Marco A Marra

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1027399/publications.pdf

Version: 2024-02-01

		120	28
513	222,497	166	446
papers	citations	h-index	g-index
533	533	533	208092
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Optimization of magnetic bead-based nucleic acid extraction for SARS-CoV-2 testing using readily available reagents. Journal of Virological Methods, 2022, 299, 114339.	1.0	4
2	Early-stage economic analysis of research-based comprehensive genomic sequencing for advanced cancer care. Journal of Community Genetics, 2022, 13, 523-538.	0.5	4
3	Integrative multiâ€omic analysis reveals neurodevelopmental gene dysregulation in <scp><i>ClC</i></scp> â€knockout and <scp><i>IDH1</i></scp> â€mutant cells. Journal of Pathology, 2022, 256, 297-309.	2.1	5
4	ICGC-ARGO precision medicine: familial matters in pancreatic cancer. Lancet Oncology, The, 2022, 23, 25-26.	5.1	6
5	A platform for oncogenomic reporting and interpretation. Nature Communications, 2022, 13, 756.	5.8	7
6	Cost-Effectiveness of Molecularly Guided Treatment in Diffuse Large B-Cell Lymphoma (DLBCL) in Patients under 60. Cancers, 2022, 14, 908.	1.7	0
7	The Neoantigen Landscape of the Coding and Noncoding Cancer Genome Space. Journal of Molecular Diagnostics, 2022, , .	1.2	O
8	Single-cell landscapes of primary glioblastomas and matched explants and cell lines show variable retention of inter- and intratumor heterogeneity. Cancer Cell, 2022, 40, 379-392.e9.	7.7	54
9	The impact of whole genome and transcriptome analysis (<scp>WGTA</scp>) on predictive biomarker discovery and diagnostic accuracy of advanced malignancies. Journal of Pathology: Clinical Research, 2022, 8, 395-407.	1.3	3
10	Combinatorial and Machine Learning Approaches for Improved Somatic Variant Calling From Formalin-Fixed Paraffin-Embedded Genome Sequence Data. Frontiers in Genetics, 2022, 13, 834764.	1.1	1
11	Whole-genome and transcriptome analysis of advanced adrenocortical cancer highlights multiple alterations affecting epigenome and DNA repair pathways Cold Spring Harbor Molecular Case Studies, 2022, 8, .	0.7	2
12	Impact of MYC and BCL2 structural variants in tumors of DLBCL morphology and mechanisms of false-negative MYC IHC. Blood, 2021, 137, 2196-2208.	0.6	18
13	Uncovering Clinically Relevant Gene Fusions with Integrated Genomic and Transcriptomic Profiling of Metastatic Cancers. Clinical Cancer Research, 2021, 27, 522-531.	3.2	14
14	Wholeâ€slide laser microdissection for tumour enrichment. Journal of Pathology, 2021, 253, 225-233.	2.1	4
15	Delving into Early-onset Pancreatic Ductal Adenocarcinoma: How Does Age Fit In?. Clinical Cancer Research, 2021, 27, 246-254.	3.2	16
16	Genome and Transcriptome Biomarkers of Response to Immune Checkpoint Inhibitors in Advanced Solid Tumors. Clinical Cancer Research, 2021, 27, 202-212.	3.2	50
17	Subtype-Discordant Pancreatic Ductal Adenocarcinoma Tumors Show Intermediate Clinical and Molecular Characteristics. Clinical Cancer Research, 2021, 27, 150-157.	3.2	24
18	Matching methods in precision oncology: An introduction and illustrative example. Molecular Genetics & Genomic Medicine, 2021, 9, e1554.	0.6	13

#	Article	lF	Citations
19	Molecular attributes underlying central nervous system and systemic relapse in diffuse large B-cell lymphoma. Haematologica, 2021, 106, 1466-1471.	1.7	9
20	Gradient of Developmental and Injury Response transcriptional states defines functional vulnerabilities underpinning glioblastoma heterogeneity. Nature Cancer, 2021, 2, 157-173.	5.7	147
21	NTRK2 Fusion driven pediatric glioblastoma: Identification of oncogenic Drivers via integrative Genome and transcriptome profiling. Clinical Case Reports (discontinued), 2021, 9, 1472-1477.	0.2	3
22	Megabase-scale methylation phasing using nanopore long reads and NanoMethPhase. Genome Biology, 2021, 22, 68.	3.8	36
23	The transcriptional landscape of Shh medulloblastoma. Nature Communications, 2021, 12, 1749.	5.8	47
24	A clinical transcriptome approach to patient stratification and therapy selection in acute myeloid leukemia. Nature Communications, 2021, 12, 2474.	5.8	49
25	Human placental cytotrophoblast epigenome dynamics over gestation and alterations in placental disease. Developmental Cell, 2021, 56, 1238-1252.e5.	3.1	29
26	A Scalable Strand-Specific Protocol Enabling Full-Length Total RNA Sequencing From Single Cells. Frontiers in Genetics, 2021, 12, 665888.	1.1	2
27	Clinical and cost outcomes following genomicsâ€informed treatment for advanced cancers. Cancer Medicine, 2021, 10, 5131-5140.	1.3	8
28	Clonal fitness inferred from time-series modelling of single-cell cancer genomes. Nature, 2021, 595, 585-590.	13.7	71
29	Tumor infiltrating neutrophils and gland formation predict overall survival and molecular subgroups in pancreatic ductal adenocarcinoma. Cancer Medicine, 2021, 10, 1155-1165.	1.3	9
30	Proteotranscriptomic classification and characterization of pancreatic neuroendocrine neoplasms. Cell Reports, 2021, 37, 109817.	2.9	14
31	Rearrangement-mediated cis-regulatory alterations in advanced patient tumors reveal interactions with therapy. Cell Reports, 2021, 37, 110023.	2.9	8
32	Clinical response to nivolumab in an INI1-deficient pediatric chordoma correlates with immunogenic recognition of brachyury. Npj Precision Oncology, 2021, 5, 103.	2.3	18
33	Altered Gene Expression along the Glycolysis–Cholesterol Synthesis Axis Is Associated with Outcome in Pancreatic Cancer. Clinical Cancer Research, 2020, 26, 135-146.	3.2	121
34	Establishing a Framework for the Clinical Translation of Germline Findings in Precision Oncology. JNCI Cancer Spectrum, 2020, 4, pkaa045.	1.4	6
35	Glioma-derived IL-33 orchestrates an inflammatory brain tumor microenvironment that accelerates glioma progression. Nature Communications, 2020, 11, 4997.	5.8	109
36	Epigenomic programming in early fetal brain development. Epigenomics, 2020, 12, 1053-1070.	1.0	9

3

#	Article	IF	Citations
37	Analysis of Ugandan cervical carcinomas identifies human papillomavirus clade–specific epigenome and transcriptome landscapes. Nature Genetics, 2020, 52, 800-810.	9.4	40
38	Single-cell analysis of ROR \hat{l} ± tracer mouse lung reveals ILC progenitors and effector ILC2 subsets. Journal of Experimental Medicine, 2020, 217, .	4.2	74
39	TRIM25 promotes Capicua degradation independently of ERK in the absence of ATXN1L. BMC Biology, 2020, 18, 154.	1.7	7
40	Evaluating genomic biomarkers associated with resistance or sensitivity to chemotherapy in patients with advanced breast and colorectal cancer. Journal of Oncology Pharmacy Practice, 2020, 27, 107815522095184.	0.5	2
41	Validation of the RHL30 digital gene expression assay as a prognostic biomarker for relapsed Hodgkin lymphoma. British Journal of Haematology, 2020, 190, 864-868.	1.2	5
42	Endogenous Retrovirus Transcript Levels Are Associated with Immunogenic Signatures in Multiple Metastatic Cancer Types. Molecular Cancer Therapeutics, 2020, 19, 1889-1897.	1.9	10
43	Genetic and evolutionary patterns of treatment resistance in relapsed B-cell lymphoma. Blood Advances, 2020, 4, 2886-2898.	2.5	59
44	Improved structural variant interpretation for hereditary cancer susceptibility using long-read sequencing. Genetics in Medicine, 2020, 22, 1892-1897.	1.1	42
45	Integration of Whole-Genome Sequencing With Circulating Tumor DNA Analysis Captures Clonal Evolution and Tumor Heterogeneity in Non-V600 BRAF Mutant Colorectal Cancer. Clinical Colorectal Cancer, 2020, 19, 132-136.e3.	1.0	1
46	TMEM30A loss-of-function mutations drive lymphomagenesis and confer therapeutically exploitable vulnerability in B-cell lymphoma. Nature Medicine, 2020, 26, 577-588.	15.2	46
47	Transcription phenotypes of pancreatic cancer are driven by genomic events during tumor evolution. Nature Genetics, 2020, 52, 231-240.	9.4	365
48	Fluorouracil sensitivity in a head and neck squamous cell carcinoma with a somatic DPYD structural variant. Journal of Physical Education and Sports Management, 2020, 6, a004713.	0.5	5
49	Patient selection for a developmental therapeutics program using whole genome and Transcriptome analysis. Investigational New Drugs, 2020, 38, 1601-1604.	1.2	0
50	Pan-cancer analysis of whole genomes. Nature, 2020, 578, 82-93.	13.7	1,966
51	Pan-cancer analysis of advanced patient tumors reveals interactions between therapy and genomic landscapes. Nature Cancer, 2020, 1, 452-468.	5 . 7	103
52	Coding and noncoding drivers of mantle cell lymphoma identified through exome and genome sequencing. Blood, 2020, 136, 572-584.	0.6	44
53	Integrative Analysis of Single-Cell RNA-Seq and ATAC-Seq Data across Treatment Time Points in Pediatric AML. Blood, 2020, 136, 29-29.	0.6	1
54	Abstract PR-009: Proteotranscriptomic classification and characterization of pancreatic neuroendocrine neoplasms. , 2020, , .		0

#	Article	IF	CITATIONS
55	Transcriptomic analysis of CIC and ATXN1L reveal a functional relationship exploited by cancer. Oncogene, 2019, 38, 273-290.	2.6	32
56	The pivotal role of sampling recurrent tumors in the precision care of patients with tumors of the central nervous system. Journal of Physical Education and Sports Management, 2019, 5, a004143.	0.5	4
57	The Genome of the Steller Sea Lion (Eumetopias jubatus). Genes, 2019, 10, 486.	1.0	4
58	Integrative genomic analysis identifies key pathogenic mechanisms in primary mediastinal large B-cell lymphoma. Blood, 2019, 134, 802-813.	0.6	96
59	High-resolution structural genomics reveals new therapeutic vulnerabilities in glioblastoma. Genome Research, 2019, 29, 1211-1222.	2.4	52
60	Therapeutic Implication of Genomic Landscape of Adult Metastatic Sarcoma. JCO Precision Oncology, 2019, 3, 1-25.	1.5	12
61	Comparative Tumor RNA Sequencing Analysis for Difficult-to-Treat Pediatric and Young Adult Patients With Cancer. JAMA Network Open, 2019, 2, e1913968.	2.8	38
62	Evaluation of protocols for rRNA depletion-based RNA sequencing of nanogram inputs of mammalian total RNA. PLoS ONE, 2019, 14, e0224578.	1.1	12
63	Identification and Analyses of Extra-Cranial and Cranial Rhabdoid Tumor Molecular Subgroups Reveal Tumors with Cytotoxic T Cell Infiltration. Cell Reports, 2019, 29, 2338-2354.e7.	2.9	74
64	Clonal Decomposition and DNA Replication States Defined by Scaled Single-Cell Genome Sequencing. Cell, 2019, 179, 1207-1221.e22.	13.5	162
65	Comprehensive genomic profiling of glioblastoma tumors, BTICs, and xenografts reveals stability and adaptation to growth environments. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 19098-19108.	3.3	42
66	Molecular and Genetic Characterization of MHC Deficiency Identifies EZH2 as Therapeutic Target for Enhancing Immune Recognition. Cancer Discovery, 2019, 9, 546-563.	7.7	213
67	A distinct neurodevelopmental syndrome with intellectual disability, autism spectrum disorder, characteristic facies, and macrocephaly is caused by defects in CHD8. Journal of Human Genetics, 2019, 64, 271-280.	1.1	35
68	Integrative genomic analysis of matched primary and metastatic pediatric osteosarcoma. Journal of Pathology, 2019, 249, 319-331.	2.1	36
69	Genomic characterization of a well-differentiated grade 3 pancreatic neuroendocrine tumor. Journal of Physical Education and Sports Management, 2019, 5, a003814.	0.5	17
70	<i>NRG1</i> Gene Fusions Are Recurrent, Clinically Actionable Gene Rearrangements in <i>KRAS</i> Wild-Type Pancreatic Ductal Adenocarcinoma. Clinical Cancer Research, 2019, 25, 4674-4681.	3.2	121
71	Capicua regulates neural stem cell proliferation and lineage specification through control of Ets factors. Nature Communications, 2019, 10, 2000.	5.8	34
72	Childhood cerebellar tumours mirror conserved fetal transcriptional programs. Nature, 2019, 572, 67-73.	13.7	293

