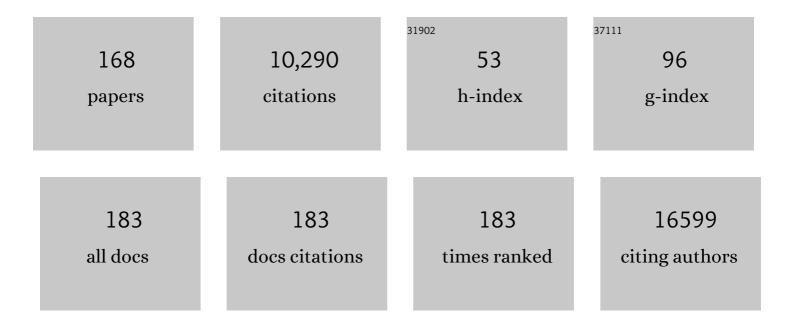
Charlotte K Y Ng

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Mutation tracking in circulating tumor DNA predicts relapse in early breast cancer. Science Translational Medicine, 2015, 7, 302ra133.	5.8	889
2	Cerebrospinal fluid-derived circulating tumour DNA better represents the genomic alterations of brain tumours than plasma. Nature Communications, 2015, 6, 8839.	5.8	605
3	Genomic characterization of metastatic breast cancers. Nature, 2019, 569, 560-564.	13.7	448
4	A framework to rank genomic alterations as targets for cancer precision medicine: the ESMO Scale for Clinical Actionability of molecular Targets (ESCAT). Annals of Oncology, 2018, 29, 1895-1902.	0.6	424
5	Spatial and Temporal Heterogeneity in High-Grade Serous Ovarian Cancer: A Phylogenetic Analysis. PLoS Medicine, 2015, 12, e1001789.	3.9	314
6	Capturing intra-tumor genetic heterogeneity by de novo mutation profiling of circulating cell-free tumor DNA: a proof-of-principle. Annals of Oncology, 2014, 25, 1729-1735.	0.6	308
7	Organoid Models of Human Liver Cancers Derived from Tumor Needle Biopsies. Cell Reports, 2018, 24, 1363-1376.	2.9	288
8	Breast cancer intra-tumor heterogeneity. Breast Cancer Research, 2014, 16, 210.	2.2	256
9	Chromosomal instability determines taxane response. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 8671-8676.	3.3	244
10	Progression from ductal carcinoma <i>in situ</i> to invasive breast cancer: Revisited. Molecular Oncology, 2013, 7, 859-869.	2.1	195
11	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
12	Hotspot activating PRKD1 somatic mutations in polymorphous low-grade adenocarcinomas of the salivary glands. Nature Genetics, 2014, 46, 1166-1169.	9.4	188
13	<i> <scp>SF3B1</scp> </i> mutations constitute a novel therapeutic target in breast cancer. Journal of Pathology, 2015, 235, 571-580.	2.1	167
14	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
15	Genomic analysis of genetic heterogeneity and evolution in high-grade serous ovarian carcinoma. Oncogene, 2010, 29, 4905-4913.	2.6	153
16	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
17	The protein histidine phosphatase LHPP is a tumour suppressor. Nature, 2018, 555, 678-682.	13.7	151
18	Genomic landscape of adenoid cystic carcinoma of the breast. Journal of Pathology, 2015, 237, 179-189.	2.1	133

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19	The Landscape of Somatic Genetic Alterations in Metaplastic Breast Carcinomas. Clinical Cancer Research, 2017, 23, 3859-3870.	3.2	129
20	The Genomic Landscape of Male Breast Cancers. Clinical Cancer Research, 2016, 22, 4045-4056.	3.2	119
21	Benchmarking mutation effect prediction algorithms using functionally validated cancer-related missense mutations. Genome Biology, 2014, 15, 484.	3.8	117
22	Whole-genome single-cell copy number profiling from formalin-fixed paraffin-embedded samples. Nature Medicine, 2017, 23, 376-385.	15.2	111
23	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
24	Recurrent hotspot mutations in HRAS Q61 and PI3K-AKT pathway genes as drivers of breast adenomyoepitheliomas. Nature Communications, 2018, 9, 1816.	5.8	105
25	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
26	<i>IDH2</i> Mutations Define a Unique Subtype of Breast Cancer with Altered Nuclear Polarity. Cancer Research, 2016, 76, 7118-7129.	0.4	99
27	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
28	Uterine adenosarcomas are mesenchymal neoplasms. Journal of Pathology, 2016, 238, 381-388.	2.1	94
29	Cancer Diagnosis Using a Liquid Biopsy: Challenges and Expectations. Diagnostics, 2018, 8, 31.	1.3	94
30	Genetic Heterogeneity in Therapy-NaÃ⁻ve Synchronous Primary Breast Cancers and Their Metastases. Clinical Cancer Research, 2017, 23, 4402-4415.	3.2	91
31	Mechanism of action of a WWTR1(TAZ)-CAMTA1 fusion oncoprotein. Oncogene, 2016, 35, 929-938.	2.6	90
