Paola Dal Cin

List of Publications by Year
in descending order

221	18,493	11651 70	13379 130
papers	citations	h-index	g-index

224	224	224	12396
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	International Consensus Classification of Myeloid Neoplasms and Acute Leukemias: integrating morphologic, clinical, and genomic data. Blood, 2022, 140, 1200-1228.	1.4	814
2	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.	1.4	793
3	Molecular profiling of diffuse large B-cell lymphoma identifies robust subtypes including one characterized by host inflammatory response. Blood, 2005, 105, 1851-1861.	1.4	778
4	Crizotinib inALK-Rearranged Inflammatory Myofibroblastic Tumor. New England Journal of Medicine, 2010, 363, 1727-1733.	27.0	769
5	TPM3-ALK and TPM4-ALK Oncogenes in Inflammatory Myofibroblastic Tumors. American Journal of Pathology, 2000, 157, 377-384.	3.8	659
6	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.	5.9	562
7	Loss of INI1 Expression is Characteristic of Both Conventional and Proximal-type Epithelioid Sarcoma. American Journal of Surgical Pathology, 2009, 33, 542-550.	3.7	538
8	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.	2.8	443
9	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
10	Predominance of beta-catenin mutations and beta-catenin dysregulation in sporadic aggressive fibromatosis (desmoid tumor). Oncogene, 1999, 18, 6615-6620.	5.9	339
11	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
12	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.	7.0	305
13	Clonal evolution in patients with chronic lymphocytic leukaemia developing resistance to BTK inhibition. Nature Communications, 2016, 7, 11589.	12.8	285
14	USP6 (Tre2) Fusion Oncogenes in Aneurysmal Bone Cyst. Cancer Research, 2004, 64, 1920-1923.	0.9	284
15	EWSR1â€CREB1 is the predominant gene fusion in angiomatoid fibrous histiocytoma. Genes Chromosomes and Cancer, 2007, 46, 1051-1060.	2.8	276
16	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.	2.8	276
17	The Clinicopathologic Features of YWHAE-FAM22 Endometrial Stromal Sarcomas. American Journal of Surgical Pathology, 2012, 36, 641-653.	3.7	265
18	Coordinated expression and amplification of theMDM2,CDK4, andHMGI-C genes in atypical lipomatous tumours. Journal of Pathology, 2000, 190, 531-536.	4.5	250

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19	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
20	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
21	MUC4 Is a Sensitive and Extremely Useful Marker for Sclerosing Epithelioid Fibrosarcoma. American Journal of Surgical Pathology, 2012, 36, 1444-1451.	3.7	230
22	Aneurysmal bone cyst variant translocations upregulate USP6 transcription by promoter swapping with the ZNF9, COL1A1, TRAP150, and OMD genes. Oncogene, 2005, 24, 3419-3426.	5.9	226
23	Combined Morphologic and Karyotypic Study of 59 Atypical Lipomatous Tumors. American Journal of Surgical Pathology, 1996, 20, 1182-1189.	3.7	200
24	NSD3â€“NUT Fusion Oncoprotein in NUT Midline Carcinoma: Implications for a Novel Oncogenic Mechanism. Cancer Discovery, 2014, 4, 928-941.	9.4	192
25	BRD4 Bromodomain Gene Rearrangement in Aggressive Carcinoma with Translocation t(15;19). American Journal of Pathology, 2001, 159, 1987-1992.	3.8	188
26	Abnormal Nuclear Shape in Solid Tumors Reflects Mitotic Instability. American Journal of Pathology, 2001, 158, 199-206.	3.8	187
27	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
28	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
29	Perivascular Epithelioid Cell Neoplasm (PEComa) of the Gynecologic Tract. American Journal of Surgical Pathology, 2014, 38, 176-188.	3.7	165
30	Cytogenetic analysis of 46 pleomorphic soft tissue sarcomas and correlation with morphologic and clinical features: A report of the CHAMP study group. Genes Chromosomes and Cancer, 1998, 22, 16-25.	2.8	161
31	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. Genes Chromosomes and Cancer, 1998, 23, 187-193.	2.8	158
32	Molecular pathogenesis of uterine smooth muscle tumors from transcriptional profiling. Genes Chromosomes and Cancer, 2004, 40, 97-108.	2.8	145
33	Translocation X;18 in synovial sarcoma. Cancer Genetics and Cytogenetics, 1986, 23, 93.	1.0	143
34	Cytogenetic characterization of peripheral nerve sheath tumours: a report of the CHAMP study group. , 2000, 190, 31-38.		141
35	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.	2.8	137
36	Immunohistochemical Detection of MYC-driven Diffuse Large B-Cell Lymphomas. PLoS ONE, 2012, 7, e33813.	2.5	137

