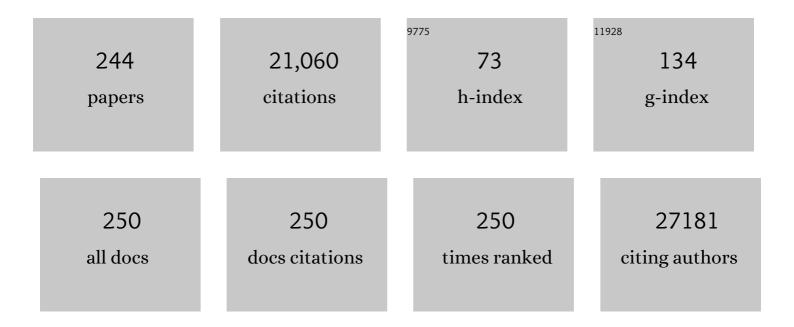
Britta Weigelt

List of Publications by Year in descending order

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#	Article	lF	CITATIONS
1	Triple-Negative Breast Cancer. New England Journal of Medicine, 2010, 363, 1938-1948.	13.9	3,233
2	Mutation tracking in circulating tumor DNA predicts relapse in early breast cancer. Science Translational Medicine, 2015, 7, 302ra133.	5.8	889
3	Clinical validity of circulating tumour cells in patients with metastatic breast cancer: a pooled analysis of individual patient data. Lancet Oncology, The, 2014, 15, 406-414.	5.1	703
4	The Genomic Landscape of Endocrine-Resistant Advanced Breast Cancers. Cancer Cell, 2018, 34, 427-438.e6.	7.7	633
5	Cerebrospinal fluid-derived circulating tumour DNA better represents the genomic alterations of brain tumours than plasma. Nature Communications, 2015, 6, 8839.	5.8	605
6	High-intensity sequencing reveals the sources of plasma circulating cell-free DNA variants. Nature Medicine, 2019, 25, 1928-1937.	15.2	485
7	Classification of endometrial carcinoma: more than two types. Lancet Oncology, The, 2014, 15, e268-e278.	5.1	479
8	Germline mutations in RAD51D confer susceptibility to ovarian cancer. Nature Genetics, 2011, 43, 879-882.	9.4	460
9	Histological and molecular types of breast cancer: is there a unifying taxonomy?. Nature Reviews Clinical Oncology, 2009, 6, 718-730.	12.5	353
10	Phyllodes tumours of the breast: a consensus review. Histopathology, 2016, 68, 5-21.	1.6	329
11	Loss of the FAT1 Tumor Suppressor Promotes Resistance to CDK4/6 Inhibitors via the Hippo Pathway. Cancer Cell, 2018, 34, 893-905.e8.	7.7	307
12	Breast cancer intra-tumor heterogeneity. Breast Cancer Research, 2014, 16, 210.	2.2	256
13	Pan-cancer analysis of intratumor heterogeneity as a prognostic determinant of survival. Oncotarget, 2016, 7, 10051-10063.	0.8	247
14	Genomic characterization of metastatic patterns from prospective clinical sequencing of 25,000 patients. Cell, 2022, 185, 563-575.e11.	13.5	223
15	Metaplastic breast carcinomas are basal-like breast cancers: a genomic profiling analysis. Breast Cancer Research and Treatment, 2009, 117, 273-280.	1.1	208
16	The clinical use of circulating tumor cells (CTCs) enumeration for staging of metastatic breast cancer (MBC): International expert consensus paper. Critical Reviews in Oncology/Hematology, 2019, 134, 39-45.	2.0	200
17	Diverse <i>BRCA1</i> and <i>BRCA2</i> Reversion Mutations in Circulating Cell-Free DNA of Therapy-Resistant Breast or Ovarian Cancer. Clinical Cancer Research, 2017, 23, 6708-6720.	3.2	194
18	Hotspot activating PRKD1 somatic mutations in polymorphous low-grade adenocarcinomas of the salivary glands. Nature Genetics, 2014, 46, 1166-1169.	9.4	188

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19	Molecular analysis reveals a genetic basis for the phenotypic diversity of metaplastic breast carcinomas. Journal of Pathology, 2010, 220, 562-573.	2.1	185
20	Diverse alterations associated with resistance to KRAS(G12C) inhibition. Nature, 2021, 599, 679-683.	13.7	183
21	Pan-cancer analysis of bi-allelic alterations in homologous recombination DNA repair genes. Nature Communications, 2017, 8, 857.	5.8	182
22	Clinicopathological analysis of endometrial carcinomas harboring somatic POLE exonuclease domain mutations. Modern Pathology, 2015, 28, 505-514.	2.9	180
23	The genetic landscape of endometrial clear cell carcinomas. Journal of Pathology, 2017, 243, 230-241.	2.1	168