32	TP53 Mutational Spectrum in Endometrioid and Serous Endometrial Cancers. International Journal of Gynecological Pathology, 2016, 35, 289-300.	0.9	89
33	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
34	Neoantigen prediction and computational perspectives towards clinical benefit: recommendations from the ESMO Precision Medicine Working Group. Annals of Oncology, 2020, 31, 978-990.	0.6	87
35	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
36	Tandem duplication of chromosomal segments is common in ovarian and breast cancer genomes. Journal of Pathology, 2012, 227, 446-455.	2.1	81

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37	Breast Cancer Genomics From Microarrays to Massively Parallel Sequencing: Paradigms and New Insights. Journal of the National Cancer Institute, 2015, 107, .	3.0	80
38	Metastatic breast carcinomas display genomic and transcriptomic heterogeneity. Modern Pathology, 2015, 28, 340-351.	2.9	80
39	Reliability of Whole-Exome Sequencing for Assessing Intratumor Genetic Heterogeneity. Cell Reports, 2018, 25, 1446-1457.	2.9	76
40	Patient-derived xenografts and organoids model therapy response in prostate cancer. Nature Communications, 2021, 12, 1117.	5.8	76
41	Genetic profiling using plasma-derived cell-free DNA in therapy-naÃ⁻ve hepatocellular carcinoma patients: a pilot study. Annals of Oncology, 2018, 29, 1286-1291.	0.6	74
42	<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
43	The Role of Long Non-Coding RNAs in Hepatocarcinogenesis. International Journal of Molecular Sciences, 2018, 19, 682.	1.8	73
44	Loss of Osteoclasts Contributes to Development of Osteosarcoma Pulmonary Metastases. Cancer Research, 2010, 70, 7063-7072.	0.4	72
45	The Tumor Profiler Study: integrated, multi-omic, functional tumor profiling for clinical decision support. Cancer Cell, 2021, 39, 288-293.	7.7	71
46	Intra-tumour genetic heterogeneity and poor chemoradiotherapy response in cervical cancer. British Journal of Cancer, 2011, 104, 361-368.	2.9	69
47	Metaplastic breast carcinoma: more than a special type. Nature Reviews Cancer, 2014, 14, 147-148.	12.8	69
48	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
49	The Genomic Landscape of Mucinous Breast Cancer. Journal of the National Cancer Institute, 2019, 111, 737-741.	3.0	68
50	Breast cancer intratumor genetic heterogeneity: causes and implications. Expert Review of Anticancer Therapy, 2012, 12, 1021-1032.	1.1	65
51	Genomic profiling of histological special types of breast cancer. Breast Cancer Research and Treatment, 2013, 142, 257-269.	1.1	64
52	Genomic and transcriptomic heterogeneity in metaplastic carcinomas of the breast. Npj Breast Cancer, 2017, 3, 48.	2.3	63
53	Comprehensive clinical and molecular analyses of neuroendocrine carcinomas of the breast. Modern Pathology, 2018, 31, 68-82.	2.9	58
54	Mutation Profiling of Key Cancer Genes in Primary Breast Cancers and Their Distant Metastases. Cancer Research, 2018, 78, 3112-3121.	0.4	57

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55	The role of tandem duplicator phenotype in tumour evolution in highâ€grade serous ovarian cancer. Journal of Pathology, 2012, 226, 703-712.	2.1	56
56	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
57	High-throughput sequencing of nodal marginal zone lymphomas identifies recurrent BRAF mutations. Leukemia, 2018, 32, 2412-2426.	3.3	53
58	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
59	The genetic landscape of breast carcinomas with neuroendocrine differentiation. Journal of Pathology, 2017, 241, 405-419.	2.1	52
60	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
61	Leiomyoma with bizarre nuclei: a morphological, immunohistochemical and molecular analysis of 31 cases. Modern Pathology, 2017, 30, 1476-1488.	2.9	51
62	High Expression of FAP in Colorectal Cancer Is Associated With Angiogenesis and Immunoregulation Processes. Frontiers in Oncology, 2020, 10, 979.	1.3	50
63	Integrative genomic and transcriptomic characterization of papillary carcinomas of the breast. Molecular Oncology, 2014, 8, 1588-1602.	2.1	49
