

Vincent A Miller

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

423
papers

18,210
citations

60
h-index

132
g-index

424
ext. papers

22,410
ext. citations

4.7
avg, IF

6.17
L-index

| # | Paper | IF | Citations |
|-----|--|------|-----------|
| 423 | EGF receptor gene mutations are common in lung cancers from "never smokers" and are associated with sensitivity of tumors to gefitinib and erlotinib. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2004 , 101, 13306-11 | 11.5 | 3659 |
| 422 | Analysis of 100,000 human cancer genomes reveals the landscape of tumor mutational burden. <i>Genome Medicine</i> , 2017 , 9, 34 | 14.4 | 1509 |
| 421 | Development and validation of a clinical cancer genomic profiling test based on massively parallel DNA sequencing. <i>Nature Biotechnology</i> , 2013 , 31, 1023-31 | 44.5 | 1353 |
| 420 | Tumor Mutational Burden as an Independent Predictor of Response to Immunotherapy in Diverse Cancers. <i>Molecular Cancer Therapeutics</i> , 2017 , 16, 2598-2608 | 6.1 | 1183 |
| 419 | Co-occurring genomic alterations define major subsets of KRAS-mutant lung adenocarcinoma with distinct biology, immune profiles, and therapeutic vulnerabilities. <i>Cancer Discovery</i> , 2015 , 5, 860-77 | 24.4 | 476 |
| 418 | Emergence of constitutively active estrogen receptor mutations in pretreated advanced estrogen receptor-positive breast cancer. <i>Clinical Cancer Research</i> , 2014 , 20, 1757-1767 | 12.9 | 415 |
| 417 | Kinase fusions are frequent in Spitz tumours and spitzoid melanomas. <i>Nature Communications</i> , 2014 , 5, 3116 | 17.4 | 394 |
| 416 | Targeted Next Generation Sequencing Identifies Markers of Response to PD-1 Blockade. <i>Cancer Immunology Research</i> , 2016 , 4, 959-967 | 12.5 | 318 |
| 415 | RAS/MAPK Activation Is Associated with Reduced Tumor-Infiltrating Lymphocytes in Triple-Negative Breast Cancer: Therapeutic Cooperation Between MEK and PD-1/PD-L1 Immune Checkpoint Inhibitors. <i>Clinical Cancer Research</i> , 2016 , 22, 1499-509 | 12.9 | 311 |
| 414 | Diverse and Targetable Kinase Alterations Drive Histiocytic Neoplasms. <i>Cancer Discovery</i> , 2016 , 6, 154-65 | 24.4 | 269 |
| 413 | Inflammatory myofibroblastic tumors harbor multiple potentially actionable kinase fusions. <i>Cancer Discovery</i> , 2014 , 4, 889-95 | 24.4 | 250 |
| 412 | Molecular profiling of cancer patients enables personalized combination therapy: the I-PREDICT study. <i>Nature Medicine</i> , 2019 , 25, 744-750 | 50.5 | 240 |
| 411 | Biliary cancer: Utility of next-generation sequencing for clinical management. <i>Cancer</i> , 2016 , 122, 3838-3847 | 17.4 | 185 |
| 410 | Integrated genomic DNA/RNA profiling of hematologic malignancies in the clinical setting. <i>Blood</i> , 2016 , 127, 3004-14 | 2.2 | 185 |
| 409 | Broad, Hybrid Capture-Based Next-Generation Sequencing Identifies Actionable Genomic Alterations in Lung Adenocarcinomas Otherwise Negative for Such Alterations by Other Genomic Testing Approaches. <i>Clinical Cancer Research</i> , 2015 , 21, 3631-9 | 12.9 | 183 |
| 408 | Genomic and functional analysis of leukemic transformation of myeloproliferative neoplasms. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014 , 111, E5401-10 | 11.5 | 183 |
| 407 | Characterization of 298 Patients with Lung Cancer Harboring MET Exon 14 Skipping Alterations. <i>Journal of Thoracic Oncology</i> , 2016 , 11, 1493-502 | 8.9 | 177 |

