

# Paola Dal Cin

## List of Publications by Year in descending order

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221  
papers

18,493  
citations

11651

70  
h-index

13379

130  
g-index

224  
all docs

224  
docs citations

224  
times ranked

12396  
citing authors

#	ARTICLE	IF	CITATIONS
1	Characteristic nuclear membrane <i>ALK</i> reactivity in chronic myelomonocytic leukemia with <i>RANBP2-ALK</i> fusion. <i>American Journal of Hematology</i> , 2023, 98, 365-367.	4.1	1
2	<i>t(4;12)(q12;p13)</i> ETV6-rearranged AML without eosinophilia does not involve PDGFRA: relevance for imatinib insensitivity. <i>Blood Advances</i> , 2022, 6, 818-827.	5.2	5
3	Atypical uterine polyps show morphologic and molecular overlap with mullerian adenocarcinoma but follow a benign clinical course. <i>Modern Pathology</i> , 2022, 35, 106-116.	5.5	4
4	Chemotherapy Resistance in B-ALL with Cryptic <i>NUP214-ABL1</i> Is Amenable to Kinase Inhibition and Immunotherapy. <i>Oncologist</i> , 2022, 27, 82-86.	3.7	5
5	<i>TP53</i> mutation defines a unique subgroup within complex karyotype de novo and therapy-related MDS/AML. <i>Blood Advances</i> , 2022, 6, 2847-2853.	5.2	87
6	N-terminus <i>DUX4</i> immunohistochemistry is a reliable methodology for the diagnosis of <i>DUX4</i> -fused B-lymphoblastic leukemia/lymphoma (N-terminus <i>DUX4</i> IHC for) Tj ET 20 0 0 rBT /Overlo	2.9	0
7	Low-grade Fibromyxoid Sarcoma of the Vulva and Vagina. <i>American Journal of Surgical Pathology</i> , 2022, 46, 1196-1206.	3.7	3
8	Guiding the global evolution of cytogenetic testing for hematologic malignancies. <i>Blood</i> , 2022, 139, 2273-2284.	1.4	29
9	International Consensus Classification of Myeloid Neoplasms and Acute Leukemias: integrating morphologic, clinical, and genomic data. <i>Blood</i> , 2022, 140, 1200-1228.	1.4	814
10	Genomic alterations in patients with somatic loss of the Y chromosome as the sole cytogenetic finding in bone marrow cells. <i>Haematologica</i> , 2021, 106, 555-564.	3.5	34
11	Pseudosarcomatous myofibroblastic proliferations of the urinary bladder are neoplasms characterized by recurrent <i>FN1-ALK</i> fusions. <i>Modern Pathology</i> , 2021, 34, 469-477.	5.5	12
12	Re-evaluating tumors of purported specialized prostatic stromal origin reveals molecular heterogeneity, including non-recurring gene fusions characteristic of uterine and soft tissue sarcoma subtypes. <i>Modern Pathology</i> , 2021, 34, 1763-1779.	5.5	8
13	Myelodysplastic/myeloproliferative neoplasms-unclassifiable with isolated isochromosome 17q represents a distinct clinico-biologic subset: a multi-institutional collaborative study from the Bone Marrow Pathology Group. <i>Modern Pathology</i> , 2021, , .	5.5	9
14	Polymorphous sweat gland carcinoma found to have MYB rearrangement. <i>Histopathology</i> , 2020, 76, 779-781.	2.9	1
15	Synovial Sarcoma of the Female Genital Tract. <i>American Journal of Surgical Pathology</i> , 2020, 44, 1487-1495.	3.7	11
16	Long: molecular tracking of CML with bilineal <i>inv(16)</i> myeloid and <i>del(9)</i> lymphoid blast crisis and durable response to CD19-directed CAR-T therapy. <i>Leukemia</i> , 2020, 34, 3050-3054.	7.2	3
17	Targeted FGFR inhibition results in a durable remission in an FGFR1-driven myeloid neoplasm with eosinophilia. <i>Blood Advances</i> , 2020, 4, 3136-3140.	5.2	28
18	Carbonic anhydrase IX (CA9) expression in multiple renal epithelial tumour subtypes. <i>Histopathology</i> , 2020, 77, 659-666.	2.9	28

