Paola Dal Cin

List of Publications by Year
in descending order

220 papers	18,493 citations	16791 66 h-index	16791 127 g-index
224 all docs	224 docs citations	224 times ranked	13198 citing authors

#	Article	IF	Citations
1	Characteristic nuclear membrane <scp>ALK</scp> reactivity in chronic myelomonocytic leukemia with <scp>RANBP2â€ALK</scp> fusion. American Journal of Hematology, 2023, 98, 365-367.	2.0	1
2	t(4;12)(q12;p13) ETV6-rearranged AML without eosinophilia does not involve PDGFRA: relevance for imatinib insensitivity. Blood Advances, 2022, 6, 818-827.	2.5	5
3	Atypical uterine polyps show morphologic and molecular overlap with mullerian adenosarcoma but follow a benign clinical course. Modern Pathology, 2022, 35, 106-116.	2.9	4
4	Chemotherapy Resistance in B-ALL with Cryptic NUP214-ABL1 Is Amenable to Kinase Inhibition and Immunotherapy. Oncologist, 2022, 27, 82-86.	1.9	5
5	TP53 mutation defines a unique subgroup within complex karyotype deÂnovo and therapy-related MDS/AML. Blood Advances, 2022, 6, 2847-2853.	2.5	87
6	Nâ€terminus <scp>DUX4</scp>â€immunohistochemistry is a reliable methodology for the diagnosis of DUX4â€fused Bâ€lymphoblastic leukemia/lymphoma (Nâ€terminus <scp>DUX4 IHC</scp> for) Tj ETQq0 0 0 rgBT /Overlo		
7	Low-grade Fibromyxoid Sarcoma of the Vulva and Vagina. American Journal of Surgical Pathology, 2022, 46, 1196-1206.	2.1	3
8	Guiding the global evolution of cytogenetic testing for hematologic malignancies. Blood, 2022, 139, 2273-2284.	0.6	29
9	International Consensus Classification of Myeloid Neoplasms and Acute Leukemias: integrating morphologic, clinical, and genomic data. Blood, 2022, 140, 1200-1228.	0.6	814
10	Genomic alterations in patients with somatic loss of the Y chromosome as the sole cytogenetic finding in bone marrow cells. Haematologica, 2021, 106, 555-564.	1.7	34
11	Pseudosarcomatous myofibroblastic proliferations of the urinary bladder are neoplasms characterized by recurrent FN1â€“ALK fusions. Modern Pathology, 2021, 34, 469-477.	2.9	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. Modern Pathology, 2021, 34, 1763-1779.	2.9	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. Modern Pathology, 2021, , .	2.9	9
14	Polymorphous sweat gland carcinoma found to have MYB rearrangement. Histopathology, 2020, 76, 779-781.	1.6	1
15	Synovial Sarcoma of the Female Genital Tract. American Journal of Surgical Pathology, 2020, 44, 1487-1495.	2.1	11
16	Long: molecular tracking of CML with bilineal inv(16) myeloid and del(9) lymphoid blast crisis and durable response to CD19-directed CAR-T therapy. Leukemia, 2020, 34, 3050-3054.	3.3	3
17	Targeted FGFR inhibition results in a durable remission in an FGFR1-driven myeloid neoplasm with eosinophilia. Blood Advances, 2020, 4, 3136-3140.	2.5	28
18	Carbonic anhydrase IX (CA9) expression in multiple renal epithelial tumour subtypes. Histopathology, 2020, 77, 659-666.	1.6	28