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| 406 | The distribution of BRAF gene fusions in solid tumors and response to targeted therapy. <i>International Journal of Cancer</i> , 2016 , 138, 881-90 | 7.5 | 168 |
| 405 | Lung Master Protocol (Lung-MAP)-A Biomarker-Driven Protocol for Accelerating Development of Therapies for Squamous Cell Lung Cancer: SWOG S1400. <i>Clinical Cancer Research</i> , 2015 , 21, 1514-24 | 12.9 | 165 |
| 404 | Targeted next-generation sequencing of head and neck squamous cell carcinoma identifies novel genetic alterations in HPV+ and HPV- tumors. <i>Genome Medicine</i> , 2013 , 5, 49 | 14.4 | 158 |
| 403 | Impact of EML4-ALK Variant on Resistance Mechanisms and Clinical Outcomes in ALK-Positive Lung Cancer. <i>Journal of Clinical Oncology</i> , 2018 , 36, 1199-1206 | 2.2 | 155 |
| 402 | Cancer Therapy Directed by Comprehensive Genomic Profiling: A Single Center Study. <i>Cancer Research</i> , 2016 , 76, 3690-701 | 10.1 | 154 |
| 401 | Prevalence of PDL1 Amplification and Preliminary Response to Immune Checkpoint Blockade in Solid Tumors. <i>JAMA Oncology</i> , 2018 , 4, 1237-1244 | 13.4 | 149 |
| 400 | Beyond microsatellite testing: assessment of tumor mutational burden identifies subsets of colorectal cancer who may respond to immune checkpoint inhibition. <i>Journal of Gastrointestinal Oncology</i> , 2018 , 9, 610-617 | 2.8 | 131 |
| 399 | Comprehensive Genomic Profiling of 282 Pediatric Low- and High-Grade Gliomas Reveals Genomic Drivers, Tumor Mutational Burden, and Hypermutation Signatures. <i>Oncologist</i> , 2017 , 22, 1478-1490 | 5.7 | 126 |
| 398 | Fluorescence in situ hybridization, immunohistochemistry, and next-generation sequencing for detection of EML4-ALK rearrangement in lung cancer. <i>Oncologist</i> , 2015 , 20, 316-22 | 5.7 | 125 |
| 397 | BRAF fusions define a distinct molecular subset of melanomas with potential sensitivity to MEK inhibition. <i>Clinical Cancer Research</i> , 2013 , 19, 6696-702 | 12.9 | 122 |
| 396 | Real-Time Targeted Genome Profile Analysis of Pancreatic Ductal Adenocarcinomas Identifies Genetic Alterations That Might Be Targeted With Existing Drugs or Used as Biomarkers. <i>Gastroenterology</i> , 2019 , 156, 2242-2253.e4 | 13.3 | 117 |
| 395 | Comprehensive genomic profiling of pancreatic acinar cell carcinomas identifies recurrent RAF fusions and frequent inactivation of DNA repair genes. <i>Cancer Discovery</i> , 2014 , 4, 1398-405 | 24.4 | 112 |
| 394 | A computational approach to distinguish somatic vs. germline origin of genomic alterations from deep sequencing of cancer specimens without a matched normal. <i>PLoS Computational Biology</i> , 2018 , 14, e1005965 | 5 | 109 |
| 393 | Advanced urothelial carcinoma: next-generation sequencing reveals diverse genomic alterations and targets of therapy. <i>Modern Pathology</i> , 2014 , 27, 271-80 | 9.8 | 108 |
| 392 | Total mutation burden (TMB) in lung cancer (LC) and relationship with response to PD-1/PD-L1 targeted therapies.. <i>Journal of Clinical Oncology</i> , 2016 , 34, 9017-9017 | 2.2 | 108 |
| 391 | Analytical Validation of a Hybrid Capture-Based Next-Generation Sequencing Clinical Assay for Genomic Profiling of Cell-Free Circulating Tumor DNA. <i>Journal of Molecular Diagnostics</i> , 2018 , 20, 686-702 | 5.1 | 103 |
| 390 | Genomic Profiling of Small-Bowel Adenocarcinoma. <i>JAMA Oncology</i> , 2017 , 3, 1546-1553 | 13.4 | 101 |
| 389 | Inactivation of Capicua drives cancer metastasis. <i>Nature Genetics</i> , 2017 , 49, 87-96 | 36.3 | 92 |