24	Massively Parallel Sequencing-Based Clonality Analysis of Synchronous Endometrioid Endometrial and Ovarian Carcinomas. Journal of the National Cancer Institute, 2015, 108, djv427.	3.0	164
25	Comprehensive Molecular Characterization of Salivary Duct Carcinoma Reveals Actionable Targets and Similarity to Apocrine Breast Cancer. Clinical Cancer Research, 2016, 22, 4623-4633.	3.2	153
26	A recurrent neomorphic mutation in MYOD1 defines a clinically aggressive subset of embryonal rhabdomyosarcoma associated with PI3K-AKT pathway mutations. Nature Genetics, 2014, 46, 595-600.	9.4	152
27	HER2-Mediated Internalization of Cytotoxic Agents in <i>ERBB2</i> Amplified or Mutant Lung Cancers. Cancer Discovery, 2020, 10, 674-687.	7.7	149
28	An approach to suppress the evolution of resistance in BRAFV600E-mutant cancer. Nature Medicine, 2017, 23, 929-937.	15.2	146
29	Adenoid cystic carcinomas constitute a genomically distinct subgroup of tripleâ€negative and basalâ€like breast cancers. Journal of Pathology, 2012, 226, 84-96.	2.1	144
30	Distinct Classes of Complex Structural Variation Uncovered across Thousands of Cancer Genome Graphs. Cell, 2020, 183, 197-210.e32.	13.5	141
31	ARID1A determines luminal identity and therapeutic response in estrogen-receptor-positive breast cancer. Nature Genetics, 2020, 52, 198-207.	9.4	140
32	Mucinous carcinoma of the breast is genomically distinct from invasive ductal carcinomas of no special type. Journal of Pathology, 2010, 222, 282-298.	2.1	139
33	In situ single-cell analysis identifies heterogeneity for PIK3CA mutation and HER2 amplification in HER2-positive breast cancer. Nature Genetics, 2015, 47, 1212-1219.	9.4	139
34	Metastasis and Immune Evasion from Extracellular cGAMP Hydrolysis. Cancer Discovery, 2021, 11, 1212-1227.	7.7	139
35	Unraveling tumor–immune heterogeneity in advanced ovarian cancer uncovers immunogenic effect of chemotherapy. Nature Genetics, 2020, 52, 582-593.	9.4	136
36	Genetic hallmarks of recurrent/metastatic adenoid cystic carcinoma. Journal of Clinical Investigation, 2019, 129, 4276-4289.	3.9	134

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37	Genomic landscape of adenoid cystic carcinoma of the breast. Journal of Pathology, 2015, 237, 179-189.	2.1	133
38	Pathogenesis of Triple-Negative Breast Cancer. Annual Review of Pathology: Mechanisms of Disease, 2022, 17, 181-204.	9.6	132
39	The Landscape of Somatic Genetic Alterations in Metaplastic Breast Carcinomas. Clinical Cancer Research, 2017, 23, 3859-3870.	3.2	129
40	Triple-negative breast cancer: the importance of molecular and histologic subtyping, and recognition of low-grade variants. Npj Breast Cancer, 2016, 2, 16036.	2.3	127
41	The Genomic Landscape of Male Breast Cancers. Clinical Cancer Research, 2016, 22, 4045-4056.	3.2	119
42	The Spectrum of Triple-Negative Breast Disease. American Journal of Pathology, 2017, 187, 2139-2151.	1.9	118
43	Benchmarking mutation effect prediction algorithms using functionally validated cancer-related missense mutations. Genome Biology, 2014, 15, 484.	3.8	117
44	Mutations in BRCA1 and BRCA2 differentially affect the tumor microenvironment and response to checkpoint blockade immunotherapy. Nature Cancer, 2020, 1, 1188-1203.	5.7	114
45	Whole-genome single-cell copy number profiling from formalin-fixed paraffin-embedded samples. Nature Medicine, 2017, 23, 376-385.	15.2	111
46	Mucinous and neuroendocrine breast carcinomas are transcriptionally distinct from invasive ductal carcinomas of no special type. Modern Pathology, 2009, 22, 1401-1414.	2.9	110
