Paul S Meltzer

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7642410/publications.pdf

Version: 2024-02-01

2509 2975 42,094 321 93 196 citations h-index g-index papers 339 339 339 46670 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Classification and diagnostic prediction of cancers using gene expression profiling and artificial neural networks. Nature Medicine, 2001, 7, 673-679.	30.7	2,352
2	Mutations of a mutS homolog in hereditary nonpolyposis colorectal cancer. Cell, 1993, 75, 1215-1225.	28.9	2,195
3	Vascular Channel Formation by Human Melanoma Cells in Vivo and in Vitro: Vasculogenic Mimicry. American Journal of Pathology, 1999, 155, 739-752.	3.8	1,705
4	Gene-Expression Profiles in Hereditary Breast Cancer. New England Journal of Medicine, 2001, 344, 539-548.	27.0	1,669
5	Rare Structural Variants Disrupt Multiple Genes in Neurodevelopmental Pathways in Schizophrenia. Science, 2008, 320, 539-543.	12.6	1,654
6	High frequency of BRAF mutations in nevi. Nature Genetics, 2003, 33, 19-20.	21.4	1,547
7	Expression profiling using cDNA microarrays. Nature Genetics, 1999, 21, 10-14.	21.4	1,529
8	AIB1, a Steroid Receptor Coactivator Amplified in Breast and Ovarian Cancer. Science, 1997, 277, 965-968.	12.6	1,514
9	High-Resolution Mapping andÂCharacterization of Open Chromatin across the Genome. Cell, 2008, 132, 311-322.	28.9	1,246
10	Mutations in the human Jagged1 gene are responsible for Alagille syndrome. Nature Genetics, 1997, 16, 235-242.	21.4	1,072
11	MicroRNA Expression, Survival, and Response to Interferon in Liver Cancer. New England Journal of Medicine, 2009, 361, 1437-1447.	27.0	778
12	Pan-cancer genome and transcriptome analyses of 1,699 paediatric leukaemias and solid tumours. Nature, 2018, 555, 371-376.	27.8	649
13	Expression profiling identifies the cytoskeletal organizer ezrin and the developmental homeoprotein Six-1 as key metastatic regulators. Nature Medicine, 2004, 10, 175-181.	30.7	480
14	Functionally defined therapeutic targets in diffuse intrinsic pontine glioma. Nature Medicine, 2015, 21, 555-559.	30.7	473
15	Mechanisms of sarcoma development. Nature Reviews Cancer, 2003, 3, 685-694.	28.4	406
16	MYC-driven accumulation of 2-hydroxyglutarate is associated with breast cancer prognosis. Journal of Clinical Investigation, 2014, 124, 398-412.	8.2	348
17	Comparative genomic hybridization using oligonucleotide microarrays and total genomic DNA. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 17765-17770.	7.1	336
18	Small RNAs with big impacts. Nature, 2005, 435, 745-746.	27.8	324

#	Article	IF	CITATIONS
19	Common Molecular Subtypes Among Asian Hepatocellular Carcinoma and Cholangiocarcinoma. Cancer Cell, 2017, 32, 57-70.e3.	16.8	324
20	Fluorescent cDNA microarray hybridization reveals complexity and heterogeneity of cellular genotoxic stress responses. Oncogene, 1999, 18, 3666-3672.	5.9	314
21	Gene Expression Profiling of Human Sarcomas: Insights into Sarcoma Biology. Cancer Research, 2005, 65, 9226-9235.	0.9	312
22	Thyroid Hormone Regulation of Hepatic Genes in Vivo Detected by Complementary DNA Microarray. Molecular Endocrinology, 2000, 14, 947-955.	3.7	303
23	Molecular Subtypes of <i>KIT/PDGFRA</i> Wild-Type Gastrointestinal Stromal Tumors. JAMA Oncology, 2016, 2, 922.	7.1	291
24	Data management and analysis for gene expression arrays. Nature Genetics, 1998, 20, 19-23.	21.4	290
25	Succinate Dehydrogenase Mutation Underlies Global Epigenomic Divergence in Gastrointestinal Stromal Tumor. Cancer Discovery, 2013, 3, 648-657.	9.4	288
26	Interaction of the Glucocorticoid Receptor with the Chromatin Landscape. Molecular Cell, 2008, 29, 611-624.	9.7	285
27	Interferon- \hat{I}^3 links ultraviolet radiation to melanomagenesis in mice. Nature, 2011, 469, 548-553.	27.8	264
28	Rapid generation of region specific probes by chromosome microdissection and their application. Nature Genetics, 1992, 1, 24-28.	21.4	261
29	Melanoma mouse model implicates metabotropic glutamate signaling in melanocytic neoplasia. Nature Genetics, 2003, 34, 108-112.	21.4	260
30	Interaction between the microbiome and TP53 in human lung cancer. Genome Biology, 2018, 19, 123.	8.8	247
31	Sunitinib in patients with chemotherapy-refractory thymoma and thymic carcinoma: an open-label phase 2 trial. Lancet Oncology, The, 2015, 16, 177-186.	10.7	240
32	The Exomes of the NCI-60 Panel: A Genomic Resource for Cancer Biology and Systems Pharmacology. Cancer Research, 2013, 73, 4372-4382.	0.9	239
33	Cloning a novel member of the human interferon-inducible gene family associated with control of tumorigenicity in a model of human melanoma. Oncogene, 1997, 15, 453-457.	5.9	238
34	High prevalence of MAP2K1 mutations in variant and IGHV4-34–expressing hairy-cell leukemias. Nature Genetics, 2014, 46, 8-10.	21.4	236
35	Integrating Global Gene Expression and Radiation Survival Parameters across the 60 Cell Lines of the National Cancer Institute Anticancer Drug Screen. Cancer Research, 2008, 68, 415-424.	0.9	226
36	Soft tissue sarcomas of adults: state of the translational science. Clinical Cancer Research, 2003, 9, 1941-56.	7.0	224