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37	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.	3.7	134
38	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.	3.7	133
39	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.	4.5	131
40	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.	2.8	129
41	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
42	PLAG1 Alterations in Lipoblastoma. American Journal of Pathology, 2001, 159, 955-962.	3.8	117
43	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
44	Correlation between Clinicopathological Features and Karyotype in Spindle Cell Sarcomas. American Journal of Pathology, 1999, 154, 1841-1847.	3.8	109
45	The structure and dynamics of ring chromosomes in human neoplastic and non-neoplastic cells. Human Genetics, 1999, 104, 315-325.	3.8	108
46	Clinical impact of molecular and cytogenetic findings in synovial sarcoma. Genes Chromosomes and Cancer, 2001, 31, 362-372.	2.8	108
47	Inflammatory Myofibroblastic Tumor of the Uterus. American Journal of Surgical Pathology, 2015, 39, 157-168.	3.7	107
48	Combined Morphologic and Karyotypic Study of 28 Myxoid Liposarcomas. American Journal of Surgical Pathology, 1996, 20, 1047-1055.	3.7	107
49	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.	2.8	106
50	Uterine Leiomyomata with t(10;17) Disrupt the Histone Acetyltransferase MORF. Cancer Research, 2004, 64, 5570-5577.	0.9	106
51	Molecular genetic characterization of theEWS/CHNandRBP56/CHNfusion genes in extraskeletal myxoid chondrosarcoma. Genes Chromosomes and Cancer, 2002, 35, 340-352.	2.8	104
52	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.	7.1	104
53	Cutaneous Syncytial Myoepithelioma. American Journal of Surgical Pathology, 2013, 37, 710-718.	3.7	103
54	Additional evidence of a variant translocation t(12;22) withEWS/CHOP fusion in myxoid liposarcoma: clinicopathological features. Journal of Pathology, 1997, 182, 437-441.	4.5	102

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55	Upper respiratory tract carcinoma with chromosomal translocation 15;19. Cancer, 2001, 92, 1195-1203.	4.1	102
56	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.	5.5	101
57	Cytogenetic, Clinical, and Morphologic Correlations in 78 Cases of Fibromatosis: A Report from the CHAMP Study Group. Modern Pathology, 2000, 13, 1080-1085.	5.5	99
58	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
59	Recurrent breakpoints at 9q31 and 22q12.2 in extraskeletal myxoid chondrosarcoma. Cancer Genetics and Cytogenetics, 1988, 30, 145-150.	1.0	97
60	Distinctive Cytogenetic Profile in Benign Metastasizing Leiomyoma: Pathogenetic Implications. American Journal of Surgical Pathology, 2007, 31, 737-743.	3.7	94
61	Variant translocations involving 16q22 and 17p13 in solid variant and extraosseous forms of aneurysmal bone cyst. Genes Chromosomes and Cancer, 2000, 28, 233-234.	2.8	88
62	TP53 mutation defines a unique subgroup within complex karyotype deÂnovo and therapy-related MDS/AML. Blood Advances, 2022, 6, 2847-2853.	5.2	87
63	Expression of ROS1 predicts ROS1 gene rearrangement in inflammatory myofibroblastic tumors. Modern Pathology, 2015, 28, 732-739.	5.5	85
64	Synovial Sarcoma of the Larynx and Hypopharynx. Annals of Otology, Rhinology and Laryngology, 1998, 107, 1080-1085.	1.1	82
65	Wellâ€differentiated and dedifferentiated liposarcomas with prominent myxoid stroma: analysis of 56 cases. Histopathology, 2013, 62, 287-293.	2.9	82
66	Clinical significance of cytogenetic abnormalities in uterine myomas. Fertility and Sterility, 1998, 69, 232-235.	1.0	80
67	Inflammatory Myofibroblastic Tumor of Bone. American Journal of Surgical Pathology, 1997, 21, 1166-1172.	3.7	80
68	Fusion transcripts involving HMGA2 are not a common molecular mechanism in uterine leiomyomata with rearrangements in 12q15. Cancer Research, 2003, 63, 1351-8.	0.9	79
69	HMGIY is the target of 6p21.3 rearrangements in various benign mesenchymal tumors. Genes Chromosomes and Cancer, 1998, 23, 279-285.	2.8	78
70	Integrative Genomic Analysis Implicates Gain of PIK3CA at 3q26 and MYC at 8q24 in Chronic Lymphocytic Leukemia. Clinical Cancer Research, 2012, 18, 3791-3802.	7.0	76
71	Endometrial polyp: Another benign tumor characterized by 12q13â€“q15 changes. Cancer Genetics and Cytogenetics, 1993, 68, 32-33.	1.0	73
72	Cytogenetic and immunohistochemical evidence that giant cell fibroblastoma is related to dermatofibrosarcoma protuberans. , 1996, 15, 73-75.		71

