Albrecht Stenzinger

List of Publications by Year in descending order

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304 papers 16,802 citations

20817 60 h-index 24258 110 g-index

319 all docs

319 docs citations

319 times ranked

22591 citing authors

#	Article	IF	Citations
1	Development of tumor mutation burden as an immunotherapy biomarker: utility for the oncology clinic. Annals of Oncology, 2019, 30, 44-56.	1.2	1,742
2	Identification of a population of blood circulating tumor cells from breast cancer patients that initiates metastasis in a xenograft assay. Nature Biotechnology, 2013, 31, 539-544.	17.5	920
3	Recommendations for the use of next-generation sequencing (NGS) for patients with metastatic cancers: a report from the ESMO Precision Medicine Working Group. Annals of Oncology, 2020, 31, 1491-1505.	1.2	658
4	The Novel Histologic International Association for the Study of Lung Cancer/American Thoracic Society/European Respiratory Society Classification System of Lung Adenocarcinoma Is a Stage-Independent Predictor of Survival. Journal of Clinical Oncology, 2012, 30, 1438-1446.	1.6	606
5	Tumor Mutational Burden as a Predictive Biomarker in Solid Tumors. Cancer Discovery, 2020, 10, 1808-1825.	9.4	388
6	SARS-CoV-2 infects and replicates in cells of the human endocrine and exocrine pancreas. Nature Metabolism, 2021, 3, 149-165.	11.9	378
7	Establishing guidelines to harmonize tumor mutational burden (TMB): in silico assessment of variation in TMB quantification across diagnostic platforms: phase I of the Friends of Cancer Research TMB Harmonization Project., 2020, 8, e000147.		329
8	Sarcoma classification by DNA methylation profiling. Nature Communications, 2021, 12, 498.	12.8	237
9	Accurate and efficient detection of gene fusions from RNA sequencing data. Genome Research, 2021, 31, 448-460.	5 . 5	215
10	Classification of Cancer at Prostate MRI: Deep Learning versus Clinical PI-RADS Assessment. Radiology, 2019, 293, 607-617.	7.3	214
11	Prognostic impact of tumour-infiltrating immune cells on biliary tract cancer. British Journal of Cancer, 2013, 109, 2665-2674.	6.4	209
12	The landscape of metastatic progression patterns across major human cancers. Oncotarget, 2015, 6, 570-583.	1.8	208
13	BRAFV600E mutant protein is expressed in cells of variable maturation in Langerhans cell histiocytosis. Blood, 2012, 120, e28-e34.	1.4	199
14	Integrative genomic and transcriptomic analysis of leiomyosarcoma. Nature Communications, 2018, 9, 144.	12.8	197
15	<i>NRG1</i> Fusions in <i>KRAS</i> Wild-Type Pancreatic Cancer. Cancer Discovery, 2018, 8, 1087-1095.	9.4	189
16	CYP3A5 mediates basal and acquired therapy resistance in different subtypes of pancreatic ductal adenocarcinoma. Nature Medicine, 2016, 22, 278-287.	30.7	184
17	Increased microtubule assembly rates influence chromosomal instability in colorectal cancer cells. Nature Cell Biology, 2014, 16, 779-791.	10.3	174
18	Tumor mutational burden standardization initiatives: Recommendations for consistent tumor mutational burden assessment in clinical samples to guide immunotherapy treatment decisions. Genes Chromosomes and Cancer, 2019, 58, 578-588.	2.8	173