47	Intra-tumor genetic heterogeneity and alternative driver genetic alterations in breast cancers with heterogeneous HER2 gene amplification. Genome Biology, 2015, 16, 107.	3.8	109
48	Recurrent hotspot mutations in HRAS Q61 and PI3K-AKT pathway genes as drivers of breast adenomyoepitheliomas. Nature Communications, 2018, 9, 1816.	5.8	105
49	FOXA1 upregulation promotes enhancer and transcriptional reprogramming in endocrine-resistant breast cancer. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 26823-26834.	3.3	103
50	Massively parallel sequencing of phyllodes tumours of the breast reveals actionable mutations, and <i><scp>TERT</scp></i> promoter hotspot mutations and <i>TERT</i> gene amplification as likely drivers of progression. Journal of Pathology, 2016, 238, 508-518.	2.1	102
51	A survey of DICER1 hotspot mutations in ovarian and testicular sex cord-stromal tumors. Modern Pathology, 2015, 28, 1603-1612.	2.9	100
52	Clinical Utility of Prospective Molecular Characterization in Advanced Endometrial Cancer. Clinical Cancer Research, 2018, 24, 5939-5947.	3.2	100
53	<i>IDH2</i> Mutations Define a Unique Subtype of Breast Cancer with Altered Nuclear Polarity. Cancer Research, 2016, 76, 7118-7129.	0.4	99
54	Precision Radiotherapy: Reduction in Radiation for Oropharyngeal Cancer in the 30 ROC Trial. Journal of the National Cancer Institute, 2021, 113, 742-751.	3.0	98

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55	HER2-Enriched Subtype and ERBB2 Expression in HER2-Positive Breast Cancer Treated with Dual HER2 Blockade. Journal of the National Cancer Institute, 2020, 112, 46-54.	3.0	97
56	Next-Generation Assessment of Human Epidermal Growth Factor Receptor 2 (ERBB2) Amplification Status. Journal of Molecular Diagnostics, 2017, 19, 244-254.	1.2	96
57	Alterations in PTEN and ESR1 promote clinical resistance to alpelisib plus aromatase inhibitors. Nature Cancer, 2020, 1, 382-393.	5.7	96
58	Genetic alterations of triple negative breast cancer by targeted next-generation sequencing and correlation with tumor morphology. Modern Pathology, 2016, 29, 476-488.	2.9	95
59	Uterine adenosarcomas are mesenchymal neoplasms. Journal of Pathology, 2016, 238, 381-388.	2.1	94
60	A novel representation of inter-site tumour heterogeneity from pre-treatment computed tomography textures classifies ovarian cancers by clinical outcome. European Radiology, 2017, 27, 3991-4001.	2.3	92
61	Loss-of-function mutations in ATP6AP1 and ATP6AP2 in granular cell tumors. Nature Communications, 2018, 9, 3533.	5.8	92
62	Genetic Heterogeneity in Therapy-NaÃ⁻ve Synchronous Primary Breast Cancers and Their Metastases. Clinical Cancer Research, 2017, 23, 4402-4415.	3.2	91
63	The value of cellâ€free DNA for molecular pathology. Journal of Pathology, 2018, 244, 616-627.	2.1	91
64	The Landscape of Somatic Genetic Alterations in Breast Cancers From ATM Germline Mutation Carriers. Journal of the National Cancer Institute, 2018, 110, 1030-1034.	3.0	90
65	An immune stratification reveals a subset of PD-1/LAG-3 double-positive triple-negative breast cancers. Breast Cancer Research, 2016, 18, 121.	2.2	89
66	TP53 Mutational Spectrum in Endometrioid and Serous Endometrial Cancers. International Journal of Gynecological Pathology, 2016, 35, 289-300.	0.9	89
67	Characterization of the genomic features and expressed fusion genes in micropapillary carcinomas of the breast. Journal of Pathology, 2014, 232, 553-565.	2.1	88
68	The molecular basis of breast cancer pathological phenotypes. Journal of Pathology, 2017, 241, 375-391.	2.1	86