#	Article	IF	CITATIONS
37	DNase-chip: a high-resolution method to identify DNase I hypersensitive sites using tiled microarrays. Nature Methods, 2006, 3, 503-509.	19.0	222
38	Activity of durvalumab plus olaparib in metastatic castration-resistant prostate cancer in men with and without DNA damage repair mutations., 2018, 6, 141.		214
39	New Horizons in the Treatment of Osteosarcoma. New England Journal of Medicine, 2021, 385, 2066-2076.	27.0	210
40	Pim-1 is up-regulated by constitutively activated FLT3 and plays a role in FLT3-mediated cell survival. Blood, 2005, 105, 1759-1767.	1.4	209
41	A specific missense mutation in GTF2I occurs at high frequency in thymic epithelial tumors. Nature Genetics, 2014, 46, 844-849.	21.4	208
42	DNA Breaks and End Resection Measured Genome-wide by End Sequencing. Molecular Cell, 2016, 63, 898-911.	9.7	206
43	Activating Signal Cointegrator 2 Belongs to a Novel Steady-State Complex That Contains a Subset of Trithorax Group Proteins. Molecular and Cellular Biology, 2003, 23, 140-149.	2.3	202
44	Point Mutations in Exon 1B of APC Reveal Gastric Adenocarcinoma and Proximal Polyposis of the Stomach as a Familial Adenomatous Polyposis Variant. American Journal of Human Genetics, 2016, 98, 830-842.	6.2	201
45	Transcription Program of Human Herpesvirus 8 (Kaposi's Sarcoma-Associated Herpesvirus). Journal of Virology, 2001, 75, 4843-4853.	3.4	198
46	Genome-Wide Identification of PAX3-FKHR Binding Sites in Rhabdomyosarcoma Reveals Candidate Target Genes Important for Development and Cancer. Cancer Research, 2010, 70, 6497-6508.	0.9	195
47	Genome-Wide Analysis of Menin Binding Provides Insights into MEN1 Tumorigenesis. PLoS Genetics, 2006, 2, e51.	3.5	193
48	Gene expression profile in multiple sclerosis patients and healthy controls: identifying pathways relevant to disease. Human Molecular Genetics, 2003, 12, 2191-2199.	2.9	191
49	A Nuclear Factor, ASC-2, as a Cancer-amplified Transcriptional Coactivator Essential for Ligand-dependent Transactivation by Nuclear Receptors in Vivo. Journal of Biological Chemistry, 1999, 274, 34283-34293.	3.4	190
50	Analyses of Resected Human Brain Metastases of Breast Cancer Reveal the Association between Up-Regulation of Hexokinase 2 and Poor Prognosis. Molecular Cancer Research, 2009, 7, 1438-1445.	3.4	185
51	A Polymorphism in IRF4 Affects Human Pigmentation through a Tyrosinase-Dependent MITF/TFAP2A Pathway. Cell, 2013, 155, 1022-1033.	28.9	184
52	Molecular classification of familial non- <i>BRCA1/BRCA2</i> breast cancer. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 2532-2537.	7.1	182
53	Specific Chromosomal Aberrations and Amplification of the AIB1 Nuclear Receptor Coactivator Gene in Pancreatic Carcinomas. American Journal of Pathology, 1999, 154, 525-536.	3.8	181
54	Genome-wide association study identifies two susceptibility loci for osteosarcoma. Nature Genetics, 2013, 45, 799-803.	21.4	181

#	Article	IF	CITATIONS
55	Immunohistochemical Loss of Succinate Dehydrogenase Subunit A (SDHA) in Gastrointestinal Stromal Tumors (GISTs) Signals SDHA Germline Mutation. American Journal of Surgical Pathology, 2013, 37, 234-240.	3.7	178
56	Differential responses of stress genes to low dose-rate gamma irradiation. Molecular Cancer Research, 2003, 1, 445-52.	3.4	177
57	Expression Profiling of Synovial Sarcoma by cDNA Microarrays. American Journal of Pathology, 2002, 161, 1587-1595.	3.8	173
58	Microarray expression profiling in melanoma reveals a BRAF mutation signature. Oncogene, 2004, 23, 4060-4067.	5.9	169
59	Evidence for an Unanticipated Relationship between Undifferentiated Pleomorphic Sarcoma and Embryonal Rhabdomyosarcoma. Cancer Cell, 2011, 19, 177-191.	16.8	167
60	Telomere capture stabilizes chromosome breakage. Nature Genetics, 1993, 4, 252-255.	21.4	160
61	Recurrent epimutation of <i>SDHC</i> in gastrointestinal stromal tumors. Science Translational Medicine, 2014, 6, 268ra177.	12.4	158
62	A Molecular Function Map of Ewing's Sarcoma. PLoS ONE, 2009, 4, e5415.	2.5	158
63	Expression profiling in cancer using cDNA microarrays. Electrophoresis, 1999, 20, 223-229.	2.4	157
64	TRAIL induces apoptosis in triple-negative breast cancer cells with a mesenchymal phenotype. Breast Cancer Research and Treatment, 2009, 113, 217-230.	2.5	157
65	Molecular determinants of human uveal melanoma invasion and metastasis. Clinical and Experimental Metastasis, 2002, 19, 233-246.	3.3	149
66	Molecular cytogenetic analysis of i(12p)-negative human male germ cell tumors. Genes Chromosomes and Cancer, 1993, 8 , $230-236$.	2.8	141
67	A genome-based strategy uncovers frequent BRAF mutations in melanoma. Cancer Cell, 2002, 2, 5-7.	16.8	139
68	Identification of cryptic sites of DNA sequence amplification in human breast cancer by chromosome microdissection. Nature Genetics, 1994, 8, 155-161.	21.4	137
69	Methylation profiling of mediastinal gray zone lymphoma reveals a distinctive signature with elements shared by classical Hodgkin's lymphoma and primary mediastinal large B-cell lymphoma. Haematologica, 2011, 96, 558-566.	3.5	135
70	Biology of childhood osteogenic sarcoma and potential targets for therapeutic development: meeting summary. Clinical Cancer Research, 2003, 9, 5442-53.	7.0	135
71	Multivariate Measurement of Gene Expression Relationships. Genomics, 2000, 67, 201-209.	2.9	133
72	Expression of multiple molecular phenotypes by aggressive melanoma tumor cells: role in vasculogenic mimicry. Critical Reviews in Oncology/Hematology, 2002, 44, 17-27.	4.4	132

#	Article	IF	Citations
73	Vorinostat Inhibits Brain Metastatic Colonization in a Model of Triple-Negative Breast Cancer and Induces DNA Double-Strand Breaks. Clinical Cancer Research, 2009, 15, 6148-6157.	7.0	132
74	Stress-specific signatures: expression profiling of p53 wild-type and -null human cells. Oncogene, 2005, 24, 4572-4579.	5.9	131
75	Toward a Drug Development Path That Targets Metastatic Progression in Osteosarcoma. Clinical Cancer Research, 2014, 20, 4200-4209.	7.0	127
76	Positively selected enhancer elements endow osteosarcoma cells with metastatic competence. Nature Medicine, 2018, 24, 176-185.	30.7	126
77	Gene-target recognition among members of the Myc superfamily and implications for oncogenesis. Nature Genetics, 2000, 24, 113-119.	21.4	125
78	Array comparative genomic hybridization-based characterization of genetic alterations in pulmonary neuroendocrine tumors. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 13040-13045.	7.1	123
79	Characterization of the 12q13-15 amplicon in soft tissue tumors. Cancer Genetics and Cytogenetics, 1995, 83, 32-36.	1.0	119
80	Expression of the cytoskeleton linker protein ezrin in human cancers. Clinical and Experimental Metastasis, 2007, 24, 69-78.	3.3	118
81	Generation of band-specific painting probes from a single microdissected chromosome. Human Molecular Genetics, 1993, 2, 1117-1121.	2.9	116
82	Rapid Generation of Whole Chromosome Painting Probes (WCPs) by Chromosome Microdissection. Genomics, 1994, 22, 101-107.	2.9	115
83	Loss-of-Function Fibroblast Growth Factor Receptor-2 Mutations in Melanoma. Molecular Cancer Research, 2009, 7, 41-54.	3.4	112
84	Genome-wide depletion of replication initiation events in highly transcribed regions. Genome Research, 2011, 21, 1822-1832.	5. 5	112
85	Effects of ligand and thyroid hormone receptor isoforms on hepatic gene expression profiles of thyroid hormone receptor knockout mice. EMBO Reports, 2003, 4, 581-587.	4.5	110
86	Nm23-H1 Suppresses Tumor Cell Motility by Down-regulating the Lysophosphatidic Acid Receptor <i>EDG2</i> . Cancer Research, 2007, 67, 7238-7246.	0.9	110
87	Genome-wide expression changes induced by HTLV-1 Tax: evidence for MLK-3 mixed lineage kinase involvement in Tax-mediated NF-κB activation. Oncogene, 2001, 20, 4484-4496.	5.9	109
88	Transcriptional activation by the thyroid hormone receptor through ligand-dependent receptor recruitment and chromatin remodelling. Nature Communications, 2015, 6, 7048.	12.8	106
89	General nonlinear framework for the analysis of gene interaction via multivariate expression arrays. Journal of Biomedical Optics, 2000, 5, 411.	2.6	104
90	An Integrated Prognostic Classifier for Stage I Lung Adenocarcinoma Based on mRNA, microRNA, and DNA Methylation Biomarkers. Journal of Thoracic Oncology, 2015, 10, 1037-1048.	1.1	103