#	Article	lF	Citations
73	Application of a Neural Network Whole Transcriptome–Based Pan-Cancer Method for Diagnosis of Primary and Metastatic Cancers. JAMA Network Open, 2019, 2, e192597.	2.8	67
74	Intratumoral Genetic and Functional Heterogeneity in Pediatric Glioblastoma. Cancer Research, 2019, 79, 2111-2123.	0.4	28
75	Base excision repair deficiency signatures implicate germline and somatic <i>MUTYH</i> aberrations in pancreatic ductal adenocarcinoma and breast cancer oncogenesis. Journal of Physical Education and Sports Management, 2019, 5, a003681.	0.5	33
76	Double-Hit Gene Expression Signature Defines a Distinct Subgroup of Germinal Center B-Cell-Like Diffuse Large B-Cell Lymphoma. Journal of Clinical Oncology, 2019, 37, 190-201.	0.8	257
77	A high-throughput protocol for isolating cell-free circulating tumor DNA from peripheral blood. BioTechniques, 2019, 66, 85-92.	0.8	13
78	Recurrent noncoding U1ÂsnRNA mutations drive cryptic splicing in SHH medulloblastoma. Nature, 2019, 574, 707-711.	13.7	129
79	Sources of erroneous sequences and artifact chimeric reads in next generation sequencing of genomic DNA from formalin-fixed paraffin-embedded samples. Nucleic Acids Research, 2019, 47, e12-e12.	6.5	50
80	Genome-wide discovery of somatic coding and noncoding mutations in pediatric endemic and sporadic Burkitt lymphoma. Blood, 2019, 133, 1313-1324.	0.6	172
81	Clinical outcomes after whole-genome sequencing in patients with metastatic non-small-cell lung cancer. Journal of Physical Education and Sports Management, 2019, 5, a002659.	0.5	3
82	Abstract B56: Endogenous retrovirus transcript levels are associated with immunogenic signatures in multiple metastatic cancer types. , $2019, \dots$		0
83	Abstract 3480: <i>TMEM30A</i> loss-of-function mutations drive lymphomagenesis and confer therapeutically exploitable vulnerability in B-cell lymphoma., 2019,,.		0
84	A Hematogenous Route for Medulloblastoma Leptomeningeal Metastases. Cell, 2018, 172, 1050-1062.e14.	13.5	85
85	Molecular characterization of <i>ERBB2</i> -amplified colorectal cancer identifies potential mechanisms of resistance to targeted therapies: a report of two instructive cases. Journal of Physical Education and Sports Management, 2018, 4, a002535.	0.5	16
86	Assessment of Capture and Amplicon-Based Approaches for the Development of a Targeted Next-Generation Sequencing Pipeline to Personalize Lymphoma Management. Journal of Molecular Diagnostics, 2018, 20, 203-214.	1.2	58
87	Application of genomics to identify therapeutic targets in recurrent pediatric papillary thyroid carcinoma. Journal of Physical Education and Sports Management, 2018, 4, a002568.	0.5	14
88	Personalized oncogenomic analysis of metastatic adenoid cystic carcinoma: using whole-genome sequencing to inform clinical decision-making. Journal of Physical Education and Sports Management, 2018, 4, a002626.	0.5	18
89	An Integrated TCGA Pan-Cancer Clinical Data Resource to Drive High-Quality Survival Outcome Analytics. Cell, 2018, 173, 400-416.e11.	13.5	2,277
90	Comprehensive Characterization of Cancer Driver Genes and Mutations. Cell, 2018, 173, 371-385.e18.	13.5	1,670

#	Article	IF	Citations
91	Cell-of-Origin Patterns Dominate the Molecular Classification of 10,000 Tumors from 33 Types of Cancer. Cell, 2018, 173, 291-304.e6.	13.5	1,718
92	A Pan-Cancer Analysis of Enhancer Expression in Nearly 9000 Patient Samples. Cell, 2018, 173, 386-399.e12.	13.5	228
93	Perspective on Oncogenic Processes at the End of the Beginning of Cancer Genomics. Cell, 2018, 173, 305-320.e10.	13.5	272
94	Machine Learning Identifies Stemness Features Associated with Oncogenic Dedifferentiation. Cell, 2018, 173, 338-354.e15.	13.5	1,417
95	Oncogenic Signaling Pathways in The Cancer Genome Atlas. Cell, 2018, 173, 321-337.e10.	13.5	2,111
96	Pathogenic Germline Variants in 10,389 Adult Cancers. Cell, 2018, 173, 355-370.e14.	13.5	620
97	Somatic Mutational Landscape of Splicing Factor Genes and Their Functional Consequences across 33 Cancer Types. Cell Reports, 2018, 23, 282-296.e4.	2.9	333
98	Driver Fusions and Their Implications in the Development and Treatment of Human Cancers. Cell Reports, 2018, 23, 227-238.e3.	2.9	407
99	Genomic, Pathway Network, and Immunologic Features Distinguishing Squamous Carcinomas. Cell Reports, 2018, 23, 194-212.e6.	2.9	245
100	Pan-Cancer Analysis of IncRNA Regulation Supports Their Targeting of Cancer Genes in Each Tumor Context. Cell Reports, 2018, 23, 297-312.e12.	2.9	205
101	The Cancer Genome Atlas Comprehensive Molecular Characterization of Renal Cell Carcinoma. Cell Reports, 2018, 23, 313-326.e5.	2.9	523
102	Spatial Organization and Molecular Correlation of Tumor-Infiltrating Lymphocytes Using Deep Learning on Pathology Images. Cell Reports, 2018, 23, 181-193.e7.	2.9	683
103	The Immune Landscape of Cancer. Immunity, 2018, 48, 812-830.e14.	6.6	3,706
104	Machine Learning Detects Pan-cancer Ras Pathway Activation in The Cancer Genome Atlas. Cell Reports, 2018, 23, 172-180.e3.	2.9	119
105	Integrated Genomic Analysis of the Ubiquitin Pathway across Cancer Types. Cell Reports, 2018, 23, 213-226.e3.	2.9	83
106	Genomic and Molecular Landscape of DNA Damage Repair Deficiency across The Cancer Genome Atlas. Cell Reports, 2018, 23, 239-254.e6.	2.9	801
107	Molecular Characterization and Clinical Relevance of Metabolic Expression Subtypes in Human Cancers. Cell Reports, 2018, 23, 255-269.e4.	2.9	204
108	Systematic Analysis of Splice-Site-Creating Mutations in Cancer. Cell Reports, 2018, 23, 270-281.e3.	2.9	177

#	Article	IF	CITATIONS
109	The Integrated Genomic Landscape of Thymic Epithelial Tumors. Cancer Cell, 2018, 33, 244-258.e10.	7.7	270
110	Scalable Open Science Approach for Mutation Calling of Tumor Exomes Using Multiple Genomic Pipelines. Cell Systems, 2018, 6, 271-281.e7.	2.9	605
111	Pan-cancer Alterations of the MYC Oncogene and Its Proximal Network across the Cancer Genome Atlas. Cell Systems, 2018, 6, 282-300.e2.	2.9	284
112	IncRNA Epigenetic Landscape Analysis Identifies EPIC1 as an Oncogenic IncRNA that Interacts with MYC and Promotes Cell-Cycle Progression in Cancer. Cancer Cell, 2018, 33, 706-720.e9.	7.7	400
113	Genomic and Functional Approaches to Understanding Cancer Aneuploidy. Cancer Cell, 2018, 33, 676-689.e3.	7.7	750
114	Comparative Molecular Analysis of Gastrointestinal Adenocarcinomas. Cancer Cell, 2018, 33, 721-735.e8.	7.7	396
115	A Comprehensive Pan-Cancer Molecular Study of Gynecologic and Breast Cancers. Cancer Cell, 2018, 33, 690-705.e9.	7.7	478
116	Whole genome and whole transcriptome genomic profiling of a metastatic eccrine porocarcinoma. Npj Precision Oncology, 2018, 2, 8.	2.3	15
117	Opposing Effects of CREBBP Mutations Govern the Phenotype of Rubinstein-Taybi Syndrome and Adult SHH Medulloblastoma. Developmental Cell, 2018, 44, 709-724.e6.	3.1	35
118	The molecular landscape of pediatric acute myeloid leukemia reveals recurrent structural alterations and age-specific mutational interactions. Nature Medicine, 2018, 24, 103-112.	15.2	525
119	Molecular characterization of metastatic pancreatic neuroendocrine tumors (PNETs) using whole-genome and transcriptome sequencing. Journal of Physical Education and Sports Management, 2018, 4, a002329.	0.5	30
120	High-resolution architecture and partner genes of MYC rearrangements in lymphoma with DLBCL morphology. Blood Advances, 2018, 2, 2755-2765.	2.5	74
121	Temporal Dynamics of Genomic Alterations in a BRCA1 Germline–Mutated Pancreatic Cancer With Low Genomic Instability Burden but Exceptional Response to Fluorouracil, Oxaliplatin, Leucovorin, and Irinotecan. JCO Precision Oncology, 2018, 2, 1-8.	1.5	1
122	Comparative RNA-Sequencing Analysis Benefits a Pediatric Patient With Relapsed Cancer. JCO Precision Oncology, 2018, 2, 1-16.	1.5	12
123	Whole-genome and transcriptome profiling of a metastatic thyroid-like follicular renal cell carcinoma. Journal of Physical Education and Sports Management, 2018, 4, a003137.	0.5	15
124	The Genome of the North American Brown Bear or Grizzly: Ursus arctos ssp. horribilis. Genes, 2018, 9, 598.	1.0	34
125	Genome-wide discovery of somatic regulatory variants in diffuse large B-cell lymphoma. Nature Communications, 2018, 9, 4001.	5.8	102
126	A Pan-Cancer Analysis Reveals High-Frequency Genetic Alterations in Mediators of Signaling by the TGF-Î ² Superfamily. Cell Systems, 2018, 7, 422-437.e7.	2.9	134

