Obi Lee Griffith

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

176	19,396	59	139
papers	citations	h-index	g-index
207	23,096 ext. citations	10.6	5.93
ext. papers		avg, IF	L-index

#	Paper	IF	Citations
176	Standards for the classification of pathogenicity of somatic variants in cancer (oncogenicity): Joint recommendations of Clinical Genome Resource (ClinGen), Cancer Genomics Consortium (CGC), and Variant Interpretation for Cancer Consortium (VICC) <i>Genetics in Medicine</i> , 2022 ,	8.1	4
175	Bam-readcount - rapid generation of basepair-resolution sequence metrics. <i>Journal of Open Source Software</i> , 2022 , 7, 3722	5.2	2
174	Hematopoietic cell transplantation donor-derived memory-like NK cells functionally persist after transfer into patients with leukemia <i>Science Translational Medicine</i> , 2022 , 14, eabm1375	17.5	2
173	Integrative genomic analysis reveals low T-cell infiltration as the primary feature of tobacco use in HPV-positive oropharyngeal cancer <i>IScience</i> , 2022 , 25, 104216	6.1	0
172	Standardized evidence-based approach for assessment of oncogenic and clinical significance of NTRK fusions <i>Cancer Genetics</i> , 2022 , 264-265, 50-59	2.3	1
171	Genomic and transcriptomic somatic alterations of hepatocellular carcinoma in non-cirrhotic livers <i>Cancer Genetics</i> , 2022 , 264-265, 90-99	2.3	1
170	A community approach to the cancer-variant-interpretation bottleneck. <i>Nature Cancer</i> , 2022 , 3, 522-525	515.4	O
169	GA4GH: International policies and standards for data sharing across genomic research and healthcare <i>Cell Genomics</i> , 2021 , 1, 100029-100029		20
168	The GA4GH Variation Representation Specification: A computational framework for variation representation and federated identification <i>Cell Genomics</i> , 2021 , 1, 100027-100027		4
167	Expert Curation of Somatic FLT3 Variants By the Clingen Somatic Hematologic Cancer Taskforce (ClinGen HCT). <i>Blood</i> , 2021 , 138, 4387-4387	2.2	
166	Cytokine-Induced Memory-like NK Cells Have a Distinct Single Cell Transcriptional Profile and Persist for Months in Adult and Pediatric Leukemia Patients after Adoptive Transfer. <i>Blood</i> , 2021 , 138, 3825-3825	2.2	
165	In Silico Epitope Prediction Analyses Highlight the Potential for Distracting Antigen Immunodominance with Allogeneic Cancer Vaccines. <i>Cancer Research Communications</i> , 2021 , 1, 115-126	5	
164	Characterization of the Genomic and Immunological Diversity of Malignant Brain Tumors Through Multi-Sector Analysis. <i>Cancer Discovery</i> , 2021 ,	24.4	4
163	Genetic Ancestry Correlations with Driver Mutations Suggest Complex Interactions between Somatic and Germline Variation in Cancer. <i>Cancer Discovery</i> , 2021 , 11, 534-536	24.4	О
162	Phase I Trial of N-803, an IL15 Receptor Agonist, with Rituximab in Patients with Indolent Non-Hodgkin Lymphoma. <i>Clinical Cancer Research</i> , 2021 , 27, 3339-3350	12.9	7
161	Epigenomic regulation of human T-cell leukemia virus by chromatin-insulator CTCF. <i>PLoS Pathogens</i> , 2021 , 17, e1009577	7.6	1
160	Multitarget Stool RNA Test for Noninvasive Detection of Colorectal Neoplasias in a Multicenter, Prospective, and Retrospective Cohort. <i>Clinical and Translational Gastroenterology</i> , 2021 , 12, e00360	4.2	O

(2020-2021)