69	HER2 Reactivation through Acquisition of the HER2 L755S Mutation as a Mechanism of Acquired Resistance to HER2-targeted Therapy in HER2+ Breast Cancer. Clinical Cancer Research, 2017, 23, 5123-5134.	3.2	85
70	Loss of 16q in high grade breast cancer is associated with estrogen receptor status: Evidence for progression in tumors with a luminal phenotype?. Genes Chromosomes and Cancer, 2009, 48, 351-365.	1.5	80
71	Breast Cancer Genomics From Microarrays to Massively Parallel Sequencing: Paradigms and New Insights. Journal of the National Cancer Institute, 2015, 107, .	3.0	80
72	Metastatic breast carcinomas display genomic and transcriptomic heterogeneity. Modern Pathology, 2015, 28, 340-351.	2.9	80

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73	Small cell carcinoma of the gynecologic tract: A multifaceted spectrum of lesions. Gynecologic Oncology, 2014, 134, 410-418.	0.6	79
74	<i>MED12</i> somatic mutations in fibroadenomas and phyllodes tumours of the breast. Histopathology, 2015, 67, 719-729.	1.6	78
75	Mesothelin Expression in Triple Negative Breast Carcinomas Correlates Significantly with Basal-Like Phenotype, Distant Metastases and Decreased Survival. PLoS ONE, 2014, 9, e114900.	1.1	77
76	Reliability of Whole-Exome Sequencing for Assessing Intratumor Genetic Heterogeneity. Cell Reports, 2018, 25, 1446-1457.	2.9	76
77	<i>MYBL1</i> rearrangements and <i>MYB</i> amplification in breast adenoid cystic carcinomas lacking the <i>MYB</i> – <i>NFIB</i> fusion gene. Journal of Pathology, 2018, 244, 143-150.	2.1	74
78	Functional screening identifies <scp>MCT4</scp> as a key regulator of breast cancer cell metabolism and survival. Journal of Pathology, 2015, 237, 152-165.	2.1	73
79	Low PTEN levels and PIK3CA mutations predict resistance to neoadjuvant lapatinib and trastuzumab without chemotherapy in patients with HER2 over-expressing breast cancer. Breast Cancer Research and Treatment, 2018, 167, 731-740.	1.1	71
80	Metaplastic breast carcinoma: more than a special type. Nature Reviews Cancer, 2014, 14, 147-148.	12.8	69
81	Genetic events in the progression of adenoid cystic carcinoma of the breast to high-grade triple-negative breast cancer. Modern Pathology, 2016, 29, 1292-1305.	2.9	68
82	The Genomic Landscape of Mucinous Breast Cancer. Journal of the National Cancer Institute, 2019, 111, 737-741.	3.0	68
83	INK4 Tumor Suppressor Proteins Mediate Resistance to CDK4/6 Kinase Inhibitors. Cancer Discovery, 2022, 12, 356-371.	7.7	68
84	FOXA1 Mutations Reveal Distinct Chromatin Profiles and Influence Therapeutic Response in Breast Cancer. Cancer Cell, 2020, 38, 534-550.e9.	7.7	67
85	Analysis of mutational signatures in primary and metastatic endometrial cancer reveals distinct patterns of DNA repair defects and shifts during tumor progression. Gynecologic Oncology, 2019, 152, 11-19.	0.6	66
86	Genomic and transcriptomic heterogeneity in metaplastic carcinomas of the breast. Npj Breast Cancer, 2017, 3, 48.	2.3	63
87	Molecular profiling and molecular classification of endometrioid ovarian carcinomas. Gynecologic Oncology, 2019, 154, 516-523.	0.6	62
88	Mutation Profiling of Key Cancer Genes in Primary Breast Cancers and Their Distant Metastases. Cancer Research, 2018, 78, 3112-3121.	0.4	57
89	Mesonephric and mesonephric-like carcinomas of the female genital tract: molecular characterization including cases with mixed histology and matched metastases. Modern Pathology, 2021, 34, 1570-1587.	2.9	57