#	Article	IF	Citations
19	A cryptic imatinib-sensitive G3BP1-PDGFRB rearrangement in a myeloid neoplasm with eosinophilia. Blood Advances, 2020, 4, 445-448.	2.5	11
20	IGH rearrangement in myeloid neoplasms. Haematologica, 2020, 105, e315-e317.	1.7	4
21	Clinical response to larotrectinib in adult Philadelphia chromosomeâ€"like ALL with cryptic ETV6-NTRK3 rearrangement. Blood Advances, 2020, 4, 106-111.	2.5	23
22	Composite chronic myeloid leukemia and essential thrombocythemia with BCRâ€ABL1 fusion and CALR mutation. American Journal of Hematology, 2019, 94, 504-505.	2.0	9
23	High NPM1 mutant allele burden at diagnosis correlates with minimal residual disease at first remission in de novo acute myeloid leukemia. American Journal of Hematology, 2019, 94, 921-928.	2.0	24
24	Clinicopathologic Features and Chromosome 12p Status of Pediatric Sacrococcygeal Teratomas: A Multi-institutional Analysis. Pediatric and Developmental Pathology, 2019, 22, 214-220.	0.5	2
25	ZMYM2-FGFR1 fusion as secondary change in acute myeloid leukemia. Leukemia and Lymphoma, 2019, 60, 556-558.	0.6	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. Blood, 2019, 134, 5371-5371.	0.6	0
27	CIC†NUTM1 fusion: A case which expands the spectrum of NUTâ€rearranged epithelioid malignancies. Genes Chromosomes and Cancer, 2018, 57, 446-451.	1.5	53
28	Outcomes after Allogeneic Stem Cell Transplantation in Patients with Double-Hit and Double-Expressor Lymphoma. Biology of Blood and Marrow Transplantation, 2018, 24, 514-520.	2.0	31
29	Expanding the spectrum of translocations in sclerosing epitheloid fibrosarcoma: A new case with EWSR1â€CREB3L3 fusion. Genes Chromosomes and Cancer, 2018, 57, 675-677.	1.5	11
30	Immunophenotypic dysplasia and aberrant T-cell antigen expression in acute myeloid leukaemia with complex karyotype and TP53 mutations. Journal of Clinical Pathology, 2018, 71, 1051-1059.	1.0	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. Leukemia and Lymphoma, 2017, 58, 569-577.	0.6	12
32	An Unusual Case of YWHAE-NUTM2A/B Endometrial Stromal Sarcoma With Confinement to the Endometrium and Lack of High-Grade Morphology. International Journal of Gynecological Pathology, 2017, 36, 165-171.	0.9	13
33	Detection of activatingMAP2K1mutations in atypical hairy cell leukemia and hairy cell leukemia variant. Leukemia and Lymphoma, 2017, 58, 233-236.	0.6	39
34	Ectopic protein interactions within BRD4â€"chromatin complexes drive oncogenic megadomain formation in NUT midline carcinoma. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E4184-E4192.	3.3	104
35	Conventional cytogenetics for myeloid neoplasms in the era of nextâ€generationâ€sequencing. American Journal of Hematology, 2017, 92, 227-229.	2.0	3
36	Ring chromosome in myeloid neoplasms is associated with complex karyotype and disease progression. Human Pathology, 2017, 68, 40-46.	1.1	5

#	Article	IF	Citations
37	Fusion of the genes BRD8 and PHF1 in endometrial stromal sarcoma. Genes Chromosomes and Cancer, 2017, 56, 841-845.	1.5	45
38	Integrated single-cell genetic and transcriptional analysis suggests novel drivers of chronic lymphocytic leukemia. Genome Research, 2017, 27, 1300-1311.	2.4	67
39	Maternal iAMP21 acute lymphoblastic leukemia detected on prenatal cell-free DNA genetic screening. Blood Advances, 2017, 1, 1491-1494.	2.5	4
40	FISHing in the dark: How the combination of FISH and conventional karyotyping improves the diagnostic yield in CpGâ€stimulated chronic lymphocytic leukemia. American Journal of Hematology, 2016, 91, 978-983.	2.0	14
41	The Dohner fluorescence inÂsitu hybridization prognostic classification of chronic lymphocytic leukaemia (<scp>CLL</scp>): the <scp>CLL</scp> Research Consortium experience. British Journal of Haematology, 2016, 173, 105-113.	1.2	66
42	Acute erythroid leukemia with <20% bone marrow blasts is clinically and biologically similar to myelodysplastic syndrome with excess blasts. Modern Pathology, 2016, 29, 1221-1231.	2.9	22
43	Clonal evolution in patients with chronic lymphocytic leukaemia developing resistance to BTK inhibition. Nature Communications, 2016, 7, 11589.	5.8	285
44	Intravenous leiomyomatosis: an unusual intermediate between benign and malignant uterine smooth muscle tumors. Modern Pathology, 2016, 29, 500-510.	2.9	65
45	MYC Immunohistochemistry to Identify MYC-Driven B-Cell Lymphomas in Clinical Practice. American Journal of Clinical Pathology, 2016, 145, 166-179.	0.4	29
46	Double-Hit and Double-Expressor Lymphomas Are Not Associated with an Adverse Outcome after Allogeneic Stem Cell Transplantation. Blood, 2016, 128, 830-830.	0.6	3
47	Molecular Pathology and Cytogenetics of Endometrial Carcinoma, Carcinosarcoma, and Uterine Sarcomas. Current Clinical Oncology, 2015, , 85-103.	0.0	0
48	Routine conventional karyotyping of lymphoma staging bone marrow samples does not contribute clinically relevant information. American Journal of Hematology, 2015, 90, 529-533.	2.0	5
49	Refractory myeloid sarcoma with a FIP1L1-PDGFRA rearrangement detected by clinical high throughput somatic sequencing. Experimental Hematology and Oncology, 2015, 4, 30.	2.0	6
50	Controversial fluorescence inÂsitu hybridization cytogenetic abnormalities in chronic lymphocytic leukaemia: new insights from a large cohort. British Journal of Haematology, 2015, 170, 694-703.	1.2	19
51	Analysis of MDM2 Amplification in 43 Endometrial Stromal Tumors. International Journal of Gynecological Pathology, 2015, 34, 576-583.	0.9	33
52	Myeloid neoplasm demonstrating a STAT5B-RARA rearrangement and genetic alterations associated with all-trans retinoic acid resistance identified by a custom next-generation sequencing assay. Journal of Physical Education and Sports Management, 2015, 1, a000307.	0.5	13
53	Expression of ROS1 predicts ROS1 gene rearrangement in inflammatory myofibroblastic tumors. Modern Pathology, 2015, 28, 732-739.	2.9	85
54	Acute myeloid leukemia in a patient with constitutional 47,XXY karyotype. Leukemia Research Reports, 2015, 4, 28-30.	0.2	0