159	Immunogenomic profiling and pathological response results from a clinical trial of docetaxel and carboplatin in triple-negative breast cancer. <i>Breast Cancer Research and Treatment</i> , 2021 , 189, 187-202	4.4	5
158	OMIC-13. THE ROLE OF COPY NUMBER ALTERATIONS IN PREDICTING SURVIVAL AND INFLUENCING TREATMENT OF CHILDHOOD BRAIN TUMORS. <i>Neuro-Oncology</i> , 2021 , 23, i40-i40	1	78
157	Integration of the Drug-Gene Interaction Database (DGIdb 4.0) with open crowdsource efforts. <i>Nucleic Acids Research</i> , 2021 , 49, D1144-D1151	20.1	71
156	Colonosight: Multitarget RNA-FIT assay for noninvasive detection of advanced colorectal neoplasia <i>Journal of Clinical Oncology</i> , 2021 , 39, 25-25	2.2	
155	Unraveling the chaotic genomic landscape of primary and metastatic canine appendicular osteosarcoma with current sequencing technologies and bioinformatic approaches. <i>PLoS ONE</i> , 2021 , 16, e0246443	3.7	О
154	Yap1 Mediates Trametinib Resistance in Head and Neck Squamous Cell Carcinomas. <i>Clinical Cancer Research</i> , 2021 , 27, 2326-2339	12.9	1
153	Exploring the Genomic Landscape of Cancer Patient Cohorts with GenVisR. <i>Current Protocols</i> , 2021 , 1, e252		2
152	Impact of a 40-Gene Targeted Panel Test on Physician Decision Making for Patients With Acute Myeloid Leukemia. <i>JCO Precision Oncology</i> , 2021 , 5,	3.6	2
151	Donor Memory-like NK cells Persist and Induce Remissions in Pediatric Patients with Relapsed AML after Transplant. <i>Blood</i> , 2021 ,	2.2	5
150	Sex- and mutation-specific p53 gain-of-function activity in gliomagenesis <i>Cancer Research Communications</i> , 2021 , 1, 148-163		О
149	The clonal evolution of metastatic colorectal cancer. <i>Science Advances</i> , 2020 , 6, eaay9691	14.3	14
148	CIViCpy: A Python Software Development and Analysis Toolkit for the CIViC Knowledgebase. <i>JCO Clinical Cancer Informatics</i> , 2020 , 4, 245-253	5.2	5
147	A harmonized meta-knowledgebase of clinical interpretations of somatic genomic variants in cancer. <i>Nature Genetics</i> , 2020 , 52, 448-457	36.3	58
146	B-Cell Acute Lymphoblastic Leukemia Arising in Patients with a Preexisting Diagnosis of Multiple Myeloma Is a Novel Cancer with High Incidence of TP53 Mutations. <i>Blood</i> , 2020 , 136, 20-20	2.2	О
145	IMMU-53. CHARACTERIZATION OF THE GENOMIC AND IMMUNOLOGICAL DIVERSITY OF MALIGNANT BRAIN TUMORS THROUGH MULTI-SECTOR ANALYSIS. <i>Neuro-Oncology</i> , 2020 , 22, ii116-ii11	6 ¹	
144	Wikidata as a knowledge graph for the life sciences. <i>ELife</i> , 2020 , 9,	8.9	24
143	Reply. <i>Gastroenterology</i> , 2020 , 158, 793-794	13.3	
142	Splicing factor SF3B1 promotes endometrial cancer progression via regulating KSR2 RNA maturation. <i>Cell Death and Disease</i> , 2020 , 11, 842	9.8	10