90	Independent realâ€world application of a clinicalâ€grade automated prostate cancer detection system. Journal of Pathology, 2021, 254, 147-158.	2.1	57

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91	Genomic and immunohistochemical analysis of adenosquamous carcinoma of the breast. Modern Pathology, 2010, 23, 951-960.	2.9	56
92	The repertoire of somatic genetic alterations of acinic cell carcinomas of the breast: an exploratory, hypothesisâ€generating study. Journal of Pathology, 2015, 237, 166-178.	2.1	53
93	Microglandular adenosis associated with tripleâ€negative breast cancer is a neoplastic lesion of tripleâ€negative phenotype harbouring <i><scp>TP53</scp></i> somatic mutations. Journal of Pathology, 2016, 238, 677-688.	2.1	52
94	The genetic landscape of breast carcinomas with neuroendocrine differentiation. Journal of Pathology, 2017, 241, 405-419.	2.1	52
95	Phyllodes tumors with and without fibroadenoma-like areas display distinct genomic features and may evolve through distinct pathways. Npj Breast Cancer, 2017, 3, 40.	2.3	52
96	Leiomyoma with bizarre nuclei: a morphological, immunohistochemical and molecular analysis of 31 cases. Modern Pathology, 2017, 30, 1476-1488.	2.9	51
97	Immunogenicity and therapeutic targeting of a public neoantigen derived from mutated PIK3CA. Nature Medicine, 2022, 28, 946-957.	15.2	50
98	Integrative genomic and transcriptomic characterization of papillary carcinomas of the breast. Molecular Oncology, 2014, 8, 1588-1602.	2.1	49
99	Activation of the IFN Signaling Pathway is Associated with Resistance to CDK4/6 Inhibitors and Immune Checkpoint Activation in ER-Positive Breast Cancer. Clinical Cancer Research, 2021, 27, 4870-4882.	3.2	49
100	Clinicopathologic and Genomic Analysis of <i>TP53</i> -Mutated Endometrial Carcinomas. Clinical Cancer Research, 2021, 27, 2613-2623.	3.2	49
101	Genetic analysis of microglandular adenosis and acinic cell carcinomas of the breast provides evidence for the existence of a low-grade triple-negative breast neoplasia family. Modern Pathology, 2017, 30, 69-84.	2.9	48
102	De-escalation of treatment in HER2-positive breast cancer: Determinants of response and mechanisms of resistance. Breast, 2017, 34, S19-S26.	0.9	46
103	Secretory carcinoma of the breast: clinicopathologic profile of 14 cases emphasising distant metastatic potential. Histopathology, 2019, 75, 213-224.	1.6	46
104	Triple-negative breast cancers — a panoply of cancer types. Nature Reviews Clinical Oncology, 2018, 15, 347-348.	12.5	45
105	Solid papillary breast carcinomas resembling the tall cell variant of papillary thyroid neoplasms (solid papillary carcinomas with reverse polarity) harbour recurrent mutations affecting <i><scp>IDH</scp>2</i> and <i><scp>PIK</scp>3<scp>CA</scp></i> a validation cohort. Histopathology, 2018, 73, 339-344.	1.6	44
106	Lobular Carcinomas <i>In Situ</i> Display Intralesion Genetic Heterogeneity and Clonal Evolution in the Progression to Invasive Lobular Carcinoma. Clinical Cancer Research, 2019, 25, 674-686.	3.2	44
107	Biâ€ellelic alterations in DNA repair genes underpin homologous recombination DNA repair defects in breast cancer. Journal of Pathology, 2017, 242, 165-177.	2.1	43
108	Whole-Exome Sequencing Analysis of the Progression from Non–Low-Grade Ductal Carcinoma <i>In Situ</i> to Invasive Ductal Carcinoma. Clinical Cancer Research, 2020, 26, 3682-3693.	3.2	42

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109	AXL-associated tumor inflammation as a poor prognostic signature in chemotherapy-treated triple-negative breast cancer patients. Npj Breast Cancer, 2016, 2, 16033.	2.3	41
