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List of Publications by Year
in descending order

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33 papers	1,493 citations	471509 17 h-index	377865 34 g-index
34 all docs	34 docs citations	34 times ranked	3082 citing authors

#	Article	IF	Citations
1	Four-Year Laboratory Performance of the First College of American Pathologists In Silico Next-Generation Sequencing Bioinformatics Proficiency Testing Surveys. Archives of Pathology and Laboratory Medicine, 2023, 147, 137-142.	2.5	3
2	Standardized evidence-based approach for assessment of oncogenic and clinical significance of NTRK fusions. Cancer Genetics, 2022, 264-265, 50-59.	0.4	5
3	PD-L1 Tumor Cell Expression in Upper Tract Urothelial Carcinomas is Associated With Higher Pathologic Stage. Applied Immunohistochemistry and Molecular Morphology, 2021, Publish Ahead of Print, 56-61.	1.2	3
4	Lorlatinib in a Child with ALK-Fusionâ€“Positive High-Grade Glioma. New England Journal of Medicine, 2021, 385, 761-763.	27.0	27
5	Female adnexal tumors of probable Wolffian origin: morphological, immunohistochemical, and molecular analysis of 15 cases. Modern Pathology, 2020, 33, 734-747.	5.5	23
6	Multi-Institutional Evaluation of Interrater Agreement of Variant Classification Based on the 2017 Association for Molecular Pathology, American Society of Clinical Oncology, and College of American Pathologists Standards and Guidelines for the Interpretation and Reporting of Sequence Variants in Cancer. Journal of Molecular Diagnostics, 2020, 22, 284-293.	2.8	10
7	Successful lung cancer EGFR sequencing from DNA extracted from TTF-1 immunohistochemistry slides: a new means to extend insufficient tissue. Human Pathology, 2020, 97, 52-59.	2.0	3
8	Metastatic HPV-Associated Oropharyngeal Versus Primary Pulmonary Squamous Cell Carcinoma: is p16 Immunostain Useful?. Head and Neck Pathology, 2020, 14, 966-973.	2.6	4
9	Tumor PD-L1 expression in malignant pleural and peritoneal mesothelioma by Dako PD-L1 22C3 pharmDx and Dako PD-L1 28-8 pharmDx assays. Human Pathology, 2019, 87, 11-17.	2.0	40
10	Cribriform-Morular Variant of Papillary Thyroid Carcinoma With Poorly Differentiated Features: A Case Report With Immunohistochemical and Molecular Genetic Analysis. International Journal of Surgical Pathology, 2019, 27, 294-304.	0.8	3
11	Clinical performance of endobronchial ultrasoundâ€guided transbronchial needle aspiration for assessing programmed death ligandâ€1 expression in nonsmall cell lung cancer. Diagnostic Cytopathology, 2018, 46, 378-383.	1.0	38
12	Feasibility of Endobronchial Ultrasound-guided Transbronchial Needle Aspiration Cytology Specimens for Next Generation Sequencing in Nonâ€“small-cell Lung Cancer. Clinical Lung Cancer, 2018, 19, 230-238.e2.	2.6	63
13	System for Informatics in the Molecular Pathology Laboratory. Journal of Molecular Diagnostics, 2018, 20, 522-532.	2.8	8
14	Treatment outcomes and HPV characteristics for an institutional cohort of patients with anal cancer receiving concurrent chemotherapy and intensity-modulated radiation therapy. PLoS ONE, 2018, 13, e0194234.	2.5	6
15	Colorectal cancer molecular profiling: from IHC to NGS in search of optimal algorithm. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2017, 471, 235-242.	2.8	6
16	High prevalence of Mi<scp>TF</scp> staining in undifferentiated pleomorphic sarcoma: caution in the use of melanocytic markers in sarcoma. Histopathology, 2017, 70, 734-745.	2.9	16
17	Clinical Validation of a Next-Generation Sequencing Genomic Oncology Panel via Cross-Platform Benchmarking against Established Amplicon Sequencing Assays. Journal of Molecular Diagnostics, 2017, 19, 43-56.	2.8	105
18	Diagnostic evaluation of RNA sequencing for the detection of genetic abnormalities associated with Ph-like acute lymphoblastic leukemia (ALL). Leukemia and Lymphoma, 2017, 58, 950-958.	1.3	18

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19	Characterization of Molecular Alterations in an Unusual Case of Lynch Syndromeâ€“Associated Adrenocortical Carcinoma. AJSP Review and Reports, 2017, 22, 164-170.	0.1	3
20	Disorders of sterol biosynthesis. Translational Science of Rare Diseases, 2016, 1, 145-182.	1.5	3
21	Polyglucosan bodies in intramuscular nerve branches are a poor predictor of GBE1 mutation and adult polyglucosan body disease. Muscle and Nerve, 2016, 53, 473-475.	2.2	3
22	Unclassified renal cell carcinoma with tubulopapillary architecture, clear cell phenotype, and chromosome 8 monosomy: a new kid on the block. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2016, 469, 81-91.	2.8	15
23	Identification of a structurally novel BTK mutation that drives ibrutinib resistance in CLL. Oncotarget, 2016, 7, 68833-68841.	1.8	67
24	Circulating Tumor DNA Testing for Liver Cancer. Cellular and Molecular Gastroenterology and Hepatology, 2015, 1, 458-459.	4.5	4
25	Do Circulating Tumor Cells, Exosomes, and Circulating Tumor Nucleic Acids Have Clinical Utility?. Journal of Molecular Diagnostics, 2015, 17, 209-224.	2.8	176
26	Amplicon Indel Hunter Is a Novel Bioinformatics Tool to Detect Large Somatic Insertion/Deletion Mutations in Amplicon-Based Next-Generation Sequencing Data. Journal of Molecular Diagnostics, 2015, 17, 635-643.	2.8	26
27	High prevalence of somatic MAP2K1 mutations in BRAF V600Eâ€“negative Langerhans cell histiocytosis. Blood, 2014, 124, 1655-1658.	1.4	311
28	Detection of MPL Mutations by a Novel Allele-Specific PCR-Based Strategy. Journal of Molecular Diagnostics, 2013, 15, 810-818.	2.8	39
29	A Multiplexed Fragment Analysis-Based Assay for Detection of JAK2 Exon 12 Mutations. Journal of Molecular Diagnostics, 2013, 15, 592-599.	2.8	20
30	Diagnostic Utility of Microsatellite Genotyping for Molar Pregnancy Testing. Archives of Pathology and Laboratory Medicine, 2013, 137, 55-63.	2.5	31
31	Characterization of large genomic deletions in the FBN1 gene using multiplex ligation-dependent probe amplification. BMC Medical Genetics, 2011, 12, 119.	2.1	24
32	Septin 9 methylated DNA is a sensitive and specific blood test for colorectal cancer. BMC Medicine, 2011, 9, 133.	5.5	361
33	A novel Xâ€linked multiple congenital anomaly syndrome associated with an EBP mutation. American Journal of Medical Genetics, Part A, 2010, 152A, 2838-2844.	1.2	28