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| 388 | Pulmonary Sarcomatoid Carcinomas Commonly Harbor Either Potentially Targetable Genomic Alterations or High Tumor Mutational Burden as Observed by Comprehensive Genomic Profiling. <i>Journal of Thoracic Oncology</i> , 2017 , 12, 932-942 | 8.9 | 85 |
| 387 | Genomic Characterization of Renal Cell Carcinoma with Sarcomatoid Dedifferentiation Pinpoints Recurrent Genomic Alterations. <i>European Urology</i> , 2016 , 70, 348-57 | 10.2 | 82 |
| 386 | Profiling of 149 Salivary Duct Carcinomas, Carcinoma Ex Pleomorphic Adenomas, and Adenocarcinomas, Not Otherwise Specified Reveals Actionable Genomic Alterations. <i>Clinical Cancer Research</i> , 2016 , 22, 6061-6068 | 12.9 | 81 |
| 385 | Diverse EGFR Exon 20 Insertions and Co-Occurring Molecular Alterations Identified by Comprehensive Genomic Profiling of NSCLC. <i>Journal of Thoracic Oncology</i> , 2018 , 13, 1560-1568 | 8.9 | 75 |
| 384 | Metastatic basal cell carcinoma with amplification of PD-L1: exceptional response to anti-PD1 therapy. <i>Npj Genomic Medicine</i> , 2016 , 1, | 6.2 | 74 |
| 383 | Triple-negative breast cancers with amplification of JAK2 at the 9p24 locus demonstrate JAK2-specific dependence. <i>Science Translational Medicine</i> , 2016 , 8, 334ra53 | 17.5 | 73 |
| 382 | A Novel Next-Generation Sequencing Approach to Detecting Microsatellite Instability and Pan-Tumor Characterization of 1000 Microsatellite Instability-High Cases in 67,000 Patient Samples. <i>Journal of Molecular Diagnostics</i> , 2019 , 21, 1053-1066 | 5.1 | 72 |
| 381 | Emergence of Preexisting MET Y1230C Mutation as a Resistance Mechanism to Crizotinib in NSCLC with MET Exon 14 Skipping. <i>Journal of Thoracic Oncology</i> , 2017 , 12, 137-140 | 8.9 | 72 |
| 380 | Identification of NTRK fusions in pediatric mesenchymal tumors. <i>Pediatric Blood and Cancer</i> , 2017 , 64, e26433 | 3 | 71 |
| 379 | High-Throughput Genomic Profiling of Adult Solid Tumors Reveals Novel Insights into Cancer Pathogenesis. <i>Cancer Research</i> , 2017 , 77, 2464-2475 | 10.1 | 71 |
| 378 | Pan-Cancer Landscape and Analysis of ERBB2 Mutations Identifies Poziotinib as a Clinically Active Inhibitor and Enhancer of T-DM1 Activity. <i>Cancer Cell</i> , 2019 , 36, 444-457.e7 | 24.3 | 69 |
| 377 | RICTOR Amplification Defines a Novel Subset of Patients with Lung Cancer Who May Benefit from Treatment with mTORC1/2 Inhibitors. <i>Cancer Discovery</i> , 2015 , 5, 1262-70 | 24.4 | 69 |
| 376 | Comprehensive Genomic Profiling of Advanced Esophageal Squamous Cell Carcinomas and Esophageal Adenocarcinomas Reveals Similarities and Differences. <i>Oncologist</i> , 2015 , 20, 1132-9 | 5.7 | 69 |
| 375 | Identification of recurrent FGFR3-TACC3 fusion oncogenes from lung adenocarcinoma. <i>Clinical Cancer Research</i> , 2014 , 20, 6551-8 | 12.9 | 67 |
| 374 | An Acquired Gatekeeper Mutation Induces Resistance to Neratinib in a Patient with HER2 Mutant-Driven Breast Cancer. <i>Cancer Discovery</i> , 2017 , 7, 575-585 | 24.4 | 66 |
| 373 | Receptor Tyrosine Kinase Fusions and BRAF Kinase Fusions are Rare but Actionable Resistance Mechanisms to EGFR Tyrosine Kinase Inhibitors. <i>Journal of Thoracic Oncology</i> , 2018 , 13, 1312-1323 | 8.9 | 65 |
| 372 | EGFR Fusions as Novel Therapeutic Targets in Lung Cancer. <i>Cancer Discovery</i> , 2016 , 6, 601-11 | 24.4 | 65 |
| 371 | Comparative analysis of primary tumour and matched metastases in colorectal cancer patients: evaluation of concordance between genomic and transcriptional profiles. <i>European Journal of Cancer</i> , 2015 , 51, 791-9 | 7.5 | 64 |