#	Article	IF	Citations
55	Inflammatory Myofibroblastic Tumor of the Uterus. American Journal of Surgical Pathology, 2015, 39, 157-168.	2.1	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.	2.9	111
57	Adult Renal Cell Carcinoma. Surgical Pathology Clinics, 2015, 8, 587-621.	0.7	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.	1.2	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.	2.9	42
60	Dedifferentiated liposarcoma and pleomorphic liposarcoma. Cancer Cytopathology, 2014, 122, 128-137.	1.4	47
61	Primary Sclerosing Epithelioid Fibrosarcoma of Bone. American Journal of Surgical Pathology, 2014, 38, 1538-1544.	2.1	64
62	Perivascular Epithelioid Cell Neoplasm (PEComa) of the Gynecologic Tract. American Journal of Surgical Pathology, 2014, 38, 176-188.	2.1	165
63	Promiscuous genes involved in recurrent chromosomal translocations in soft tissue tumours. Pathology, 2014, 46, 105-112.	0.3	40
64	Nuclear Protein in Testis Midline Carcinoma Misdiagnosed As Adamantinoma. Journal of Clinical Oncology, 2014, 32, e57-e60.	0.8	13
65	NSD3â€“NUT Fusion Oncoprotein in NUT Midline Carcinoma: Implications for a Novel Oncogenic Mechanism. Cancer Discovery, 2014, 4, 928-941.	7.7	192
66	Multiple EWSR1-WT1 and WT1-EWSR1 copies in two cases of desmoplastic round cell tumor. Cancer Genetics, 2013, 206, 387-392.	0.2	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.	0.2	51
68	Wellâ€differentiated and dedifferentiated liposarcomas with prominent myxoid stroma: analysis of 56 cases. Histopathology, 2013, 62, 287-293.	1.6	82
69	Cutaneous Syncytial Myoepithelioma. American Journal of Surgical Pathology, 2013, 37, 710-718.	2.1	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.	3.3	239
72	MUC4 Is a Sensitive and Extremely Useful Marker for Sclerosing Epithelioid Fibrosarcoma. American Journal of Surgical Pathology, 2012, 36, 1444-1451.	2.1	230

#	Article	IF	Citations
73	The Clinicopathologic Features of YWHAE-FAM22 Endometrial Stromal Sarcomas. American Journal of Surgical Pathology, 2012, 36, 641-653.	2.1	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.	2.1	184
75	Integrative Genomic Analysis Implicates Gain of PIK3CA at 3q26 and MYC at 8q24 in Chronic Lymphocytic Leukemia. Clinical Cancer Research, 2012, 18, 3791-3802.	3.2	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.	1.1	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.	0.5	17
79	Identification of a novel, recurrent HEY1â€NCOA2 fusion in mesenchymal chondrosarcoma based on a genomeâ€wide screen of exonâ€level expression data. Genes Chromosomes and Cancer, 2012, 51, 127-139.	1.5	276
80	MUC4 Is a Highly Sensitive and Specific Marker for Low-grade Fibromyxoid Sarcoma. American Journal of Surgical Pathology, 2011, 35, 733-741.	2.1	358
81	Consistent t(1;10) with rearrangements of TGFBR3 and MGEA5 in both myxoinflammatory fibroblastic sarcoma and hemosiderotic fibrolipomatous tumor. Genes Chromosomes and Cancer, 2011, 50, 757-764.	1.5	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 CBFB-MYH11 Fusion Transcripts in Patients (Pts) with Acute Myeloid Leukemia (AML) and inv(16)/t(16;16). Blood, 2011, 118, 2514-2514.	0.6	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.	0.6	0
85	Recurrent t(2;2) and t(2;8) translocations in rhabdomyosarcoma without the canonical PAXâ€FOXO1 fuse PAX3 to members of the nuclear receptor transcriptional coactivator family. Genes Chromosomes and Cancer, 2010, 49, 224-236.	1.5	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.	2.1	327
87	Dedifferentiated Liposarcoma With â€œHomologousâ€•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.	2.1	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.	2.1	67
89	Pericentric inversion (12)(p12q13â¼q14) as the sole chromosomal abnormality in a leiomyoma of the vulva. Cancer Genetics and Cytogenetics, 2010, 199, 21-23.	1.0	11
90	HMGA1 and HMGA2 rearrangements in massâ€forming endometriosis. Genes Chromosomes and Cancer, 2010, 49, 630-634.	1.5	15