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19	A cryptic imatinib-sensitive G3BP1-PDGFRB rearrangement in a myeloid neoplasm with eosinophilia. <i>Blood Advances</i> , 2020, 4, 445-448.	5.2	11
20	IGH rearrangement in myeloid neoplasms. <i>Haematologica</i> , 2020, 105, e315-e317.	3.5	4
21	Clinical response to larotrectinib in adult Philadelphia chromosome-like ALL with cryptic ETV6-NTRK3 rearrangement. <i>Blood Advances</i> , 2020, 4, 106-111.	5.2	23
22	Composite chronic myeloid leukemia and essential thrombocythemia with <i>BCR-ABL1</i> fusion and <i>CALR</i> mutation. <i>American Journal of Hematology</i> , 2019, 94, 504-505.	4.1	9
23	High <i>NPM1</i> mutant allele burden at diagnosis correlates with minimal residual disease at first remission in de novo acute myeloid leukemia. <i>American Journal of Hematology</i> , 2019, 94, 921-928.	4.1	24
24	Clinicopathologic Features and Chromosome 12p Status of Pediatric Sacrococcygeal Teratomas: A Multi-institutional Analysis. <i>Pediatric and Developmental Pathology</i> , 2019, 22, 214-220.	1.0	2
25	<i>ZMYM2-FGFR1</i> fusion as secondary change in acute myeloid leukemia. <i>Leukemia and Lymphoma</i> , 2019, 60, 556-558.	1.3	0
26	Targeted FGFR Inhibition Results in Hematologic and Cytogenetic Remission in a Myeloid Neoplasm Driven By a Novel PCM1-FGFR1 Fusion: Data from an Expanded Access Program. <i>Blood</i> , 2019, 134, 5371-5371.	1.4	0
27	<i>CIC-NUTM1</i> fusion: A case which expands the spectrum of <i>NUT</i> -rearranged epithelioid malignancies. <i>Genes Chromosomes and Cancer</i> , 2018, 57, 446-451.	2.8	53
28	Outcomes after Allogeneic Stem Cell Transplantation in Patients with Double-Hit and Double-Expressor Lymphoma. <i>Biology of Blood and Marrow Transplantation</i> , 2018, 24, 514-520.	2.0	31
29	Expanding the spectrum of translocations in sclerosing epithelioid fibrosarcoma: A new case with <i>EWSR1-CREB3L3</i> fusion. <i>Genes Chromosomes and Cancer</i> , 2018, 57, 675-677.	2.8	11
30	Immunophenotypic dysplasia and aberrant T-cell antigen expression in acute myeloid leukaemia with complex karyotype and <i>TP53</i> mutations. <i>Journal of Clinical Pathology</i> , 2018, 71, 1051-1059.	2.0	6
31	Clinicopathologic evaluation of cytopenic patients with isolated trisomy 8: a detailed comparison between idiopathic cytopenia of unknown significance and low-grade myelodysplastic syndrome. <i>Leukemia and Lymphoma</i> , 2017, 58, 569-577.	1.3	12
32	An Unusual Case of YWHAE-NUTM2A/B Endometrial Stromal Sarcoma With Confinement to the Endometrium and Lack of High-Grade Morphology. <i>International Journal of Gynecological Pathology</i> , 2017, 36, 165-171.	1.4	13
33	Detection of activating <i>MAP2K1</i> mutations in atypical hairy cell leukemia and hairy cell leukemia variant. <i>Leukemia and Lymphoma</i> , 2017, 58, 233-236.	1.3	39
34	Ectopic protein interactions within BRD4 chromatin complexes drive oncogenic megadomain formation in NUT midline carcinoma. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, E4184-E4192.	7.1	104
35	Conventional cytogenetics for myeloid neoplasms in the era of next-generation sequencing. <i>American Journal of Hematology</i> , 2017, 92, 227-229.	4.1	3
36	Ring chromosome in myeloid neoplasms is associated with complex karyotype and disease progression. <i>Human Pathology</i> , 2017, 68, 40-46.	2.0	5