141	Neoadjuvant and Adjuvant Pembrolizumab in Resectable Locally Advanced, Human Papillomavirus-Unrelated Head and Neck Cancer: A Multicenter, Phase II Trial. <i>Clinical Cancer Research</i> , 2020 , 26, 5140-5152	12.9	71
140	Collaborative, Multidisciplinary Evaluation of Cancer Variants Through Virtual Molecular Tumor Boards Informs Local Clinical Practices. <i>JCO Clinical Cancer Informatics</i> , 2020 , 4, 602-613	5.2	11
139	pVACtools: A Computational Toolkit to Identify and Visualize Cancer Neoantigens. <i>Cancer Immunology Research</i> , 2020 , 8, 409-420	12.5	53
138	Targeting the Mevalonate Pathway to Overcome Acquired Anti-HER2 Treatment Resistance in Breast Cancer. <i>Molecular Cancer Research</i> , 2019 , 17, 2318-2330	6.6	22
137	Rapid progression of adult T-cell leukemia/lymphoma as tumor-infiltrating Tregs after PD-1 blockade. <i>Blood</i> , 2019 , 134, 1406-1414	2.2	50
136	Best practices for bioinformatic characterization of neoantigens for clinical utility. <i>Genome Medicine</i> , 2019 , 11, 56	14.4	73
135	A Spontaneous Aggressive ER⊞ Mammary Tumor Model Is Driven by Kras Activation. <i>Cell Reports</i> , 2019 , 28, 1526-1537.e4	10.6	10
134	Noninvasive Detection of High-Risk Adenomas Using Stool-Derived Eukaryotic RNA Sequences as Biomarkers. <i>Gastroenterology</i> , 2019 , 157, 884-887.e3	13.3	6
133	Open-Sourced CIViC Annotation Pipeline to Identify and Annotate Clinically Relevant Variants Using Single-Molecule Molecular Inversion Probes. <i>JCO Clinical Cancer Informatics</i> , 2019 , 3, 1-12	5.2	4
132	Immunological ignorance is an enabling feature of the oligo-clonal T cell response to melanoma neoantigens. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019 , 116, 23662-23670	11.5	19
131	Standard operating procedure for curation and clinical interpretation of variants in cancer. <i>Genome Medicine</i> , 2019 , 11, 76	14.4	6
130	Text-mining clinically relevant cancer biomarkers for curation into the CIViC database. <i>Genome Medicine</i> , 2019 , 11, 78	14.4	19
129	Accounting for proximal variants improves neoantigen prediction. <i>Nature Genetics</i> , 2019 , 51, 175-179	36.3	23
128	Standard operating procedure for somatic variant refinement of sequencing data with paired tumor and hormal samples. <i>Genetics in Medicine</i> , 2019 , 21, 972-981	8.1	31
127	DGIdb 3.0: a redesign and expansion of the drug-gene interaction database. <i>Nucleic Acids Research</i> , 2018 , 46, D1068-D1073	20.1	414
126	Single-agent ibrutinib in relapsed or refractory follicular lymphoma: a phase 2 consortium trial. <i>Blood</i> , 2018 , 131, 182-190	2.2	92
125	Oral Cavity Squamous Cell Carcinoma Xenografts Retain Complex Genotypes and Intertumor Molecular Heterogeneity. <i>Cell Reports</i> , 2018 , 24, 2167-2178	10.6	17
124	ClinGen Cancer Somatic Working Group - standardizing and democratizing access to cancer molecular diagnostic data to drive translational research. <i>Pacific Symposium on Biocomputing Pacific Symposium on Biocomputing</i> , 2018 , 23, 247-258	1.3	13