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19	A field guide for cancer diagnostics using cellâ€free DNA: From principles to practice and clinical applications. Genes Chromosomes and Cancer, 2018, 57, 123-139.	2.8	155
20	Implementing tumor mutational burden (TMB) analysis in routine diagnostics—a primer for molecular pathologists and clinicians. Translational Lung Cancer Research, 2018, 7, 703-715.	2.8	152
21	Precision oncology based on omics data: The NCT Heidelberg experience. International Journal of Cancer, 2017, 141, 877-886.	5.1	133
22	Automated sample preparation with <scp>SP</scp> 3 for lowâ€input clinical proteomics. Molecular Systems Biology, 2020, 16, e9111.	7.2	133
23	Colorectal mixed adenoneuroendocrine carcinomas and neuroendocrine carcinomas are genetically closely related to colorectal adenocarcinomas. Modern Pathology, 2017, 30, 610-619.	5 . 5	131
24	Size matters: Dissecting key parameters for panelâ€based tumor mutational burden analysis. International Journal of Cancer, 2019, 144, 848-858.	5.1	131
25	Three molecular pathways model colorectal carcinogenesis in <scp>L</scp> ynch syndrome. International Journal of Cancer, 2018, 143, 139-150.	5.1	129
26	Comprehensive Genomic and Transcriptomic Analysis for Guiding Therapeutic Decisions in Patients with Rare Cancers. Cancer Discovery, 2021, 11, 2780-2795.	9.4	125
27	Optimizing panel-based tumor mutational burden (TMB) measurement. Annals of Oncology, 2019, 30, 1496-1506.	1.2	123
28	Tumour cell proliferation (Ki-67) in non-small cell lung cancer: a critical reappraisal of its prognostic role. British Journal of Cancer, 2014, 111, 1222-1229.	6.4	114
29	The presence of circulating tumor cells (CTCs) correlates with lymph node metastasis in nonresectable squamous cell carcinoma of the head and neck region (SCCHN). Annals of Oncology, 2011, 22, 1878-1885.	1.2	112
30	Molecular Diagnostic Profiling of Lung Cancer Specimens with a Semiconductor-Based Massive Parallel Sequencing Approach. Journal of Molecular Diagnostics, 2013, 15, 765-775.	2.8	107
31	Correlation of radio- and histomorphological pattern of pulmonary adenocarcinoma. European Respiratory Journal, 2013, 41, 943-951.	6.7	105
32	Interobserver variability in the application of the novel IASLC/ATS/ERS classification for pulmonary adenocarcinomas. European Respiratory Journal, 2012, 40, 1221-1227.	6.7	97
33	EGFR, KRAS, BRAF and ALK gene alterations in lung adenocarcinomas: patient outcome, interplay with morphology and immunophenotype. European Respiratory Journal, 2014, 43, 872-883.	6.7	97
34	Global alterations of DNA methylation in cholangiocarcinoma target the Wnt signaling pathway. Hepatology, 2014, 59, 544-554.	7.3	97
35	BRAF V600E-specific immunohistochemistry reveals low mutation rates in biliary tract cancer and restriction to intrahepatic cholangiocarcinoma. Modern Pathology, 2014, 27, 1028-1034.	5. 5	96
36	<i><scp>ROS</scp>1</i> expression and translocations in nonâ€small ell lung cancer: clinicopathological analysis of 1478 cases. Histopathology, 2014, 65, 187-194.	2.9	96