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37	Fusion of the genes <i>BRD8</i> and <i>PHF1</i> in endometrial stromal sarcoma. <i>Genes Chromosomes and Cancer</i> , 2017, 56, 841-845.	2.8	45
38	Integrated single-cell genetic and transcriptional analysis suggests novel drivers of chronic lymphocytic leukemia. <i>Genome Research</i> , 2017, 27, 1300-1311.	5.5	67
39	Maternal <i>iAMP21</i> acute lymphoblastic leukemia detected on prenatal cell-free DNA genetic screening. <i>Blood Advances</i> , 2017, 1, 1491-1494.	5.2	4
40	FISHing in the dark: How the combination of FISH and conventional karyotyping improves the diagnostic yield in CpG-methylated chronic lymphocytic leukemia. <i>American Journal of Hematology</i> , 2016, 91, 978-983.	4.1	14
41	The Dohner fluorescence <i>in situ</i> hybridization prognostic classification of chronic lymphocytic leukaemia (CLL): the CLL Research Consortium experience. <i>British Journal of Haematology</i> , 2016, 173, 105-113.	2.5	66
42	Acute erythroid leukemia with <20% bone marrow blasts is clinically and biologically similar to myelodysplastic syndrome with excess blasts. <i>Modern Pathology</i> , 2016, 29, 1221-1231.	5.5	22
43	Clonal evolution in patients with chronic lymphocytic leukaemia developing resistance to BTK inhibition. <i>Nature Communications</i> , 2016, 7, 11589.	12.8	285
44	Intravenous leiomyomatosis: an unusual intermediate between benign and malignant uterine smooth muscle tumors. <i>Modern Pathology</i> , 2016, 29, 500-510.	5.5	65
45	MYC Immunohistochemistry to Identify MYC-Driven B-Cell Lymphomas in Clinical Practice. <i>American Journal of Clinical Pathology</i> , 2016, 145, 166-179.	0.7	29
46	Double-Hit and Double-Expressor Lymphomas Are Not Associated with an Adverse Outcome after Allogeneic Stem Cell Transplantation. <i>Blood</i> , 2016, 128, 830-830.	1.4	3
47	Molecular Pathology and Cytogenetics of Endometrial Carcinoma, Carcinosarcoma, and Uterine Sarcomas. <i>Current Clinical Oncology</i> , 2015, , 85-103.	0.0	0
48	Routine conventional karyotyping of lymphoma staging bone marrow samples does not contribute clinically relevant information. <i>American Journal of Hematology</i> , 2015, 90, 529-533.	4.1	5
49	Refractory myeloid sarcoma with a <i>FIP1L1-PDGFR</i> rearrangement detected by clinical high throughput somatic sequencing. <i>Experimental Hematology and Oncology</i> , 2015, 4, 30.	5.0	6
50	Controversial fluorescence <i>in situ</i> hybridization cytogenetic abnormalities in chronic lymphocytic leukaemia: new insights from a large cohort. <i>British Journal of Haematology</i> , 2015, 170, 694-703.	2.5	19
51	Analysis of <i>MDM2</i> Amplification in 43 Endometrial Stromal Tumors. <i>International Journal of Gynecological Pathology</i> , 2015, 34, 576-583.	1.4	33
52	Myeloid neoplasm demonstrating a <i>STAT5B-RARA</i> rearrangement and genetic alterations associated with all-trans retinoic acid resistance identified by a custom next-generation sequencing assay. <i>Journal of Physical Education and Sports Management</i> , 2015, 1, a000307.	1.2	13
53	Expression of <i>ROS1</i> predicts <i>ROS1</i> gene rearrangement in inflammatory myofibroblastic tumors. <i>Modern Pathology</i> , 2015, 28, 732-739.	5.5	85
54	Acute myeloid leukemia in a patient with constitutional 47,XXY karyotype. <i>Leukemia Research Reports</i> , 2015, 4, 28-30.	0.4	0