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73	Intravenous leiomyomatosis is characterized by a der(14)t(12;14)(q15;q24). Genes Chromosomes and Cancer, 2003, 36, 205-206.	2.8	69
74	Physical Mapping of Chromosome 12q Breakpoints in Lipoma, Pleomorphic Salivary Gland Adenoma, Uterine Leiomyoma, and Myxoid Liposarcoma. Genomics, 1994, 20, 210-222.	2.9	68
75	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
76	Disseminated peritoneal leiomyomatosis after laparoscopic supracervical hysterectomy with characteristic molecular cytogenetic findings of uterine leiomyoma. Genes Chromosomes and Cancer, 2010, 49, 1152-1160.	2.8	67
77	Integrated single-cell genetic and transcriptional analysis suggests novel drivers of chronic lymphocytic leukemia. Genome Research, 2017, 27, 1300-1311.	5.5	67
78	Further Evidence that Renal Oncocytoma has Malignant Potential. Journal of Urology, 1988, 139, 585-587.	0.4	66
79	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.	2.5	66
80	Three possible cytogenetic subgroups of leiomyosarcoma. Cancer Genetics and Cytogenetics, 1989, 43, 39-49.	1.0	65
81	New discriminative chromosomal marker in adipose tissue tumors. Cancer Genetics and Cytogenetics, 1994, 78, 232-235.	1.0	65
82	Intravenous leiomyomatosis: an unusual intermediate between benign and malignant uterine smooth muscle tumors. Modern Pathology, 2016, 29, 500-510.	5.5	65
83	Primary Sclerosing Epithelioid Fibrosarcoma of Bone. American Journal of Surgical Pathology, 2014, 38, 1538-1544.	3.7	64
84	Uterine leiomyomata with deletions of 1p represent a distinct cytogenetic subgroup associated with unusual histologic features. Genes Chromosomes and Cancer, 2006, 45, 304-312.	2.8	62
85	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
86	Comparative Cytogenetic Study of Spindle Cell and Pleomorphic Leiomyosarcomas of Soft Tissues. Cancer Genetics and Cytogenetics, 2000, 116, 66-73.	1.0	60
87	HMGA2 Rearrangement in a Case of Vulvar Aggressive Angiomyxoma. International Journal of Gynecological Pathology, 2006, 25, 403-407.	1.4	60
88	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
89	Recurrent chromosome translocations in liposarcoma. Cancer Genetics and Cytogenetics, 1986, 22, 93-94.	1.0	57
90	Trisomies 8 and 20 in desmoid tumors. Cancer Genetics and Cytogenetics, 1996, 92, 147-149.	1.0	56