64	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
65	Integrative proteogenomic characterization of hepatocellular carcinoma across etiologies and stages. Nature Communications, 2022, 13, 2436.	5.8	45
66	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
67	Biâ€allelic alterations in DNA repair genes underpin homologous recombination DNA repair defects in breast cancer. Journal of Pathology, 2017, 242, 165-177.	2.1	43
68	PI3K Pathway Activation in High-Grade Ductal Carcinoma <i>In Situ</i> —Implications for Progression to Invasive Breast Carcinoma. Clinical Cancer Research, 2014, 20, 2326-2337.	3.2	41
69	HMGA1 Expression in Human Hepatocellular Carcinoma Correlates with Poor Prognosis and Promotes Tumor Growth and Migration in in vitro Models. Neoplasia, 2016, 18, 724-731.	2.3	41
70	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
71	High expression of HOXA13 correlates with poorly differentiated hepatocellular carcinomas and modulates sorafenib response in in vitro models. Laboratory Investigation, 2018, 98, 95-105.	1.7	41
72	A tumor DNA complex aberration index is an independent predictor of survival in breast and ovarian cancer. Molecular Oncology, 2015, 9, 115-127.	2.1	38

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73	Circulating Cell-Free DNA in Hepatocellular Carcinoma: Current Insights and Outlook. Frontiers in Medicine, 2018, 5, 78.	1.2	38
74	The repertoire of genetic alterations in salivary duct carcinoma including a novel HNRNPH3-ALK rearrangement. Human Pathology, 2019, 88, 66-77.	1.1	38
75	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
76	Are acinic cell carcinomas of the breast and salivary glands distinct diseases?. Histopathology, 2015, 67, 529-537.	1.6	37
77	Contralateral breast cancers: Independent cancers or metastases?. International Journal of Cancer, 2018, 142, 347-356.	2.3	37
78	SCIM: universal single-cell matching with unpaired feature sets. Bioinformatics, 2020, 36, i919-i927.	1.8	37
79	Genetic heterogeneity and actionable mutations in HER2-positive primary breast cancers and their brain metastases. Oncotarget, 2018, 9, 20617-20630.	0.8	36
80	LATS1 but not LATS2 represses autophagy by a kinase-independent scaffold function. Nature Communications, 2019, 10, 5755.	5.8	36
81	Infiltration by myeloperoxidase-positive neutrophils is an independent prognostic factor in breast cancer. Breast Cancer Research and Treatment, 2019, 177, 581-589.	1.1	34
82	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
83	Structural analysis of the genome of breast cancer cell line ZR-75-30 identifies twelve expressed fusion genes. BMC Genomics, 2012, 13, 719.	1.2	32
84	Gene expression profiling of lobular carcinoma in situ reveals candidate precursor genes for invasion. Molecular Oncology, 2015, 9, 772-782.	2.1	32
85	Genomic Analysis Revealed New Oncogenic Signatures in TP53-Mutant Hepatocellular Carcinoma. Frontiers in Genetics, 2018, 9, 2.	1.1	32
86	Infiltrating epitheliosis of the breast: characterization of histological features, immunophenotype and genomic profile. Histopathology, 2016, 68, 1030-1039.	1.6	31
87	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
88	NGS-pipe: a flexible, easily extendable and highly configurable framework for NGS analysis. Bioinformatics, 2018, 34, 107-108.	1.8	25
89	Hepatocellular Carcinoma Xenografts Established From Needle Biopsies Preserve the Characteristics of the Originating Tumors. Hepatology Communications, 2019, 3, 971-986.	2.0	24
90	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€ŧype classic polymorphous lowâ€grade adenocarcinomas of the salivary gland. Histopathology, 2016, 68, 1055-1062.	1.6	23

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91	Resolving quandaries: basaloid adenoid cystic carcinoma or breast cylindroma? The role of massively parallel sequencing. Histopathology, 2016, 68, 262-271.	1.6	22
