

Michael R Clay

List of Publications by Year in descending order

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Version: 2024-02-01

40
papers

1,492
citations

567281

15
h-index

361022

35
g-index

46
all docs

46
docs citations

46
times ranked

3455
citing authors

#	ARTICLE	IF	CITATIONS
1	Clinical and Functional Significance of TP53 Exon 4 Intron 4 Splice Junction Variants. <i>Molecular Cancer Research</i> , 2022, 20, 207-216.	3.4	4
2	Pathological and Genetic Stratification for Management of Adrenocortical Carcinoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, 1159-1169.	3.6	7
3	TERT Expression in Wilms Tumor Is Regulated by Promoter Mutation or Hypermethylation, WT1, and N-MYC. <i>Cancers</i> , 2022, 14, 1655.	3.7	3
4	The myogenesis program drives clonal selection and drug resistance in rhabdomyosarcoma. <i>Developmental Cell</i> , 2022, 57, 1226-1240.e8.	7.0	24
5	Evaluating bone biopsy quality by technique in an animal model. , 2022, 2, 100008.		1
6	The Common Germline TP53-R337H Mutation Is Hypomorphic and Confers Incomplete Penetrance and Late Tumor Onset in a Mouse Model. <i>Cancer Research</i> , 2021, 81, 2442-2456.	0.9	9
7	Genomes for Kids: The Scope of Pathogenic Mutations in Pediatric Cancer Revealed by Comprehensive DNA and RNA Sequencing. <i>Cancer Discovery</i> , 2021, 11, 3008-3027.	9.4	88
8	Abstract 642: Genomes for Kids: Comprehensive DNA and RNA sequencing defining the scope of actionable mutations in pediatric cancer. , 2021, , .		0
9	SLFN11 is Widely Expressed in Pediatric Sarcoma and Induces Variable Sensitization to Replicative Stress Caused By DNA-Damaging Agents. <i>Molecular Cancer Therapeutics</i> , 2021, 20, 2151-2165.	4.1	6
10	A genetic mouse model with postnatal Nf1 and p53 loss recapitulates the histology and transcriptome of human malignant peripheral nerve sheath tumor. <i>Neuro-Oncology Advances</i> , 2021, 3, vdab129.	0.7	3
11	An update on the central nervous system manifestations of Fraumeni syndrome. <i>Acta Neuropathologica</i> , 2020, 139, 669-687.	7.7	44
12	Partial response to carboplatin, etoposide phosphate, and atezolizumab in a pediatric patient with high-grade metastatic tumor with rhabdoid and focal neuroendocrine features. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28048.	1.5	5
13	A phase I trial of talazoparib and irinotecan with and without temozolomide in children and young adults with recurrent or refractory solid malignancies. <i>European Journal of Cancer</i> , 2020, 137, 204-213.	2.8	39
14	Dasatinib induces a dramatic response in a child with refractory juvenile xanthogranuloma with a novel MRC1-PDGFRB fusion. <i>Blood Advances</i> , 2020, 4, 2991-2995.	5.2	10
15	Route of 41BB/41BBL Costimulation Determines Effector Function of B7-H3-CAR.CD28 T Cells. <i>Molecular Therapy - Oncolytics</i> , 2020, 18, 202-214.	4.4	37
16	A single-cell and single-nucleus RNA-Seq toolbox for fresh and frozen human tumors. <i>Nature Medicine</i> , 2020, 26, 792-802.	30.7	381
17	Benign infiltrative myofibroblastic neoplasms of childhood with USP6 gene rearrangement. <i>Histopathology</i> , 2020, 77, 760-768.	2.9	15
18	XAF1 as a modifier of p53 function and cancer susceptibility. <i>Science Advances</i> , 2020, 6, eaba3231.	10.3	37