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37	Measurement of tumor mutational burden (TMB) in routine molecular diagnostics: ⟨i⟩in silico⟨/i⟩ and realâ€ife analysis of three larger gene panels. International Journal of Cancer, 2019, 144, 2303-2312.	5.1	95
38	KRAS Genotyping of Paraffin-Embedded Colorectal Cancer Tissue in Routine Diagnostics. Journal of Molecular Diagnostics, 2010, 12, 35-42.	2.8	94
39	<i>EML4â€ALK</i> fusion variant V3 is a highâ€risk feature conferring accelerated metastatic spread, early treatment failure and worse overall survival in ALK ⁺ nonâ€small cell lung cancer. International Journal of Cancer, 2018, 142, 2589-2598.	5.1	93
40	Classical pathology and mutational load of breast cancer – integration of two worlds. Journal of Pathology: Clinical Research, 2015, 1, 225-238.	3.0	91
41	mTOR expression and activity patterns in gastroenteropancreatic neuroendocrine tumours. Endocrine-Related Cancer, 2011, 18, 181-192.	3.1	90
42	Aligning tumor mutational burden (TMB) quantification across diagnostic platforms: phase II of the Friends of Cancer Research TMB Harmonization Project. Annals of Oncology, 2021, 32, 1626-1636.	1.2	86
43	Expression of Amphiregulin and EGFRvIII Affect Outcome of Patients with Squamous Cell Carcinoma of the Head and Neck Receiving Cetuximab–Docetaxel Treatment. Clinical Cancer Research, 2011, 17, 5197-5204.	7.0	85
44	Combined targeted DNA and RNA sequencing of advanced NSCLC in routine molecular diagnostics: Analysis of the first 3,000 Heidelberg cases. International Journal of Cancer, 2019, 145, 649-661.	5.1	85
45	Prognostic Impact and Clinicopathological Correlations of the Cribriform Pattern in Pulmonary Adenocarcinoma. Journal of Thoracic Oncology, 2015, 10, 638-644.	1.1	83
46	Co-expression of MET and CD47 is a novel prognosticator for survival of luminal-type breast cancer patients. Oncotarget, 2014, 5, 8147-8160.	1.8	83
47	Targeted ultra-deep sequencing reveals recurrent and mutually exclusive mutations of cancer genes in blastic plasmacytoid dendritic cell neoplasm. Oncotarget, 2014, 5, 6404-6413.	1.8	82
48	Pancreatic Ductal Adenocarcinoma Subtyping Using the Biomarkers Hepatocyte Nuclear Factor-1A and Cytokeratin-81 Correlates with Outcome and Treatment Response. Clinical Cancer Research, 2018, 24, 351-359.	7.0	81
49	Harmonization and Standardization of Panel-Based Tumor Mutational Burden Measurement: Real-World Results and Recommendations ofÂtheÂQuality in Pathology Study. Journal of Thoracic Oncology, 2020, 15, 1177-1189.	1.1	81
50	Panâ€cancer analysis of copy number changes in programmed deathâ€ligand 1 (PD‣1, CD274) – associations with gene expression, mutational load, and survival. Genes Chromosomes and Cancer, 2016, 55, 626-639.	S 2.8	80
51	Postoperative Complications Deteriorate Long-Term Outcome in Pancreatic Cancer Patients. Annals of Surgical Oncology, 2012, 19, 856-863.	1.5	78
52	Potential clinical implications of <i>BRAF</i> mutations in histiocytic proliferations. Oncotarget, 2014, 5, 4060-4070.	1.8	78
53	Variant classification in precision oncology. International Journal of Cancer, 2019, 145, 2996-3010.	5.1	76
54	Contribution of human papilloma virus to the incidence of squamous cell carcinoma of the head and neck in a European population with high smoking prevalence. European Journal of Cancer, 2015, 51, 514-521.	2.8	75