#	Article	IF	Citations
91	Separate amplified regions encompassing CDK4 and MDM2 in human sarcomas., 1996, 17, 254-259.		102
92	An Unliganded Thyroid Hormone \hat{l}^2 Receptor Activates the Cyclin D1/Cyclin-Dependent Kinase/Retinoblastoma/E2F Pathway and Induces Pituitary Tumorigenesis. Molecular and Cellular Biology, 2005, 25, 124-135.	2.3	100
93	Oncogenic ETS fusions deregulate E2F3 target genes in Ewing sarcoma and prostate cancer. Genome Research, 2013, 23, 1797-1809.	5.5	99
94	DNA Methylation Profiling Identifies Global Methylation Differences and Markers of Adrenocortical Tumors. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E1004-E1013.	3.6	98
95	Clinical differentiation between proteus syndrome and hemihyperplasia: Description of a distinct form of hemihyperplasia., 1998, 79, 311-318.		97
96	Archival Fine-Needle Aspiration Cytopathology (FNAC) Samples. Journal of Molecular Diagnostics, 2010, 12, 739-745.	2.8	97
97	Harnessing synthetic lethality to predict the response to cancer treatment. Nature Communications, 2018, 9, 2546.	12.8	97
98	Repeat expansions confer WRN dependence in microsatellite-unstable cancers. Nature, 2020, 586, 292-298.	27.8	95
99	Data analysis and integration: of steps and arrows. Nature Genetics, 1999, 22, 213-215.	21.4	93
100	Disease fingerprinting with cDNA microarrays reveals distinct gene expression profiles in lethal typeâ€1 and typeâ€2 cytokineâ€mediated inflammatory reactions. FASEB Journal, 2001, 15, 2545-2547.	0.5	92
101	A p21-ZEB1 Complex Inhibits Epithelial-Mesenchymal Transition through the MicroRNA 183-96-182 Cluster. Molecular and Cellular Biology, 2014, 34, 533-550.	2.3	92
102	Imputation and subset-based association analysis across different cancer types identifies multiple independent risk loci in the TERT-CLPTM1L region on chromosome 5p15.33. Human Molecular Genetics, 2014, 23, 6616-6633.	2.9	90
103	Focus on sarcomas. Cancer Cell, 2002, 2, 175-178.	16.8	89
104	<i>SLFN11</i> Is a Transcriptional Target of EWS-FLI1 and a Determinant of Drug Response in Ewing Sarcoma. Clinical Cancer Research, 2015, 21, 4184-4193.	7.0	89
105	Characterization of the metastatic phenotype of a panel of established osteosarcoma cells. Oncotarget, 2015, 6, 29469-29481.	1.8	89
106	A Genome-Wide Scan Identifies Variants in <i>NFIB</i> Associated with Metastasis in Patients with Osteosarcoma. Cancer Discovery, 2015, 5, 920-931.	9.4	88
107	Separate and variably shaped chromosome arm domains are disclosed by chromosome arm painting in human cell nuclei. Chromosome Research, 1998, 6, 25-33.	2.2	87
108	Lineage of origin in rhabdomyosarcoma informs pharmacological response. Genes and Development, 2014, 28, 1578-1591.	5.9	87

#	Article	IF	CITATIONS
109	Chromothripsis and Focal Copy Number Alterations Determine Poor Outcome in Malignant Melanoma. Cancer Research, 2013, 73, 1454-1460.	0.9	86
110	Provocative questions in osteosarcoma basic and translational biology: A report from the Children's Oncology Group. Cancer, 2019, 125, 3514-3525.	4.1	86
111	SCLC-CellMiner: A Resource for Small Cell Lung Cancer Cell Line Genomics and Pharmacology Based on Genomic Signatures. Cell Reports, 2020, 33, 108296.	6.4	86
112	The Synovial Sarcoma–Associated SS18-SSX2 Fusion Protein Induces Epigenetic Gene (De)Regulation. Cancer Research, 2006, 66, 9474-9482.	0.9	85
113	Antagonistic Cross-Regulation between Sox9 and Sox10 Controls an Anti-tumorigenic Program in Melanoma. PLoS Genetics, 2015, 11, e1004877.	3 . 5	85
114	Lipid defect underlies selective skin barrier impairment of an epidermal-specific deletion of Gata-3. Journal of Cell Biology, 2006, 175, 661-670.	5. 2	80
115	Unfavorable prognosis of <i>CRTC1â€MAML2</i> positive mucoepidermoid tumors with <i>CDKN2A</i> deletions. Genes Chromosomes and Cancer, 2010, 49, 59-69.	2.8	80
116	Nonrandom chromosome alterations in rhabdomyosarcoma. Cancer Genetics and Cytogenetics, 1985, 16, 189-197.	1.0	79
117	CDC91L1 (PIG-U) is a newly discovered oncogene in human bladder cancer. Nature Medicine, 2004, 10, 374-381.	30.7	79
118	Suppressor role of activating transcription factor 2 (ATF2) in skin cancer. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 1674-1679.	7.1	78
119	Twelve amplified and expressed genes localized in a single domain in glioma. Human Genetics, 1996, 98, 625-628.	3.8	77
120	A chromatin structureâ€based model accurately predicts <scp>DNA</scp> replication timing in human cells. Molecular Systems Biology, 2014, 10, 722.	7.2	77
121	The gene expression response of breast cancer to growth regulators: patterns and correlation with tumor expression profiles. Cancer Research, 2003, 63, 7158-66.	0.9	75
122	Thyroid Hormone Responsive Genes in Cultured Human Fibroblasts. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 936-943.	3.6	74
123	Translocation 1;7 in hematologic disorders: A brief review of 22 cases. Cancer Genetics and Cytogenetics, 1985, 18, 207-213.	1.0	72
124	Rapid generation of region-specific genomic clones by chromosome microdissection: Isolation of DNA from a region frequently deleted in malignant melanoma. Genomics, 1992, 14, 680-684.	2.9	71
125	Generation and analysis of melanoma SAGE libraries: SAGE advice on the melanoma transcriptome. Oncogene, 2004, 23, 2264-2274.	5.9	71
126	Alterations in genomic profiles during tumor progression in a mouse model of follicular thyroid carcinoma. Carcinogenesis, 2003, 24, 1467-1479.	2.8	68