#	Article	IF	CITATIONS
127	Integrative Molecular Characterization of Malignant Pleural Mesothelioma. Cancer Discovery, 2018, 8, 1548-1565.	7.7	422
128	Comprehensive Molecular Characterization of the Hippo Signaling Pathway in Cancer. Cell Reports, 2018, 25, 1304-1317.e5.	2.9	329
129	The genetic basis and cell of origin of mixed phenotype acute leukaemia. Nature, 2018, 562, 373-379.	13.7	236
130	ABT-888 restores sensitivity in temozolomide resistant glioma cells and xenografts. PLoS ONE, 2018, 13, e0202860.	1.1	28
131	Comprehensive Analysis of Alternative Splicing Across Tumors from 8,705 Patients. Cancer Cell, 2018, 34, 211-224.e6.	7.7	623
132	Spectrum and prevalence of genetic predisposition in medulloblastoma: a retrospective genetic study and prospective validation in a clinical trial cohort. Lancet Oncology, The, 2018, 19, 785-798.	5.1	268
133	Integrated Molecular Characterization of Testicular Germ Cell Tumors. Cell Reports, 2018, 23, 3392-3406.	2.9	324
134	Integrated genomic and molecular characterization of cervical cancer. Nature, 2017, 543, 378-384.	13.7	1,158
135	Comprehensive Molecular Characterization of Pheochromocytoma and Paraganglioma. Cancer Cell, 2017, 31, 181-193.	7.7	532
136	Spatial heterogeneity in medulloblastoma. Nature Genetics, 2017, 49, 780-788.	9.4	112
136	Spatial heterogeneity in medulloblastoma. Nature Genetics, 2017, 49, 780-788. Genomic consequences of aberrant DNA repair mechanisms stratify ovarian cancer histotypes. Nature Genetics, 2017, 49, 856-865.	9.4	112 220
	Genomic consequences of aberrant DNA repair mechanisms stratify ovarian cancer histotypes. Nature		
137	Genomic consequences of aberrant DNA repair mechanisms stratify ovarian cancer histotypes. Nature Genetics, 2017, 49, 856-865. Comprehensive and Integrative Genomic Characterization of Hepatocellular Carcinoma. Cell, 2017, 169,	9.4	220
137	Genomic consequences of aberrant DNA repair mechanisms stratify ovarian cancer histotypes. Nature Genetics, 2017, 49, 856-865. Comprehensive and Integrative Genomic Characterization of Hepatocellular Carcinoma. Cell, 2017, 169, 1327-1341.e23. Genomic profiling of pelvic genital type leiomyosarcoma in a woman with a germline <i>CHEK2</i> i>c.1100delC mutation and a concomitant diagnosis of metastatic invasive ductal breast	9.4	220 1,794
137 138 139	Genomic consequences of aberrant DNA repair mechanisms stratify ovarian cancer histotypes. Nature Genetics, 2017, 49, 856-865. Comprehensive and Integrative Genomic Characterization of Hepatocellular Carcinoma. Cell, 2017, 169, 1327-1341.e23. Genomic profiling of pelvic genital type leiomyosarcoma in a woman with a germline <i>CHEK2</i> Sc.1100delC mutation and a concomitant diagnosis of metastatic invasive ductal breast carcinoma. Journal of Physical Education and Sports Management, 2017, 3, a001628. Pyruvate Kinase Inhibits Proliferation during Postnatal Cerebellar Neurogenesis and Suppresses	9.4 13.5 0.5	220 1,794 8
137 138 139	Genomic consequences of aberrant DNA repair mechanisms stratify ovarian cancer histotypes. Nature Genetics, 2017, 49, 856-865. Comprehensive and Integrative Genomic Characterization of Hepatocellular Carcinoma. Cell, 2017, 169, 1327-1341.e23. Genomic profiling of pelvic genital type leiomyosarcoma in a woman with a germline <i>CHEK2 </i> CHEK2 Comprehensive and a concomitant diagnosis of metastatic invasive ductal breast carcinoma. Journal of Physical Education and Sports Management, 2017, 3, a001628. Pyruvate Kinase Inhibits Proliferation during Postnatal Cerebellar Neurogenesis and Suppresses Medulloblastoma Formation. Cancer Research, 2017, 77, 3217-3230. Whole-genome analysis reveals unexpected dynamics of mutant subclone development in a patient	9.4 13.5 0.5	220 1,794 8 45
137 138 139 140	Genomic consequences of aberrant DNA repair mechanisms stratify ovarian cancer histotypes. Nature Genetics, 2017, 49, 856-865. Comprehensive and Integrative Genomic Characterization of Hepatocellular Carcinoma. Cell, 2017, 169, 1327-1341.e23. Genomic profiling of pelvic genital type leiomyosarcoma in a woman with a germline ⟨i>CHEK2 ⟨i>c.1100delC mutation and a concomitant diagnosis of metastatic invasive ductal breast carcinoma. Journal of Physical Education and Sports Management, 2017, 3, a001628. Pyruvate Kinase Inhibits Proliferation during Postnatal Cerebellar Neurogenesis and Suppresses Medulloblastoma Formation. Cancer Research, 2017, 77, 3217-3230. Whole-genome analysis reveals unexpected dynamics of mutant subclone development in a patient with JAK2-V617F-positive chronic myeloid leukemia. Experimental Hematology, 2017, 53, 48-58. Genetic profiling of MYC and BCL2 in diffuse large B-cell lymphoma determines cell-of-origin–specific	9.4 13.5 0.5 0.4	220 1,794 8 45

#	Article	IF	Citations
145	Integrative Genomic Analysis of Cholangiocarcinoma Identifies Distinct IDH-Mutant Molecular Profiles. Cell Reports, 2017, 18, 2780-2794.	2.9	416
146	The cost and cost trajectory of wholeâ€genome analysis guiding treatment of patients with advanced cancers. Molecular Genetics & Denomic Medicine, 2017, 5, 251-260.	0.6	40
147	Integrated genomic characterization of oesophageal carcinoma. Nature, 2017, 541, 169-175.	13.7	1,448
148	Comprehensive Molecular Characterization of Muscle-Invasive Bladder Cancer. Cell, 2017, 171, 540-556.e25.	13.5	1,742
149	A Children's Oncology Group and TARGET initiative exploring the genetic landscape of Wilms tumor. Nature Genetics, 2017, 49, 1487-1494.	9.4	255
150	Clonal expansion and epigenetic reprogramming following deletion or amplification of mutant $\langle i \rangle$ IDH1 $\langle i \rangle$. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 10743-10748.	3.3	109
151	Identification of GPC2 as an Oncoprotein and Candidate Immunotherapeutic Target in High-Risk Neuroblastoma. Cancer Cell, 2017, 32, 295-309.e12.	7.7	148
152	Detection and genomic characterization of a mammary-like adenocarcinoma. Journal of Physical Education and Sports Management, 2017, 3, a002170.	0.5	13
153	The whole-genome landscape of medulloblastoma subtypes. Nature, 2017, 547, 311-317.	13.7	787
154	Integrative Analysis Identifies Four Molecular and Clinical Subsets in Uveal Melanoma. Cancer Cell, 2017, 32, 204-220.e15.	7.7	642
155	Characterization of the human thyroid epigenome. Journal of Endocrinology, 2017, 235, 153-165.	1.2	8
156	Integrated Genomic Characterization of Pancreatic Ductal Adenocarcinoma. Cancer Cell, 2017, 32, 185-203.e13.	7.7	1,428
157	Homologous Recombination Deficiency and Platinum-Based Therapy Outcomes in Advanced Breast Cancer. Clinical Cancer Research, 2017, 23, 7521-7530.	3.2	144
158	Comprehensive and Integrated Genomic Characterization of Adult Soft Tissue Sarcomas. Cell, 2017, 171, 950-965.e28.	13.5	738
159	Comprehensive whole genome sequence analyses yields novel genetic and structural insights for Intellectual Disability. BMC Genomics, 2017, 18, 403.	1.2	15
160	Successful targeting of the NRG1 pathway indicates novel treatment strategy for metastatic cancer. Annals of Oncology, 2017, 28, 3092-3097.	0.6	83
161	Increasing quality, throughput and speed of sample preparation for strand-specific messenger RNA sequencing. BMC Genomics, 2017, 18, 515.	1.2	8
162	The Genome of the Beluga Whale (Delphinapterus leucas). Genes, 2017, 8, 378.	1.0	39

#	Article	IF	CITATIONS
163	The Genome of the Northern Sea Otter (Enhydra lutris kenyoni). Genes, 2017, 8, 379.	1.0	24
164	Carcinoma Ex Pleomorphic Adenoma: Case Report and Options for Systemic Therapy. Current Oncology, 2017, 24, 251-254.	0.9	13
165	MicroRNA Expression-Based Model Indicates Event-Free Survival in Pediatric Acute Myeloid Leukemia. Journal of Clinical Oncology, 2017, 35, 3964-3977.	0.8	49
166	Automated high throughput nucleic acid purification from formalin-fixed paraffin-embedded tissue samples for next generation sequence analysis. PLoS ONE, 2017, 12, e0178706.	1.1	18
167	miR-509-3p is clinically significant and strongly attenuates cellular migration and multi-cellular spheroids in ovarian cancer. Oncotarget, 2016, 7, 25930-25948.	0.8	49
168	Small molecule epigenetic screen identifies novel EZH2 and HDAC inhibitors that target glioblastoma brain tumor-initiating cells. Oncotarget, 2016, 7, 59360-59376.	0.8	34
169	Immunohistochemistry for NF2, LATS1/2, and YAP/TAZ Fails to Separate Benign From Malignant Mesothelial Proliferations. Archives of Pathology and Laboratory Medicine, 2016, 140, 391-391.	1.2	16
170	Tumour-suppressor microRNAs regulate ovarian cancer cell physical properties and invasive behaviour. Open Biology, 2016, 6, 160275.	1.5	29
171	Tracking of Normal and Malignant Progenitor Cell Cycle Transit in a Defined Niche. Scientific Reports, 2016, 6, 23885.	1.6	7
172	Response to angiotensin blockade with irbesartan in a patient with metastatic colorectal cancer. Annals of Oncology, 2016, 27, 801-806.	0.6	39
173	Divergent modes of clonal spread and intraperitoneal mixing in high-grade serous ovarian cancer. Nature Genetics, 2016, 48, 758-767.	9.4	287
174	Molecular etiology of an indolent lymphoproliferative disorder determined by whole-genome sequencing. Journal of Physical Education and Sports Management, 2016, 2, a000679.	0.5	3
175	Comprehensive Pan-Genomic Characterization of Adrenocortical Carcinoma. Cancer Cell, 2016, 29, 723-736.	7.7	482
176	Distinct patterns of somatic genome alterations in lung adenocarcinomas and squamous cell carcinomas. Nature Genetics, 2016, 48, 607-616.	9.4	933
177	Clinical impact of molecular features in diffuse large B-cell lymphoma and follicular lymphoma. Blood, 2016, 127, 181-186.	0.6	21
178	Comprehensive characterization of programmed death ligand structural rearrangements in B-cell non-Hodgkin lymphomas. Blood, 2016, 128, 1206-1213.	0.6	47
179	CSF3R mutations have a high degree of overlap with CEBPA mutations in pediatric AML. Blood, 2016, 127, 3094-3098.	0.6	49
180	Investigation of PD-L1 Biomarker Testing Methods for PD-1 Axis Inhibition in Non-squamous Non–small Cell Lung Cancer. Journal of Histochemistry and Cytochemistry, 2016, 64, 587-600.	1.3	30