Standardizing And Democratizing Access To Cancer Molecular Diagnostic Test Data From Patients 123 To Drive Translational Research. AMIA Summits on Translational Science Proceedings, **2018**, 2017, 152-15 $9^{1.1}$ A case of acute myeloid leukemia with promyelocytic features characterized by expression of a 18 122 7.8 novel - fusion. *Blood Advances*, **2018**, 2, 1295-1299 A deep learning approach to automate refinement of somatic variant calling from cancer 121 36.3 38 sequencing data. Nature Genetics, 2018, 50, 1735-1743 Adapting crowdsourced clinical cancer curation in CIVIC to the ClinGen minimum variant level data 120 11 community-driven standards. Human Mutation, 2018, 39, 1721-1732 Recurrent WNT pathway alterations are frequent in relapsed small cell lung cancer. Nature 119 63 17.4 Communications, 2018, 9, 3787 The prognostic effects of somatic mutations in ER-positive breast cancer. Nature Communications, 118 17.4 **2018**, 9, 3476 CIViC is a community knowledgebase for expert crowdsourcing the clinical interpretation of 308 117 36.3 variants in cancer. Nature Genetics, 2017, 49, 170-174 NeoPalAna: Neoadjuvant Palbociclib, a Cyclin-Dependent Kinase 4/6 Inhibitor, and Anastrozole for 116 160 Clinical Stage 2 or 3 Estrogen Receptor-Positive Breast Cancer. Clinical Cancer Research, 2017, 23, 4055-4065 Melorheostosis: Exome sequencing of an associated dermatosis implicates postzygotic mosaicism 115 4.7 25 of mutated KRAS. Bone, 2017, 101, 145-155 Genomic characterization of HER2-positive breast cancer and response to neoadjuvant trastuzumab and chemotherapy-results from the ACOSOG Z1041 (Alliance) trial. Annals of Oncology 114 10.3 35 , **2017**, 28, 1070-1077 Recurrent somatic mutations affecting B-cell receptor signaling pathway genes in follicular 113 2.2 98 lymphoma. Blood, 2017, 129, 473-483 Clinical implications of neoepitope landscapes for adult and pediatric cancers. Genome Medicine, 112 14.4 4 **2017**, 9, 77 A Phase II Trial of Neoadjuvant MK-2206, an AKT Inhibitor, with Anastrozole in Clinical Stage II or III 111 12.9 51 -Mutant ER-Positive and HER2-Negative Breast Cancer. Clinical Cancer Research, 2017, 23, 6823-6832 Comprehensive discovery of noncoding RNAs in acute myeloid leukemia cell transcriptomes. 110 8 3.1 Experimental Hematology, 2017, 55, 19-33 Neoantigens in immunotherapy and personalized vaccines: Implications for head and neck 109 13 4.4 squamous cell carcinoma. Oral Oncology, 2017, 71, 169-176 Aromatase inhibition remodels the clonal architecture of estrogen-receptor-positive breast 108 17.4 47 cancers. Nature Communications, 2016, 7, 12498 A genomic analysis of Philadelphia chromosome-negative AML arising in patients with CML. Blood 107 12 Cancer Journal, 2016, 6, e413 Comprehensive genomic analysis reveals FLT3 activation and a therapeutic strategy for a patient 106 33 with relapsed adult B-lymphoblastic leukemia. Experimental Hematology, 2016, 44, 603-13