#	Article	IF	CITATIONS
127	Identification and Cloning of the Human Homolog (JAG1) of the RatJagged1Gene from the Alagille Syndrome Critical Region at 20p12. Genomics, 1997, 43, 376-379.	2.9	67
128	Biological indicators for the identification of ionizing radiation exposure in humans. Expert Review of Molecular Diagnostics, 2001, 1, 211-219.	3.1	66
129	NUP98–PHF23 Is a Chromatin-Modifying Oncoprotein That Causes a Wide Array of Leukemias Sensitive to Inhibition of PHD Histone Reader Function. Cancer Discovery, 2014, 4, 564-577.	9.4	66
130	Hypoxia Modulates EWS-FLI1 Transcriptional Signature and Enhances the Malignant Properties of Ewing's Sarcoma Cells <i>In vitro</i> Cancer Research, 2010, 70, 4015-4023.	0.9	65
131	Melanoblast transcriptome analysis reveals pathways promoting melanoma metastasis. Nature Communications, 2020, 11, 333.	12.8	65
132	Phenol oxidase activation in Drosophila: A cascade of reactions. Biochemical Genetics, 1975, 13, 85-108.	1.7	64
133	Genomic aberrations in pediatric diffuse intrinsic pontine gliomas. Neuro-Oncology, 2012, 14, 326-332.	1.2	62
134	Discovery and validation of methylation markers for endometrial cancer. International Journal of Cancer, 2014, 135, 1860-1868.	5.1	62
135	Microarray analysis of knockout mice identifies cyclin D2 as a possible mediator for the action of thyroid hormone during the postnatal development of the cerebellum. Developmental Biology, 2003, 254, 188-199.	2.0	61
136	Expression and Mutational Status of c-kit in Thymic Epithelial Tumors. Journal of Thoracic Oncology, 2010, 5, 1447-1453.	1.1	61
137	Mutant thyroid hormone receptor beta represses the expression and transcriptional activity of peroxisome proliferator-activated receptor gamma during thyroid carcinogenesis. Cancer Research, 2003, 63, 5274-80.	0.9	61
138	Molecular Grading of Ductal Carcinoma <i>In situ</i> of the Breast. Clinical Cancer Research, 2008, 14, 8244-8252.	7.0	60
139	RNA Sequencing of the NCI-60: Integration into CellMiner and CellMiner CDB. Cancer Research, 2019, 79, 3514-3524.	0.9	58
140	Differentially Painting Human Chromosome Arms with Combined Binary Ratio-labeling Fluorescence In Situ Hybridization. Genome Research, 2000, 10, 861-865.	5.5	56
141	ATP11B mediates platinum resistance in ovarian cancer. Journal of Clinical Investigation, 2013, 123, 2119-2130.	8.2	56
142	Genome-Wide Methylation Patterns in Papillary Thyroid Cancer Are Distinct Based on Histological Subtype and Tumor Genotype. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E329-E337.	3.6	55
143	The second European interdisciplinary Ewing sarcoma research summit - A joint effort to deconstructing the multiple layers of a complex disease. Oncotarget, 2016, 7, 8613-8624.	1.8	55
144	Assessment of Automated Image Analysis of Breast Cancer Tissue Microarrays for Epidemiologic Studies. Cancer Epidemiology Biomarkers and Prevention, 2010, 19, 992-999.	2.5	54

#	Article	IF	CITATIONS
145	Molecular cytogenetic characterization and physical mapping of 12q13–15 amplification in human cancers. Genes Chromosomes and Cancer, 1996, 17, 205-214.	2.8	53
146	Spotting the target: microarrays for disease gene discovery. Current Opinion in Genetics and Development, 2001, 11, 258-263.	3.3	51
147	Related subunits of NF-κB map to two distinct loci associated with translocations in leukemia, NFKB1 and NFKB2. Genomics, 1992, 13, 287-292.	2.9	50
148	Complete sequence analysis of a gene (OS-9) ubiquitously expressed in human tissues and amplified in sarcomas. Molecular Carcinogenesis, 1996, 15, 270-275.	2.7	50
149	Transgenic Targeting of a Dominant Negative Corepressor to Liver Blocks Basal Repression by Thyroid Hormone Receptor and Increases Cell Proliferation. Journal of Biological Chemistry, 2001, 276, 15066-15072.	3.4	49
150	The NCI-60 Methylome and Its Integration into CellMiner. Cancer Research, 2017, 77, 601-612.	0.9	48
151	Preferential Localization of Human Origins of DNA Replication at the 5′-Ends of Expressed Genes and at Evolutionarily Conserved DNA Sequences. PLoS ONE, 2011, 6, e17308.	2.5	47
152	Alpha-globulins suppress human leukocyte tumor necrosis factor secretion. European Journal of Immunology, 1989, 19, 939-942.	2.9	46
153	Verapamil suppresses the emergence of P-glycoprotein-mediated multi-drug resistance. , 1996, 66, 520-525.		46
154	Large-Scale Profiling of Archival Lymph Nodes Reveals Pervasive Remodeling of the Follicular Lymphoma Methylome. Cancer Research, 2009, 69, 758-764.	0.9	46
155	Imprints and <i>DPPA3</i> are bypassed during pluripotency- and differentiation-coupled methylation reprogramming in testicular germ cell tumors. Genome Research, 2016, 26, 1490-1504.	5.5	44
156	Characterization of a highly conserved gene (OS4) amplified with CDK4 in human sarcomas. Oncogene, 1997, 15, 1289-1294.	5.9	43
157	A small protein encoded by a putative lncRNA regulates apoptosis and tumorigenicity in human colorectal cancer cells. ELife, 2020, 9, .	6.0	43
158	Bromodomain and Extraterminal Protein Inhibitor JQ1 Suppresses Thyroid Tumor Growth in a Mouse Model. Clinical Cancer Research, 2017, 23, 430-440.	7.0	42
159	Biomarker significance of plasma and tumor miR-21, miR-221, and miR-106a in osteosarcoma. Oncotarget, 2017, 8, 96738-96752.	1.8	41
160	Role of glutathione and its associated enzymes in multidrug-resistant human myeloma cells. Biochemical Pharmacology, 1989, 38, 787-793.	4.4	40
161	Detection of Antisense and Ribozyme Accessible Sites on Native mRNAs: Application to NCOA3 mRNA. Molecular Therapy, 2001, 4, 454-460.	8.2	39
162	Inhibition of Polo-like kinase 1 prevents the growth of metastatic breast cancer cells in the brain. Clinical and Experimental Metastasis, 2011, 28, 899-908.	3.3	39