#	Article	IF	Citations
181	Significance of <i>TP53</i> Mutation in Wilms Tumors with Diffuse Anaplasia: A Report from the Children's Oncology Group. Clinical Cancer Research, 2016, 22, 5582-5591.	3.2	82
182	A somatic reference standard for cancer genome sequencing. Scientific Reports, 2016, 6, 24607.	1.6	64
183	The International Human Epigenome Consortium: A Blueprint for Scientific Collaboration and Discovery. Cell, 2016, 167, 1145-1149.	13.5	404
184	Analysis of Normal Human Mammary Epigenomes Reveals Cell-Specific Active Enhancer States and Associated Transcription Factor Networks. Cell Reports, 2016, 17, 2060-2074.	2.9	90
185	Genomic analyses identify recurrent MEF2D fusions in acute lymphoblastic leukaemia. Nature Communications, 2016, 7, 13331.	5.8	218
186	Intragenic CNVs for epigenetic regulatory genes in intellectual disability: Survey identifies pathogenic and benign single exon changes. American Journal of Medical Genetics, Part A, 2016, 170, 2916-2926.	0.7	14
187	ETV6-NTRK3 Is Expressed in a Subset of ALK-Negative Inflammatory Myofibroblastic Tumors. American Journal of Surgical Pathology, 2016, 40, 1051-1061.	2.1	139
188	ADAR1 Activation Drives Leukemia Stem Cell Self-Renewal by Impairing Let-7 Biogenesis. Cell Stem Cell, 2016, 19, 177-191.	5.2	182
189	DNA methylation in adult diffuse gliomas. Briefings in Functional Genomics, 2016, 15, elw019.	1.3	11
190	Divergent clonal selection dominates medulloblastoma at recurrence. Nature, 2016, 529, 351-357.	13.7	266
191	Large-scale profiling of microRNAs for The Cancer Genome Atlas. Nucleic Acids Research, 2016, 44, e3-e3.	6.5	125
192	Toward Personalized Lymphoma Immunotherapy: Identification of Common Driver Mutations Recognized by Patient CD8+ T Cells. Clinical Cancer Research, 2016, 22, 2226-2236.	3.2	26
193	Genome-Wide Profiles of Extra-cranial Malignant Rhabdoid Tumors Reveal Heterogeneity and Dysregulated Developmental Pathways. Cancer Cell, 2016, 29, 394-406.	7.7	105
194	PutativeBRAFactivating fusion in a medullary thyroid cancer. Journal of Physical Education and Sports Management, 2016, 2, a000729.	0.5	14
195	Molecular Profiling Reveals Biologically Discrete Subsets and Pathways of Progression in Diffuse Glioma. Cell, 2016, 164, 550-563.	13.5	1,695
196	Comprehensive Molecular Characterization of Papillary Renal-Cell Carcinoma. New England Journal of Medicine, 2016, 374, 135-145.	13.9	1,040
197	Burkitt Lymphoma Genome Sequencing Project (BLGSP): Introduction. Blood, 2016, 128, 1760-1760.	0.6	1
198	Sequencing Strategies to Guide Decision Making in Cancer Treatment. PLoS Medicine, 2016, 13, e1002189.	3.9	4

#	Article	IF	Citations
199	Histological Transformation and Progression in Follicular Lymphoma: A Clonal Evolution Study. PLoS Medicine, 2016, 13, e1002197.	3.9	185
200	MEF2 transcription factors: developmental regulators and emerging cancer genes. Oncotarget, 2016, 7, 2297-2312.	0.8	132
201	Medulloblastoma-associated DDX3 variant selectively alters the translational response to stress. Oncotarget, 2016, 7, 28169-28182.	0.8	62
202	The genomic and transcriptomic landscape of anaplastic thyroid cancer: implications for therapy. BMC Cancer, 2015, 15, 984.	1.1	55
203	An RCOR1 loss–associated gene expression signature identifies a prognostically significant DLBCL subgroup. Blood, 2015, 125, 959-966.	0.6	24
204	Cell of origin of transformed follicular lymphoma. Blood, 2015, 126, 2118-2127.	0.6	91
205	GLI2 inhibition abrogates human leukemia stem cell dormancy. Journal of Translational Medicine, 2015, 13, 98.	1.8	80
206	Recurrent genomic rearrangements in primary testicular lymphoma. Journal of Pathology, 2015, 236, 136-141.	2.1	47
207	Multifocal endometriotic lesions associated with cancer are clonal and carry a high mutation burden. Journal of Pathology, 2015, 236, 201-209.	2.1	131
208	The <i>Drosophila</i> TIPE family member Sigmar interacts with the Ste20-like kinase Misshapen and modulates JNK signaling, cytoskeletal remodeling and autophagy. Biology Open, 2015, 4, 672-684.	0.6	10
209	MLLT1 YEATS domain mutations in clinically distinctive Favourable Histology Wilms tumours. Nature Communications, 2015, 6, 10013.	5.8	64
210	Integrative genomic analysis of ghost cell odontogenic carcinoma. Oral Oncology, 2015, 51, e71-e75.	0.8	20
211	Lessons learned from the application of whole-genome analysis to the treatment of patients with advanced cancers. Journal of Physical Education and Sports Management, 2015, 1, a000570.	0.5	92
212	Recurrent DGCR8, DROSHA, and SIX Homeodomain Mutations in Favorable Histology Wilms Tumors. Cancer Cell, 2015, 27, 286-297.	7.7	244
213	The Release 6 reference sequence of the <i>Drosophila melanogaster</i> genome. Genome Research, 2015, 25, 445-458.	2.4	359
214	Loss of the Notch effector RBPJ promotes tumorigenesis. Journal of Experimental Medicine, 2015, 212, 37-52.	4.2	52
215	Intermediate DNA methylation is a conserved signature of genome regulation. Nature Communications, 2015, 6, 6363.	5.8	91
216	Integrative analysis of 111 reference human epigenomes. Nature, 2015, 518, 317-330.	13.7	5,653

#	Article	IF	Citations
217	MEN1 Mutations in Hürthle Cell (Oncocytic) Thyroid Carcinoma. Journal of Clinical Endocrinology and Metabolism, 2015, 100, E611-E615.	1.8	20
218	Comprehensive genomic characterization of head and neck squamous cell carcinomas. Nature, 2015, 517, 576-582.	13.7	3,209
219	Epigenetic and transcriptional determinants of the human breast. Nature Communications, 2015, 6, 6351.	5.8	56
220	Comprehensive miRNA sequence analysis reveals survival differences in diffuse large B-cell lymphoma patients. Genome Biology, 2015, 16, 18.	3.8	107
221	Genomic Classification of Cutaneous Melanoma. Cell, 2015, 161, 1681-1696.	13.5	2,562
222	Cross-cancer profiling of molecular alterations within the human autophagy interaction network. Autophagy, 2015, 11, 1668-1687.	4.3	107
223	Precursor States of Brain Tumor Initiating Cell Lines Are Predictive of SurvivalÂin Xenografts and Associated with Glioblastoma Subtypes. Stem Cell Reports, 2015, 5, 1-9.	2.3	72
224	Comprehensive, Integrative Genomic Analysis of Diffuse Lower-Grade Gliomas. New England Journal of Medicine, 2015, 372, 2481-2498.	13.9	2,582
225	MEF2B mutations in non-Hodgkin lymphoma dysregulate cell migration by decreasing MEF2B target gene activation. Nature Communications, 2015, 6, 7953.	5.8	50
226	SOX9 modulates the expression of key transcription factors required for heart valve development. Development (Cambridge), 2015, 142, 4340-50.	1.2	49
227	The Molecular Taxonomy of Primary Prostate Cancer. Cell, 2015, 163, 1011-1025.	13.5	2,435
228	Combined serial analysis of gene expression and transcription factor binding site prediction identifies novel-candidate-target genes of Nr2e1 in neocortex development. BMC Genomics, 2015, 16, 545.	1.2	9
229	Comprehensive Molecular Portraits of Invasive Lobular Breast Cancer. Cell, 2015, 163, 506-519.	13.5	1,485
230	Reversion to an embryonic alternative splicing program enhances leukemia stem cell self-renewal. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 15444-15449.	3.3	36
231	Dynamics of genomic clones in breast cancer patient xenografts at single-cell resolution. Nature, 2015, 518, 422-426.	13.7	545
232	Comprehensive Sequence Analysis of Relapse and Refractory Pediatric Acute Myeloid Leukemia Identifies miRNA and mRNA Transcripts Associated with Treatment Resistance - a Report from the COG/NCI-Target AML Initiative. Blood, 2015, 126, 687-687.	0.6	2
233	Discovery and Functional Validation of Novel Pediatric Specific FLT3 Activating Mutations in Acute Myeloid Leukemia: Results from the COG/NCI Target Initiative. Blood, 2015, 126, 87-87.	0.6	19
234	Personalized Oncogenomics: Clinical Experience with Malignant Peritoneal Mesothelioma Using Whole Genome Sequencing. PLoS ONE, 2015, 10, e0119689.	1.1	36

#	Article	IF	CITATIONS
235	Next-Generation Sequencing Approaches in Cancer: Where Have They Brought Us and Where Will They Take Us?. Cancers, 2015, 7, 1925-1958.	1.7	51
236	The expression level of small nonâ€coding <scp>RNA</scp> s derived from the first exon of proteinâ€coding genes is predictive of cancer status. EMBO Reports, 2014, 15, 402-410.	2.0	13
237	Analysis of the Genome and Transcriptome of Cryptococcus neoformans var. grubii Reveals Complex RNA Expression and Microevolution Leading to Virulence Attenuation. PLoS Genetics, 2014, 10, e1004261.	1.5	336
238	TITAN: inference of copy number architectures in clonal cell populations from tumor whole-genome sequence data. Genome Research, 2014, 24, 1881-1893.	2.4	322
239	Integrated Genomic Characterization of Papillary Thyroid Carcinoma. Cell, 2014, 159, 676-690.	13.5	2,318
240	Regulatory network decoded from epigenomes of surface ectoderm-derived cell types. Nature Communications, 2014, 5, 5442.	5.8	25
241	Diagnostic Value of Next-Generation Sequencing in an Unusual Sphenoid Tumor. Oncologist, 2014, 19, 623-630.	1.9	20
242	Desmosterolosis: an illustration of diagnostic ambiguity of cholesterol synthesis disorders. Orphanet Journal of Rare Diseases, 2014, 9, 94.	1.2	19
243	Hippo Signaling Influences HNF4A and FOXA2 Enhancer Switching during Hepatocyte Differentiation. Cell Reports, 2014, 9, 261-271.	2.9	89
244	Second-Generation Sequencing for Cancer Genome Analysis. , 2014, , 13-30.		2
244		13.7	2,496
	Second-Generation Sequencing for Cancer Genome Analysis. , 2014, , 13-30.	13.7	
245	Second-Generation Sequencing for Cancer Genome Analysis., 2014, , 13-30. Comprehensive molecular characterization of urothelial bladder carcinoma. Nature, 2014, 507, 315-322. Single exon-resolution targeted chromosomal microarray analysis of known and candidate		2,496
245 246	Second-Generation Sequencing for Cancer Genome Analysis., 2014, , 13-30. Comprehensive molecular characterization of urothelial bladder carcinoma. Nature, 2014, 507, 315-322. Single exon-resolution targeted chromosomal microarray analysis of known and candidate intellectual disability genes. European Journal of Human Genetics, 2014, 22, 792-800. Mutational Analysis Reveals the Origin and Therapy-Driven Evolution of Recurrent Glioma. Science,	1.4	2,496 35
245 246 247	Second-Generation Sequencing for Cancer Genome Analysis., 2014, , 13-30. Comprehensive molecular characterization of urothelial bladder carcinoma. Nature, 2014, 507, 315-322. Single exon-resolution targeted chromosomal microarray analysis of known and candidate intellectual disability genes. European Journal of Human Genetics, 2014, 22, 792-800. Mutational Analysis Reveals the Origin and Therapy-Driven Evolution of Recurrent Glioma. Science, 2014, 343, 189-193. Characterization of HPV and host genome interactions in primary head and neck cancers. Proceedings	1.4 6.0	2,496 35 1,147
245 246 247 248	Second-Generation Sequencing for Cancer Genome Analysis., 2014, , 13-30. Comprehensive molecular characterization of urothelial bladder carcinoma. Nature, 2014, 507, 315-322. Single exon-resolution targeted chromosomal microarray analysis of known and candidate intellectual disability genes. European Journal of Human Genetics, 2014, 22, 792-800. Mutational Analysis Reveals the Origin and Therapy-Driven Evolution of Recurrent Glioma. Science, 2014, 343, 189-193. Characterization of HPV and host genome interactions in primary head and neck cancers. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 15544-15549. A Notchâ€dependent transcriptional hierarchy promotes mesenchymal transdifferentiation in the	1.4 6.0 3.3	2,496 35 1,147 317
245 246 247 248	Second-Generation Sequencing for Cancer Genome Analysis., 2014, , 13-30. Comprehensive molecular characterization of urothelial bladder carcinoma. Nature, 2014, 507, 315-322. Single exon-resolution targeted chromosomal microarray analysis of known and candidate intellectual disability genes. European Journal of Human Genetics, 2014, 22, 792-800. Mutational Analysis Reveals the Origin and Therapy-Driven Evolution of Recurrent Glioma. Science, 2014, 343, 189-193. Characterization of HPV and host genome interactions in primary head and neck cancers. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 15544-15549. A Notchâ€dependent transcriptional hierarchy promotes mesenchymal transdifferentiation in the cardiac cushion. Developmental Dynamics, 2014, 243, 894-905. Recurrent somatic mutations of PTPN1 in primary mediastinal B cell lymphoma and Hodgkin lymphoma.	1.4 6.0 3.3	2,496 35 1,147 317 21