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55	The landscape of chromothripsis across adult cancer types. Nature Communications, 2020, 11, 2320.	12.8	7 5
56	Reliable Entity Subtyping in Non-small Cell Lung Cancer by Matrix-assisted Laser Desorption/Ionization Imaging Mass Spectrometry on Formalin-fixed Paraffin-embedded Tissue Specimens. Molecular and Cellular Proteomics, 2016, 15, 3081-3089.	3.8	72
57	Morphological and molecular breast cancer profiling through explainable machine learning. Nature Machine Intelligence, 2021, 3, 355-366.	16.0	72
58	Survival of Patients with Oral Cavity Cancer in Germany. PLoS ONE, 2013, 8, e53415.	2.5	69
59	Spatial and Temporal Heterogeneity of Panel-Based Tumor Mutational Burden in Pulmonary Adenocarcinoma: Separating Biology From Technical Artifacts. Journal of Thoracic Oncology, 2019, 14, 1935-1947.	1.1	69
60	Homologous Recombination Deficiency: Concepts, Definitions, and Assays. Oncologist, 2022, 27, 167-174.	3.7	69
61	Identification of a highly lethal V3 ⁺ TP53 ⁺ subset in ALK ⁺ lung adenocarcinoma. International Journal of Cancer, 2019, 144, 190-199.	5.1	67
62	Highâ€throughput diagnostic profiling of clinically actionable gene fusions in lung cancer. Genes Chromosomes and Cancer, 2016, 55, 30-44.	2.8	65
63	Defective homologous recombination DNA repair as therapeutic target in advanced chordoma. Nature Communications, 2019, 10, 1635.	12.8	64
64	High SIRT1 expression is a negative prognosticator in pancreatic ductal adenocarcinoma. BMC Cancer, 2013, 13, 450.	2.6	63
65	Integrative Analysis Defines Distinct Prognostic Subgroups of Intrahepatic Cholangiocarcinoma. Hepatology, 2019, 69, 2091-2106.	7.3	63
66	Deep Learning for the Classification of Small-Cell and Non-Small-Cell Lung Cancer. Cancers, 2020, 12, 1604.	3.7	63
67	Patients Resistant Against PSMA-Targeting α-Radiation Therapy Often Harbor Mutations in DNA Damage-Repair–Associated Genes. Journal of Nuclear Medicine, 2020, 61, 683-688.	5.0	61
68	Phenotyping of pulmonary carcinoids and a Ki-67-based grading approach. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2012, 460, 299-308.	2.8	60
69	Loss of SOX2 expression induces cell motility via vimentin upâ€regulation and is an unfavorable risk factor for survival ofÂhead and neck squamous cell carcinoma. Molecular Oncology, 2015, 9, 1704-1719.	4.6	60
70	Integrated analysis of the immunological and genetic status in and across cancer types: impact of mutational signatures beyond tumor mutational burden. Oncolmmunology, 2018, 7, e1526613.	4.6	60
71	Position of a panel of international lung cancer experts on the approval decision for use of durvalumab in stage III non-small-cell lung cancer (NSCLC) by the Committee for Medicinal Products for Human Use (CHMP). Annals of Oncology, 2019, 30, 161-165.	1.2	60
72	Associations of Pathogenic Variants in MLH1, MSH2, and MSH6 With Risk of Colorectal Adenomas and Tumors and With Somatic Mutations in Patients With Lynch Syndrome. Gastroenterology, 2020, 158, 1326-1333.	1.3	60

