	16
29	Multiple linear combination (MLC) regression tests for common variants adapted to linkage disequilibrium structure. Genetic Epidemiology, 2017, 41, 108-121.	1.3	14
30	Tumoral BRD4 expression in lymph node-negative breast cancer: association with T-bet+ tumor-infiltrating lymphocytes and disease-free survival. BMC Cancer, 2018, 18, 750.	2.6	13
31	Genome-wide association analyses of North American Rheumatoid Arthritis Consortium and Framingham Heart Study data utilizing genome-wide linkage results. BMC Proceedings, 2009, 3, S103.	1.6	12
32	Resampling to Address the Winner's Curse in Genetic Association Analysis of Time to Event. Genetic Epidemiology, 2015, 39, 518-528.	1.3	11
33	Region-based analysis in genome-wide association study of Framingham Heart Study blood lipid phenotypes. BMC Proceedings, 2009, 3, S127.	1.6	10
34	New Locus for Skin Intrinsic Fluorescence in Type 1 Diabetes Also Associated With Blood and Skin Glycated Proteins. Diabetes, 2016, 65, 2060-2071.	0.6	10
35	Fine mapping by linkage and association in nuclear family and case-control designs. Genetic Epidemiology, 2005, 29, S48-S58.	1.3	9
36	Gene-based multiple regression association testing for combined examination of common and low frequency variants in quantitative trait analysis. Frontiers in Genetics, 2013, 4, 233.	2.3	9

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37	Clique-Based Clustering of Correlated SNPs in a Gene Can Improve Performance of Gene-Based Multi-Bin Linear Combination Test. BioMed Research International, 2015, 2015, 1-11.	1.9	9
38	Regression models for allele sharing: analysis of accumulating data in affected sib pair studies. Statistics in Medicine, 2002, 21, 431-444.	1.6	8
39	Are quantitative trait-dependent sampling designs cost-effective for analysis of rare and common variants?. BMC Proceedings, 2011, 5, S111.	1.6	8
40	Does Familial Clustering of Risk Factors for Long-Term Diabetic Complications Leave Any Place for Genes That Act independently?. Journal of Cardiovascular Translational Research, 2012, 5, 388-398.	2.4	8
41	Statistical power in COVID-19 case-control host genomic study design. Genome Medicine, 2020, 12, 115.	8.2	7
42	Heterogeneity in IBD Allele Sharing among Covariate-Defined Subgroups: Issues and Findings for Affected Relatives. Human Heredity, 2003, 56, 94-106.	0.8	6
43	Regression models, scan statistics and reappearance probabilities to detect regions of association between gene expression and copy number. Statistics in Medicine, 2011, 30, 1157-1178.	1.6	6
44	Twoâ€phase designs for joint quantitativeâ€traitâ€dependent and genotypeâ€dependent sampling in postâ€GW regional sequencing. Genetic Epidemiology, 2018, 42, 104-116.	/AS 1.3	6
45	The Efficiency of Multinomial Logistic Regression Compared With Multiple Group Discriminant Analysis. Journal of the American Statistical Association, 1987, 82, 1118.	3.1	6
46	Twoâ€Phase Stratified Sampling Designs for Regional Sequencing. Genetic Epidemiology, 2012, 36, 320-332.	1.3	5
47	A 2-step strategy for detecting pleiotropic effects on multiple longitudinal traits. Frontiers in Genetics, 2014, 5, 357.	2.3	5
48	Efficiency of reduced logistic regression models. Canadian Journal of Statistics, 1994, 22, 319-334.	0.9	4
49	Non-invasive Electroarthrography Measures Load-Induced Cartilage Streaming Potentials via Electrodes Placed on Skin Surrounding an Articular Joint. Cartilage, 2020, , 194760352092858.	2.7	3
50	Sequence kernel association test for survival outcomes in the presence of a non-susceptible fraction. Biostatistics, 2020, 21, 518-530.	1.5	3
51	Disease-marker associations: Power and heterogeneity in independent population samples. Genetic Epidemiology, 1999, 17, S509-S514.	1.3	2
52	Modeling Complex Disease with Demographic and Environmental Covariates and a Candidate Gene Marker. Genetic Epidemiology, 2001, 21, S423-S428.	1.3	2
53	A Note on the Efficiencies of Sampling Strategies in Twoâ€Stage Bayesian Regional Fine Mapping of a Quantitative Trait. Genetic Epidemiology, 2014, 38, 599-609.	1.3	2
54	Multiphase analysis by linkage, quantitative transmission disequilibrium, and measured genotype: systolic blood pressure in complex Mexican American pedigrees. BMC Proceedings, 2014, 8, S108.	1.6	2

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#	Article	IF	CITATIONS
55	Twoâ€phase sample selection strategies for design and analysis in postâ€genomeâ€wide association fineâ€mapping studies. Statistics in Medicine, 2021, 40, 6792-6817.	1.6	2
56	Comparative Expression of the Mitotic Regulators SAK and PLK in Colorectal Cancer. Annals of Surgical Oncology, 2001, 8, 729-740.	1.5	2
57	Uncovering the Contribution of Moderate-Penetrance Susceptibility Genes to Breast Cancer by Whole-Exome Sequencing and Targeted Enrichment Sequencing of Candidate Genes in Women of European Ancestry. Cancers, 2022, 14, 3363.	3.7	2
58	Comparison of evidence for linkage from different analytic methods. Genetic Epidemiology, 1997, 14, 965-970.	1.3	1
59	Does Simultaneous Consideration of Multiple Regions Improve Disease Gene Localization?. Genetic Epidemiology, 2001, 21, S504-S509.	1.3	1
60	An exploration of heterogeneity in genetic analysis of complex pedigrees: linkage and association using whole genome sequencing data in the MAP4 region. BMC Proceedings, 2014, 8, S107.	1.6	1
61	Geneâ€based and pathwayâ€based testing for rareâ€variant association in affected sib pairs. Genetic Epidemiology, 2020, 44, 368-381.	1.3	1
62	Statistical challenges in highâ€dimensional molecular and genetic epidemiology. Canadian Journal of Statistics, 2018, 46, 24-40.	0.9	0