#	Article	IF	CITATIONS
253	Comprehensive molecular characterization of gastric adenocarcinoma. Nature, 2014, 513, 202-209.	13.7	5,055
254	The Molecular Landscape of Pediatric Brain Tumors in the Next-Generation Sequencing Era. Current Neurology and Neuroscience Reports, 2014, 14, 474.	2.0	11
255	The Somatic Genomic Landscape of Chromophobe Renal Cell Carcinoma. Cancer Cell, 2014, 26, 319-330.	7.7	665
256	Comprehensive molecular profiling of lung adenocarcinoma. Nature, 2014, 511, 543-550.	13.7	4,572
257	Multiplatform Analysis of 12 Cancer Types Reveals Molecular Classification within and across Tissues of Origin. Cell, 2014, 158, 929-944.	13.5	1,242
258	Quiescent Sox2+ Cells Drive Hierarchical Growth and Relapse in Sonic Hedgehog Subgroup Medulloblastoma. Cancer Cell, 2014, 26, 33-47.	7.7	241
259	A transgenic mouse model demonstrating the oncogenic role of mutations in the polycomb-group gene EZH2 in lymphomagenesis. Blood, 2014, 123, 3914-3924.	0.6	69
260	Clinical Significance of Genetic Aberrations in Diffuse Large B Cell Lymphoma. Blood, 2014, 124, 703-703.	0.6	5
261	Mutations in CIC and IDH1 cooperatively regulate 2-hydroxyglutarate levels and cell clonogenicity. Oncotarget, 2014, 5, 7960-7979.	0.8	35
262	Aberrant patterns of H3K4 and H3K27 histone lysine methylation occur across subgroups in medulloblastoma. Acta Neuropathologica, 2013, 125, 373-384.	3.9	169
263	A Clinically Validated Diagnostic Second-Generation Sequencing Assay for Detection of Hereditary BRCA1 and BRCA2 Mutations. Journal of Molecular Diagnostics, 2013, 15, 796-809.	1.2	29
264	The Somatic Genomic Landscape of Glioblastoma. Cell, 2013, 155, 462-477.	13.5	3,979
265	Estimating absolute methylation levels at single-CpG resolution from methylation enrichment and restriction enzyme sequencing methods. Genome Research, 2013, 23, 1541-1553.	2.4	138
266	Novel mRNA isoforms and mutations of uridine monophosphate synthetase and 5-fluorouracil resistance in colorectal cancer. Pharmacogenomics Journal, 2013, 13, 148-158.	0.9	29
267	The Cancer Genome Atlas Pan-Cancer analysis project. Nature Genetics, 2013, 45, 1113-1120.	9.4	6,265
268	Genomic testing to determine drug response: measuring preferences of the public and patients using Discrete Choice Experiment (DCE). BMC Health Services Research, 2013, 13, 454.	0.9	28
269	A Pan-BCL2 Inhibitor Renders Bone-Marrow-Resident Human Leukemia Stem Cells Sensitive to Tyrosine Kinase Inhibition. Cell Stem Cell, 2013, 12, 316-328.	5.2	167
270	Analysis of FOXO1 mutations in diffuse large B-cell lymphoma. Blood, 2013, 121, 3666-3674.	0.6	139

#	Article	IF	CITATIONS
271	Cancer genome-sequencing study design. Nature Reviews Genetics, 2013, 14, 321-332.	7.7	100
272	Complete genomic landscape of a recurring sporadic parathyroid carcinoma. Journal of Pathology, 2013, 230, 249-260.	2.1	57
273	Integrated genomic characterization of endometrial carcinoma. Nature, 2013, 497, 67-73.	13.7	4,075
274	Genomic and Epigenomic Landscapes of Adult De Novo Acute Myeloid Leukemia. New England Journal of Medicine, 2013, 368, 2059-2074.	13.9	4,139
275	DNA hypomethylation within specific transposable element families associates with tissue-specific enhancer landscape. Nature Genetics, 2013, 45, 836-841.	9.4	207
276	The genetic landscape of high-risk neuroblastoma. Nature Genetics, 2013, 45, 279-284.	9.4	990
277	Transcriptomics in the Age of Ultra High-Throughput Sequencing. , 2013, , 145-154.		0
278	Genome-wide microRNA and messenger RNA profiling in rodent liver development implicates mir302b and mir20a in repressing transforming growth factor-beta signaling. Hepatology, 2013, 57, 2491-2501.	3.6	17
279	A Cancer Stem Cell Model for Studying Brain Metastases From Primary Lung Cancer. Journal of the National Cancer Institute, 2013, 105, 551-562.	3.0	50
280	Comprehensive molecular characterization of clear cell renal cell carcinoma. Nature, 2013, 499, 43-49.	13.7	2,839
281	Cell culture and (i>Drosophila / i>model systems define three classes of anaplastic lymphoma kinase mutations in neuroblastoma. DMM Disease Models and Mechanisms, 2013, 6, 373-82.	1.2	59
282	Functional DNA methylation differences between tissues, cell types, and across individuals discovered using the M&M algorithm. Genome Research, 2013, 23, 1522-1540.	2.4	162
283	ADAR1 promotes malignant progenitor reprogramming in chronic myeloid leukemia. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 1041-1046.	3.3	148
284	Distinct evolutionary trajectories of primary highâ€grade serous ovarian cancers revealed through spatial mutational profiling. Journal of Pathology, 2013, 231, 21-34.	2.1	357
285	The E3 ubiquitin ligase UBR5 is recurrently mutated in mantle cell lymphoma. Blood, 2013, 121, 3161-3164.	0.6	124
286	Mutational and structural analysis of diffuse large B-cell lymphoma using whole-genome sequencing. Blood, 2013, 122, 1256-1265.	0.6	349
287	MicroRNA dysregulation in B-cell non-Hodgkin lymphoma. Blood and Lymphatic Cancer: Targets and Therapy, 2013, , 25.	1.2	0
288	A Phosphoproteomics Approach to Identify Candidate Kinase Inhibitor Pathway Targets in Lymphoma-Like Primary Cell Lines. Current Drug Discovery Technologies, 2013, 10, 283-304.	0.6	2

#	Article	IF	CITATIONS
289	Next-Generation Sequencing of Prostate Tumors Provides Independent Evidence of Xenotropic Murine Leukemia Virus-Related Gammaretrovirus Contamination. Journal of Clinical Microbiology, 2012, 50, 536-537.	1.8	2
290	Interaction of Cyclin-Dependent Kinase 12/CrkRS with Cyclin K1 Is Required for the Phosphorylation of the C-Terminal Domain of RNA Polymerase II. Molecular and Cellular Biology, 2012, 32, 4691-4704.	1.1	93
291	Next Generation Sequencing of Prostate Cancer from a Patient Identifies a Deficiency of Methylthioadenosine Phosphorylase, an Exploitable Tumor Target. Molecular Cancer Therapeutics, 2012, 11, 775-783.	1.9	34
292	Recurrent Somatic <i>DICER1</i> Mutations in Nonepithelial Ovarian Cancers. New England Journal of Medicine, 2012, 366, 234-242.	13.9	401
293	Integrative analysis of genome-wide loss of heterozygosity and monoallelic expression at nucleotide resolution reveals disrupted pathways in triple-negative breast cancer. Genome Research, 2012, 22, 1995-2007.	2.4	237
294	Feature-based classifiers for somatic mutation detection in tumour–normal paired sequencing data. Bioinformatics, 2012, 28, 167-175.	1.8	130
295	14-3-3 fusion oncogenes in high-grade endometrial stromal sarcoma. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 929-934.	3.3	239
296	Penetrance of biallelic SMARCAL1 mutations is associated with environmental and genetic disturbances of gene expression. Human Molecular Genetics, 2012, 21, 2572-2587.	1.4	57
297	Subgroup-specific structural variation across 1,000 medulloblastoma genomes. Nature, 2012, 488, 49-56.	13.7	761
298	JointSNVMix: a probabilistic model for accurate detection of somatic mutations in normal/tumour paired next-generation sequencing data. Bioinformatics, 2012, 28, 907-913.	1.8	159
299	Allelic Ratios and the Mutational Landscape Reveal Biologically Significant Heterozygous SNVs. Genetics, 2012, 190, 1225-1233.	1.2	12
300	Whole transcriptome sequencing reveals recurrent NOTCH1 mutations in mantle cell lymphoma. Blood, 2012, 119, 1963-1971.	0.6	313
301	TBL1XR1/TP63: a novel recurrent gene fusion in B-cell non-Hodgkin lymphoma. Blood, 2012, 119, 4949-4952.	0.6	60
302	BreakFusion: targeted assembly-based identification of gene fusions in whole transcriptome paired-end sequencing data. Bioinformatics, 2012, 28, 1923-1924.	1.8	54
303	Rapid, reliable, and reproducible molecular sub-grouping of clinical medulloblastoma samples. Acta Neuropathologica, 2012, 123, 615-626.	3.9	318
304	Genetic Alterations Activating Kinase and Cytokine Receptor Signaling in High-Risk Acute Lymphoblastic Leukemia. Cancer Cell, 2012, 22, 153-166.	7.7	621
305	Hive plotsrational approach to visualizing networks. Briefings in Bioinformatics, 2012, 13, 627-644.	3.2	187
306	Lifeâ€history chronicle for a patient with the recently described chromosome 4q21 microdeletion syndrome. American Journal of Medical Genetics, Part A, 2012, 158A, 2606-2609.	0.7	5