#	Article	IF	CITATIONS
163	EWS-FLI1 employs an E2F switch to drive target gene expression. Nucleic Acids Research, 2015, 43, 2780-2789.	14.5	39
164	Copy Number Aberrations of Genes Regulating Normal Thymus Development in Thymic Epithelial Tumors. Clinical Cancer Research, 2013, 19, 1960-1971.	7.0	38
165	Distinct methylation profiles characterize fusion-positive and fusion-negative rhabdomyosarcoma. Modern Pathology, 2015, 28, 1214-1224.	5.5	38
166	Current state of pediatric sarcoma biology and opportunities for future discovery: A report from the sarcoma translational research workshop. Cancer Genetics, 2016, 209, 182-194.	0.4	38
167	Epigenome-wide DNA methylation analysis of small cell lung cancer cell lines suggests potential chemotherapy targets. Clinical Epigenetics, 2020, 12, 93.	4.1	38
168	NCI-60 Whole Exome Sequencing and Pharmacological CellMiner Analyses. PLoS ONE, 2014, 9, e101670.	2.5	38
169	Chromosome arm-specific multicolor FISH. Genes Chromosomes and Cancer, 2001, 30, 105-109.	2.8	37
170	Advanced Bone Formation in Mice with a Dominant-negative Mutation in the Thyroid Hormone Receptor \hat{l}^2 Gene due to Activation of Wnt/ \hat{l}^2 -Catenin Protein Signaling. Journal of Biological Chemistry, 2012, 287, 17812-17822.	3.4	37
171	G-Cimp Status Prediction Of Glioblastoma Samples Using mRNA Expression Data. PLoS ONE, 2012, 7, e47839.	2.5	37
172	Transcript Mapping in a 46-kb Sequenced Region at the Core of 12q13.3 Amplification in Human Cancers. Genomics, 1997, 42, 295-301.	2.9	35
173	A combined approach identifies a limited number of new thyroid hormone target genes in post-natal mouse cerebellum. Journal of Molecular Endocrinology, 2007, 39, 17-28.	2.5	35
174	Identification of RECQ1-regulated transcriptome uncovers a role of RECQ1 in regulation of cancer cell migration and invasion. Cell Cycle, 2014, 13, 2431-2445.	2.6	35
175	Targeting loss of the Hippo signaling pathway in <i>NF2</i> deficient papillary kidney cancers. Oncotarget, 2018, 9, 10723-10733.	1.8	35
176	Lucky draw in the gene raffle. Nature, 2002, 417, 906-907.	27.8	34
177	Constitutive Fmsâ€like tyrosine kinase 3 activation results in specific changes in gene expression in myeloid leukaemic cells. British Journal of Haematology, 2007, 138, 603-615.	2.5	34
178	Targeting MYC as a Therapeutic Intervention for Anaplastic Thyroid Cancer. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 2268-2280.	3.6	34
179	Rapid and reversible suppression of ALT by DAXX in osteosarcoma cells. Scientific Reports, 2019, 9, 4544.	3.3	34
180	Alleles of the Estrogen Receptor \hat{l}_{\pm} -Gene and an Estrogen Receptor Cotranscriptional Activator Gene, Amplified in Breast Cancer-1 (AlB1), Are Associated with Quantitative Calcaneal Ultrasound. Journal of Bone and Mineral Research, 2000, 15, 2231-2239.	2.8	33

#	Article	IF	Citations
181	Modulation of transcriptional sensitivity of mineralocorticoid and estrogen receptors. Journal of Steroid Biochemistry and Molecular Biology, 2004, 91, 197-210.	2.5	33
182	Post-Transcriptional Dysregulation by miRNAs Is Implicated in the Pathogenesis of Gastrointestinal Stromal Tumor [GIST]. PLoS ONE, 2013, 8, e64102.	2.5	33
183	Micro-Environment Causes Reversible Changes in DNA Methylation and mRNA Expression Profiles in Patient-Derived Glioma Stem Cells. PLoS ONE, 2014, 9, e94045.	2.5	33
184	Detection of Gains and Losses in 18 Meningiomas by Comparative Genomic Hybridization. Cancer Genetics and Cytogenetics, 1998, 103, 95-100.	1.0	32
185	Negative Regulation of TSHα Target Gene by Thyroid Hormone Involves Histone Acetylation and Corepressor Complex Dissociation. Molecular Endocrinology, 2009, 23, 600-609.	3.7	31
186	Coverage of chromosome 6 by chromosome microdissection: generation of 14 subregion-specific probes. Human Genetics, 1995, 95, 637-40.	3.8	30
187	Effects of HIV-1 nef on cellular gene expression profiles. Journal of Biomedical Science, 2002, 9, 82-96.	7.0	30
188	Association of steroid receptor coactivator AIB1 with estrogen receptor- \hat{l}_{\pm} in breast cancer cells. Breast Cancer Research and Treatment, 2001, 70, 89-101.	2.5	29
189	Mutation-targeted therapy with sunitinib or everolimus in patients with advanced low-grade or intermediate-grade neuroendocrine tumours of the gastrointestinal tract and pancreas with or without cytoreductive surgery: protocol for a phase II clinical trial. BMJ Open, 2015, 5, e008248-e008248.	1.9	29
190	M6A RNA Methylation Regulates Histone Ubiquitination to Support Cancer Growth and Progression. Cancer Research, 2022, 82, 1872-1889.	0.9	29
191	Molecular mapping of the oncogeneMYB and rearrangements in malignant melanoma. Genes Chromosomes and Cancer, 1989, 1, 88-94.	2.8	28
192	Identification of genes differentially regulated by the P210 BCR/ABL1 fusion oncogene using cDNA microarrays. Experimental Hematology, 2004, 32, 476-482.	0.4	28
193	Epigenetic and genetic inactivation of tyrosyl-DNA-phosphodiesterase 1 (TDP1) in human lung cancer cells from the NCI-60 panel. DNA Repair, 2014, 13, 1-9.	2.8	28
194	miR-23a impairs bone differentiation in osteosarcoma via down-regulation of GJA1. Frontiers in Genetics, 2015, 6, 233.	2.3	28
195	Integrated genome-wide analysis of genomic changes and gene regulation in human adrenocortical tissue samples. Nucleic Acids Research, 2015, 43, 9327-9339.	14.5	28
196	Skp, Cullin, F-box (SCF)-Met30 and SCF-Cdc4-Mediated Proteolysis of CENP-A Prevents Mislocalization of CENP-A for Chromosomal Stability in Budding Yeast. PLoS Genetics, 2020, 16, e1008597.	3.5	28
197	Isolation of genes amplified in human cancers by microdissection mediated cDNA capture. Human Molecular Genetics, 1996, 5, 595-600.	2.9	27
198	Aneuploidy, <i>TP53</i> mutation, and amplification of <i>MYC</i> correlate with increased intratumor heterogeneity and poor prognosis of breast cancer patients. Genes Chromosomes and Cancer, 2018, 57, 165-175.	2.8	27