92	Phosphoprotein enriched in diabetes (PED/PEA15) promotes migration in hepatocellular carcinoma and confers resistance to sorafenib. Cell Death and Disease, 2017, 8, e3138-e3138.	2.7	22
93	Nectin-4 Expression Is an Independent Prognostic Biomarker and Associated With Better Survival in Triple-Negative Breast Cancer. Frontiers in Medicine, 2019, 6, 200.	1.2	22
94	Predictive Performance of Microarray Gene Signatures: Impact of Tumor Heterogeneity and Multiple Mechanisms of Drug Resistance. Cancer Research, 2014, 74, 2946-2961.	0.4	20
95	The Dilemma of HER2 Double-equivocal Breast Carcinomas. American Journal of Surgical Pathology, 2018, 42, 1190-1200.	2.1	20
96	Preoperative plasma fatty acid metabolites inform risk of prostate cancer progression and may be used for personalized patient stratification. BMC Cancer, 2019, 19, 1216.	1.1	20
97	Adenylosuccinate lyase is oncogenic in colorectal cancer by causing mitochondrial dysfunction and independent activation of NRF2 and mTOR-MYC-axis. Theranostics, 2021, 11, 4011-4029.	4.6	19
98	DNA Copy Number Aberrations, and Human Papillomavirus Status in Penile Carcinoma. Clinico-Pathological Correlations and Potential Driver Genes. PLoS ONE, 2016, 11, e0146740.	1.1	19
99	Response to dual HER2 blockade in a patient with HER3-mutant metastatic breast cancer. Annals of Oncology, 2015, 26, 1704-1709.	0.6	18
100	Therapeutic Targeting of CD146/MCAM Reduces Bone Metastasis in Prostate Cancer. Molecular Cancer Research, 2019, 17, 1049-1062.	1.5	17
101	Expression of RET is associated with Oestrogen receptor expression but lacks prognostic significance in breast cancer. BMC Cancer, 2019, 19, 41.	1.1	16
102	PIKing the type and pattern of PI3K pathway mutations in endometrioid endometrial carcinomas. Gynecologic Oncology, 2015, 137, 321-328.	0.6	15
103	Vascular endothelial growth factor A amplification in colorectal cancer is associated with reduced M1 and M2 macrophages and diminished PD-1-expressing lymphocytes. PLoS ONE, 2017, 12, e0175563.	1.1	15
104	Diagnostic Targeted Sequencing Panel for Hepatocellular Carcinoma Genomic Screening. Journal of Molecular Diagnostics, 2018, 20, 836-848.	1.2	15
105	Infiltration by IL22-Producing T Cells Promotes Neutrophil Recruitment and Predicts Favorable Clinical Outcome in Human Colorectal Cancer. Cancer Immunology Research, 2020, 8, 1452-1462.	1.6	15
106	Establishing standardized immune phenotyping of metastatic melanoma by digital pathology. Laboratory Investigation, 2021, 101, 1561-1570.	1.7	15
107	Transcriptional Enhancer Factor Domain Family member 4 Exerts an Oncogenic Role in Hepatocellular Carcinoma by Hippoâ€Independent Regulation of Heat Shock Protein 70 Family Members. Hepatology Communications, 2021, 5, 661-674.	2.0	13
108	Interferon lambda 4 impairs hepatitis C viral antigen presentation and attenuates T cell responses. Nature Communications, 2021, 12, 4882.	5.8	13

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109	Alterations in homologous recombination repair genes in prostate cancer brain metastases. Nature Communications, 2022, 13, 2400.	5.8	13
110	Multi-omics data integration reveals novel drug targets in hepatocellular carcinoma. BMC Genomics, 2021, 22, 592.	1.2	12
111	Cenetic analysis of uterine adenosarcomas and phyllodes tumors of the breast. Molecular Oncology, 2017, 11, 913-926.	2.1	11
112	Molecular classification of hepatocellular carcinoma: The view from metabolic zonation. Hepatology, 2017, 66, 1377-1380.	3.6	11
113	Pregnancy at early age is associated with a reduction of progesterone-responsive cells and epithelial Wnt signaling in human breast tissue. Oncotarget, 2017, 8, 22353-22360.	0.8	11
114	PipelT. Journal of Molecular Diagnostics, 2019, 21, 884-894.	1.2	11
115	ESR1 and endocrine therapy resistance: more than just mutations. Annals of Oncology, 2018, 29, 787-789.	0.6	10
116	Genetic Alterations in Benign Breast Biopsies of Subsequent Breast Cancer Patients. Frontiers in Medicine, 2019, 6, 166.	1.2	10