#	Article	IF	Citations
307	Polyâ€gene fusion transcripts and chromothripsis in prostate cancer. Genes Chromosomes and Cancer, 2012, 51, 1144-1153.	1.5	46
308	Comprehensive molecular characterization of human colon and rectal cancer. Nature, 2012, 487, 330-337.	13.7	7,168
309	Comprehensive molecular portraits of human breast tumours. Nature, 2012, 490, 61-70.	13.7	10,282
310	Barriers to integrating personalized medicine into clinical practice: a best–worst scaling choice experiment. Genetics in Medicine, 2012, 14, 520-526.	1.1	34
311	Comprehensive genomic characterization of squamous cell lung cancers. Nature, 2012, 489, 519-525.	13.7	3,483
312	Twist1 Transcriptional Targets in the Developing Atrio-Ventricular Canal of the Mouse. PLoS ONE, 2012, 7, e40815.	1.1	10
313	Mutation Discovery in Regions of Segmental Cancer Genome Amplifications with CoNAn-SNV: A Mixture Model for Next Generation Sequencing of Tumors. PLoS ONE, 2012, 7, e41551.	1.1	4
314	Recurrent targets of aberrant somatic hypermutation in lymphoma. Oncotarget, 2012, 3, 1308-1319.	0.8	127
315	Concurrent <i>CIC</i> mutations, <i>IDH</i> mutations, and 1p/19q loss distinguish oligodendrogliomas from other cancers. Journal of Pathology, 2012, 226, 7-16.	2.1	272
316	Integrated genome and transcriptome sequencing identifies a novel form of hybrid and aggressive prostate cancer. Journal of Pathology, 2012, 227, 53-61.	2.1	63
317	From sequence to molecular pathology, and a mechanism driving the neuroendocrine phenotype in prostate cancer. Journal of Pathology, 2012, 227, 286-297.	2.1	161
318	Applications of High-Throughput Sequencing. , 2012, , 27-53.		1
319	DNA hypermethylation and 1p Loss silence <i>NHEâ€1</i> i> in oligodendroglioma. Annals of Neurology, 2012, 71, 845-849.	2.8	22
320	The clonal and mutational evolution spectrum of primary triple-negative breast cancers. Nature, 2012, 486, 395-399.	13.7	1,778
321	Subgroup-specific alternative splicing in medulloblastoma. Acta Neuropathologica, 2012, 123, 485-499.	3.9	28
322	Mutations in EZH2 Cause Weaver Syndrome. American Journal of Human Genetics, 2012, 90, 110-118.	2.6	253
323	High-throughput microfluidic single-cell RT-qPCR. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 13999-14004.	3.3	406
324	Frequent mutation of histone-modifying genes in non-Hodgkin lymphoma. Nature, 2011, 476, 298-303.	13.7	1,428

#	Article	IF	Citations
325	Notch Initiates the Endothelial-to-Mesenchymal Transition in the Atrioventricular Canal through Autocrine Activation of Soluble Guanylyl Cyclase. Developmental Cell, 2011, 21, 288-300.	3.1	144
326	Comprehensive analysis of mammalian miRNA* species and their role in myeloid cells. Blood, 2011, 118, 3350-3358.	0.6	90
327	Genome-wide identification of human microRNAs located in leukemia-associated genomic alterations. Blood, 2011, 117, 595-607.	0.6	105
328	Somatic mutations at EZH2 Y641 act dominantly through a mechanism of selectively altered PRC2 catalytic activity, to increase H3K27 trimethylation. Blood, 2011, 117, 2451-2459.	0.6	556
329	MHC class II transactivator CIITA is a recurrent gene fusion partner in lymphoid cancers. Nature, 2011, 471, 377-381.	13.7	551
330	Retinoblastoma-binding proteins 4 and 9 are important for human pluripotent stem cell maintenance. Experimental Hematology, 2011, 39, 866-879.e1.	0.2	30
331	Cell of Origin in AML: Susceptibility to MN1-Induced Transformation Is Regulated by the MEIS1/AbdB-like HOX Protein Complex. Cancer Cell, 2011, 20, 39-52.	7.7	76
332	Comparison of genome-wide array genomic hybridization platforms for the detection of copy number variants in idiopathic mental retardation. BMC Medical Genomics, 2011, 4, 25.	0.7	26
333	Using nextâ€generation sequencing for the diagnosis of rare disorders: a family with retinitis pigmentosa and skeletal abnormalities. Journal of Pathology, 2011, 225, 12-18.	2.1	29
334	Deep annotation of <i>Drosophila melanogaster</i> microRNAs yields insights into their processing, modification, and emergence. Genome Research, 2011, 21, 203-215.	2.4	207
335	Extensive relationship between antisense transcription and alternative splicing in the human genome. Genome Research, 2011, 21, 1203-1212.	2.4	68
336	Genome and transcriptome analyses of the mountain pine beetle-fungal symbiont <i>Grosmannia clavigera</i> , a lodgepole pine pathogen. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 2504-2509.	3.3	218
337	deFuse: An Algorithm for Gene Fusion Discovery in Tumor RNA-Seq Data. PLoS Computational Biology, 2011, 7, e1001138.	1.5	477
338	Characterization of the Contradictory Chromatin Signatures at the $3\hat{a}\in^2$ Exons of Zinc Finger Genes. PLoS ONE, 2011, 6, e17121.	1.1	64
339	Next generation sequencing based approaches to epigenomics. Briefings in Functional Genomics, 2010, 9, 455-465.	1.3	60
340	Hypomorphic Temperature-Sensitive Alleles of NSDHL Cause CK Syndrome. American Journal of Human Genetics, 2010, 87, 905-914.	2.6	64
341	Genomic sequence of a mutant strain of Caenorhabditis elegans with an altered recombination pattern. BMC Genomics, 2010, 11, 131.	1.2	9
342	Selective targeting of neuroblastoma tumourâ€initiating cells by compounds identified in stem cellâ€based small molecule screens. EMBO Molecular Medicine, 2010, 2, 371-384.	3.3	62

#	Article	IF	CITATIONS
343	High resolution analysis of follicular lymphoma genomes reveals somatic recurrent sites of copyâ€neutral loss of heterozygosity and copy number alterations that target single genes. Genes Chromosomes and Cancer, 2010, 49, 669-681.	1.5	51
344	LNCaP Atlas: Gene expression associated with in vivoprogression to castration-recurrent prostate cancer. BMC Medical Genomics, 2010, 3, 43.	0.7	73
345	Convergent Genesis of an Adult Neural Crest-Like Dermal Stem Cell from Distinct Developmental Origins. Stem Cells, 2010, 28, 2027-2040.	1.4	100
346	International network of cancer genome projects. Nature, 2010, 464, 993-998.	13.7	2,114
347	Conserved role of intragenic DNA methylation in regulating alternative promoters. Nature, 2010, 466, 253-257.	13.7	1,568
348	Comparison of sequencing-based methods to profile DNA methylation and identification of monoallelic epigenetic modifications. Nature Biotechnology, 2010, 28, 1097-1105.	9.4	647
349	The NIH Roadmap Epigenomics Mapping Consortium. Nature Biotechnology, 2010, 28, 1045-1048.	9.4	1,705
350	Somatic mutations altering EZH2 (Tyr641) in follicular and diffuse large B-cell lymphomas of germinal-center origin. Nature Genetics, 2010, 42, 181-185.	9.4	1,504
351	Identification of miR-145 and miR-146a as mediators of the 5q– syndrome phenotype. Nature Medicine, 2010, 16, 49-58.	15.2	588
352	Alternative expression analysis by RNA sequencing. Nature Methods, 2010, 7, 843-847.	9.0	283
352 353	Alternative expression analysis by RNA sequencing. Nature Methods, 2010, 7, 843-847. De novo assembly and analysis of RNA-seq data. Nature Methods, 2010, 7, 909-912.	9.0	283 886
353	De novo assembly and analysis of RNA-seq data. Nature Methods, 2010, 7, 909-912.	9.0	886
353 354	De novo assembly and analysis of RNA-seq data. Nature Methods, 2010, 7, 909-912. Whole-Genome Profiling of Mutagenesis in < i> Caenorhabditis elegans < / i> . Genetics, 2010, 185, 431-441. Acquired < i> TNFRSF14 < / i> Mutations in Follicular Lymphoma Are Associated with Worse Prognosis.	9.0	135
353 354 355	De novo assembly and analysis of RNA-seq data. Nature Methods, 2010, 7, 909-912. Whole-Genome Profiling of Mutagenesis in < i>Caenorhabditis elegans < / i>. Genetics, 2010, 185, 431-441. Acquired < i>TNFRSF14 < / i> Mutations in Follicular Lymphoma Are Associated with Worse Prognosis. Cancer Research, 2010, 70, 9166-9174. LaneRuler: Automated Lane Tracking for DNA Electrophoresis Gel Images. IEEE Transactions on	9.0 1.2 0.4	886 135 160
353 354 355 356	De novo assembly and analysis of RNA-seq data. Nature Methods, 2010, 7, 909-912. Whole-Genome Profiling of Mutagenesis in <i>Caenorhabditis elegans </i> Caenothabditis elegans Cancer Research, 2010, 70, 9166-9174. LaneRuler: Automated Lane Tracking for DNA Electrophoresis Gel Images. IEEE Transactions on Automation Science and Engineering, 2010, 7, 706-708. System-Level Analysis of Neuroblastoma Tumor†Initiating Cells Implicates AURKB as a Novel Drug	9.0 1.2 0.4 3.4	886 135 160 12
353 354 355 356	De novo assembly and analysis of RNA-seq data. Nature Methods, 2010, 7, 909-912. Whole-Genome Profiling of Mutagenesis in ⟨i⟩ Caenorhabditis elegans ⟨li⟩. Genetics, 2010, 185, 431-441. Acquired ⟨i⟩ TNFRSF14⟨li⟩ Mutations in Follicular Lymphoma Are Associated with Worse Prognosis. Cancer Research, 2010, 70, 9166-9174. LaneRuler: Automated Lane Tracking for DNA Electrophoresis Gel Images. IEEE Transactions on Automation Science and Engineering, 2010, 7, 706-708. System-Level Analysis of Neuroblastoma Tumor–Initiating Cells Implicates AURKB as a Novel Drug Target for Neuroblastoma. Clinical Cancer Research, 2010, 16, 4572-4582. Genomic analysis distinguishes phases of early development of the mouse atrio-ventricular canal.	9.0 1.2 0.4 3.4	135 160 12 43

#	Article	IF	Citations
361	MicroRNA transcriptome in the newborn mouse ovaries determined by massive parallel sequencing. Molecular Human Reproduction, 2010, 16, 463-471.	1.3	122
362	Identification of Functional Elements and Regulatory Circuits by <i>Drosophila</i> modENCODE. Science, 2010, 330, 1787-1797.	6.0	1,124
363	ATR-X Syndrome Protein Targets Tandem Repeats and Influences Allele-Specific Expression in a Size-Dependent Manner. Cell, 2010, 143, 367-378.	13.5	365
364	<i>ARID1A</i> Mutations in Endometriosis-Associated Ovarian Carcinomas. New England Journal of Medicine, 2010, 363, 1532-1543.	13.9	1,460
365	Evolution of an adenocarcinoma in response to selection by targeted kinase inhibitors. Genome Biology, 2010, 11, R82.	13.9	159
366	BCL6 repression of EP300 in human diffuse large B cell lymphoma cells provides a basis for rational combinatorial therapy. Journal of Clinical Investigation, 2010, 120, 4569-4582.	3.9	101
367	Genetic Variation in Healthy Oldest-Old. PLoS ONE, 2009, 4, e6641.	1.1	42
368	The completion of the Mammalian Gene Collection (MGC). Genome Research, 2009, 19, 2324-2333.	2.4	125
369	High-resolution profiling and discovery of planarian small RNAs. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 11546-11551.	3.3	128
370	Massively parallel sequencing of the polyadenylated transcriptome of <i>C. elegans</i> . Genome Research, 2009, 19, 657-666.	2.4	169
371	An Integrated Strategy to Study Muscle Development and Myofilament Structure in Caenorhabditis elegans. PLoS Genetics, 2009, 5, e1000537.	1.5	89
372	Sequence Variant Discovery in DNA Repair Genes from Radiosensitive and Radiotolerant Prostate Brachytherapy Patients. Clinical Cancer Research, 2009, 15, 5008-5016.	3.2	42
373	<i>De novo</i> transcriptome assembly with ABySS. Bioinformatics, 2009, 25, 2872-2877.	1.8	371
374	Identification of genes expressed in the hermaphrodite germ line of C. elegans using SAGE. BMC Genomics, 2009, 10, 213.	1.2	105
375	Identification of novel androgen-responsive genes by sequencing of LongSAGE libraries. BMC Genomics, 2009, 10, 476.	1.2	75
376	Detection of pathogenic copy number variants in children with idiopathic intellectual disability using 500 K SNP array genomic hybridization. BMC Genomics, 2009, 10, 526.	1.2	30
377	A novel de novo 1.1 Mb duplication of 17q21.33 associated with cognitive impairment and other anomalies. American Journal of Medical Genetics, Part A, 2009, 149A, 1257-1262.	0.7	14
378	Mutational evolution in a lobular breast tumour profiled at single nucleotide resolution. Nature, 2009, 461, 809-813.	13.7	984