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| 370 | Genetic hallmarks of recurrent/metastatic adenoid cystic carcinoma. <i>Journal of Clinical Investigation</i> , 2019 , 129, 4276-4289 | 15.9 | 64 |
| 369 | Enrichment of Targetable Mutations in the Relapsed Neuroblastoma Genome. <i>PLoS Genetics</i> , 2016 , 12, e1006501 | 6 | 64 |
| 368 | ROS1 Fusions Rarely Overlap with Other Oncogenic Drivers in Non-Small Cell Lung Cancer. <i>Journal of Thoracic Oncology</i> , 2017 , 12, 872-877 | 8.9 | 63 |
| 367 | Emergence of RET rearrangement co-existing with activated EGFR mutation in EGFR-mutated NSCLC patients who had progressed on first- or second-generation EGFR TKI. <i>Lung Cancer</i> , 2015 , 89, 357-9 | 5.9 | 63 |
| 366 | Characterization of Clinical Cases of Collecting Duct Carcinoma of the Kidney Assessed by Comprehensive Genomic Profiling. <i>European Urology</i> , 2016 , 70, 516-21 | 10.2 | 61 |
| 365 | Comprehensive Genomic Profiling Facilitates Implementation of the National Comprehensive Cancer Network Guidelines for Lung Cancer Biomarker Testing and Identifies Patients Who May Benefit From Enrollment in Mechanism-Driven Clinical Trials. <i>Oncologist</i> , 2016 , 21, 684-91 | 5.7 | 61 |
| 364 | Genomic Profiling of a Large Set of Diverse Pediatric Cancers Identifies Known and Novel Mutations across Tumor Spectra. <i>Cancer Research</i> , 2017 , 77, 509-519 | 10.1 | 60 |
| 363 | HER2 Transmembrane Domain (TMD) Mutations (V659/G660) That Stabilize Homo- and Heterodimerization Are Rare Oncogenic Drivers in Lung Adenocarcinoma That Respond to Afatinib. <i>Journal of Thoracic Oncology</i> , 2017 , 12, 446-457 | 8.9 | 59 |
| 362 | The Genomic Landscape of Merkel Cell Carcinoma and Clinicogenomic Biomarkers of Response to Immune Checkpoint Inhibitor Therapy. <i>Clinical Cancer Research</i> , 2019 , 25, 5961-5971 | 12.9 | 58 |
| 361 | Comprehensive genomic profiling of inflammatory breast cancer cases reveals a high frequency of clinically relevant genomic alterations. <i>Breast Cancer Research and Treatment</i> , 2015 , 154, 155-62 | 4.4 | 57 |
| 360 | STUMP un"stumped": anti-tumor response to anaplastic lymphoma kinase (ALK) inhibitor based targeted therapy in uterine inflammatory myofibroblastic tumor with myxoid features harboring DCTN1-ALK fusion. <i>Journal of Hematology and Oncology</i> , 2015 , 8, 66 | 22.4 | 56 |
| 359 | Comprehensive Genomic Profiling of Advanced Penile Carcinoma Suggests a High Frequency of Clinically Relevant Genomic Alterations. <i>Oncologist</i> , 2016 , 21, 33-9 | 5.7 | 55 |
| 358 | Effect of the RET Inhibitor Vandetanib in a Patient With RET Fusion-Positive Metastatic Non-Small-Cell Lung Cancer. <i>Journal of Clinical Oncology</i> , 2016 , 34, e141-4 | 2.2 | 55 |
| 357 | Oncogenic alterations in ERBB2/HER2 represent potential therapeutic targets across tumors from diverse anatomic sites of origin. <i>Oncologist</i> , 2015 , 20, 7-12 | 5.7 | 54 |
| 356 | Characterization of Clinical Cases of Advanced Papillary Renal Cell Carcinoma via Comprehensive Genomic Profiling. <i>European Urology</i> , 2018 , 73, 71-78 | 10.2 | 54 |
| 355 | BRAFV600E Mutations in High-Grade Colorectal Neuroendocrine Tumors May Predict Responsiveness to BRAF-MEK Combination Therapy. <i>Cancer Discovery</i> , 2016 , 6, 594-600 | 24.4 | 52 |
| 354 | On-target Resistance to the Mutant-Selective EGFR Inhibitor Osimertinib Can Develop in an Allele-Specific Manner Dependent on the Original EGFR-Activating Mutation. <i>Clinical Cancer Research</i> , 2019 , 25, 3341-3351 | 12.9 | 51 |
| 353 | TP53 Alterations Correlate with Response to VEGF/VEGFR Inhibitors: Implications for Targeted Therapeutics. <i>Molecular Cancer Therapeutics</i> , 2016 , 15, 2475-2485 | 6.1 | 49 |