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73	Mutational profiles in triple-negative breast cancer defined by ultradeep multigene sequencing show high rates of PI3K pathway alterations and clinically relevant entity subgroup specific differences. Oncotarget, 2014, 5, 9952-9965.	1.8	58
74	Integration of genomics and histology revises diagnosis and enables effective therapy of refractory cancer of unknown primary with $\langle i \rangle$ PDL1 $\langle i \rangle$ amplification. Journal of Physical Education and Sports Management, 2016, 2, a001180.	1.2	57
75	Marker chromosomes can arise from chromothripsis and predict adverse prognosis in acute myeloid leukemia. Blood, 2017, 129, 1333-1342.	1.4	57
76	Mutations in POLE and survival of colorectal cancer patients $\hat{a} \in \text{``link to disease stage and treatment.}$ Cancer Medicine, 2014, 3, 1527-1538.	2.8	56
77	Training increases concordance in classifying pulmonary adenocarcinomas according to the novel IASLC/ATS/ERS classification. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2012, 461, 185-193.	2.8	55
78	Distribution of <i>MED12</i> mutations in fibroadenomas and phyllodes tumors of the breastâe"implications for tumor biology and pathological diagnosis. Genes Chromosomes and Cancer, 2015, 54, 444-452.	2.8	55
79	Standards for the classification of pathogenicity of somatic variants in cancer (oncogenicity): Joint recommendations of Clinical Genome Resource (ClinGen), Cancer Genomics Consortium (CGC), and Variant Interpretation for Cancer Consortium (VICC). Genetics in Medicine, 2022, 24, 986-998.	2.4	55
80	Major histocompatibility complex class I expression impacts on patient survival and type and density of immune cells in biliary tract cancer. British Journal of Cancer, 2015, 113, 1343-1349.	6.4	54
81	Biomarker testing in non-small cell lung cancer in routine care: Analysis of the first 3,717 patients in the German prospective, observational, nation-wide CRISP Registry (AIO-TRK-0315). Lung Cancer, 2021, 152, 174-184.	2.0	53
82	Who Is at Risk for Diagnostic Discrepancies? Comparison of Pre- and Postmortal Diagnoses in 1800 Patients of 3 Medical Decades in East and West Berlin. PLoS ONE, 2012, 7, e37460.	2.5	53
83	Testing <i>NTRK</i> testing: Wet″ab and in silico comparison of RNAâ€based targeted sequencing assays. Genes Chromosomes and Cancer, 2020, 59, 178-188.	2.8	52
84	MiR-200b and miR-155 as predictive biomarkers for the efficacy of chemoradiation in locally advanced head and neck squamous cell carcinoma. European Journal of Cancer, 2017, 77, 3-12.	2.8	51
85	Recurrent YAP1 and MAML2 Gene Rearrangements in Retiform and Composite Hemangioendothelioma. American Journal of Surgical Pathology, 2020, 44, 1677-1684.	3.7	51
86	PD-L1 (CD274) copy number gain, expression, and immune cell infiltration as candidate predictors for response to immune checkpoint inhibitors in soft-tissue sarcoma. Oncolmmunology, 2017, 6, e1279777.	4.6	50
87	Role of <i>TP53</i> mutations in triple negative and HER2-positive breast cancer treated with neoadjuvant anthracycline/taxane-based chemotherapy. Oncotarget, 2016, 7, 67686-67698.	1.8	50
88	EWSR1/FUS–CREB fusions define a distinctive malignant epithelioid neoplasm with predilection for mesothelial-lined cavities. Modern Pathology, 2020, 33, 2233-2243.	5 . 5	49
89	The combinatorial complexity of cancer precision medicine. Oncoscience, 2014, 1, 504-509.	2.2	48
90	Molecular driver alterations and their clinical relevance in cancer of unknown primary site. Oncotarget, 2016, 7, 44322-44329.	1.8	47