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55	Inflammatory Myofibroblastic Tumor of the Uterus. American Journal of Surgical Pathology, 2015, 39, 157-168.	3.7	107
56	Targeted genomic profiling reveals recurrent KRAS mutations and gain of chromosome 1q in mesonephric carcinomas of the female genital tract. Modern Pathology, 2015, 28, 1504-1514.	5.5	111
57	Adult Renal Cell Carcinoma. Surgical Pathology Clinics, 2015, 8, 587-621.	1.7	33
58	Molecular Classification of MYC-Driven B-Cell Lymphomas by Targeted Gene Expression Profiling of Fixed Biopsy Specimens. Journal of Molecular Diagnostics, 2015, 17, 19-30.	2.8	25
59	High p53 protein expression in therapy-related myeloid neoplasms is associated with adverse karyotype and poor outcome. Modern Pathology, 2015, 28, 552-563.	5.5	42
60	Dedifferentiated liposarcoma and pleomorphic liposarcoma. Cancer Cytopathology, 2014, 122, 128-137.	2.4	47
61	Primary Sclerosing Epithelioid Fibrosarcoma of Bone. American Journal of Surgical Pathology, 2014, 38, 1538-1544.	3.7	64
62	Perivascular Epithelioid Cell Neoplasm (PEComa) of the Gynecologic Tract. American Journal of Surgical Pathology, 2014, 38, 176-188.	3.7	165
63	Promiscuous genes involved in recurrent chromosomal translocations in soft tissue tumours. Pathology, 2014, 46, 105-112.	0.6	40
64	Nuclear Protein in Testis Midline Carcinoma Misdiagnosed As Adamantinoma. Journal of Clinical Oncology, 2014, 32, e57-e60.	1.6	13
65	NSD3-“NUT Fusion Oncoprotein in NUT Midline Carcinoma: Implications for a Novel Oncogenic Mechanism. Cancer Discovery, 2014, 4, 928-941.	9.4	192
66	Multiple EWSR1-WT1 and WT1-EWSR1 copies in two cases of desmoplastic round cell tumor. Cancer Genetics, 2013, 206, 387-392.	0.4	19
67	ALK as a paradigm of oncogenic promiscuity: different mechanisms of activation and different fusion partners drive tumors of different lineages. Cancer Genetics, 2013, 206, 357-373.	0.4	51
68	Well-differentiated and dedifferentiated liposarcomas with prominent myxoid stroma: analysis of 56 cases. Histopathology, 2013, 62, 287-293.	2.9	82
69	Cutaneous Syncytial Myoepithelioma. American Journal of Surgical Pathology, 2013, 37, 710-718.	3.7	103
70	Molecular Pathology of Soft Tissue and Bone Tumors. , 2013, , 325-356.		0
71	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.	7.1	239
72	MUC4 Is a Sensitive and Extremely Useful Marker for Sclerosing Epithelioid Fibrosarcoma. American Journal of Surgical Pathology, 2012, 36, 1444-1451.	3.7	230

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73	The Clinicopathologic Features of YWHAE-FAM22 Endometrial Stromal Sarcomas. American Journal of Surgical Pathology, 2012, 36, 641-653.	3.7	265
74	Cyclin D1 as a Diagnostic Immunomarker for Endometrial Stromal Sarcoma With YWHAE-FAM22 Rearrangement. American Journal of Surgical Pathology, 2012, 36, 1562-1570.	3.7	184
75	Integrative Genomic Analysis Implicates Gain of <i>PIK3CA</i> at 3q26 and <i>MYC</i> at 8q24 in Chronic Lymphocytic Leukemia. Clinical Cancer Research, 2012, 18, 3791-3802.	7.0	76
76	Metaphase Harvest and Cytogenetic Analysis of Malignant Hematological Specimens. Current Protocols in Human Genetics, 2012, 73, Unit 10.2.1-15.	3.5	0
77	Immunohistochemical Detection of MYC-driven Diffuse Large B-Cell Lymphomas. PLoS ONE, 2012, 7, e33813.	2.5	137
78	Angiomatoid fibrous histiocytoma a series of five cytologic cases with literature review and emphasis on diagnostic pitfalls. Diagnostic Cytopathology, 2012, 40, E86-93.	1.0	17
79	Identification of a novel, recurrent <i>HEY1</i> - <i>NCOA2</i> fusion in mesenchymal chondrosarcoma based on a genome-wide screen of exon-level expression data. Genes Chromosomes and Cancer, 2012, 51, 127-139.	2.8	276
80	MUC4 Is a Highly Sensitive and Specific Marker for Low-grade Fibromyxoid Sarcoma. American Journal of Surgical Pathology, 2011, 35, 733-741.	3.7	358
81	Consistent t(1;10) with rearrangements of <i>TGFBR3</i> and <i>MGEA5</i> in both myxoinflammatory fibroblastic sarcoma and hemosiderotic fibrolipomatous tumor. Genes Chromosomes and Cancer, 2011, 50, 757-764.	2.8	137
82	Validation of a TFE3 Break-apart FISH Assay for Xp11.2 Translocation Renal Cell Carcinomas. Diagnostic Molecular Pathology, 2011, 20, 129-137.	2.1	60
83	Cytogenetic, Molecular and Clinical Features Associated with Rare CFBF-MYH11 Fusion Transcripts in Patients (Pts) with Acute Myeloid Leukemia (AML) and inv(16)/t(16;16). Blood, 2011, 118, 2514-2514.	1.4	0
84	Rearrangement of 14q32 in the Absence of t(14;18) Is Associated with Short Time to First Treatment in Chronic Lymphocytic Leukemia. Blood, 2011, 118, 1438-1438.	1.4	0
85	Recurrent t(2;2) and t(2;8) translocations in rhabdomyosarcoma without the canonical <i>PAX</i> - <i>FOXO1</i> fuse <i>PAX3</i> to members of the nuclear receptor transcriptional coactivator family. Genes Chromosomes and Cancer, 2010, 49, 224-236.	2.8	129
86	B-cell Lymphomas With Concurrent IGH-BCL2 and MYC Rearrangements Are Aggressive Neoplasms With Clinical and Pathologic Features Distinct From Burkitt Lymphoma and Diffuse Large B-cell Lymphoma. American Journal of Surgical Pathology, 2010, 34, 327-340.	3.7	327
87	Dedifferentiated Liposarcoma With "Homologous" Lipoblastic (Pleomorphic Liposarcoma-like) Differentiation: Clinicopathologic and Molecular Analysis of a Series Suggesting Revised Diagnostic Criteria. American Journal of Surgical Pathology, 2010, 34, 1122-1131.	3.7	134
88	Hybrid Myxoinflammatory Fibroblastic Sarcoma/Hemosiderotic Fibrolipomatous Tumor: Report of a Case Providing Further Evidence for a Pathogenetic Link. American Journal of Surgical Pathology, 2010, 34, 1723-1727.	3.7	67
89	Pericentric inversion (12)(p12q13~1/4q14) as the sole chromosomal abnormality in a leiomyoma of the vulva. Cancer Genetics and Cytogenetics, 2010, 199, 21-23.	1.0	11
90	<i>HMGA1</i> and <i>HMGA2</i> rearrangements in mass-forming endometriosis. Genes Chromosomes and Cancer, 2010, 49, 630-634.	2.8	15