105	High-performance web services for querying gene and variant annotation. <i>Genome Biology</i> , 2016 , 17, 91	18.3	111
104	pVAC-Seq: A genome-guided in silico approach to identifying tumor neoantigens. <i>Genome Medicine</i> , 2016 , 8, 11	14.4	221
103	GenVisR: Genomic Visualizations in R. <i>Bioinformatics</i> , 2016 , 32, 3012-4	7.2	146
102	ORegAnno 3.0: a community-driven resource for curated regulatory annotation. <i>Nucleic Acids Research</i> , 2016 , 44, D126-32	20.1	89
101	A Phase I Trial of BKM120 (Buparlisib) in Combination with Fulvestrant in Postmenopausal Women with Estrogen Receptor-Positive Metastatic Breast Cancer. <i>Clinical Cancer Research</i> , 2016 , 22, 1583-91	12.9	77
100	Exome Sequencing of Hodgkins and Non-Hodgkin Composite Lymphomas Identifies Shared Somatic Mutations Indicative of Common Founding Precursors. <i>Blood</i> , 2016 , 128, 5285-5285	2.2	
99	Clonal Evolution Revealed By Exome Sequencing in a Case of Primary Myelofibrosis Associated with Subsequent Development of Aggressive Systemic Mastocytosis/Mast Cell Leukemia. <i>Blood</i> , 2016 , 128, 5496-5496	2.2	
98	Cancer Immunogenomics: Computational Neoantigen Identification and Vaccine Design. <i>Cold Spring Harbor Symposia on Quantitative Biology</i> , 2016 , 81, 105-111	3.9	15
97	A common founding clone with TP53 and PTEN mutations gives rise to a concurrent germ cell tumor and acute megakaryoblastic leukemia. <i>Journal of Physical Education and Sports Management</i> , 2016 , 2, a000687	2.8	12
96	DGIdb 2.0: mining clinically relevant drug-gene interactions. <i>Nucleic Acids Research</i> , 2016 , 44, D1036-44	20.1	222
95	A genomic case study of mixed fibrolamellar hepatocellular carcinoma. <i>Annals of Oncology</i> , 2016 , 27, 1148-1154	10.3	13
94	DoCM: a database of curated mutations in cancer. <i>Nature Methods</i> , 2016 , 13, 806-7	21.6	63
93	Truncating Prolactin Receptor Mutations Promote Tumor Growth in Murine Estrogen Receptor-Alpha Mammary Carcinomas. <i>Cell Reports</i> , 2016 , 17, 249-260	10.6	14
92	FOXA1 overexpression mediates endocrine resistance by altering the ER transcriptome and IL-8 expression in ER-positive breast cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016 , 113, E6600-E6609	11.5	91
91	U2AF1 mutations alter sequence specificity of pre-mRNA binding and splicing. <i>Leukemia</i> , 2015 , 29, 909-	17 0.7	79
90	Epigenetic and transcriptional determinants of the human breast. <i>Nature Communications</i> , 2015 , 6, 635	1 17.4	44
89	RNA-sequencing reveals oligodendrocyte and neuronal transcripts in microglia relevant to central nervous system disease. <i>Glia</i> , 2015 , 63, 531-548	9	26
88	Statistically identifying tumor suppressors and oncogenes from pan-cancer genome-sequencing data. <i>Bioinformatics</i> , 2015 , 31, 3561-8	7.2	30

(2014-2015)

87	Mutant U2AF1 Expression Alters Hematopoiesis and Pre-mRNA Splicing In Vivo. <i>Cancer Cell</i> , 2015 , 27, 631-43	24.3	186
86	Association Between Mutation Clearance After Induction Therapy and Outcomes in Acute Myeloid Leukemia. <i>JAMA - Journal of the American Medical Association</i> , 2015 , 314, 811-22	27.4	242
85	RNA Sequencing of Tumor-Associated Microglia Reveals Ccl5 as a Stromal Chemokine Critical for Neurofibromatosis-1 Glioma Growth. <i>Neoplasia</i> , 2015 , 17, 776-88	6.4	62
84	Convergent loss of PTEN leads to clinical resistance to a PI(3)K nhibitor. <i>Nature</i> , 2015 , 518, 240-4	50.4	366
83	Optimizing cancer genome sequencing and analysis. <i>Cell Systems</i> , 2015 , 1, 210-223	10.6	135
82	Genome Modeling System: A Knowledge Management Platform for Genomics. <i>PLoS Computational Biology</i> , 2015 , 11, e1004274	5	59
81	Informatics for RNA Sequencing: A Web Resource for Analysis on the Cloud. <i>PLoS Computational Biology</i> , 2015 , 11, e1004393	5	50
8o	Recurrent Somatic Genomic Alterations in Follicular NHL (FL) Revealed By Exome and Custom-Capture Next Generation Sequencing. <i>Blood</i> , 2015 , 126, 574-574	2.2	2
79	Decoupling of the PI3K Pathway via Mutation Necessitates Combinatorial Treatment in HER2+ Breast Cancer. <i>PLoS ONE</i> , 2015 , 10, e0133219	3.7	16
78	Single exon-resolution targeted chromosomal microarray analysis of known and candidate intellectual disability genes. <i>European Journal of Human Genetics</i> , 2014 , 22, 792-800	5.3	30
77	Organizing knowledge to enable personalization of medicine in cancer. <i>Genome Biology</i> , 2014 , 15, 438	18.3	53
76	Identification of gene regulation patterns underlying both oestrogen- and tamoxifen-stimulated cell growth through global gene expression profiling in breast cancer cells. <i>European Journal of Cancer</i> , 2014 , 50, 2877-86	7.5	12
75	cDNA hybrid capture improves transcriptome analysis on low-input and archived samples. <i>Journal of Molecular Diagnostics</i> , 2014 , 16, 440-51	5.1	35
74	Clonal architectures and driver mutations in metastatic melanomas. <i>PLoS ONE</i> , 2014 , 9, e111153	3.7	53
73	SciClone: inferring clonal architecture and tracking the spatial and temporal patterns of tumor evolution. <i>PLoS Computational Biology</i> , 2014 , 10, e1003665	5	301
72	Hippo signaling influences HNF4A and FOXA2 enhancer switching during hepatocyte differentiation. <i>Cell Reports</i> , 2014 , 9, 261-271	10.6	59
71	Functional heterogeneity of genetically defined subclones in acute myeloid leukemia. <i>Cancer Cell</i> , 2014 , 25, 379-92	24.3	273
70	The cure: design and evaluation of a crowdsourcing game for gene selection for breast cancer survival prediction. <i>JMIR Serious Games</i> , 2014 , 2, e7	3.4	16