117	Radiogenomics Analysis of Intratumor Heterogeneity in a Patient With High-Grade Serous Ovarian Cancer. JCO Precision Oncology, 2019, 3, 1-9.	1.5	10
118	Circulating Cell-Free DNA Captures the Intratumor Heterogeneity in Multinodular Hepatocellular Carcinoma. JCO Precision Oncology, 2022, 6, e2100335.	1.5	9
119	RNASeq analysis reveals biological processes governing the clinical behaviour of endometrioid and serous endometrial cancers. European Journal of Cancer, 2016, 64, 149-158.	1.3	8
120	Stroma Transcriptomic and Proteomic Profile of Prostate Cancer Metastasis Xenograft Models Reveals Prognostic Value of Stroma Signatures. Cancers, 2020, 12, 3786.	1.7	8
121	GATA3 and MDM2 are synthetic lethal in estrogen receptor-positive breast cancers. Communications Biology, 2022, 5, 373.	2.0	7
122	High-resolution genomic profiling of thyroid lesions uncovers preferential copy number gains affecting mitochondrial biogenesis loci in the oncocytic variants. American Journal of Cancer Research, 2015, 5, 1954-71.	1.4	6
123	Nestin and CD34 expression in colorectal cancer predicts improved overall survival. Acta Oncológica, 2021, 60, 727-734.	0.8	5
124	Genomic analysis of focal nodular hyperplasia with associated hepatocellular carcinoma unveils its malignant potential: a case report. Communications Medicine, 2022, 2, .	1.9	5
125	Prognostic signatures in breast cancer: correlation does not imply causation. Breast Cancer Research, 2012, 14, 313.	2.2	4
126	Systematic identification of novel cancer genes through analysis of deep shRNA perturbation screens. Nucleic Acids Research, 2021, 49, 8488-8504.	6.5	4

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127	Abstract S4-03: A functional assay for homologous recombination (HR) DNA repair and whole exome sequencing reveal that HR-defective sporadic breast cancers are enriched for genetic alterations in DNA repair genes. , 2016, , .		4
128	The Genomic Landscape of Serrated Lesion of the Colorectum: Similarities and Differences With Tubular and Tubulovillous Adenomas. Frontiers in Oncology, 2021, 11, 668466.	1.3	4
129	Hepatocellular Carcinoma: Pathology and Genetics. , 2018, , 198-198.		3
130	Genomic evolutionary trajectory of metastatic squamous cell carcinoma of the lung. Translational Lung Cancer Research, 2021, 10, 1792-1803.	1.3	3
131	Epigenetic priming in chronic liver disease impacts the transcriptional and genetic landscapes of hepatocellular carcinoma. Molecular Oncology, 2022, 16, 665-682.	2.1	3
132	Fibroepithelial Breast Lesion: When Sequencing Can Help to Make a Clinical Decision. A Case Report. Clinical Breast Cancer, 2019, 19, e1-e6.	1.1	2
133	Discovery of heterozygous <i>KRT10</i> alterations in MAUIE cases underlines the importance of regular skin cancer screening in ichthyosis with confetti. British Journal of Dermatology, 2020, 183, 954-955.	1.4	2
134	Identification of Somatic Mutations in Thirty-year-old Serum Cell-free DNA From Patients With Breast Cancer: A Feasibility Study. Clinical Breast Cancer, 2020, 20, 413-421.e1.	1.1	2
135	Abstract 927: Targeted capture next generation sequencing of fresh frozen lobular carcinoma in situ and invasive lobular cancer identifies a common repertoire of mutations. , 2014, , .		2
136	Reliability of Whole-Exome Sequencing for Assessing Intratumor Genetic Heterogeneity. SSRN Electronic Journal, 0, , .	0.4	2
137	Abstract 930: Analysis of cell-free tumor DNA in cerebrospinal fluid to characterize and monitor the genetic alterations of brain tumors. Cancer Research, 2015, 75, 930-930.	0.4	2
138	The Role of Chronic Liver Diseases in the Emergence and Recurrence of Hepatocellular Carcinoma: An Omics Perspective. Frontiers in Medicine, 0, 9, .	1.2	2
139	Combining two antibodies to define <scp>E</scp> â€cadherin loss of expression in nonâ€lobular breast carcinomas: when less is more. Histopathology, 2013, 63, 439-440.	1.6	1