#	Article	IF	Citations
199	ATP11B mediates platinum resistance in ovarian cancer. Journal of Clinical Investigation, 2018, 128, 3199-3199.	8.2	27
200	Epidermal growth factor receptor overexpression and trisomy 7 in a case of Barrett's esophagus. Digestive Diseases and Sciences, 1990, 35, 1115-1120.	2.3	26
201	Outcomes of Children and Adolescents with Advanced Hereditary Medullary Thyroid Carcinoma Treated with Vandetanib. Clinical Cancer Research, 2018, 24, 753-765.	7.0	26
202	Immuno-transcriptomic profiling of extracranial pediatric solid malignancies. Cell Reports, 2021, 37, 110047.	6.4	26
203	Gene expression profiles in a panel of childhood leukemia cell lines mirror critical features of the disease. Molecular Cancer Therapeutics, 2003, 2, 671-7.	4.1	26
204	The role of mutation of metabolism-related genes in genomic hypermethylation. Biochemical and Biophysical Research Communications, 2014, 455, 16-23.	2.1	25
205	Is Ewing's Sarcoma a Stem Cell Tumor?. Cell Stem Cell, 2007, 1, 13-15.	11.1	24
206	Integrated high-resolution array CGH and SKY analysis of homozygous deletions and other genomic alterations present in malignant mesothelioma cell lines. Cancer Genetics, 2013, 206, 191-205.	0.4	23
207	Deposition of Centromeric Histone H3 Variant CENP-A/Cse4 into Chromatin Is Facilitated by Its C-Terminal Sumoylation. Genetics, 2020, 214, 839-854.	2.9	23
208	A Novel Chordoma Xenograft Allows In Vivo Drug Testing and Reveals the Importance of NF-κB Signaling in Chordoma Biology. PLoS ONE, 2013, 8, e79950.	2.5	23
209	Predicting continuous values of prognostic markers in breast cancer from microarray gene expression profiles. Molecular Cancer Therapeutics, 2004, 3, 161-8.	4.1	23
210	Molecular cloning of human hic-5, a potential regulator involved in signal transduction and cellular senescence. Molecular Carcinogenesis, 2000, 27, 177-183.	2.7	22
211	SAGE Identification and Fluorescence Imaging Analysis of Genes and Transcripts in Melanomas and Precursor Lesions. Cancer Biology and Therapy, 2004, 3, 104-109.	3.4	22
212	Zoom-in comparative genomic hybridisation arrays for the characterisation of variable breakpoint contiguous gene syndromes. Journal of Medical Genetics, 2006, 44, e59-e59.	3.2	22
213	Targeting Notch1 and IKKα Enhanced NF-κB Activation in CD133+ Skin Cancer Stem Cells. Molecular Cancer Therapeutics, 2018, 17, 2034-2048.	4.1	22
214	Dynamics of Genome Alterations in Crohn's Disease–Associated Colorectal Carcinogenesis. Clinical Cancer Research, 2018, 24, 4997-5011.	7.0	22
215	Recurrent PTPRT/JAK2 mutations in lung adenocarcinoma among African Americans. Nature Communications, 2019, 10, 5735.	12.8	22
216	A PAC Containing the Human Mitochondrial DNA Polymerase Gamma Gene (POLG) Maps to Chromosome 15q25. Genomics, 1997, 40, 376-378.	2.9	21

#	Article	IF	CITATIONS
217	Application of molecular cytogenetic techniques in a case study of human cutaneous metastatic melanoma. Cancer Genetics and Cytogenetics, 2001, 131, 97-103.	1.0	21
218	Loss of 18q22.3 Involving the Carboxypeptidase of Glutamate-like Gene Is Associated with Poor Prognosis in Resected Pancreatic Cancer. Clinical Cancer Research, 2012, 18, 524-533.	7.0	21
219	A Circular RNA from the <i>MDM2</i> Locus Controls Cell Cycle Progression by Suppressing p53 Levels. Molecular and Cellular Biology, 2020, 40, .	2.3	21
220	Whole Genome Sequencing of Newly Established Pancreatic Cancer Lines Identifies Novel Somatic Mutation (c.2587G>A) in Axon Guidance Receptor Plexin A1 as Enhancer of Proliferation and Invasion. PLoS ONE, 2016, 11, e0149833.	2.5	21
221	Isolation of a Cosmid Sublibrary for a Region of Chromosome 12 Frequently Amplified in Human Cancers Using a Complex Chromosome Microdissection Probe. Genomics, 1996, 31, 343-347.	2.9	20
222	Structure and Promoter Analysis of the Humanunc-33-like Phosphoprotein Gene. Journal of Biological Chemistry, 2000, 275, 16560-16568.	3.4	20
223	A Genome-Wide Screen Reveals a Role for the HIR Histone Chaperone Complex in Preventing Mislocalization of Budding Yeast CENP-A. Genetics, 2018, 210, 203-218.	2.9	20
224	CDC73 Germline Mutation in a Family With Mixed Epithelial and Stromal Tumors. Urology, 2019, 124, 91-97.	1.0	20
225	Qualitative changes in the basic protein fraction of developing embryos. Developmental Biology, 1970, 21, 73-86.	2.0	19
226	Development and utilization of a somatic cell hybrid mapping panel to assign Notl linking probes to the long arm of human chromosome 6. Genomics, 1992, 12, 542-548.	2.9	19
227	Cloning and tissue expression of the mouse ortholog of AIM1, a $\hat{l}^2\hat{l}^3$ -crystallin superfamily member. Mammalian Genome, 1998, 9, 715-720.	2.2	19
228	Progenitor B-1 B-cell acute lymphoblastic leukemia is associated with collaborative mutations in 3 critical pathways. Blood Advances, 2017, 1, 1749-1759.	5.2	19
229	Immunohistochemical detection of PAX-FOXO1 fusion proteins in alveolar rhabdomyosarcoma using breakpoint specific monoclonal antibodies. Modern Pathology, 2021, 34, 748-757.	5.5	19
230	Detection of Novel Amplicons in Prostate Cancer by Comprehensive Genomic Profiling of Prostate Cancer Cell Lines Using Oligonucleotide-Based ArrayCGH. PLoS ONE, 2007, 2, e769.	2.5	18
231	Patterns of somatic uniparental disomy identify novel tumor suppressor genes in colorectal cancer. Carcinogenesis, 2015, 36, 1103-1110.	2.8	18
232	Genomic profiling of multiple sequentially acquired tumor metastatic sites from an "exceptional responder―lung adenocarcinoma patient reveals extensive genomic heterogeneity and novel somatic variants driving treatment response. Journal of Physical Education and Sports Management, 2016, 2, a001263.	1,2	18
233	Genomic structure of SAS, a member of the transmembrane 4 superfamily amplified in human sarcomas. Genomics, 1995, 25, 501-506.	2.9	17
234	Modeling Synovial Sarcoma: Timing Is Everything. Cancer Cell, 2007, 11, 305-307.	16.8	17