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|-----|--|------|----|
| 352 | Clinical Actionability of Comprehensive Genomic Profiling for Management of Rare or Refractory Cancers. <i>Oncologist</i> , 2016 , 21, 1315-1325 | 5.7 | 49 |
| 351 | Systemic and CNS activity of the RET inhibitor vandetanib combined with the mTOR inhibitor everolimus in KIF5B-RET re-arranged non-small cell lung cancer with brain metastases. <i>Lung Cancer</i> , 2015 , 89, 76-9 | 5.9 | 48 |
| 350 | Real-time genomic profiling of histiocytoses identifies early-kinase domain BRAF alterations while improving treatment outcomes. <i>JCI Insight</i> , 2017 , 2, e89473 | 9.9 | 48 |
| 349 | Analysis of DNA Damage Response Gene Alterations and Tumor Mutational Burden Across 17,486 Tubular Gastrointestinal Carcinomas: Implications for Therapy. <i>Oncologist</i> , 2019 , 24, 1340-1347 | 5.7 | 43 |
| 348 | Pediatric, Adolescent, and Young Adult Thyroid Carcinoma Harbors Frequent and Diverse Targetable Genomic Alterations, Including Kinase Fusions. <i>Oncologist</i> , 2017 , 22, 255-263 | 5.7 | 42 |
| 347 | Checkpoint inhibitor is active against large cell neuroendocrine carcinoma with high tumor mutation burden 2017 , 5, 75 | | 42 |
| 346 | Comprehensive genomic profiling of extrahepatic cholangiocarcinoma reveals a long tail of therapeutic targets. <i>Journal of Clinical Pathology</i> , 2016 , 69, 403-8 | 3.9 | 40 |
| 345 | Pan-Cancer Analysis of and Genomic Alterations and Their Association With Genomic Instability as Measured by Genome-Wide Loss of Heterozygosity. <i>JCO Precision Oncology</i> , 2020 , 4, 442-465 | 3.6 | 38 |
| 344 | Response of an ERBB2-mutated inflammatory breast carcinoma to human epidermal growth factor receptor 2-targeted therapy. <i>Journal of Clinical Oncology</i> , 2014 , 32, e88-91 | 2.2 | 38 |
| 343 | Acquired FGFR and FGF Alterations Confer Resistance to Estrogen Receptor (ER) Targeted Therapy in ER Metastatic Breast Cancer. <i>Clinical Cancer Research</i> , 2020 , 26, 5974-5989 | 12.9 | 37 |
| 342 | Hybrid Capture-Based Genomic Profiling of Circulating Tumor DNA from Patients with Advanced Cancers of the Gastrointestinal Tract or Anus. <i>Clinical Cancer Research</i> , 2018 , 24, 1881-1890 | 12.9 | 36 |
| 341 | Combined Blockade of Activating Mutations and ER Results in Synthetic Lethality of ER+/HER2 Mutant Breast Cancer. <i>Clinical Cancer Research</i> , 2019 , 25, 277-289 | 12.9 | 35 |
| 340 | Clinical Benefit in Response to Palbociclib Treatment in Refractory Uterine Leiomyosarcomas with a Common Alteration. <i>Oncologist</i> , 2017 , 22, 416-421 | 5.7 | 33 |
| 339 | Detection of Known and Novel FGFR Fusions in Non-Small Cell Lung Cancer by Comprehensive Genomic Profiling. <i>Journal of Thoracic Oncology</i> , 2019 , 14, 54-62 | 8.9 | 33 |
| 338 | Detection of clonal hematopoiesis of indeterminate potential in clinical sequencing of solid tumor specimens. <i>Blood</i> , 2018 , 131, 2501-2505 | 2.2 | 32 |
| 337 | Comprehensive genomic profiling of malignant phyllodes tumors of the breast. <i>Breast Cancer Research and Treatment</i> , 2017 , 162, 597-602 | 4.4 | 31 |
| 336 | Comprehensive genomic profiling of different subtypes of nasopharyngeal carcinoma reveals similarities and differences to guide targeted therapy. <i>Cancer</i> , 2017 , 123, 3628-3637 | 6.4 | 30 |
| 335 | Identification of a novel TMEM106B-ROS1 fusion variant in lung adenocarcinoma by comprehensive genomic profiling. <i>Lung Cancer</i> , 2015 , 88, 352-4 | 5.9 | 30 |