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91	Downâ€regulation of the microRNA processing enzyme Dicer is a prognostic factor in human colorectal cancer. Histopathology, 2012, 61, 552-561.	2.9	44
92	Appendiceal goblet cell carcinoids and adenocarcinomas ex-goblet cell carcinoid are genetically distinct from primary colorectal-type adenocarcinoma of the appendix. Modern Pathology, 2018, 31, 829-839.	5 . 5	44
93	Copy number changes of clinically actionable genes in melanoma, nonâ€small cell lung cancer and colorectal cancer—A survey across 822 routine diagnostic cases. Genes Chromosomes and Cancer, 2016, 55, 821-833.	2.8	43
94	KRAS G12C-mutated advanced non-small cell lung cancer: A real-world cohort from the German prospective, observational, nation-wide CRISP Registry (AIO-TRK-0315). Lung Cancer, 2021, 154, 51-61.	2.0	43
95	Genomic Characterization of Cholangiocarcinoma in Primary Sclerosing Cholangitis Reveals Therapeutic Opportunities. Hepatology, 2020, 72, 1253-1266.	7.3	42
96	Evaluation of a Hybrid Capture–Based Pan-Cancer Panel for Analysis of Treatment Stratifying Oncogenic Aberrations and Processes. Journal of Molecular Diagnostics, 2020, 22, 757-769.	2.8	42
97	Artificial intelligence and pathology: From principles to practice and future applications in histomorphology and molecular profiling. Seminars in Cancer Biology, 2022, 84, 129-143.	9.6	41
98	A gene expression signature associated with B cells predicts benefit from immune checkpoint blockade in lung adenocarcinoma. Oncolmmunology, 2021, 10, 1860586.	4.6	40
99	Combined Clinical Parameters and Multiparametric Magnetic Resonance Imaging for the Prediction of Extraprostatic Disease—A Risk Model for Patient-tailored Risk Stratification When Planning Radical Prostatectomy. European Urology Focus, 2020, 6, 1205-1212.	3.1	39
100	Targeted next-generation sequencing identifies molecular subgroups in squamous cell carcinoma of the head and neck with distinct outcome after concurrent chemoradiation. Annals of Oncology, 2016, 27, 2262-2268.	1.2	38
101	The BRCA2 mutation status shapes the immune phenotype of prostate cancer. Cancer Immunology, Immunotherapy, 2019, 68, 1621-1633.	4.2	38
102	Label-Free Enrichment and Molecular Characterization of Viable Circulating Tumor Cells from Diagnostic Leukapheresis Products. Clinical Chemistry, 2019, 65, 549-558.	3.2	37
103	Cancer beyond organ and tissue specificity: Nextâ€generationâ€sequencing gene mutation data reveal complex genetic similarities across major cancers. International Journal of Cancer, 2014, 135, 2362-2369.	5.1	36
104	KRAS Mutations in Codon 12 or 13 Are Associated With Worse Prognosis in Pancreatic Ductal Adenocarcinoma. Pancreas, 2014, 43, 578-583.	1.1	36
105	Allelic Ratio of <i>KRAS</i> Mutations in Pancreatic Cancer. Oncologist, 2015, 20, e8-e9.	3.7	36
106	ALK-FISH borderline cases in non-small cell lung cancer: Implications for diagnostics and clinical decision making. Lung Cancer, 2015, 90, 465-471.	2.0	36
107	A non-controlled, single arm, open label, phase II study of intravenous and intratumoral administration of ParvOryx in patients with metastatic, inoperable pancreatic cancer: ParvOryx02 protocol. BMC Cancer, 2017, 17, 576.	2.6	36
108	Detection of TP53 Mutations in Tissue or Liquid Rebiopsies at Progression Identifies ALK+ Lung Cancer Patients with Poor Survival. Cancers, 2019, 11, 124.	3.7	36

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109	Hidden Variables in Deep Learning Digital Pathology and Their Potential to Cause Batch Effects: Prediction Model Study. Journal of Medical Internet Research, 2021, 23, e23436.	4.3	36
110	Distinctive Spatiotemporal Stability of Somatic Mutations in Metastasized Microsatellite-stable Colorectal Cancer. American Journal of Surgical Pathology, 2015, 39, 1140-1147.	3.7	35
111	Establishment of a patient-derived orthotopic osteosarcoma mouse model. Journal of Translational Medicine, 2015, 13, 136.	4.4	35
112	Mutation patterns in genes encoding interferon signaling and antigen presentation: A panâ \in cancer survey with implications for the use of immune checkpoint inhibitors. Genes Chromosomes and Cancer, 2017, 56, 651-659.	2.8	35
113	Tumor Mutational Burden as a Pan-cancer Biomarker for Immunotherapy: The Limits and Potential for Convergence. Cancer Cell, 2020, 38, 624-625.	16.8	35
114	Association of the advanced lung cancer inflammation index (ALI) with immune checkpoint inhibitor efficacy in patients with advanced non-small-cell lung cancer. ESMO Open, 2021, 6, 100254.	4.5	35
115	Defining molecular risk in ALK+ NSCLC. Oncotarget, 2019, 10, 3093-3103.	1.8	35
116	Genotyping of colorectal cancer for cancer precision medicine: Results from the IPH Center for Molecular Pathology. Genes Chromosomes and Cancer, 2016, 55, 505-521.	2.8	34
117	The Value of Prostate-specific Antigen Density for Prostate Imaging-Reporting and Data System 3 Lesions on Multiparametric Magnetic Resonance Imaging: A Strategy to Avoid Unnecessary Prostate Biopsies. European Urology Focus, 2021, 7, 325-331.	3.1	34
118	Cadherin-6 is a putative tumor suppressor and target of epigenetically dysregulated miR-429 in cholangiocarcinoma. Epigenetics, 2016, 11, 780-790.	2.7	33
119	Therapeutic and Prognostic Implications of Immune-Related Adverse Events in Advanced Non-Small-Cell Lung Cancer. Frontiers in Oncology, 2021, 11, 703893.	2.8	33
120	Simultaneous whole-body 18F–PSMA-1007-PET/MRI with integrated high-resolution multiparametric imaging of the prostatic fossa for comprehensive oncological staging of patients with prostate cancer: a pilot study. European Journal of Nuclear Medicine and Molecular Imaging, 2018, 45, 340-347.	6.4	32
121	RNA-Based Detection of Gene Fusions in Formalin-Fixed and Paraffin-Embedded Solid Cancer Samples. Cancers, 2019, 11, 1309.	3.7	32
122	Mutational Diversity and Therapy Response in Breast Cancer: A Sequencing Analysis in the Neoadjuvant GeparSepto Trial. Clinical Cancer Research, 2019, 25, 3986-3995.	7.0	32
123	Longitudinal therapy monitoring of ALK-positive lung cancer by combined copy number and targeted mutation profiling of cell-free DNA. EBioMedicine, 2020, 62, 103103.	6.1	32
124	Endometrial stromal sarcomas with <i>BCOR</i> â€rearrangement harbor <i>MDM2</i> amplifications. Journal of Pathology: Clinical Research, 2020, 6, 178-184.	3.0	32
125	Efficacy of Immune Checkpoint Inhibitors Alone or in Combination With Chemotherapy in NSCLC Harboring ERBB2 Mutations. Journal of Thoracic Oncology, 2021, 16, 1952-1958.	1.1	32
126	Fusionâ€positive nonâ€small cell lung carcinoma: Biological principles, clinical practice, and diagnostic implications. Genes Chromosomes and Cancer, 2022, 61, 244-260.	2.8	32