#	Article	IF	Citations
235	Genetic Diversity in Melanoma. New England Journal of Medicine, 2005, 353, 2104-2107.	27.0	15
236	T7-based linear amplification of low concentration mRNA samples using beads and microfluidics for global gene expression measurements. Lab on A Chip, 2009, 9, 917-924.	6.0	15
237	A Methyl-Deviator Epigenotype of Estrogen Receptor–Positive Breast Carcinoma Is Associated with Malignant Biology. American Journal of Pathology, 2011, 179, 55-65.	3.8	15
238	Broader utilization of origins of DNA replication in cancer cell lines along a 78 kb region of human chromosome 2q34. Journal of Cellular Biochemistry, 2012, 113, 132-140.	2.6	15
239	Positional Cloning Utilizing Genomic DNA Microarrays: The Niemann–Pick Type C Gene as a Model System. Molecular Genetics and Metabolism, 2000, 70, 10-18.	1.1	14
240	Genomic and metabolic characterization of a chromophobe renal cell carcinoma cell line model (UOK276). Genes Chromosomes and Cancer, 2017, 56, 719-729.	2.8	14
241	Gene amplification in cancer: A molecular cytogenetic approach. Cancer Genetics and Cytogenetics, 1986, 19, 93-99.	1.0	13
242	Gene amplification elucidated by combined chromosomal microdissection and comparative genomic hybridization. Cancer Genetics and Cytogenetics, 1995, 80, 55-59.	1.0	13
243	A Phase II Trial of Vandetanib in Children and Adults with Succinate Dehydrogenase–Deficient Gastrointestinal Stromal Tumor. Clinical Cancer Research, 2019, 25, 6302-6308.	7.0	13
244	Genome-Wide Analysis of the FOXA1 Transcriptional Network Identifies Novel Protein-Coding and Long Noncoding RNA Targets in Colorectal Cancer Cells. Molecular and Cellular Biology, 2020, 40, .	2.3	13
245	Detection of Chromosome 6 Abnormalities in Melanoma Cell Lines by Chromosome Arm Painting Probes. Cancer Genetics and Cytogenetics, 1998, 107, 89-92.	1.0	12
246	Chimeric Negative Regulation ofp14ARFandTBX1by a t(9;22) Translocation Associated with Melanoma, Deafness, and DNA Repair Deficiency. Human Mutation, 2013, 34, 1250-1259.	2.5	11
247	Translational Predictive Biomarker Analysis of the Phase 1b Sorafenib and Bevacizumab Study Expansion Cohort. Molecular and Cellular Proteomics, 2013, 12, 1621-1631.	3.8	11
248	Engineered Bcor mutations lead to acute leukemia of progenitor B-1 lymphocyte origin in a sensitized background. Blood, 2019, 133, 2610-2614.	1.4	11
249	Generation of Novel Genetic Models to Dissect Resistance to Thyroid Hormone Receptor \hat{l}_{\pm} in Zebrafish. Thyroid, 2020, 30, 314-328.	4.5	11
250	Alu-PCR: Characterization of a chromosome 6-specific hybrid mapping panel and cloning of chromosome-specific markers. Genomics, 1992, 12, 549-554.	2.9	10
251	Isolation of YAC insert sequences by representational difference analysis. Nucleic Acids Research, 1995, 23, 4127-4133.	14.5	10
252	<i>In Vivo</i> Role of Alternative Splicing and Serine Phosphorylation of the Microphthalmia-Associated Transcription Factor. Genetics, 2012, 191, 133-144.	2.9	10

#	Article	IF	CITATIONS
253	Prohibitin expression is associated with high grade breast cancer but is not a driver of amplification at 17q21.33. Pathology, 2013, 45, 629-636.	0.6	10
254	Replicon: a software to accurately predict DNA replication timing in metazoan cells. Frontiers in Genetics, 2014, 5, 378.	2.3	10
255	Relationship of DNA methylation to mutational changes and transcriptional organization in fusionâ€positive and fusionâ€negative rhabdomyosarcoma. International Journal of Cancer, 2019, 144, 2707-2717.	5.1	10
256	Characterization of genetically defined sporadic and hereditary type 1 papillary renal cell carcinoma cell lines. Genes Chromosomes and Cancer, 2021 , 60 , 434 - 446 .	2.8	10
257	Characterization of Genomic Alterations in Radiation-Associated Breast Cancer among Childhood Cancer Survivors, Using Comparative Genomic Hybridization (CGH) Arrays. PLoS ONE, 2015, 10, e0116078.	2.5	10
258	Telomere Length and Survival of Patients with Hepatocellular Carcinoma in the United States. PLoS ONE, 2016, 11, e0166828.	2. 5	10
259	Localization of the gene encoding RκB (NFRKB), a tissue-specific DNA binding protein, to chromosome 11q24–q25. Genomics, 1992, 14, 270-274.	2.9	9
260	Identification of region specific genes by chromosome microdissection. Cancer Genetics and Cytogenetics, 1997, 93, 29-32.	1.0	9
261	Production and Characterization of Monoclonal Antibodies to the Steroid Receptor Coactivator AIB1. Hybridoma, 1999, 18, 281-287.	0.6	9
262	Complex temporal changes in TGF \hat{l}^2 oncogenic signaling drive thyroid carcinogenesis in a mouse model. Carcinogenesis, 2013, 34, 2389-2400.	2.8	9
263	Somatic mutations in murine models of leukemia and lymphoma: Disease specificity and clinical relevance. Genes Chromosomes and Cancer, 2017, 56, 472-483.	2.8	9
264	Oncogene Panel Sequencing Analysis Identifies Candidate Actionable Genes in Advanced Well-Differentiated Gastroenteropancreatic Neuroendocrine Tumors. Endocrine Practice, 2019, 25, 580-588.	2.1	9
265	Low-penetrant RB allele in small-cell cancer shows geldanamycin instability and discordant expression with mutant ras. Cell Cycle, 2008, 7, 2384-2391.	2.6	8
266	CD8+Lymphocytes Suppress Human Immunodeficiency Virus 1 Replication by Secreting Type I Interferons. Journal of Interferon and Cytokine Research, 2013, 33, 632-645.	1.2	8
267	Impact of telomere length on survival in classic and variant hairy cell leukemia. Leukemia Research, 2015, 39, 1360-1366.	0.8	8
268	Leukocyte DNA methylation and colorectal cancer among male smokers. World Journal of Gastrointestinal Oncology, 2012, 4, 193.	2.0	8
269	Exclusion of the 750â€kb genetically unstable region at Xq27 as a candidate locus for prostate malignancy in HPCX1â€linked families. Genes Chromosomes and Cancer, 2012, 51, 933-948.	2.8	7
270	Integrated analysis of DNA methylation, immunohistochemistry and mRNA expression, data identifies a methylation expression index (MEI) robustly associated with survival of ER-positive breast cancer patients. Breast Cancer Research and Treatment, 2015, 150, 457-466.	2.5	7

#	Article	IF	Citations
271	Novel renal medullary carcinoma cell lines, <scp>UOK353</scp> and <scp>UOK360</scp> , provide preclinical tools to identify new therapeutic treatments. Genes Chromosomes and Cancer, 2020, 59, 472-483.	2.8	7
272	Mutant <i>Idh2</i> Cooperates with a <i>NUP98-HOXD13</i> Fusion to Induce Early Immature Thymocyte Precursor ALL. Cancer Research, 2021, 81, 5033-5046.	0.9	7
273	Analysis of dihydrofolate reductase gene amplification in a methotrexate-resistant human tumor cell line. Cancer Genetics and Cytogenetics, 1985, 17, 289-300.	1.0	6
274	Microdissection and microcloning of chromosomal alterations in human breast cancer. Breast Cancer Research and Treatment, 1995, 33, 95-102.	2.5	6
275	Chromosome breakpoint at $17q11.2$ and insertion of DNA from three different chromosomes in a glioblastoma with exceptional glial fibrillary acidic protein expression. Cancer Genetics and Cytogenetics, 1996, 87, 41-47.	1.0	6
276	Comparison of Eight Technologies to Determine Genotype at the UGT1A1 (TA)n Repeat Polymorphism: Potential Clinical Consequences of Genotyping Errors?. International Journal of Molecular Sciences, 2020, 21, 896.	4.1	6
277	LRRC15 antibodyâ€drug conjugates show promise as osteosarcoma therapeutics in preclinical studies. Pediatric Blood and Cancer, 2021, 68, e28771.	1.5	6
278	Regulation of cancer stem cell activity by thyroid hormone receptor \hat{l}^2 . Oncogene, 2022, 41, 2315-2325.	5.9	6
279	Isolation of a yeast artificial chromosome clone that spans the (12;16) translocation breakpoint characteristic of myxoid liposarcoma. Cancer Genetics and Cytogenetics, 1992, 62, 166-170.	1.0	5
280	The last shall be first. Nature Genetics, 1993, 3, 101-102.	21.4	5
281	Ewing's sarcoma: General insights from a rare model. Cancer Cell, 2006, 9, 331-332.	16.8	5
282	Caspase-8 expression is predictive of tumour response to death receptor 5 agonist antibody in Ewing's sarcoma. British Journal of Cancer, 2015, 113, 894-901.	6.4	5
283	Somatic VHL Mutation in a Patient With MEN1-Associated Metastatic Pancreatic Neuroendocrine Tumor Responding to Sunitinib Treatment: A Case Report. Journal of the Endocrine Society, 2017, 1, 1124-1134.	0.2	5
284	Expression of the muscle-associated gene MYF6 in hairy cell leukemia. PLoS ONE, 2020, 15, e0227586.	2.5	5
285	Multifocal Renal Cell Carcinomas With Somatic IDH2 Mutation: Report of a Previously Undescribed Neoplasm. American Journal of Surgical Pathology, 2021, 45, 137-142.	3.7	5
286	Targeting Epigenetic Misregulation in Synovial Sarcoma. Cancer Cell, 2012, 21, 323-324.	16.8	4
287	Avalanching mutations in biallelic mismatch repair deficiency syndrome. Nature Genetics, 2015, 47, 194-196.	21.4	4
288	caOmicsV: an R package for visualizing multidimensional cancer genomic data. BMC Bioinformatics, 2016, 17, 141.	2.6	4