110	Targeted capture massively parallel sequencing analysis of LCIS and invasive lobular cancer: Repertoire of somatic genetic alterations and clonal relationships. Molecular Oncology, 2016, 10, 360-370.	2.1	41
111	Massively parallel sequencing analysis of mucinous ovarian carcinomas: genomic profiling and differential diagnoses. Gynecologic Oncology, 2018, 150, 127-135.	0.6	41
112	Targeting the Mevalonate Pathway to Overcome Acquired Anti-HER2 Treatment Resistance in Breast Cancer. Molecular Cancer Research, 2019, 17, 2318-2330.	1.5	41
113	Homologous recombination DNA repair defects in PALB2-associated breast cancers. Npj Breast Cancer, 2019, 5, 23.	2.3	39
114	Histologic Classification and Molecular Signature of Polymorphous Adenocarcinoma (PAC) and Cribriform Adenocarcinoma of Salivary Gland (CASG). American Journal of Surgical Pathology, 2020, 44, 545-552.	2.1	39
115	Translating neoadjuvant therapy into survival benefits: one size does not fit all. Nature Reviews Clinical Oncology, 2016, 13, 566-579.	12.5	38
116	PAX8–GLIS3 gene fusion is a pathognomonic genetic alteration of hyalinizing trabecular tumors of the thyroid. Modern Pathology, 2019, 32, 1734-1743.	2.9	38
117	The repertoire of genetic alterations in salivary duct carcinoma including a novel HNRNPH3-ALK rearrangement. Human Pathology, 2019, 88, 66-77.	1.1	38
118	Genomic profiling of primary and recurrent adult granulosa cell tumors of the ovary. Modern Pathology, 2020, 33, 1606-1617.	2.9	38
119	Pathogenic <i>ATM</i> Mutations in Cancer and a Genetic Basis for Radiotherapeutic Efficacy. Journal of the National Cancer Institute, 2021, 113, 266-273.	3.0	38
120	Poor response to neoadjuvant chemotherapy in metaplastic breast carcinoma. Npj Breast Cancer, 2021, 7, 96.	2.3	38
121	Establishing the origin of metastatic deposits in the setting of multiple primary malignancies: The role of massively parallel sequencing. Molecular Oncology, 2014, 8, 150-158.	2.1	37
122	Are acinic cell carcinomas of the breast and salivary glands distinct diseases?. Histopathology, 2015, 67, 529-537.	1.6	37
123	Breast lesions of uncertain malignant nature and limited metastatic potential: proposals to improve their recognition and clinical management. Histopathology, 2016, 68, 45-56.	1.6	37
124	Contralateral breast cancers: Independent cancers or metastases?. International Journal of Cancer, 2018, 142, 347-356.	2.3	37
125	Genetic heterogeneity and actionable mutations in HER2-positive primary breast cancers and their brain metastases. Oncotarget, 2018, 9, 20617-20630.	0.8	36
126	Clinical outcomes of patients with POLE mutated endometrioid endometrial cancer. Gynecologic Oncology, 2020, 156, 194-202.	0.6	35

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127	Immunohistochemical analysis of IDH2 R172 hotspot mutations in breast papillary neoplasms: applications in the diagnosis of tall cell carcinoma with reverse polarity. Modern Pathology, 2020, 33, 1056-1064.	2.9	35
128	Machine learning-based prediction of microsatellite instability and high tumor mutation burden from contrast-enhanced computed tomography in endometrial cancers. Scientific Reports, 2020, 10, 17769.	1.6	35
129	Identification of recurrent FHL2-GLI2 oncogenic fusion in sclerosing stromal tumors of the ovary. Nature Communications, 2020, 11, 44.	5.8	34
130	Endometrial Carcinomas with a "Serous―Component in Young Women Are Enriched for DNA Mismatch Repair Deficiency, Lynch Syndrome, and POLE Exonuclease Domain Mutations. American Journal of Surgical Pathology, 2020, 44, 641-648.	2.1	34