#	Article	IF	Citations
91	EWSR1â€POU5F1 fusion in soft tissue myoepithelial tumors. A molecular analysis of sixtyâ€six cases, including soft tissue, bone, and visceral lesions, showing common involvement of the EWSR1 gene. Genes Chromosomes and Cancer, 2010, 49, 1114-1124.	1.5	443
92	Disseminated peritoneal leiomyomatosis after laparoscopic supracervical hysterectomy with characteristic molecular cytogenetic findings of uterine leiomyoma. Genes Chromosomes and Cancer, 2010, 49, 1152-1160.	1.5	67
93	Crizotinib inALK-Rearranged Inflammatory Myofibroblastic Tumor. New England Journal of Medicine, 2010, 363, 1727-1733.	13.9	769
94	Standardization of fluorescence in situ hybridization studies on chronic lymphocytic leukemia (CLL) blood and marrow cells by the CLL Research Consortium. Cancer Genetics and Cytogenetics, 2010, 203, 141-148.	1.0	20
95	Molecular Profiling of Extramedullary and Medullary Plasmacytomas Compared to Multiple Myeloma. Blood, 2010, 116, 4042-4042.	0.6	0
96	Cytogenetics Abnormalities Predict the Outcome of Allogeneic Transplantation In AML: A CIBMTR Study. Blood, 2010, 116, 680-680.	0.6	0
97	Cytogenetics of Mesenchymal Tumors of the Female Genital Tract. Surgical Pathology Clinics, 2009, 2, 813-821.	0.7	3
98	Loss of INI1 Expression is Characteristic of Both Conventional and Proximal-type Epithelioid Sarcoma. American Journal of Surgical Pathology, 2009, 33, 542-550.	2.1	538
99	Molecular Profiling of Extramedullary and Medullary Plasmacytomas.. Blood, 2009, 114, 1806-1806.	0.6	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.. Blood, 2009, 114, 1614-1614.	0.6	0
102	Fluorescence in situ hybridization is a useful ancillary diagnostic tool for extraskeletal myxoid chondrosarcoma. Modern Pathology, 2008, 21, 1303-1310.	2.9	44
103	Molecular and Cytogenetic Characterization of Plexiform Leiomyomata Provide Further Evidence for Genetic Heterogeneity Underlying Uterine Fibroids. American Journal of Pathology, 2008, 172, 1403-1410.	1.9	24
104	Molecular Testing for Solid Tumors. , 2008, , 467-495.		1
105	Distinctive Cytogenetic Profile in Benign Metastasizing Leiomyoma: Pathogenetic Implications. American Journal of Surgical Pathology, 2007, 31, 737-743.	2.1	94
106	Molecular Analysis of the JAZF1-JJAZ1 Gene Fusion by RT-PCR and Fluorescence In Situ Hybridization in Endometrial Stromal Neoplasms. American Journal of Surgical Pathology, 2007, 31, 65-70.	2.1	133
107	EWSR1â€CREB1 is the predominant gene fusion in angiomatoid fibrous histiocytoma. Genes Chromosomes and Cancer, 2007, 46, 1051-1060.	1.5	276
108	A Phase 2 Study of Fludarabine and Rituximab for the Treatment of Marginal Zone Lymphomas.. Blood, 2007, 110, 1358-1358.	0.6	2