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| 334 | Correlation Between Molecular Subclassifications of Clear Cell Renal Cell Carcinoma and Targeted Therapy Response. <i>European Urology Focus</i> , 2016 , 2, 204-209 | 5.1 | 30 |
| 333 | Clonal diversity predicts adverse outcome in chronic lymphocytic leukemia. <i>Leukemia</i> , 2019 , 33, 390-402 | 10.7 | 30 |
| 332 | Genomic landscape of advanced basal cell carcinoma: Implications for precision treatment with targeted and immune therapies. <i>Oncolmmunology</i> , 2018 , 7, e1404217 | 7.2 | 30 |
| 331 | Prospective Comprehensive Genomic Profiling of Primary and Metastatic Prostate Tumors. <i>JCO Precision Oncology</i> , 2019 , 3, | 3.6 | 29 |
| 330 | Emergence of FGFR3-TACC3 fusions as a potential by-pass resistance mechanism to EGFR tyrosine kinase inhibitors in EGFR mutated NSCLC patients. <i>Lung Cancer</i> , 2017 , 111, 61-64 | 5.9 | 29 |
| 329 | Presence of both alterations in FGFR/FGF and PI3K/AKT/mTOR confer improved outcomes for patients with metastatic breast cancer treated with PI3K/AKT/mTOR inhibitors. <i>Oncoscience</i> , 2016 , 3, 164-72 | 0.8 | 29 |
| 328 | Hybrid Capture-Based Genomic Profiling of Circulating Tumor DNA from Patients with Advanced Non-Small Cell Lung Cancer. <i>Journal of Thoracic Oncology</i> , 2019 , 14, 255-264 | 8.9 | 28 |
| 327 | Phosphatidylinositol 3-kinase pathway genomic alterations in 60,991 diverse solid tumors informs targeted therapy opportunities. <i>Cancer</i> , 2019 , 125, 1185-1199 | 6.4 | 27 |
| 326 | Use of comprehensive genomic profiling to direct point-of-care management of patients with gynecologic cancers. <i>Gynecologic Oncology</i> , 2016 , 141, 2-9 | 4.9 | 26 |
| 325 | Comprehensive Genomic Profiling of Esthesioneuroblastoma Reveals Additional Treatment Options. <i>Oncologist</i> , 2017 , 22, 834-842 | 5.7 | 26 |
| 324 | Biomarker-driven therapies for previously treated squamous non-small-cell lung cancer (Lung-MAP SWOG S1400): a biomarker-driven master protocol. <i>Lancet Oncology, The</i> , 2020 , 21, 1589-1601 | 21.7 | 26 |
| 323 | A Case of Metastatic Atypical Neuroendocrine Tumor with Translocation and Diffuse Brain Metastases. <i>Oncologist</i> , 2017 , 22, 768-773 | 5.7 | 25 |
| 322 | Targeted genomic landscape of metastases compared to primary tumours in clear cell metastatic renal cell carcinoma. <i>British Journal of Cancer</i> , 2018 , 118, 1238-1242 | 8.7 | 25 |
| 321 | BRCA2 Reversion Mutation Associated With Acquired Resistance to Olaparib in Estrogen Receptor-positive Breast Cancer Detected by Genomic Profiling of Tissue and Liquid Biopsy. <i>Clinical Breast Cancer</i> , 2018 , 18, 184-188 | 3 | 25 |
| 320 | Durable Response to Crizotinib in a MET-Amplified, KRAS-Mutated Carcinoma of Unknown Primary. <i>Case Reports in Oncology</i> , 2014 , 7, 503-8 | 1 | 25 |
| 319 | Mutation of MET Y1230 as an Acquired Mechanism of Crizotinib Resistance in NSCLC with MET Exon 14 Skipping. <i>Journal of Thoracic Oncology</i> , 2017 , 12, e89-e90 | 8.9 | 25 |
| 318 | Tumor mutational burden as a potential biomarker for PD1/PD-L1 therapy in colorectal cancer.. <i>Journal of Clinical Oncology</i> , 2016 , 34, 3587-3587 | 2.2 | 24 |
| 317 | Profiling of 3,634 cholangiocarcinomas (CCA) to identify genomic alterations (GA), tumor mutational burden (TMB), and genomic loss of heterozygosity (gLOH).. <i>Journal of Clinical Oncology</i> , 2019 , 37, 4087-4087 | 2.2 | 24 |