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91	CIC–NUTM1 fusion: A case which expands the spectrum of NUT–rearranged epithelioid malignancies. Genes Chromosomes and Cancer, 2018, 57, 446-451.	2.8	53
92	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
93	Some desmoid tumors are characterized by trisomy 8. Genes Chromosomes and Cancer, 1994, 10, 131-135.	2.8	49
94	Amplification ofAML1 in childhood acute lymphoblastic leukemias. Genes Chromosomes and Cancer, 2001, 30, 407-409.	2.8	49
95	Renal Cortical Tumors: Cytogenetic Characterization. American Journal of Clinical Pathology, 1989, 92, 408-414.	0.7	48
96	Dedifferentiated liposarcoma and pleomorphic liposarcoma. Cancer Cytopathology, 2014, 122, 128-137.	2.4	47
97	Localization and expression of the human estrogen receptor beta gene in uterine leiomyomata. , 1998, 23, 361-366.		45
98	Fusion of the genes BRD8 and PHF1 in endometrial stromal sarcoma. Genes Chromosomes and Cancer, 2017, 56, 841-845.	2.8	45
99	Rearrangement of 12q14-15 in pulmonary chondroid hamartoma. Genes Chromosomes and Cancer, 1993, 8, 131-133.	2.8	44
100	Fluorescence in situ hybridization is a useful ancillary diagnostic tool for extraskeletal myxoid chondrosarcoma. Modern Pathology, 2008, 21, 1303-1310.	5.5	44
101	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
102	Promiscuous genes involved in recurrent chromosomal translocations in soft tissue tumours. Pathology, 2014, 46, 105-112.	0.6	40
103	Detection of activatingMAP2K1mutations in atypical hairy cell leukemia and hairy cell leukemia variant. Leukemia and Lymphoma, 2017, 58, 233-236.	1.3	39
104	Breakpoints in benign lipoma may be at 12q13 or 12q14. Cancer Genetics and Cytogenetics, 1988, 36, 131-135.	1.0	38
105	Consistent chromosome changes in leiomyosarcoma. Cancer Genetics and Cytogenetics, 1988, 35, 47-50.	1.0	37
106	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.	7.0	37
107	Trisomy 21 in solitary fibrous tumor. Cancer Genetics and Cytogenetics, 1996, 86, 58-60.	1.0	36
108	Molecular cytogenetic definition of three distinct chromosome arm 14q deletion intervals in gastrointestinal stromal tumors. Genes Chromosomes and Cancer, 2001, 32, 26-32.	2.8	36

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109	Inflammatory leiomyosarcoma may be characterized by specific near-haploid chromosome changes. , 1998, 185, 112-115.		35
110	Amplification and expression of theHMGIC gene in a benign endometrial polyp. Genes Chromosomes and Cancer, 1998, 22, 95-99.	2.8	35
111	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
112	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.	3.5	34
113	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.	2.8	33
114	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.	1.0	33
115	Analysis of MDM2 Amplification in 43 Endometrial Stromal Tumors. International Journal of Gynecological Pathology, 2015, 34, 576-583.	1.4	33
116	Adult Renal Cell Carcinoma. Surgical Pathology Clinics, 2015, 8, 587-621.	1.7	33
117	Chromosome abnormalities in benign prostatic hyperplasia. Genes Chromosomes and Cancer, 1994, 9, 227-233.	2.8	31
118	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
119	MYC Immunohistochemistry to Identify MYC-Driven B-Cell Lymphomas in Clinical Practice. American Journal of Clinical Pathology, 2016, 145, 166-179.	0.7	29
120	Guiding the global evolution of cytogenetic testing for hematologic malignancies. Blood, 2022, 139, 2273-2284.	1.4	29
121	Targeted FGFR inhibition results in a durable remission in an FGFR1-driven myeloid neoplasm with eosinophilia. Blood Advances, 2020, 4, 3136-3140.	5.2	28
122	Carbonic anhydrase IX (CA9) expression in multiple renal epithelial tumour subtypes. Histopathology, 2020, 77, 659-666.	2.9	28
123	Inflammatory Myofibroblastic Tumor with HMGIC Rearrangement. Cancer Genetics and Cytogenetics, 1999, 112, 156-160.	1.0	27
124	Chromosome changes in a case of hibernoma. Genes Chromosomes and Cancer, 1992, 5, 178-180.	2.8	26
125	The 6p21 chromosome region is nonrandomly involved in endometrial polyps. Gynecologic Oncology, 1992, 46, 393-396.	1.4	25
126	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