#	Article	IF	Citations
109	Detecting Apoptotic Blocks and Sensitvity to ABT-737 and Conventional Chemotherapy Via BH3 Profiling.. Blood, 2007, 110, 4523-4523.	0.6	0
110	HMGA2 Rearrangement in a Case of Vulvar Aggressive Angiomyxoma. International Journal of Gynecological Pathology, 2006, 25, 403-407.	0.9	60
111	Uterine leiomyomata with deletions of Ip represent a distinct cytogenetic subgroup associated with unusual histologic features. Genes Chromosomes and Cancer, 2006, 45, 304-312.	1.5	62
112	FAS Death Domain Deletions and Cellular FADD-like Interleukin 1Î² Converting Enzyme Inhibitory Protein (Long) Overexpression: Alternative Mechanisms for Deregulating the Extrinsic Apoptotic Pathway in Diffuse Large B-Cell Lymphoma Subtypes. Clinical Cancer Research, 2006, 12, 3265-3271.	3.2	37
113	EWS-CREB1: A Recurrent Variant Fusion in Clear Cell Sarcomaâ€"Association with Gastrointestinal Location and Absence of Melanocytic Differentiation. Clinical Cancer Research, 2006, 12, 5356-5362.	3.2	305
114	Hodgkinâ€™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.. Blood, 2006, 108, 469-469.	0.6	1
115	Aneurysmal bone cyst variant translocations upregulate USP6 transcription by promoter swapping with the ZNF9, COL1A1, TRAP150, and OMD genes. Oncogene, 2005, 24, 3419-3426.	2.6	226
116	Molecular profiling of diffuse large B-cell lymphoma identifies robust subtypes including one characterized by host inflammatory response. Blood, 2005, 105, 1851-1861.	0.6	778
117	Uterine Leiomyomata with t(10;17) Disrupt the Histone Acetyltransferase MORF. Cancer Research, 2004, 64, 5570-5577.	0.4	106
118	USP6 (Tre2) Fusion Oncogenes in Aneurysmal Bone Cyst. Cancer Research, 2004, 64, 1920-1923.	0.4	284
119	An intragenic rearrangement of HMGA2 is not necessary for lipoma formation. Cancer Genetics and Cytogenetics, 2004, 149, 178-179.	1.0	6
120	Molecular pathogenesis of uterine smooth muscle tumors from transcriptional profiling. Genes Chromosomes and Cancer, 2004, 40, 97-108.	1.5	145
121	Urachal inflammatory myofibroblastic tumor with ALK gene rearrangement: a study of urachal remnants. Urology, 2004, 64, 140-144.	0.5	21
122	Molecular Profiling of Diffuse Large B-Cell Lymphoma Identifies Robust Subtypes Including One Characterized by Host Inflammatory Response.. Blood, 2004, 104, 25-25.	0.6	1
123	Intravenous leiomyomatosis is characterized by a der(14)t(12;14)(q15;q24). Genes Chromosomes and Cancer, 2003, 36, 205-206.	1.5	69
124	Genetics in renal cell carcinoma. Current Opinion in Urology, 2003, 13, 463-466.	0.9	8
125	Metaphase Harvest and Cytogenetic Analysis of Malignant Hematological Specimens. Current Protocols in Human Genetics, 2003, 36, Unit 10.2.	3.5	6
126	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. Blood, 2003, 102, 3871-3879.	0.6	793

#	Article	IF	Citations
127	Fusion transcripts involving HMGA2 are not a common molecular mechanism in uterine leiomyomata with rearrangements in 12q15. Cancer Research, 2003, 63, 1351-8.	0.4	79
128	Chromosome changes in sarcomatoid renal carcinomas are different from those in renal cell carcinomas. Cancer Genetics and Cytogenetics, 2002, 134, 38-40.	1.0	21
129	Molecular genetic characterization of theEWS/CHNandRBP56/CHNfusion genes in extraskeletal myxoid chondrosarcoma. Genes Chromosomes and Cancer, 2002, 35, 340-352.	1.5	104
130	Correlation between clinicopathological features and karyotype in 100 cartilaginous and chordoid tumours. A report from the Chromosomes and Morphology (CHAMP) Collaborative Study Group. Journal of Pathology, 2002, 196, 194-203.	2.1	131
131	PLAG1 Alterations in Lipoblastoma. American Journal of Pathology, 2001, 159, 955-962.	1.9	117
132	BRD4 Bromodomain Gene Rearrangement in Aggressive Carcinoma with Translocation t(15;19). American Journal of Pathology, 2001, 159, 1987-1992.	1.9	188
133	Abnormal Nuclear Shape in Solid Tumors Reflects Mitotic Instability. American Journal of Pathology, 2001, 158, 199-206.	1.9	187
134	Upper respiratory tract carcinoma with chromosomal translocation 15;19. Cancer, 2001, 92, 1195-1203.	2.0	102
135	Clinical impact of molecular and cytogenetic findings in synovial sarcoma. Genes Chromosomes and Cancer, 2001, 31, 362-372.	1.5	108
136	Molecular cytogenetic definition of three distinct chromosome arm 14q deletion intervals in gastrointestinal stromal tumors. Genes Chromosomes and Cancer, 2001, 32, 26-32.	1.5	36
137	Amplification ofAML1 in childhood acute lymphoblastic leukemias. Genes Chromosomes and Cancer, 2001, 30, 407-409.	1.5	49
138	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. Oncogene, 2001, 20, 48-57.	2.6	562
139	Cytogenetic characterization of peripheral nerve sheath tumours: a report of the CHAMP study group. , 2000, 190, 31-38.		141
140	Coordinated expression and amplification of theMDM2,CDK4, andHMGI-C genes in atypical lipomatous tumours. , 2000, 190, 531-536.		250
141	Variant translocations involving 16q22 and 17p13 in solid variant and extraosseous forms of aneurysmal bone cyst. , 2000, 28, 233-234.		88
142	Comparative Cytogenetic Study of Spindle Cell and Pleomorphic Leiomyosarcomas of Soft Tissues. Cancer Genetics and Cytogenetics, 2000, 116, 66-73.	1.0	60
143	Cytogenetic-Morphologic Correlations in Aneurysmal Bone Cyst, Giant Cell Tumor of Bone and Combined Lesions. A Report from the CHAMP Study Group. Modern Pathology, 2000, 13, 1206-1210.	2.9	101
144	Precancerous Lesions in the Kidney. Scandinavian Journal of Urology and Nephrology, 2000, 34, 136-165.	1.4	20