131	Proteasome Addiction Defined in Ewing Sarcoma Is Effectively Targeted by a Novel Class of 19S Proteasome Inhibitors. Cancer Research, 2016, 76, 4525-4534.	0.4	33
132	Genomic landscape of endometrial carcinomas of no specific molecular profile. Modern Pathology, 2022, 35, 1269-1278.	2.9	33
133	Gene expression profiling of lobular carcinoma in situ reveals candidate precursor genes for invasion. Molecular Oncology, 2015, 9, 772-782.	2.1	32
134	Infiltrating epitheliosis of the breast: characterization of histological features, immunophenotype and genomic profile. Histopathology, 2016, 68, 1030-1039.	1.6	31
135	Evaluation of the Predictive Role of Tumor Immune Infiltrate in Patients with HER2-Positive Breast Cancer Treated with Neoadjuvant Anti-HER2 Therapy without Chemotherapy. Clinical Cancer Research, 2020, 26, 738-745.	3.2	31
136	Somatic mutations in leukocytes infiltrating primary breast cancers. Npj Breast Cancer, 2015, 1, 15005.	2.3	30
137	Genomic Profiling Aids Classification of Diagnostically Challenging Uterine Mesenchymal Tumors With Myomelanocytic Differentiation. American Journal of Surgical Pathology, 2021, 45, 77-92.	2.1	30
138	Functional and topographic effects on DNA methylation in IDH1/2 mutant cancers. Scientific Reports, 2019, 9, 16830.	1.6	29
139	Histologic spectrum of polymorphous adenocarcinoma of the salivary gland harbor genetic alterations affecting PRKD genes. Modern Pathology, 2020, 33, 65-73.	2.9	29
140	<i>BRCA</i> Mutations, Homologous DNA Repair Deficiency, Tumor Mutational Burden, and Response to Immune Checkpoint Inhibition in Recurrent Ovarian Cancer. JCO Precision Oncology, 2020, 4, 665-679.	1.5	29
141	Mutant FOXL2C134W Hijacks SMAD4 and SMAD2/3 to Drive Adult Granulosa Cell Tumors. Cancer Research, 2020, 80, 3466-3479.	0.4	29
142	A P53-Independent DNA Damage Response Suppresses Oncogenic Proliferation and Genome Instability. Cell Reports, 2020, 30, 1385-1399.e7.	2.9	29
143	Massively parallel sequencing analysis of synchronous fibroepithelial lesions supports the concept of progression from fibroadenoma to phyllodes tumor. Npj Breast Cancer, 2016, 2, 16035.	2.3	28
144	Massively parallel sequencing analysis of 68 gastric-type cervical adenocarcinomas reveals mutations in cell cycle-related genes and potentially targetable mutations. Modern Pathology, 2021, 34, 1213-1225.	2.9	28

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145	Lobular Neoplasia. Surgical Oncology Clinics of North America, 2014, 23, 487-503.	0.6	27
146	V211D Mutation in MEK1 Causes Resistance to MEK Inhibitors in Colon Cancer. Cancer Discovery, 2019, 9, 1182-1191.	7.7	27
147	The genomic landscape of metastatic histologic special types of invasive breast cancer. Npj Breast Cancer, 2020, 6, 53.	2.3	27
148	Ultraviolet radiation drives mutations in a subset of mucosal melanomas. Nature Communications, 2021, 12, 259.	5.8	27
149	The molecular genetic make-up of male breast cancer. Endocrine-Related Cancer, 2019, 26, 779-794.	1.6	27
150	Myxoid fibroadenomas differ from conventional fibroadenomas: a hypothesisâ€generating study. Histopathology, 2017, 71, 626-634.	1.6	26
151	Mixed Mesonephric Adenocarcinoma and High-grade Neuroendocrine Carcinoma of the Uterine Cervix: Case Description of a Previously Unreported Entity With Insights Into Its Molecular Pathogenesis. International Journal of Gynecological Pathology, 2017, 36, 76-89.	0.9	26