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91	<i>EWSR1</i> – <i>POU5F1</i> fusion in soft tissue myoepithelial tumors. A molecular analysis of sixty-six cases, including soft tissue, bone, and visceral lesions, showing common involvement of the <i>EWSR1</i> gene. <i>Genes Chromosomes and Cancer</i> , 2010, 49, 1114-1124.	2.8	443
92	Disseminated peritoneal leiomyomatosis after laparoscopic supracervical hysterectomy with characteristic molecular cytogenetic findings of uterine leiomyoma. <i>Genes Chromosomes and Cancer</i> , 2010, 49, 1152-1160.	2.8	67
93	Crizotinib in <i>ALK</i> -Rearranged Inflammatory Myofibroblastic Tumor. <i>New England Journal of Medicine</i> , 2010, 363, 1727-1733.	27.0	769
94	Standardization of fluorescence in situ hybridization studies on chronic lymphocytic leukemia (CLL) blood and marrow cells by the CLL Research Consortium. <i>Cancer Genetics and Cytogenetics</i> , 2010, 203, 141-148.	1.0	20
95	Molecular Profiling of Extramedullary and Medullary Plasmacytomas Compared to Multiple Myeloma. <i>Blood</i> , 2010, 116, 4042-4042.	1.4	0
96	Cytogenetics Abnormalities Predict the Outcome of Allogeneic Transplantation In AML: A CIBMTR Study. <i>Blood</i> , 2010, 116, 680-680.	1.4	0
97	Cytogenetics of Mesenchymal Tumors of the Female Genital Tract. <i>Surgical Pathology Clinics</i> , 2009, 2, 813-821.	1.7	3
98	Loss of INI1 Expression is Characteristic of Both Conventional and Proximal-type Epithelioid Sarcoma. <i>American Journal of Surgical Pathology</i> , 2009, 33, 542-550.	3.7	538
99	Molecular Profiling of Extramedullary and Medullary Plasmacytomas.. <i>Blood</i> , 2009, 114, 1806-1806.	1.4	1
100	Molecular Pathology and Cytogenetics of Endometrial Carcinoma, Carcinosarcoma, and Uterine Sarcomas. , 2009, , 87-104.		0
101	Karyotype Results From CpG Oligodeoxynucleotide Stimulated Chronic Lymphocytic Leukemia (CLL) Cultures Are Consistent Among Laboratories: a CLL Research Consortium (CRC) Study.. <i>Blood</i> , 2009, 114, 1614-1614.	1.4	0
102	Fluorescence in situ hybridization is a useful ancillary diagnostic tool for extraskeletal myxoid chondrosarcoma. <i>Modern Pathology</i> , 2008, 21, 1303-1310.	5.5	44
103	Molecular and Cytogenetic Characterization of Plexiform Leiomyomata Provide Further Evidence for Genetic Heterogeneity Underlying Uterine Fibroids. <i>American Journal of Pathology</i> , 2008, 172, 1403-1410.	3.8	24
104	Molecular Testing for Solid Tumors. , 2008, , 467-495.		1
105	Distinctive Cytogenetic Profile in Benign Metastasizing Leiomyoma: Pathogenetic Implications. <i>American Journal of Surgical Pathology</i> , 2007, 31, 737-743.	3.7	94
106	Molecular Analysis of the JAZF1-JAZ1 Gene Fusion by RT-PCR and Fluorescence In Situ Hybridization in Endometrial Stromal Neoplasms. <i>American Journal of Surgical Pathology</i> , 2007, 31, 65-70.	3.7	133
107	<i>EWSR1</i> – <i>CREB1</i> is the predominant gene fusion in angiomatoid fibrous histiocytoma. <i>Genes Chromosomes and Cancer</i> , 2007, 46, 1051-1060.	2.8	276
108	A Phase 2 Study of Fludarabine and Rituximab for the Treatment of Marginal Zone Lymphomas.. <i>Blood</i> , 2007, 110, 1358-1358.	1.4	2