#	Article	IF	Citations
145	Cytogenetic, Clinical, and Morphologic Correlations in 78 Cases of Fibromatosis: A Report from the CHAMP Study Group. Modern Pathology, 2000, 13, 1080-1085.	2.9	99
146	TPM3-ALK and TPM4-ALK Oncogenes in Inflammatory Myofibroblastic Tumors. American Journal of Pathology, 2000, 157, 377-384.	1.9	659
147	Coordinated expression and amplification of the MDM2, CDK4, and HMGI-C genes in atypical lipomatous tumours. , 2000, 190, 531.		4
148	Predominance of beta-catenin mutations and beta-catenin dysregulation in sporadic aggressive fibromatosis (desmoid tumor). Oncogene, 1999, 18, 6615-6620.	2.6	339
149	The structure and dynamics of ring chromosomes in human neoplastic and non-neoplastic cells. Human Genetics, 1999, 104, 315-325.	1.8	108
150	Embryonal Rhabdomyosarcoma with Only Numerical Chromosome Changes. Cancer Genetics and Cytogenetics, 1999, 109, 161-165.	1.0	21
151	Inflammatory Myofibroblastic Tumor with HMGIC Rearrangement. Cancer Genetics and Cytogenetics, 1999, 112, 156-160.	1.0	27
152	Involvement of 12q12-13 is a nonrandom chromosome change in renal oncocytoma. , 1999, 24, 94-94.		7
153	Clonal chromosome abnormalities in a so-called Dupuytren's subungual exostosis. , 1999, 24, 162-164.		24
154	Involvement of theHMGI(Y) gene in a microfollicular adenoma of the thyroid. Genes Chromosomes and Cancer, 1999, 24, 286-289.	1.5	12
155	HMG1 is not rearranged by 13q12 aberrations in lipomas. , 1999, 24, 290-292.		3
156	Correlation between Clinicopathological Features and Karyotype in Spindle Cell Sarcomas. American Journal of Pathology, 1999, 154, 1841-1847.	1.9	109
157	Comparison of Chromosomal Patterns with Clinical Features in 165 Lipomas: A Report of the CHAMP Study Group. Cancer Genetics and Cytogenetics, 1998, 102, 46-49.	1.0	60
158	Inflammatory leiomyosarcoma may be characterized by specific near-haploid chromosome changes. , 1998, 185, 112-115.		35
159	Cytogenetic analysis of 46 pleomorphic soft tissue sarcomas and correlation with morphologic and clinical features: A report of the CHAMP study group. , 1998, 22, 16-25.		161
160	Amplification and expression of theHMGIC gene in a benign endometrial polyp. Genes Chromosomes and Cancer, 1998, 22, 95-99.	1.5	35
161	Various regions within the alpha-helical domain of theCOL1A1 gene are fused to the second exon of thePDGFB gene in dermatofibrosarcomas and giant-cell fibroblastomas. , 1998, 23, 187-193.		158
162	HMGIY is the target of 6p21.3 rearrangements in various benign mesenchymal tumors. Genes Chromosomes and Cancer, 1998, 23, 279-285.	1.5	78