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127	Ten years of the cytogenetics of soft tissue tumors. Cancer Genetics and Cytogenetics, 1997, 95, 59-66.	1.0	24
128	Hamartoma of the breast with involvement of 6p21 and rearrangement ofHMGIY. , 1997, 20, 90-92.		24
129	Clonal chromosome abnormalities in a so-called Dupuytren's subungual exostosis. , 1999, 24, 162-164.		24
130	Molecular and Cytogenetic Characterization of Plexiform Leiomyomata Provide Further Evidence for Genetic Heterogeneity Underlying Uterine Fibroids. American Journal of Pathology, 2008, 172, 1403-1410.	3.8	24
131	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.	4.1	24
132	Clinical response to larotrectinib in adult Philadelphia chromosomeâ€“like ALL with cryptic ETV6-NTRK3 rearrangement. Blood Advances, 2020, 4, 106-111.	5.2	23
133	Is the chromosome band 1p36 another hot-spot for rearrangements in uterine leiomyoma?. Genes Chromosomes and Cancer, 1990, 2, 255-256.	2.8	22
134	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.	5.5	22
135	Embryonal Rhabdomyosarcoma with Only Numerical Chromosome Changes. Cancer Genetics and Cytogenetics, 1999, 109, 161-165.	1.0	21
136	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
137	Urachal inflammatory myofibroblastic tumor with ALK gene rearrangement: a study of urachal remnants. Urology, 2004, 64, 140-144.	1.0	21
138	Is t(6;20)(p21;q13) a characteristic chromosome change in endometrial polyps?. Genes Chromosomes and Cancer, 1991, 3, 318-319.	2.8	20
139	Ring chromosome 6 as the only change in a thymoma. Genes Chromosomes and Cancer, 1993, 6, 243-244.	2.8	20
140	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
141	Precancerous Lesions in the Kidney. Scandinavian Journal of Urology and Nephrology, 2000, 34, 136-165.	1.4	20
142	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
143	Multiple EWSR1-WT1 and WT1-EWSR1 copies in two cases of desmoplastic round cell tumor. Cancer Genetics, 2013, 206, 387-392.	0.4	19
144	Controversial fluorescence inÂsitu hybridization cytogenetic abnormalities in chronic lymphocytic leukaemia: new insights from a large cohort. British Journal of Haematology, 2015, 170, 694-703.	2.5	19

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145	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.	2.8	17
146	Molecular cytogenetic characterization of del(7q) in two uterine leiomyoma-derived cell lines. , 1997, 18, 155-161.		17
147	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
148	Involvement of 6p in an endometrial polyp. Cancer Genetics and Cytogenetics, 1991, 51, 279-280.	1.0	16
149	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.	2.8	16
150	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
151	Minor myeloid component in Ph chromosomeâ€positive acute lymphoblastic leukaemia: correlation with cytogenetic pattern and implication for poor response to therapy. British Journal of Haematology, 1994, 87, 515-522.	2.5	15
152	Multiple chromosome rearrangements in a fibrosarcoma. Cancer Genetics and Cytogenetics, 1996, 87, 176-178.	1.0	15
153	HMGA1 and HMGA2 rearrangements in massâ€forming endometriosis. Genes Chromosomes and Cancer, 2010, 49, 630-634.	2.8	15
154	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.	4.1	14
155	Nuclear Protein in Testis Midline Carcinoma Misdiagnosed As Adamantinoma. Journal of Clinical Oncology, 2014, 32, e57-e60.	1.6	13
156	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.	1.2	13
157	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.	1.4	13
158	Rearrangements of immunoglobulin and TCR genes in lymphoid blast crisis of Ph + chronic myeloid leukaemia. British Journal of Haematology, 1990, 74, 414-419.	2.5	12
159	Identification of an enriched CD4+ CD8Î±++ CD8Î²+ T-cell subset among tumor-infiltrating lymphocytes in human renal cell carcinoma. , 1997, 71, 178-182.		12
160	Involvement of theHMGI(Y) gene in a microfollicular adenoma of the thyroid. Genes Chromosomes and Cancer, 1999, 24, 286-289.	2.8	12
161	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.	1.3	12
162	Pseudosarcomatous myofibroblastic proliferations of the urinary bladder are neoplasms characterized by recurrent FN1â€“ALK fusions. Modern Pathology, 2021, 34, 469-477.	5.5	12

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163	Nonrandom translocation in extraskeletal myxoid chondrosarcoma. Cancer Genetics and Cytogenetics, 1987, 26, 377.	1.0	11
164	Clinical Review on Features and Cytogenetic Patterns in Adult Acute Myeloid Leukemia with Lymphoid Markers. Leukemia and Lymphoma, 1993, 9, 285-291.	1.3	11
165	Pericentric inversion (12)(p12q13âˆ¼q14) as the sole chromosomal abnormality in a leiomyoma of the vulva. Cancer Genetics and Cytogenetics, 2010, 199, 21-23.	1.0	11
166	Expanding the spectrum of translocations in sclerosing epitheloid fibrosarcoma: A new case with EWSR1â€CREB3L3 fusion. Genes Chromosomes and Cancer, 2018, 57, 675-677.	2.8	11
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