140	Genetic profiling using plasma-derived cell-free DNA in therapy-naÃ⁻ve hepatocellular carcinoma patients: A pilot study. Digestive and Liver Disease, 2018, 50, 27.	0.4	1
141	Cell-free DNA in hepatocellular carcinoma. , 2020, , 199-209.		1
142	Abstract PD05-08: Genomic characterisation of invasive breast cancers with heterogeneous HER2 gene amplification. , 2012, , .		1
143	Abstract 100: The landscape of somatic genetic alterations in BRCA1 and BRCA2 breast cancers. , 2016, , .		1
144	Abstract S6-06: The genomic landscape of male breast cancers. , 2015, , .		1

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145	Abstract PD3-4: Reliability of whole exome sequencing for assessing intratumor heterogeneity from breast tumor biopsies. , 2015, , .		1
146	Abstract 2989: Intra-tumor heterogeneity and clonal changes in the progression of DCIS to invasiveness: Combined tumor bulk and single cell analysis. , 2015, , .		1
147	Abstract 138: Solid papillary carcinoma with reverse polarization are driven by IDH2 and PI3K pathway mutations. , 2016, , .		1
148	575 SF3B1 mutations are associated with alternative splicing in ER-positive breast cancer. European Journal of Cancer, 2014, 50, 186.	1.3	0
149	SU30ANALYSIS OF WGS DATA FROM 108 INDIVIDUALS OF 8 SPANISH FAMILIES AFFECTED WITH BIPOLAR DISORDER. European Neuropsychopharmacology, 2019, 29, S1283-S1284.	0.3	Ο
150	Abstract 4029: The impact of multiple drug resistance mechanisms on microarray predictive gene signature performance , 2013, , .		0
151	Abstract P4-04-05: Molecular subtyping reveals the heterogeneity of metaplastic breast cancers. , 2013, , .		Ο
152	Abstract P4-04-08: Genomic and transcriptomic characterization of papillary carcinomas of the breast. , 2013, , .		0
153	Abstract 1544: The clinical behavior of endometrioid and serous endometrial carcinomas is governed by distinct biological processes. , 2014, , .		Ο
154	Abstract 4258: Benchmarking algorithms for mutation impact prediction using functionally validated missense mutations. , 2014, , .		0
155	Abstract P2-03-09: Benchmarking mutation function prediction algorithms using validated cancer driver and passenger mutations. , 2015, , .		0
156	Abstract P2-03-08: Mutational landscape of metaplastic breast carcinomas. , 2015, , .		0
157	Abstract 3889: Genetic heterogeneity and distinct driver mutations in synchronous primary and metastatic breast cancers from therapy-naÃ ⁻ ve patients. , 2015, , .		Ο
158	Abstract 2971: Whole exome sequencing reveals heterogeneity within lobular carcinomain situ(LCIS) and clonal selection in the progression to malignant lesions. , 2015, , .		0
159	Abstract 4817: Microsatellite instability status in endometrioid endometrial carcinomas is associated with distinct types and patterns of PI3K pathway mutations. , 2015, , .		Ο
160	Abstract S4-04: Lobular carcinoma in situ displays intra-lesion genetic heterogeneity and its progression to invasive disease involves clonal selection and variations in mutational processes. , 2016, , .		0
161	Abstract P6-07-04: Distinct repertoires of somatic mutations affecting driver genes in mucinous and neuroendocrine carcinomas of the breast. , 2016, , .		0
162	Abstract P2-01-02: Capturing intra-tumor genetic heterogeneity in cell-free plasma DNA from patients with oligometastatic breast cancer. , 2016, , .		0

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163	Abstract P6-06-02: GermlineCDH1mutations in lobular carcinomain situ. , 2016, , .		Ο
164	Abstract P6-03-10: Genomic and transcriptomic heterogeneity in metaplastic breast carcinomas. , 2016, ,		0
165	Abstract 91: The mutational landscape of mucinous carcinomas of the breast. , 2016, , .		Ο
166	Abstract 134: Mutational landscape of breast cancers from PALB2 germline mutation carriers. , 2016, , .		0
167	Abstract 4612: HOXA13 drives hepatocytes proliferation and liver tumorigenesis in mice. , 2019, , .		О
168	Abstract 2821: Gut commensal bacteria modulate functions of tumor-associated neutrophils in human colorectal cancer. , 2019, , .		0