#	Article	IF	Citations
163	Localization and expression of the human estrogen receptor beta gene in uterine leiomyomata. , 1998, 23, 361-366.		45
164	Intracranial germ cell tumour (embryonal carcinoma with teratoma) with complex karyotype including isochromosome 12p. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 1998, 433, 571-574.	1.4	16
165	Clinical significance of cytogenetic abnormalities in uterine myomas. Fertility and Sterility, 1998, 69, 232-235.	0.5	80
166	Synovial Sarcoma of the Larynx and Hypopharynx. Annals of Otology, Rhinology and Laryngology, 1998, 107, 1080-1085.	0.6	82
167	Ten years of the cytogenetics of soft tissue tumors. Cancer Genetics and Cytogenetics, 1997, 95, 59-66.	1.0	24
168	The role of cytogenetics in the classification of soft tissue tumours. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 1997, 431, 83-94.	1.4	106
169	Additional evidence of a variant translocation t(12;22) withEWS/CHOP fusion in myxoid liposarcoma: clinicopathological features. , 1997, 182, 437-441.		102
170	Molecular delineation of the commonly deleted segment in mature B-cell lymphoid neoplasias with deletion of 7q. Genes Chromosomes and Cancer, 1997, 18, 147-150.	1.5	17
171	Molecular cytogenetic characterization of del(7q) in two uterine leiomyoma-derived cell lines. , 1997, 18, 155-161.		17
172	Hamartoma of the breast with involvement of 6p21 and rearrangement ofHMGIY. , 1997, 20, 90-92.		24
173	Identification of an enriched CD4+ CD8Î±++ CD8Î²+ T-cell subset among tumor-infiltrating lymphocytes in human renal cell carcinoma. , 1997, 71, 178-182.		12
174	Inflammatory Myofibroblastic Tumor of Bone. American Journal of Surgical Pathology, 1997, 21, 1166-1172.	2.1	80
175	Analysis of the karyotype and desoxyribonucleic acid content of uterine myomas in premenopausal, menopausal, and gonadotropin-releasing hormone agonist-treated females. Fertility and Sterility, 1996, 66, 376-379.	0.5	33
176	Mapping of the translocation breakpoints of primary pleomorphic adenomas and lipomas within a common region of chromosome 12. Cancer Genetics and Cytogenetics, 1996, 86, 39-45.	1.0	20
177	Trisomy 21 in solitary fibrous tumor. Cancer Genetics and Cytogenetics, 1996, 86, 58-60.	1.0	36
178	Multiple chromosome rearrangements in a fibrosarcoma. Cancer Genetics and Cytogenetics, 1996, 87, 176-178.	1.0	15
179	Trisomies 8 and 20 in desmoid tumors. Cancer Genetics and Cytogenetics, 1996, 92, 147-149.	1.0	56
180	Duplication of chromosome segment 12q15-24 is associated with atypical lipomatous tumors. A report of the CHAMP collaborative study group. , 1996, 67, 632-635.		34

#	Article	IF	Citations
181	Cytogenetic and immunohistochemical evidence that giant cell fibroblastoma is related to dermatofibrosarcoma protuberans. , 1996, 15, 73-75.		71
182	Observation of a deletion of the long arm of chromosome 6 in benign fibrocystic disease of the breast constitutes a challenging problem. , 1996, 16, 68-71.		4
183	Cytogenetic and immunohistochemical evidence that giant cell fibroblastoma is related to dermatofibrosarcoma protuberans. , 1996, 15, 73.		1
184	Combined Morphologic and Karyotypic Study of 28 Myxoid Liposarcomas. American Journal of Surgical Pathology, 1996, 20, 1047-1055.	2.1	107
185	Combined Morphologic and Karyotypic Study of 59 Atypical Lipomatous Tumors. American Journal of Surgical Pathology, 1996, 20, 1182-1189.	2.1	200
186	A new cytogenetic subgroup in uterine leiomyoma is characterized by a deletion of the long arm of chromosome 3. Genes Chromosomes and Cancer, 1995, 13, 219-220.	1.5	33
187	Chromosome abnormalities in benign prostatic hyperplasia. Genes Chromosomes and Cancer, 1994, 9, 227-233.	1.5	31
188	Some desmoid tumors are characterized by trisomy 8. Genes Chromosomes and Cancer, 1994, 10, 131-135.	1.5	49
189	New discriminative chromosomal marker in adipose tissue tumors. Cancer Genetics and Cytogenetics, 1994, 78, 232-235.	1.0	65
190	Minor myeloid component in Ph chromosome-positive acute lymphoblastic leukaemia: correlation with cytogenetic pattern and implication for poor response to therapy. British Journal of Haematology, 1994, 87, 515-522.	1.2	15
191	Physical Mapping of Chromosome 12q Breakpoints in Lipoma, Pleomorphic Salivary Gland Adenoma, Uterine Leiomyoma, and Myxoid Liposarcoma. Genomics, 1994, 20, 210-222.	1.3	68
192	Cytogenetics of tumours of the female reproductive tract. Reproductive Medicine Review, 1994, 3, 1-10.	0.3	0
193	Ring chromosome 6 as the only change in a thymoma. Genes Chromosomes and Cancer, 1993, 6, 243-244.	1.5	20
194	Rearrangement of 12q14-15 in pulmonary chondroid hamartoma. Genes Chromosomes and Cancer, 1993, 8, 131-133.	1.5	44
195	Cytogenetic and fluorescence in situ hybridization investigation of ring chromosomes characterizing a specific pathologic subgroup of adipose tissue tumors. Cancer Genetics and Cytogenetics, 1993, 68, 85-90.	1.0	167
196	Endometrial polyp: Another benign tumor characterized by 12q13â€“q15 changes. Cancer Genetics and Cytogenetics, 1993, 68, 32-33.	1.0	73
197	Clinical Review on Features and Cytogenetic Patterns in Adult Acute Myeloid Leukemia with Lymphoid Markers. Leukemia and Lymphoma, 1993, 9, 285-291.	0.6	11
198	The 6p21 chromosome region is nonrandomly involved in endometrial polyps. Gynecologic Oncology, 1992, 46, 393-396.	0.6	25