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127	High nuclear polyâ€(ADPâ€ribose)â€polymerase expression is prognostic of improved survival in pancreatic cancer. Histopathology, 2012, 61, 409-416.	2.9	31
128	High extracellular matrix metalloproteinase inducer/CD147 expression is strongly and independently associated with poor prognosis in colorectal cancer. Human Pathology, 2012, 43, 1471-1481.	2.0	30
129	Mutant KIT as imatinib-sensitive target in metastatic sinonasal carcinoma. Annals of Oncology, 2017, 28, 142-148.	1.2	30
130	Mutations in genes encoding <scp>Pl3Kâ€AKT</scp> and <scp>MAPK</scp> signaling define anogenital papillary hidradenoma. Genes Chromosomes and Cancer, 2016, 55, 113-119.	2.8	29
131	EGFR T790M mutation testing of non-small cell lung cancer tissue and blood samples artificially spiked with circulating cell-free tumor DNA: results of a round robin trial. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2017, 471, 509-520.	2.8	29
132	<i>RSPO2</i> gene rearrangement: a powerful driver of \hat{I}^2 -catenin activation in liver tumours. Gut, 2019, 68, 1287-1296.	12.1	29
133	Genetic heterogeneity in synchronous colorectal cancers impacts genotyping approaches and therapeutic strategies. Genes Chromosomes and Cancer, 2016, 55, 268-277.	2.8	28
134	Quantifying potential confounders of panel-based tumor mutational burden (TMB) measurement. Lung Cancer, 2020, 142, 114-119.	2.0	28
135	Standardized Magnetic Resonance Imaging Reporting Using the Prostate Cancer Radiological Estimation of Change in Sequential Evaluation Criteria and Magnetic Resonance Imaging/Transrectal Ultrasound Fusion with Transperineal Saturation Biopsy to Select Men on Active Surveillance. European Urology Focus, 2021, 7, 102-110.	3.1	28
136	p53 partial loss-of-function mutations sensitize to chemotherapy. Oncogene, 2022, 41, 1011-1023.	5.9	28
137	Tubular, lactating, and ductal adenomas are devoid of MED12 Exon2 mutations, and ductal adenomas show recurrent mutations in GNAS and the PI3Kâ \in AKT pathway. Genes Chromosomes and Cancer, 2017, 56, 11-17.	2.8	27
138	Targeted molecular profiling reveals genetic heterogeneity of poromas and porocarcinomas. Pathology, 2018, 50, 327-332.	0.6	27
139	In-house Implementation of Tumor Mutational Burden Testing to Predict Durable Clinical Benefit in Non-small Cell Lung Cancer and Melanoma Patients. Cancers, 2019, 11, 1271.	3.7	27
140	<scp>NTRK</scp> testing: First results of the <scp>QuiPâ€EQA</scp> scheme and a comprehensive map of <scp><i>NTRK</i></scp> fusion variants and their diagnostic coverage by targeted <scp>RNA</scp> â€based <scp>NGS</scp> assays. Genes Chromosomes and Cancer, 2020, 59, 445-453.	2.8	27
141	Novel GATA6-FOXO1 fusions in a subset of epithelioid hemangioma. Modern Pathology, 2021, 34, 934-941.	5. 5	27
142	Fully Automatic Deep Learning in Bi-institutional Prostate Magnetic Resonance Imaging. Investigative Radiology, 2021, 56, 799-808.	6.2	27
143	The Different Immune Profiles of Normal Colonic Mucosa in Cancer-Free Lynch Syndrome Carriers and Lynch Syndrome Colorectal Cancer Patients. Gastroenterology, 2022, 162, 907-919.e10.	1.3	27
144	The novel protein PTPIP51 exhibits tissue- and cell-specific expression. Histochemistry and Cell Biology, 2005, 123, 19-28.	1.7	26