69	DGIdb: mining the druggable genome. <i>Nature Methods</i> , 2013 , 10, 1209-10	21.6	317
68	Novel mRNA isoforms and mutations of uridine monophosphate synthetase and 5-fluorouracil resistance in colorectal cancer. <i>Pharmacogenomics Journal</i> , 2013 , 13, 148-58	3.5	18
67	A robust prognostic signature for hormone-positive node-negative breast cancer. <i>Genome Medicine</i> , 2013 , 5, 92	14.4	17
66	Endocrine-therapy-resistant ESR1 variants revealed by genomic characterization of breast-cancer-derived xenografts. <i>Cell Reports</i> , 2013 , 4, 1116-30	10.6	447
65	c-Src modulates estrogen-induced stress and apoptosis in estrogen-deprived breast cancer cells. <i>Cancer Research</i> , 2013 , 73, 4510-20	10.1	65
64	Modeling precision treatment of breast cancer. <i>Genome Biology</i> , 2013 , 14, R110	18.3	204
63	F11R is a novel monocyte prognostic biomarker for malignant glioma. <i>PLoS ONE</i> , 2013 , 8, e77571	3.7	32
62	Cross-platform pathway-based analysis identifies markers of response to the PARP inhibitor olaparib. <i>Breast Cancer Research and Treatment</i> , 2012 , 135, 505-17	4.4	54
61	Lectin chromatography/mass spectrometry discovery workflow identifies putative biomarkers of aggressive breast cancers. <i>Journal of Proteome Research</i> , 2012 , 11, 2508-20	5.6	45
60	Genomic landscape of non-small cell lung cancer in smokers and never-smokers. <i>Cell</i> , 2012 , 150, 1121-3	456.2	860
59	Identification of PADI2 as a potential breast cancer biomarker and therapeutic target. <i>BMC Cancer</i> , 2012 , 12, 500	4.8	79
58	The clonal and mutational evolution spectrum of primary triple-negative breast cancers. <i>Nature</i> , 2012 , 486, 395-9	50.4	1417
57	On the Deep Order-Preserving Submatrix Problem: A Best Effort Approach. <i>IEEE Transactions on Knowledge and Data Engineering</i> , 2012 , 24, 309-325	4.2	16
56	Loss of cell-surface laminin anchoring promotes tumor growth and is associated with poor clinical outcomes. <i>Cancer Research</i> , 2012 , 72, 2578-88	10.1	37
55	Interaction of cyclin-dependent kinase 12/CrkRS with cyclin K1 is required for the phosphorylation of the C-terminal domain of RNA polymerase II. <i>Molecular and Cellular Biology</i> , 2012 , 32, 4691-704	4.8	71
54	Frequent mutation of histone-modifying genes in non-Hodgkin lymphoma. <i>Nature</i> , 2011 , 476, 298-303	50.4	1180
53	Immunophenotyping of thyroid tumors identifies molecular markers altered during transformation of differentiated into anaplastic carcinoma. <i>American Journal of Surgery</i> , 2011 , 201, 580-6	2.7	13
52	Somic approaches to preventing or managing metastatic breast cancer. <i>Breast Cancer Research</i> , 2011 , 13, 230	8.3	19