#	Article	IF	Citations
199	Numerical Chromosome Aberrations in Fibrothecoma. Tumori, 1992, 78, 140-142.	0.6	8
200	Chromosome changes in a case of hibernoma. Genes Chromosomes and Cancer, 1992, 5, 178-180.	1.5	26
201	Involvement of 6p in an endometrial polyp. Cancer Genetics and Cytogenetics, 1991, 51, 279-280.	1.0	16
202	Leiomyoma cells with 12q15 aberrations can be transformed in vitro and show a relatively stable karyotype during precrisis period. Cancer Genetics and Cytogenetics, 1991, 54, 223-228.	1.0	15
203	Trisomy 8 as the only chromosome change in an epithelioid smooth muscle tumor. Genes Chromosomes and Cancer, 1991, 3, 235-237.	1.5	8
204	Is t(6;20)(p21;q13) a characteristic chromosome change in endometrial polyps?. Genes Chromosomes and Cancer, 1991, 3, 318-319.	1.5	20
205	Rearrangements of immunoglobulin and TCR genes in lymphoid blast crisis of Ph + chronic myeloid leukaemia. British Journal of Haematology, 1990, 74, 414-419.	1.2	12
206	Is the chromosome band 1p36 another hot-spot for rearrangements in uterine leiomyoma?. Genes Chromosomes and Cancer, 1990, 2, 255-256.	1.5	22
207	Three possible cytogenetic subgroups of leiomyosarcoma. Cancer Genetics and Cytogenetics, 1989, 43, 39-49.	1.0	65
208	Chromosome Changes in Soft Tissue Tumors: Benign and Malignant. Cancer Investigation, 1989, 7, 63-76.	0.6	5
209	Renal Cortical Tumors: Cytogenetic Characterization. American Journal of Clinical Pathology, 1989, 92, 408-414.	0.4	48
210	Normal blood lymphocytes from patients with adipose tissue tumors with rearrangements at 12q13-q14 do not express the fragile site fra(12)(q13.1). Cancer Genetics and Cytogenetics, 1988, 31, 35-39.	1.0	3
211	Breakpoints in benign lipoma may be at 12q13 or 12q14. Cancer Genetics and Cytogenetics, 1988, 36, 131-135.	1.0	38
212	Recurrent breakpoints at 9q31 and 22q12.2 in extraskeletal myxoid chondrosarcoma. Cancer Genetics and Cytogenetics, 1988, 30, 145-150.	1.0	97
213	Consistent chromosome changes in leiomyosarcoma. Cancer Genetics and Cytogenetics, 1988, 35, 47-50.	1.0	37
214	Further Evidence that Renal Oncocytoma has Malignant Potential. Journal of Urology, 1988, 139, 585-587.	0.2	66
215	Nonrandom translocation in extraskeletal myxoid chondrosarcoma. Cancer Genetics and Cytogenetics, 1987, 26, 377.	1.0	11
216	Cytogenetic studies of adipose tissue tumors. I. A benign lipoma with reciprocal translocation t(3;12)(q28;q14). Cancer Genetics and Cytogenetics, 1986, 23, 283-289.	1.0	117

#	Article	IF	Citations
217	Cytogenetic studies of adipose tissue tumors. II. Recurrent reciprocal translocation t(12;16)(q13;p11) in myxoid liposarcomas. Cancer Genetics and Cytogenetics, 1986, 23, 291-299.	1.0	235
218	Recurrent chromosome translocations in liposarcoma. Cancer Genetics and Cytogenetics, 1986, 22, 93-94.	1.0	57
219	Translocations involving the X chromosome in solid tumors: Presentation of two sarcomas with t(X;18)(q13;p11). Cancer Genetics and Cytogenetics, 1986, 23, 87-91.	1.0	98
220	Translocation X;18 in synovial sarcoma. Cancer Genetics and Cytogenetics, 1986, 23, 93.	1.0	143