152	High-grade transformation of low-grade endometrial stromal sarcomas lacking YWHAE and BCOR genetic abnormalities. Modern Pathology, 2020, 33, 1861-1870.	2.9	26
153	Invasion in breast lesions: the role of the epithelial–stroma barrier. Histopathology, 2018, 72, 1075-1083.	1.6	25
154	Pleomorphic adenomas and mucoepidermoid carcinomas of the breast are underpinned by fusion genes. Npj Breast Cancer, 2020, 6, 20.	2.3	25
155	Problematic breast tumors reassessed in light of novel molecular data. Modern Pathology, 2021, 34, 38-47.	2.9	25
156	The clinical behavior and genomic features of the so-called adenoid cystic carcinomas of the solid variant with basaloid features. Modern Pathology, 2022, 35, 193-201.	2.9	25
157	MAPK Pathway Genetic Alterations Are Associated with Prolonged Overall Survival in Low-Grade Serous Ovarian Carcinoma. Clinical Cancer Research, 2022, 28, 4456-4465.	3.2	25
158	Ovarian carcinoma patient derived xenografts reproduce their tumor of origin and preserve an oligoclonal structure. Oncotarget, 2015, 6, 28327-28340.	0.8	24
159	Lack of <i><scp>PRKD</scp>2</i> and <i><scp>PRKD</scp>3</i> kinase domain somatic mutations in <i><scp>PRKD</scp>1</i> wildâ€type classic polymorphous lowâ€grade adenocarcinomas of the salivary gland. Histopathology, 2016, 68, 1055-1062.	1.6	23
160	Generation of conditional oncogenic chromosomal translocations using <scp>CRISPR</scp> –Cas9 genomic editing and homologyâ€directed repair. Journal of Pathology, 2017, 242, 102-112.	2.1	23
161	Clinical and pathologic features associated with PD-L1 (SP142) expression in stromal tumor-infiltrating immune cells of triple-negative breast carcinoma. Modern Pathology, 2020, 33, 2221-2232.	2.9	23
162	Fundamental immune–oncogenicity trade-offs define driver mutationÂfitness. Nature, 2022, 606, 172-179.	13.7	23

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163	Resolving quandaries: basaloid adenoid cystic carcinoma or breast cylindroma? The role of massively parallel sequencing. Histopathology, 2016, 68, 262-271.	1.6	22
164	Micropapillary variant of mucinous carcinoma of the breast shows genetic alterations intermediate between those of mucinous carcinoma and micropapillary carcinoma. Histopathology, 2019, 75, 139-145.	1.6	22
165	The role of a monoclonal antibody 11C8B1 as a diagnostic marker of IDH2-mutated sinonasal undifferentiated carcinoma. Modern Pathology, 2019, 32, 205-215.	2.9	22
166	Recurrent <i>MED12</i> exon 2 mutations in benign breast fibroepithelial lesions in adolescents and young adults. Journal of Clinical Pathology, 2019, 72, 258-262.	1.0	22
167	Histopathologic features and molecular genetic landscape of HER2-amplified endometrial carcinomas. Modern Pathology, 2022, 35, 962-971.	2.9	22
168	Molecular Subclasses of Clear Cell Ovarian Carcinoma and Their Impact on Disease Behavior and Outcomes. Clinical Cancer Research, 2022, 28, 4947-4956.	3.2	22
169	Assessment of HMGA2 and PLAG1 rearrangements in breast adenomyoepitheliomas. Npj Breast Cancer, 2019, 5, 6.	2.3	21
170	Solid pseudopapillary neoplasms of the pancreas are dependent on the Wnt pathway. Molecular Oncology, 2019, 13, 1684-1692.	2.1	21
171	The genetic landscape of metaplastic breast cancers and uterine carcinosarcomas. Molecular Oncology, 2021, 15, 1024-1039.	2.1	21
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173	The Landscape of Somatic Genetic Alterations in Breast Cancers from CHEK2 Germline Mutation Carriers. JNCI Cancer Spectrum, 2019, 3, pkz027.	1.4	20
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