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109	Philadelphia Chromosome-Positive Acute Myeloid Leukemia: A Rare Aggressive Leukemia With Clinicopathologic Features Distinct From Chronic Myeloid Leukemia in Myeloid Blast Crisis. <i>American Journal of Clinical Pathology</i> , 2007, 127, 642-650.	0.7	1
110	Detecting Apoptotic Blocks and Sensitivity to ABT-737 and Conventional Chemotherapy Via BH3 Profiling. <i>Blood</i> , 2007, 110, 4523-4523.	1.4	0
111	HMGA2 Rearrangement in a Case of Vulvar Aggressive Angiomyxoma. <i>International Journal of Gynecological Pathology</i> , 2006, 25, 403-407.	1.4	60
112	Uterine leiomyomata with deletions of 1p represent a distinct cytogenetic subgroup associated with unusual histologic features. <i>Genes Chromosomes and Cancer</i> , 2006, 45, 304-312.	2.8	62
113	FAS Death Domain Deletions and Cellular FADD-like Interleukin 1 $\beta$ Converting Enzyme Inhibitory Protein (Long) Overexpression: Alternative Mechanisms for Deregulating the Extrinsic Apoptotic Pathway in Diffuse Large B-Cell Lymphoma Subtypes. <i>Clinical Cancer Research</i> , 2006, 12, 3265-3271.	7.0	37
114	EWS-CREB1: A Recurrent Variant Fusion in Clear Cell Sarcoma Association with Gastrointestinal Location and Absence of Melanocytic Differentiation. <i>Clinical Cancer Research</i> , 2006, 12, 5356-5362.	7.0	305
115	Hodgkin's Lymphoma Reed Sternberg Cells over Express the T-Cell Inhibitory Carbohydrate-Binding Lectin, Galectin 1: Role of AP-1 and Likely Mechanism of Tumor Immune Escape. <i>Blood</i> , 2006, 108, 469-469.	1.4	1
116	Aneurysmal bone cyst variant translocations upregulate USP6 transcription by promoter swapping with the ZNF9, COL1A1, TRAP150, and OMD genes. <i>Oncogene</i> , 2005, 24, 3419-3426.	5.9	226
117	Molecular profiling of diffuse large B-cell lymphoma identifies robust subtypes including one characterized by host inflammatory response. <i>Blood</i> , 2005, 105, 1851-1861.	1.4	778
118	Uterine Leiomyomata with t(10;17) Disrupt the Histone Acetyltransferase MORF. <i>Cancer Research</i> , 2004, 64, 5570-5577.	0.9	106
119	USP6 (Tre2) Fusion Oncogenes in Aneurysmal Bone Cyst. <i>Cancer Research</i> , 2004, 64, 1920-1923.	0.9	284
120	An intragenic rearrangement of HMGA2 is not necessary for lipoma formation. <i>Cancer Genetics and Cytogenetics</i> , 2004, 149, 178-179.	1.0	6
121	Molecular pathogenesis of uterine smooth muscle tumors from transcriptional profiling. <i>Genes Chromosomes and Cancer</i> , 2004, 40, 97-108.	2.8	145
122	Urachal inflammatory myofibroblastic tumor with ALK gene rearrangement: a study of urachal remnants. <i>Urology</i> , 2004, 64, 140-144.	1.0	21
123	Molecular Profiling of Diffuse Large B-Cell Lymphoma Identifies Robust Subtypes Including One Characterized by Host Inflammatory Response. <i>Blood</i> , 2004, 104, 25-25.	1.4	1
124	Intravenous leiomyomatosis is characterized by a der(14)t(12;14)(q15;q24). <i>Genes Chromosomes and Cancer</i> , 2003, 36, 205-206.	2.8	69
125	Genetics in renal cell carcinoma. <i>Current Opinion in Urology</i> , 2003, 13, 463-466.	1.8	8
126	Metaphase Harvest and Cytogenetic Analysis of Malignant Hematological Specimens. <i>Current Protocols in Human Genetics</i> , 2003, 36, Unit 10.2.	3.5	6