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145	EML4-ALK V3, treatment resistance, and survival: refining the diagnosis of ALK+ NSCLC. Journal of Thoracic Disease, 2018, 10, S1989-S1991.	1.4	26
146	Risk stratification of EGFR+ lung cancer diagnosed with panel-based next-generation sequencing. Lung Cancer, 2020, 148, 105-112.	2.0	26
147	Chapter 6 Cell and Molecular Biology of the Novel Protein Tyrosineâ€Phosphataseâ€Interacting Protein 51. International Review of Cell and Molecular Biology, 2009, 275, 183-246.	3.2	25
148	Fibroblast Growth Factor Receptor 1 as a Putative Therapy Target in Colorectal Cancer. Digestion, 2013, 88, 172-181.	2.3	25
149	Prevalence of somatic mitochondrial mutations and spatial distribution of mitochondria in non-small cell lung cancer. British Journal of Cancer, 2017, 117, 220-226.	6.4	25
150	NGS-based BRCA1/2 mutation testing of high-grade serous ovarian cancer tissue: results and conclusions of the first international round robin trial. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2016, 468, 697-705.	2.8	24
151	Comparative genetic profiling aids diagnosis and clinical decision making in challenging cases of CUP syndrome. International Journal of Cancer, 2019, 145, 2963-2973.	5.1	24
152	Histopathological to multiparametric MRI spatial mapping of extended systematic sextant and MR/TRUS-fusion-targeted biopsy of the prostate. European Radiology, 2019, 29, 1820-1830.	4.5	24
153	Simulated clinical deployment of fully automatic deep learning for clinical prostate MRI assessment. European Radiology, 2021, 31, 302-313.	4.5	24
154	Real-world implementation of sequential targeted therapies for EGFR-mutated lung cancer. Therapeutic Advances in Medical Oncology, 2021, 13, 175883592199650.	3.2	24
155	Combined Immunohistochemistry after Mass Spectrometry Imaging for Superior Spatial Information. Proteomics - Clinical Applications, 2019, 13, e1800035.	1.6	23
156	Conceptual framework for precision cancer medicine in Germany: Consensus statement of the Deutsche Krebshilfe working group â€~Molecular Diagnostics and Therapy'. European Journal of Cancer, 2020, 135, 1-7.	2.8	23
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