#	Article	IF	Citations
289	Analysis of the 9p21.3 sequence associated with coronary artery disease reveals a tendency for duplication in a CAD patient. Oncotarget, 2018, 9, 15275-15291.	1.8	4
290	1q21.1 deletion and a rare functional polymorphism in siblings with thrombocytopenia-absent radius–like phenotypes. Journal of Physical Education and Sports Management, 2019, 5, a004564.	1.2	4
291	A Large Ribosomal Subunit Protein Abnormality in Diamond-Blackfan Anemia (DBA) Blood, 2007, 110, 422-422.	1.4	4
292	Leukemia Driven By a NUP98-Phd Domain Fusion Is Highly Sensitive To Disruption Of H3K4me3-Phd Domain Binding By a Small Molecule Inhibitor. Blood, 2013, 122, 3759-3759.	1.4	4
293	A unique mutator phenotype reveals complementary oncogenic lesions leading to acute leukemia. JCI Insight, 2019, 4, .	5.0	4
294	Genetic and epigenetic analysis of monozygotic twins discordant for testicular cancer. International Journal of Molecular Epidemiology and Genetics, 2014, 5, 135-9.	0.4	4
295	Activation of integrin-ERBB2 signaling in undifferentiated thyroid cancer. American Journal of Cancer Research, 2014, 4, 776-88.	1.4	4
296	Restriction Mapping of Recombinant Cosmid Clones Using \hat{l} » Terminase and Field Inversion Gel Electrophoresis. Analytical Biochemistry, 1997, 245, 105-107.	2.4	2
297	Genomic Investigation of Dedifferentiated Liposarcoma Suggests a Role for Therapeutic Targeting of the Tumor Epigenome. Cancer Discovery, 2011, 1, 555-556.	9.4	2
298	Structure and promoter analysis of the humanunc-33-like phosphoprotein gene: E-box required for maximal expression in neuroblastoma and myoblasts Journal of Biological Chemistry, 2000, 275, 25052-25053.	3.4	2
299	Novel near-diploid ovarian cancer cell line derived from a highly aneuploid metastatic ovarian tumor. PLoS ONE, 2017, 12, e0182610.	2.5	2
300	Wasting disease associated with Epstein-Barr virus infection. Pediatric Infectious Disease Journal, 1988, 7, 719-723.	2.0	1
301	Gene Expression Profiling in Breast Cancer Research. Breast Disease, 2004, 19, 23-27.	0.8	1
302	Building through Breaking: The Development of Cancer Neochromosomes. Cancer Cell, 2014, 26, 593-595.	16.8	1
303	P2.01-041 Integrated Proteo-Genomics Analyses Reveal Extensive Tumor Heterogeneity and Novel Somatic Variants in Lung Adenocarcinoma. Journal of Thoracic Oncology, 2017, 12, S810-S811.	1.1	1
304	A Non-canonical Polycomb Dependency in Synovial Sarcoma. Cancer Cell, 2018, 33, 344-346.	16.8	1
305	CNVScope: Visually Exploring Copy Number Aberrations in Cancer Genomes. Cancer Informatics, 2019, 18, 117693511989029.	1.9	1
306	Genomic Composition of Howell-Jolly Bodies Blood, 2006, 108, 1570-1570.	1.4	1

#	Article	IF	CITATIONS
307	Enforced Expression of Lin28b Drives Development of Peripheral T Cell Lymphoma In Vivo. Blood, 2011, 118, 1392-1392.	1.4	1
308	A NUP98-PHF23 Transgenic Mouse Model Develops AML and T-ALL. Blood, 2011, 118, 2467-2467.	1.4	1
309	Spontaneous Mutations of Bcor and Jak $1/2$ genes Lead to an Aggressive Leukemia of B-1 Progenitor B Cells. Blood, 2014, 124, 3573-3573.	1.4	1
310	Inflammation suppression prevents tumor cell proliferation in a mouse model of thyroid cancer. American Journal of Cancer Research, 2020, 10, 1857-1870.	1.4	1
311	Selective amplification of chromosome 6 specific DNA sequences by Alu-based PCR. Cancer Genetics and Cytogenetics, 1991, 56, 95-96.	1.0	0
312	Isolation and regional assignment of chromosome 6 specific HNCDNA clones to analyze human malignant melanoma. Cancer Genetics and Cytogenetics, 1994, 77, 182.	1.0	0
313	Large-Scale Genome Analysis. Methods of Biochemical Analysis, 2002, 43, 393-412.	0.2	0
314	The role of the vasculogenic phenotype and its associated extracellular matrix in tumor progression Implications for immune surveillance. Clinical and Applied Immunology Reviews, 2003, 3, 263-276.	0.4	0
315	Application of gene set enrichment method to ChIP-chip data analysis. , 2008, , .		0
316	Constitutive FLT3 Activation Results in Specific Changes in Gene Expression in Myeloid Leukemic Cells Blood, 2004, 104, 1115-1115.	1.4	0
317	Methylation Profiling of Mediastinal Gray Zone Lymphoma Reveals a Distinctive Signature with Elements Shared by Classical Hodgkin's Lymphoma and Mediastinal Large B-Cell Lymphoma. Blood, 2010, 116, 747-747.	1.4	0
318	Co-Expression of NUP98-HOXD13 and Mutant IDH2 Triggers an Early T-Cell Precursor-like Leukemia in Mice. Blood, 2015, 126, 904-904.	1.4	0
319	Increased Expression of Myc in Hairy Cell Leukemia, and Cell-Sensitivity to JQ1. Blood, 2016, 128, 5112-5112.	1.4	0
320	Engineered Bcor Mutations Lead to Acute Lymphoblastic Leukemia of Progenitor B-1 Lymphocyte Origin in a Sensitized Background. Blood, 2018, 132, 1331-1331.	1.4	0
321	Mcm2 Deficiency Leads to Bone Marrow Failure and Lymphoid Malignancies Dependent on Age and Genetic Background. Blood, 2021, 138, 2223-2223.	1.4	O