#	Article	IF	Citations
379	A first look at entire human methylomes. Nature Biotechnology, 2009, 27, 1130-1132.	9.4	14
380	Massively Parallel Sequencing: The Next Big Thing in Genetic Medicine. American Journal of Human Genetics, 2009, 85, 142-154.	2.6	308
381	Next-generation tag sequencing for cancer gene expression profiling. Genome Research, 2009, 19, 1825-1835.	2.4	306
382	The Genome Sequence of Taurine Cattle: A Window to Ruminant Biology and Evolution. Science, 2009, 324, 522-528.	6.0	1,038
383	A characteristic syndrome associated with microduplication of 8q12, inclusive of CHD7. European Journal of Medical Genetics, 2009, 52, 436-439.	0.7	21
384	Epigenetics and human disease. International Journal of Biochemistry and Cell Biology, 2009, 41, 136-146.	1.2	99
385	SKPs Derive from Hair Follicle Precursors and Exhibit Properties of Adult Dermal Stem Cells. Cell Stem Cell, 2009, 5, 610-623.	5.2	335
386	ELT-2 is the predominant transcription factor controlling differentiation and function of the C. elegans intestine, from embryo to adult. Developmental Biology, 2009, 327, 551-565.	0.9	129
387	Mutation of (i>FOXL2ii>in Granulosa-Cell Tumors of the Ovary. New England Journal of Medicine, 2009, 360, 2719-2729.	13.9	706
388	De novo genome sequence assembly of a filamentous fungus using Sanger, 454 and Illumina sequence data. Genome Biology, 2009, 10, R94.	13.9	130
389	Applications of New Sequencing Technologies for Transcriptome Analysis. Annual Review of Genomics and Human Genetics, 2009, 10, 135-151.	2.5	457
390	Circos: An information aesthetic for comparative genomics. Genome Research, 2009, 19, 1639-1645.	2.4	9,003
391	Prospective isolation and molecular characterization of hematopoietic stem cells with durable self-renewal potential. Blood, 2009, 113, 6342-6350.	0.6	300
392	Transcriptome analysis for Caenorhabditis elegansbased on novel expressed sequence tags. BMC Biology, 2008, 6, 30.	1.7	46
393	ALEXA: a microarray design platform for alternative expression analysis. Nature Methods, 2008, 5, 118-118.	9.0	19
394	A conifer genomics resource of 200,000 spruce (Picea spp.) ESTs and 6,464 high-quality, sequence-finished full-length cDNAs for Sitka spruce (Picea sitchensis). BMC Genomics, 2008, 9, 484.	1.2	113
395	Analysis of 4,664 high-quality sequence-finished poplar full-length cDNA clones and their utility for the discovery of genes responding to insect feeding. BMC Genomics, 2008, 9, 57.	1.2	68
396	Identification of a set of genes showing regionally enriched expression in the mouse brain. BMC Neuroscience, 2008, 9, 66.	0.8	25

#	Article	IF	Citations
397	Molecular profiling reveals similarities and differences between primitive subsets of hematopoietic cells generated in vitro from human embryonic stem cells and in vivo during embryogenesis. Experimental Hematology, 2008, 36, 1377-1389.	0.2	17
398	From cytogenetics to next-generation sequencing technologies: advances in the detection of genome rearrangements in tumorsThis paper is one of a selection of papers published in this Special Issue, entitled CSBMCB — Systems and Chemical Biology, and has undergone the Journal's usual peer review process Biochemistry and Cell Biology, 2008, 86, 81-91.	0.9	35
399	Identification of transcripts with enriched expression in the developing and adult pancreas. Genome Biology, 2008, 9, R99.	13.9	33
400	Applications of next-generation sequencing technologies in functional genomics. Genomics, 2008, 92, 255-264.	1.3	1,013
401	Transcriptome Analysis of the Normal Human Mammary Cell Commitment and Differentiation Process. Cell Stem Cell, 2008, 3, 109-118.	5.2	310
402	In-depth characterization of the microRNA transcriptome in a leukemia progression model. Genome Research, 2008, 18, 1787-1797.	2.4	162
403	Application of massively parallel sequencing to microRNA profiling and discovery in human embryonic stem cells. Genome Research, 2008, 18, 610-621.	2.4	964
404	Global analysis of in vivo Foxa2-binding sites in mouse adult liver using massively parallel sequencing. Nucleic Acids Research, 2008, 36, 4549-4564.	6.5	137
405	Profiling the HeLa S3 transcriptome using randomly primed cDNA and massively parallel short-read sequencing. BioTechniques, 2008, 45, 81-94.	0.8	355
406	Genome-wide relationship between histone H3 lysine 4 mono- and tri-methylation and transcription factor binding. Genome Research, 2008, 18, 1906-1917.	2.4	163
407	Use of Affymetrix Mapping Arrays in the Diagnosis of Gene Copy Number Variation. Current Protocols in Human Genetics, 2008, 59, Unit 8.13.	3.5	7
408	A Seriation Approach for Visualization-Driven Discovery of Co-Expression Patterns in Serial Analysis of Gene Expression (SAGE) Data. PLoS ONE, 2008, 3, e3205.	1.1	1
409	Largeâ€Scale BAC Clone Restriction Digest Fingerprinting. Current Protocols in Human Genetics, 2007, 53, Unit 5.19.	3.5	5
410	High-Throughput In Vivo Analysis of Gene Expression in Caenorhabditis elegans. PLoS Biology, 2007, 5, e237.	2.6	346
411	Novel deletions of 14q11.2 associated with developmental delay, cognitive impairment and similar minor anomalies in three children. Journal of Medical Genetics, 2007, 44, 556-561.	1.5	68
412	Osteopoikilosis, short stature and mental retardation as key features of a new microdeletion syndrome on 12q14. Journal of Medical Genetics, 2007, 44, 264-268.	1.5	58
413	The molecular signature and <i>cis</i> -regulatory architecture of a <i>C. elegans</i> gustatory neuron. Genes and Development, 2007, 21, 1653-1674.	2.7	151
414	Identification and analysis of internal promoters in <i>Caenorhabditis elegans</i> operons. Genome Research, 2007, 17, 1478-1485.	2.4	41

#	Article	IF	CITATIONS
415	The ELT-2 GATA-factor and the global regulation of transcription in the C. elegans intestine. Developmental Biology, 2007, 302, 627-645.	0.9	165
416	Generation of ESTs in Vitis vinifera wine grape (Cabernet Sauvignon) and table grape (Muscat) Tj ETQq0 0 0 rgBT 402, 40-50.	Overlock 1.0	10 Tf 50 707 45
417	Epidermal growth factor receptor (EGFR) is transcriptionally induced by the Y-box binding protein-1 (YB-1) and can be inhibited with Iressa in basal-like breast cancer, providing a potential target for therapy. Breast Cancer Research, 2007, 9, R61.	2.2	126
418	A BAC clone fingerprinting approach to the detection of human genome rearrangements. Genome Biology, 2007, 8, R224.	13.9	10
419	A physical map of the bovine genome. Genome Biology, 2007, 8, R165.	13.9	73
420	LongSAGE profiling of nine human embryonic stem cell lines. Genome Biology, 2007, 8, R113.	13.9	21
421	DiscoverySpace: an interactive data analysis application. Genome Biology, 2007, 8, R6.	13.9	41
422	Evolutionary and Biomedical Insights from the Rhesus Macaque Genome. Science, 2007, 316, 222-234.	6.0	1,283
423	Analyses of deep mammalian sequence alignments and constraint predictions for 1% of the human genome. Genome Research, 2007, 17, 760-774.	2.4	184
424	Novel expressed sequences identified in a model of androgen independent prostate cancer. BMC Genomics, 2007, 8, 32.	1.2	3
425	A Modified Polymerase Chain Reaction-Long Serial Analysis of Gene Expression Protocol Identifies Novel Transcripts in Human CD34+Bone Marrow Cells. Stem Cells, 2007, 25, 1681-1689.	1.4	8
426	A systematic screen for genes expressed in definitive endoderm by Serial Analysis of Gene Expression (SAGE). BMC Developmental Biology, 2007, 7, 92.	2.1	52
427	Genome-wide profiles of STAT1 DNA association using chromatin immunoprecipitation and massively parallel sequencing. Nature Methods, 2007, 4, 651-657.	9.0	1,254
428	Genome of the marsupial Monodelphis domestica reveals innovation in non-coding sequences. Nature, 2007, 447, 167-177.	13.7	661
429	A physical map of the highly heterozygous Populus genome: integration with the genome sequence and genetic map and analysis of haplotype variation. Plant Journal, 2007, 50, 1063-1078.	2.8	70
430	Generation of a wheat leaf rust, Puccinia triticina, EST database from stage-specific cDNA libraries. Molecular Plant Pathology, 2007, 8, 451-467.	2.0	49
431	Correlations of EGFR mutations and increases in EGFR and HER2 copy number to gefitinib response in a retrospective analysis of lung cancer patients. BMC Cancer, 2007, 7, 128.	1.1	36
432	Assessment of algorithms for high throughput detection of genomic copy number variation in oligonucleotide microarray data. BMC Bioinformatics, 2007, 8, 368.	1.2	49