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| 316 | Antitumor Response of VEGFR2- and VEGFR3-Amplified Angiosarcoma to Pazopanib. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2016 , 14, 499-502 | 7.3 | 23 |
| 315 | Pan-Cancer Analysis of Loss-of-Function Alterations and Their Association with the Focal Tandem-Duplicator Phenotype. <i>Oncologist</i> , 2019 , 24, 1526-1533 | 5.7 | 23 |
| 314 | Comprehensive Genomic Profiling Identifies Frequent Drug-Sensitive EGFR Exon 19 Deletions in NSCLC not Identified by Prior Molecular Testing. <i>Clinical Cancer Research</i> , 2016 , 22, 3281-5 | 12.9 | 22 |
| 313 | Comprehensive genomic profiling identifies novel NTRK fusions in neuroendocrine tumors. <i>Oncotarget</i> , 2018 , 9, 35809-35812 | 3.3 | 22 |
| 312 | RET Fusion Lung Carcinoma: Response to Therapy and Clinical Features in a Case Series of 14 Patients. <i>Clinical Lung Cancer</i> , 2017 , 18, e223-e232 | 4.9 | 21 |
| 311 | Acquired ALK L1152R Mutation Confers Resistance to Ceritinib and Predicts Response to Alectinib. <i>Journal of Thoracic Oncology</i> , 2016 , 11, e87-8 | 8.9 | 21 |
| 310 | Mutational Landscapes of Smoking-Related Cancers in Caucasians and African Americans: Precision Oncology Perspectives at Wake Forest Baptist Comprehensive Cancer Center. <i>Theranostics</i> , 2017 , 7, 2914-2923 ^{12-1, 20} | 12.1 | 20 |
| 309 | Impact of next-generation sequencing (NGS) on diagnostic and therapeutic options in soft-tissue and bone sarcoma.. <i>Journal of Clinical Oncology</i> , 2017 , 35, 11001-11001 | 2.2 | 20 |
| 308 | Comprehensive Genomic Profiling of Upper-tract and Bladder Urothelial Carcinoma. <i>European Urology Focus</i> , 2021 , 7, 1339-1346 | 5.1 | 20 |
| 307 | Exceptional Response on Addition of Everolimus to Taxane in Urothelial Carcinoma Bearing an NF2 Mutation. <i>European Urology</i> , 2015 , 67, 1195-1196 | 10.2 | 19 |
| 306 | Comprehensive Assessment of Immuno-oncology Biomarkers in Adenocarcinoma, Urothelial Carcinoma, and Squamous-cell Carcinoma of the Bladder. <i>European Urology</i> , 2020 , 77, 548-556 | 10.2 | 19 |
| 305 | Genomic profiling of cell-free circulating tumor DNA in patients with colorectal cancer and its fidelity to the genomics of the tumor biopsy. <i>Journal of Gastrointestinal Oncology</i> , 2019 , 10, 831-840 | 2.8 | 18 |
| 304 | Antitumor response of an ERBB2 amplified inflammatory breast carcinoma with EGFR mutation to the EGFR-TKI erlotinib. <i>Clinical Breast Cancer</i> , 2014 , 14, e14-6 | 3 | 18 |
| 303 | Personalized Treatment for a Patient With a BRAF V600E Mutation Using Dabrafenib and a Tumor Treatment Fields Device in a High-Grade Glioma Arising From Ganglioglioma. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2016 , 14, 1345-1350 | 7.3 | 18 |
| 302 | Exceptional durable response to everolimus in a patient with biphenotypic breast cancer harboring an variant. <i>Journal of Physical Education and Sports Management</i> , 2017 , 3, | 2.8 | 17 |
| 301 | C2 Domain Deletions Hyperactivate Phosphoinositide 3-kinase (PI3K), Generate Oncogene Dependence, and Are Exquisitely Sensitive to PI3K Inhibitors. <i>Clinical Cancer Research</i> , 2018 , 24, 1426-1435 ¹²⁻⁹ | 12.9 | 17 |
| 300 | Comprehensive Genomic Profiling of Hodgkin Lymphoma Reveals Recurrently Mutated Genes and Increased Mutation Burden. <i>Oncologist</i> , 2019 , 24, 219-228 | 5.7 | 17 |
| 299 | Comprehensive genetic alteration profiling in primary and recurrent glioblastoma. <i>Journal of Neuro-Oncology</i> , 2019 , 142, 111-118 | 4.8 | 17 |