(2008-2011)

51	Detection and management of hypothyroidism following thyroid lobectomy: evaluation of a clinical algorithm. <i>Annals of Surgical Oncology</i> , 2011 , 18, 2548-54	3.1	37
50	Lrig1 is an estrogen-regulated growth suppressor and correlates with longer relapse-free survival in EREpositive breast cancer. <i>Molecular Cancer Research</i> , 2011 , 9, 1406-17	6.6	51
49	Extensive relationship between antisense transcription and alternative splicing in the human genome. <i>Genome Research</i> , 2011 , 21, 1203-12	9.7	63
48	deFuse: an algorithm for gene fusion discovery in tumor RNA-Seq data. <i>PLoS Computational Biology</i> , 2011 , 7, e1001138	5	409
47	Somatic mutations altering EZH2 (Tyr641) in follicular and diffuse large B-cell lymphomas of germinal-center origin. <i>Nature Genetics</i> , 2010 , 42, 181-5	36.3	1273
46	Alternative expression analysis by RNA sequencing. <i>Nature Methods</i> , 2010 , 7, 843-7	21.6	227
45	De novo assembly and analysis of RNA-seq data. <i>Nature Methods</i> , 2010 , 7, 909-12	21.6	701
44	Diagnostic utility of galectin-3 in thyroid cancer. American Journal of Pathology, 2010 , 176, 2067-81	5.8	118
43	Evolution of an adenocarcinoma in response to selection by targeted kinase inhibitors. <i>Genome Biology</i> , 2010 , 11, R82	18.3	144
42	Personalized oncogenomics 2010 , 11, I5		78
41	Genomic analysis of a rare human tumor. <i>BMC Bioinformatics</i> , 2010 , 11,	3.6	78
40	Annotating the regulatory genome. <i>Methods in Molecular Biology</i> , 2010 , 674, 313-49	1.4	2
39	Thyroid Cancer: Identification of Gene Expression Markers for Diagnosis 2010 , 353-377		
38	The completion of the Mammalian Gene Collection (MGC). Genome Research, 2009, 19, 2324-33	9.7	98
37	KiWi: A Scalable Subspace Clustering Algorithm for Gene Expression Analysis 2009,		6
36	ALEXA: a microarray design platform for alternative expression analysis. <i>Nature Methods</i> , 2008 , 5, 118	21.6	19
35	Biomarker panel diagnosis of thyroid cancer: a critical review. <i>Expert Review of Anticancer Therapy</i> , 2008 , 8, 1399-413	3.5	35
	2000 , 0, 1399-413		