#	Article	IF	Citations
433	Genes that may modulate longevity in C. elegans in both dauer larvae and long-lived daf-2 adults. Experimental Gerontology, 2007, 42, 825-839.	1.2	26
434	Identification of ciliary and ciliopathy genes in Caenorhabditis elegans through comparative genomics. Genome Biology, 2006, 7, R126.	13.9	86
435	The Genome of the Sea Urchin Strongylocentrotus purpuratus. Science, 2006, 314, 941-952.	6.0	1,018
436	Oligonucleotide Microarray Analysis of Genomic Imbalance in Children with Mental Retardation. American Journal of Human Genetics, 2006, 79, 500-513.	2.6	261
437	The complete genome of Rhodococcus sp. RHA1 provides insights into a catabolic powerhouse. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 15582-15587.	3.3	586
438	Mating factor linkage and genome evolution in basidiomycetous pathogens of cereals. Fungal Genetics and Biology, 2006, 43, 655-666.	0.9	59
439	Identification of Novel Lung Genes in Bronchial Epithelium by Serial Analysis of Gene Expression. American Journal of Respiratory Cell and Molecular Biology, 2006, 35, 651-661.	1.4	51
440	Genomics of hybrid poplar (Populus trichocarpa× deltoides) interacting with forest tent caterpillars (Malacosoma disstria): normalized and full-length cDNA libraries, expressed sequence tags, and a cDNA microarray for the study of insect-induced defences. Molecular Ecology, 2006, 15, 1275-1297.	2.0	183
441	Conifer defence against insects: microarray gene expression profiling of Sitka spruce (Picea) Tj ETQq1 1 0.784314 transcriptome. Plant, Cell and Environment, 2006, 29, 1545-1570.	4 rgBT /Ov 2.8	verlock 10 Tf 221
442	Analysis of the prostate cancer cell line LNCaP transcriptome using a sequencing-by-synthesis approach. BMC Genomics, 2006, 7, 246.	1.2	173
443	Sequence biases in large scale gene expression profiling data. Nucleic Acids Research, 2006, 34, e83-e83.	6.5	51
444	Physical map-assisted whole-genome shotgun sequence assemblies. Genome Research, 2006, 16, 768-775.	2.4	27
445	Large-scale production of SAGE libraries from microdissected tissues, flow-sorted cells, and cell lines. Genome Research, 2006, 17, 108-116.	2.4	34
446	Sequencing and analysis of 10,967 full-length cDNA clones from Xenopus laevis and Xenopus tropicalis reveals post-tetraploidization transcriptome remodeling. Genome Research, 2006, 16, 796-803.	2.4	73
447	Generation and annotation of the DNA sequences of human chromosomes 2 and 4. Nature, 2005, 434, 724-731.	13.7	85
448	Functional Genomics of the Cilium, a Sensory Organelle. Current Biology, 2005, 15, 935-941.	1.8	245
449	Generation, annotation, analysis and database integration of 16,500 white spruce EST clusters. BMC Genomics, 2005, 6, 144.	1.2	119
450	High-throughput sequencing: a failure mode analysis. BMC Genomics, 2005, 6, 2.	1.2	10

#	Article	IF	CITATIONS
451	Management and visualization of whole genome shotgun assemblies using SAM. BioTechniques, 2005, 38, 715-720.	0.8	7
452	Simple, robust methods for high-throughput nanoliter-scale DNA sequencing. Genome Research, 2005, 15, 1447-1450.	2.4	13
453	Analysis of long-lived C. elegans daf-2 mutants using serial analysis of gene expression. Genome Research, 2005, 15, 603-615.	2.4	180
454	A high-resolution whole-genome cattle-human comparative map reveals details of mammalian chromosome evolution. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 18526-18531.	3.3	115
455	Serial Analysis of Gene Expression Reveals Conserved Links between Protein Kinase A, Ribosome Biogenesis, and Phosphate Metabolism in Ustilago maydis. Eukaryotic Cell, 2005, 4, 2029-2043.	3.4	25
456	A mouse atlas of gene expression: Large-scale digital gene-expression profiles from precisely defined developing C57BL/6J mouse tissues and cells. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 18485-18490.	3.3	112
457	Identification by full-coverage array CGH of human DNA copy number increases relative to chimpanzee and gorilla. Genome Research, 2005, 16, 173-181.	2.4	48
458	The Genome of the Kinetoplastid Parasite, Leishmania major. Science, 2005, 309, 436-442.	6.0	1,237
459	The Genome of the Basidiomycetous Yeast and Human Pathogen Cryptococcus neoformans. Science, 2005, 307, 1321-1324.	6.0	664
460	A physical map of the genome of Atlantic salmon, Salmo salar. Genomics, 2005, 86, 396-404.	1.3	97
461	Development and Application of a Salmonid EST Database and cDNA Microarray: Data Mining and Interspecific Hybridization Characteristics. Genome Research, 2004, 14, 478-490.	2.4	279
462	Functional Characterization of a Catabolic Plasmid from Polychlorinated-Biphenyl-Degrading Rhodococcus sp. Strain RHA1. Journal of Bacteriology, 2004, 186, 7783-7795.	1.0	65
463	Integrated and Sequence-Ordered BAC- and YAC-Based Physical Maps for the Rat Genome. Genome Research, 2004, 14, 766-779.	2.4	44
464	Systematic Recovery and Analysis of Full-ORF Human cDNA Clones. Genome Research, 2004, 14, 2083-2092.	2.4	28
465	The Status, Quality, and Expansion of the NIH Full-Length cDNA Project: The Mammalian Gene Collection (MGC). Genome Research, 2004, 14, 2121-2127.	2.4	486
466	Differential expression of a novel ankyrin containing E3 ubiquitin-protein ligase, Hace1, in sporadic Wilms' tumor versus normal kidney. Human Molecular Genetics, 2004, 13, 2061-2074.	1.4	100
467	Automated ordering of fingerprinted clones. Bioinformatics, 2004, 20, 1264-1271.	1.8	18
468	Iron-regulated transcription and capsule formation in the fungal pathogen Cryptococcus neoformans. Molecular Microbiology, 2004, 55, 1452-1472.	1.2	90

#	Article	IF	CITATIONS
469	A tiling resolution DNA microarray with complete coverage of the human genome. Nature Genetics, 2004, 36, 299-303.	9.4	597
470	Genome sequence of the Brown Norway rat yields insights into mammalian evolution. Nature, 2004, 428, 493-521.	13.7	1,943
471	Effect of TERT and ATM on gene expression profiles in human fibroblasts. Genes Chromosomes and Cancer, 2004, 39, 298-310.	1.5	11
472	Delineation of a minimal region of deletion at 6q16.3 in follicular lymphoma and construction of a bacterial artificial chromosome contig spanning a 6-megabase region of 6q16-q21. Genes Chromosomes and Cancer, 2004, 40, 60-65.	1.5	26
473	A set of BAC clones spanning the human genome. Nucleic Acids Research, 2004, 32, 3651-3660.	6.5	119
474	New Genomic Tools for Molecular Studies of Evolutionary Change in Threespine Sticklebacks. Behaviour, 2004, 141, 1331-1344.	0.4	64
475	Serial Analysis of Gene Expression Profiles of Developmental Stages in Non-small Cell Lung Carcinoma. Chest, 2004, 125, 97S.	0.4	6
476	Novel Avian Influenza H7N3 Strain Outbreak, British Columbia. Emerging Infectious Diseases, 2004, 10, 2192-2195.	2.0	182
477	A SAGE Approach to Discovery of Genes Involved in Autophagic Cell Death. Current Biology, 2003, 13, 358-363.	1.8	198
478	The DNA sequence of human chromosome 7. Nature, 2003, 424, 157-164.	13.7	236
479	Comparison of medulloblastoma and normal neural transcriptomes identifies a restricted set of activated genes. Oncogene, 2003, 22, 7687-7694.	2.6	72
480	The Genome Sequence of the SARS-Associated Coronavirus. Science, 2003, 300, 1399-1404.	6.0	1,842
481	Software for Automated Analysis of DNA Fingerprinting Gels. Genome Research, 2003, 13, 940-953.	2.4	35
482	Internet Contig Explorer (iCE)A Tool for Visualizing Clone Fingerprint Maps. Genome Research, 2003, 13, 1244-1249.	2.4	22
483	Assessment of SAGE in Transcript Identification. Genome Research, 2003, 13, 1203-1215.	2.4	61
484	The Genome Sequence of Caenorhabditis briggsae: A Platform for Comparative Genomics. PLoS Biology, 2003, 1, e45.	2.6	812
485	Physical Maps for Genome Analysis of Serotype A and D Strains of the Fungal Pathogen Cryptococcus neoformans. Genome Research, 2002, 12, 1445-1453.	2.4	38
486	Systematic sequencing of cDNA clones using the transposon Tn5. Nucleic Acids Research, 2002, 30, 2469-2477.	6.5	55

#	Article	IF	CITATIONS
487	An efficient strategy for large-scale high-throughput transposon-mediated sequencing of cDNA clones. Nucleic Acids Research, 2002, 30, 2460-2468.	6.5	29
488	Temperature-Regulated Transcription in the Pathogenic Fungus Cryptococcus neoformans. Genome Research, 2002, 12, 1386-1400.	2.4	84
489	Transferrin receptor 2 (TfR2) and HFE mutational analysis in non-C282Y iron overload: identification of a novel TfR2 mutation. Blood, 2002, 100, 1075-1077.	0.6	97
490	Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences. Proceedings of the National Academy of Sciences of the United States of America, 2002, 99, 16899-16903.	3.3	1,610
491	A physical map of the mouse genome. Nature, 2002, 418, 743-750.	13.7	316
492	Dissection of the promoter region of the inositol 1,4,5-trisphosphate receptor gene, itr-1, in C. elegans: a molecular basis for cell-specific expression of IP3R isoforms11Edited by J. Karn. Journal of Molecular Biology, 2001, 306, 145-157.	2.0	38
493	An SSLP marker–anchored BAC framework map of the mouse genome. Nature Genetics, 2001, 29, 133-134.	9.4	14
494	Initial sequencing and analysis of the human genome. Nature, 2001, 409, 860-921.	13.7	21,074
495	Changes in Gene Expression Associated with Developmental Arrest and Longevity in Caenorhabditis elegans. Genome Research, 2001, 11, 1346-1352.	2.4	202
496	An Oligonucleotide Fingerprint Normalized and Expressed Sequence Tag Characterized Zebrafish cDNA Library. Genome Research, 2001, 11, 1594-1602.	2.4	64
497	Identifying Potential Tumor Markers and Antigens by Database Mining and Rapid Expression Screening. Genome Research, 2000, 10, 1393-1402.	2.4	92
498	Conservation of sequence and function of the pag-3 genes from C. elegans and C. briggsae. Gene, 2000, 243, 67-74.	1.0	7
499	Molecular Profiling of Clinical Tissue Specimens. Journal of Molecular Diagnostics, 2000, 2, 60-66.	1.2	54
500	A map for sequence analysis of the Arabidopsis thaliana genome. Nature Genetics, 1999, 22, 265-270.	9.4	134
501	An encyclopedia of mouse genes. Nature Genetics, 1999, 21, 191-194.	9.4	114
502	Genetic Definition and Sequence Analysis of Arabidopsis Centromeres. Science, 1999, 286, 2468-2474.	6.0	417
503	Functional Genomics in <i>Caenorhabditis elegans</i> Sequences from Related Nematodes. Genome Research, 1999, 9, 348-359.	2.4	45
504	Expressed sequence tags â€" ESTablishing bridges between genomes. Trends in Genetics, 1998, 14, 4-7.	2.9	121

#	Article	IF	CITATIONS
505	Gene Discovery by EST Sequencing in∢i>Toxoplasma gondii∢/i>Reveals Sequences Restricted to the Apicomplexa. Genome Research, 1998, 8, 18-28.	2.4	179
506	High Throughput Fingerprint Analysis of Large-Insert Clones. Genome Research, 1997, 7, 1072-1084.	2.4	405
507	Recovery of duplications by drug resistance selection in <i>Caenorhabditis elegans</i> . Genome, 1994, 37, 701-705.	0.9	2
508	The Caenorhabditis elegans unc-60 gene encodes proteins homologous to a family of actin-binding proteins. Molecular Genetics and Genomics, 1994, 242, 346-357.	2.4	113
509	Molecular analysis of two genes between let-653 and let-56 in the unc 22(IV) region of Caenorhabditis elegans. Molecular Genetics and Genomics, 1993, 236-236, 289-298.	2.4	32
510	The use of deficiencies to determine essential gene content in the let-56–unc-22 region of Caenorhabditis elegans. Genome, 1993, 36, 1148-1156.	0.9	5
511	Exceptional response to combination ipilimumab and nivolumab in metastatic uveal melanoma: Insights from genomic analysis. Melanoma Research, 0, Publish Ahead of Print, .	0.6	4
512	Automated Library Construction and Analysis for High-throughput Nanopore Sequencing of SARS-CoV-2. journal of applied laboratory medicine, The, 0, , .	0.6	0
513	Genome-wide detection of imprinted differentially methylated regions using nanopore sequencing. ELife, 0, 11 , .	2.8	21