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127	The molecular signature of mediastinal large B-cell lymphoma differs from that of other diffuse large B-cell lymphomas and shares features with classical Hodgkin lymphoma. <i>Blood</i> , 2003, 102, 3871-3879.	1.4	793
128	Fusion transcripts involving HMGA2 are not a common molecular mechanism in uterine leiomyomata with rearrangements in 12q15. <i>Cancer Research</i> , 2003, 63, 1351-8.	0.9	79
129	Chromosome changes in sarcomatoid renal carcinomas are different from those in renal cell carcinomas. <i>Cancer Genetics and Cytogenetics</i> , 2002, 134, 38-40.	1.0	21
130	Molecular genetic characterization of the EWS/CHN and RBP56/CHN fusion genes in extraskeletal myxoid chondrosarcoma. <i>Genes Chromosomes and Cancer</i> , 2002, 35, 340-352.	2.8	104
131	Correlation between clinicopathological features and karyotype in 100 cartilaginous and chordoid tumours. A report from the Chromosomes and Morphology (CHAMP) Collaborative Study Group. <i>Journal of Pathology</i> , 2002, 196, 194-203.	4.5	131
132	PLAG1 Alterations in Lipoblastoma. <i>American Journal of Pathology</i> , 2001, 159, 955-962.	3.8	117
133	BRD4 Bromodomain Gene Rearrangement in Aggressive Carcinoma with Translocation t(15;19). <i>American Journal of Pathology</i> , 2001, 159, 1987-1992.	3.8	188
134	Abnormal Nuclear Shape in Solid Tumors Reflects Mitotic Instability. <i>American Journal of Pathology</i> , 2001, 158, 199-206.	3.8	187
135	Upper respiratory tract carcinoma with chromosomal translocation 15;19. <i>Cancer</i> , 2001, 92, 1195-1203.	4.1	102
136	Clinical impact of molecular and cytogenetic findings in synovial sarcoma. <i>Genes Chromosomes and Cancer</i> , 2001, 31, 362-372.	2.8	108
137	Molecular cytogenetic definition of three distinct chromosome arm 14q deletion intervals in gastrointestinal stromal tumors. <i>Genes Chromosomes and Cancer</i> , 2001, 32, 26-32.	2.8	36
138	Amplification of AML1 in childhood acute lymphoblastic leukemias. <i>Genes Chromosomes and Cancer</i> , 2001, 30, 407-409.	2.8	49
139	The der(17)t(X;17)(p11;q25) of human alveolar soft part sarcoma fuses the TFE3 transcription factor gene to ASPL, a novel gene at 17q25. <i>Oncogene</i> , 2001, 20, 48-57.	5.9	562
140	Cytogenetic characterization of peripheral nerve sheath tumours: a report of the CHAMP study group. <i>Journal of Pathology</i> , 2000, 190, 31-38.		141
141	Coordinated expression and amplification of the MDM2, CDK4, and HMGI-C genes in atypical lipomatous tumours. <i>Journal of Pathology</i> , 2000, 190, 531-536.	4.5	250
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