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| 298 | Clinical utility of tumor genomic profiling in patients with high plasma circulating tumor DNA burden or metabolically active tumors. <i>Journal of Hematology and Oncology</i> , 2018 , 11, 129 | 22.4 | 17 |
| 297 | Detection of an Fusion in Colorectal Carcinoma by Hybrid Capture-Based Assay of Circulating Tumor DNA. <i>Oncologist</i> , 2017 , 22, 774-779 | 5.7 | 14 |
| 296 | Next Generation Sequencing Reveals Potentially Actionable Alterations in the Majority of Patients with Lymphoid Malignancies. <i>JCO Precision Oncology</i> , 2017 , 1, | 3.6 | 14 |
| 295 | Retrospective analysis of real-world data to determine clinical outcomes of patients with advanced non-small cell lung cancer following cell-free circulating tumor DNA genomic profiling. <i>Lung Cancer</i> , 2020 , 148, 69-78 | 5.9 | 14 |
| 294 | Genomic Profiling of T-Cell Neoplasms Reveals Frequent and Mutations With Clonal Evasion From Targeted Therapies. <i>JCO Precision Oncology</i> , 2018 , 2018, | 3.6 | 14 |
| 293 | General paucity of genomic alteration and low tumor mutation burden in refractory and metastatic hepatoblastoma: comprehensive genomic profiling study. <i>Human Pathology</i> , 2017 , 70, 84-91 | 3.7 | 13 |
| 292 | Phenotypic and Genomic Determinants of Immunotherapy Response Associated with Squamousness. <i>Cancer Immunology Research</i> , 2019 , 7, 866-873 | 12.5 | 13 |
| 291 | Clinically advanced and metastatic pure mucinous carcinoma of the breast: a comprehensive genomic profiling study. <i>Breast Cancer Research and Treatment</i> , 2016 , 155, 405-13 | 4.4 | 13 |
| 290 | Comprehensive genomic profiling of biliary tract cancers to reveal tumor-specific differences and frequency of clinically relevant genomic alterations.. <i>Journal of Clinical Oncology</i> , 2015 , 33, 4009-4009 | 2.2 | 13 |
| 289 | MDM2 amplification (Amp) to mediate cabozantinib resistance in patients (Pts) with advanced RET-rearranged lung cancers.. <i>Journal of Clinical Oncology</i> , 2016 , 34, 9068-9068 | 2.2 | 13 |
| 288 | Landscape of genomic alterations (GA) and tumor mutational burden (TMB) in different metastatic melanoma (MM) subtypes.. <i>Journal of Clinical Oncology</i> , 2017 , 35, 9536-9536 | 2.2 | 13 |
| 287 | Approach to evaluating tumor mutational burden in routine clinical practice. <i>Translational Lung Cancer Research</i> , 2018 , 7, 678-681 | 4.4 | 13 |
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