33	Clinical utility of type 1 growth factor receptor expression in colon cancer. <i>American Journal of Surgery</i> , 2008 , 195, 604-10	2.7	16
32	Evaluation of type 1 growth factor receptor family expression in benign and malignant thyroid lesions. <i>American Journal of Surgery</i> , 2008 , 195, 667-73; discussion 673	2.7	15
31	In-depth characterization of the microRNA transcriptome in a leukemia progression model. <i>Genome Research</i> , 2008 , 18, 1787-97	9.7	148
30	Application of massively parallel sequencing to microRNA profiling and discovery in human embryonic stem cells. <i>Genome Research</i> , 2008 , 18, 610-21	9.7	879
29	Impact of whole genome amplification on analysis of copy number variants. <i>Nucleic Acids Research</i> , 2008 , 36, e80	20.1	70
28	Phosphorylated caveolin-1 regulates Rho/ROCK-dependent focal adhesion dynamics and tumor cell migration and invasion. <i>Cancer Research</i> , 2008 , 68, 8210-20	10.1	200
27	Meta-analysis of colorectal cancer gene expression profiling studies identifies consistently reported candidate biomarkers. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2008 , 17, 543-52	4	120
26	Molecular phenotyping of thyroid tumors identifies a marker panel for differentiated thyroid cancer diagnosis. <i>Annals of Surgical Oncology</i> , 2008 , 15, 2811-26	3.1	43
25	ORegAnno: an open-access community-driven resource for regulatory annotation. <i>Nucleic Acids Research</i> , 2008 , 36, D107-13	20.1	199
24	Genome-wide profiles of STAT1 DNA association using chromatin immunoprecipitation and massively parallel sequencing. <i>Nature Methods</i> , 2007 , 4, 651-7	21.6	1077
23	Cell cycle regulators show diagnostic and prognostic utility for differentiated thyroid cancer. <i>Annals of Surgical Oncology</i> , 2007 , 14, 3403-11	3.1	38
22	A survey of genomic properties for the detection of regulatory polymorphisms. <i>PLoS Computational Biology</i> , 2007 , 3, e106	5	22
21	Identification of molecular markers altered during transformation of differentiated into anaplastic thyroid carcinoma. <i>Archives of Surgery</i> , 2007 , 142, 717-27; discussion 727-9		72
20	Analysis of the prostate cancer cell line LNCaP transcriptome using a sequencing-by-synthesis approach. <i>BMC Genomics</i> , 2006 , 7, 246	4.5	139
19	Sequence biases in large scale gene expression profiling data. <i>Nucleic Acids Research</i> , 2006 , 34, e83	20.1	44
18	Discovering significant OPSM subspace clusters in massive gene expression data 2006,		19
17	Sequencing and analysis of 10,967 full-length cDNA clones from Xenopus laevis and Xenopus tropicalis reveals post-tetraploidization transcriptome remodeling. <i>Genome Research</i> , 2006 , 16, 796-803	9.7	59
16	Meta-analysis and meta-review of thyroid cancer gene expression profiling studies identifies important diagnostic biomarkers. <i>Journal of Clinical Oncology</i> , 2006 , 24, 5043-51	2.2	232

LIST OF PUBLICATIONS

15	Assessment and integration of publicly available SAGE, cDNA microarray, and oligonucleotide microarray expression data for global coexpression analyses. <i>Genomics</i> , 2005 , 86, 476-88	4.3	40
14	Systematic recovery and analysis of full-ORF human cDNA clones. <i>Genome Research</i> , 2004 , 14, 2083-92	9.7	22
13	The status, quality, and expansion of the NIH full-length cDNA project: the Mammalian Gene Collection (MGC). <i>Genome Research</i> , 2004 , 14, 2121-7	9.7	404
12	Novel avian influenza H7N3 strain outbreak, British Columbia. <i>Emerging Infectious Diseases</i> , 2004 , 10, 2192-5	10.2	154
11	Escherichia coli endA deletion strain for use in two-hybrid shuttle vector selection. <i>BioTechniques</i> , 2003 , 35, 272-4, 276, 278	2.5	6
10	The Genome sequence of the SARS-associated coronavirus. <i>Science</i> , 2003 , 300, 1399-404	33.3	1632
9	Clingen Cancer Somatic Working Group Istandardizing and democratizing access to cancer molecular diagnostic data to drive translational research		1
8	CIViC: A knowledgebase for expert-crowdsourcing the clinical interpretation of variants in cancer		2
7	DGIdb 3.0: a redesign and expansion of the drug-gene interaction database		5
6	Integration of the Drug-Gene Interaction Database (DGIdb) with open crowdsource efforts		1
5	A harmonized meta-knowledgebase of clinical interpretations of cancer genomic variants		5
4	RegTools: Integrated analysis of genomic and transcriptomic data for the discovery of splicing variants in cancer		25
3	pVACtools: a computational toolkit to identify and visualize cancer neoantigens		5
2	Wikidata as a FAIR knowledge graph for the life sciences		1
1	The GA4GH Variation Representation Specification (VRS): a Computational Framework for the Precise Representation and Federated Identification of Molecular Variation		3