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19	MYCN amplification and ATRX mutations are incompatible in neuroblastoma. <i>Nature Communications</i> , 2020, 11, 913.	12.8	66
20	An addition to the evolving spectrum of lipofibromatosis and lipofibromatosis-like neural tumor: Molecular findings in an unusual phenotype aid in accurate classification. <i>Pathology Research and Practice</i> , 2020, 216, 152942.	2.3	7
21	Dedifferentiation in SDH-Deficient Gastrointestinal Stromal Tumor: A Report With Histologic, Immunophenotypic, and Molecular Characterization. <i>Pediatric and Developmental Pathology</i> , 2019, 22, 492-498.	1.0	15
22	Forty-five patient-derived xenografts capture the clinical and biological heterogeneity of Wilms tumor. <i>Nature Communications</i> , 2019, 10, 5806.	12.8	27
23	DNA Methylation Profiling Reveals Prognostically Significant Groups in Pediatric Adrenocortical Tumors: A Report From the International Pediatric Adrenocortical Tumor Registry. <i>JCO Precision Oncology</i> , 2019, 3, 1-21.	3.0	6
24	OR02-1 DNA Methylation Profiling in Pediatric Adrenocortical Tumors Reveals Distinct Methylation Signatures with Prognostic Significance: A Report from the International Pediatric Adrenocortical Tumor Registry. <i>Journal of the Endocrine Society</i> , 2019, 3, .	0.2	0
25	SAT-LB058 Effect of a Genetic Modifier of Cancer Risk in TP53 Mutation Carriers. <i>Journal of the Endocrine Society</i> , 2019, 3, .	0.2	0
26	Identification of Therapeutic Targets in Rhabdomyosarcoma through Integrated Genomic, Epigenomic, and Proteomic Analyses. <i>Cancer Cell</i> , 2018, 34, 411-426.e19.	16.8	106
27	Genitopatellar syndrome and neuroblastoma: The multidisciplinary management of a previously unreported association. <i>Pediatric Blood and Cancer</i> , 2018, 65, e27373.	1.5	1
28	Recurrent BCOR internal tandem duplication and BCOR or BCL6 expression distinguish primitive myxoid mesenchymal tumor of infancy from congenital infantile fibrosarcoma. <i>Modern Pathology</i> , 2017, 30, 884-891.	5.5	60
29	Clear cell sarcoma of kidney involving a horseshoe kidney and harboring <i>EGFR</i> internal tandem duplication. <i>Pediatric Blood and Cancer</i> , 2017, 64, e26602.	1.5	14
30	Calcifying nested stromal-epithelial tumor (CNSET) of the liver in Beckwith-Wiedemann syndrome. <i>European Journal of Medical Genetics</i> , 2017, 60, 136-139.	1.3	22
31	Orthotopic patient-derived xenografts of paediatric solid tumours. <i>Nature</i> , 2017, 549, 96-100.	27.8	223
32	Bioinformatics Education in Pathology Training: Current Scope and Future Direction. <i>Cancer Informatics</i> , 2017, 16, 117693511770338.	1.9	9
33	Identification of Clinical and Biologic Correlates Associated With Outcome in Children With Adrenocortical Tumors Without Germline TP53 Mutations: A St Jude Adrenocortical Tumor Registry and Children's Oncology Group Study. <i>Journal of Clinical Oncology</i> , 2017, 35, 3956-3963.	1.6	33
34	Bullous Kaposiform Hemangioendothelioma Masquerading as Aplasia Cutis Congenita. <i>Pediatric Dermatology</i> , 2016, 33, e267-9.	0.9	0
35	MDM2 and CDK4 Immunohistochemistry: Should It Be Used in Problematic Differentiated Lipomatous Tumors?. <i>American Journal of Surgical Pathology</i> , 2016, 40, 1647-1652.	3.7	67
36	MDM2 Amplification in Problematic Lipomatous Tumors. <i>American Journal of Surgical Pathology</i> , 2015, 39, 1433-1439.	3.7	70

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37	Risk of Secondary Malignancy (Including Breast) in Patients With Mismatch-repair Protein Deficiency. American Journal of Surgical Pathology, 2014, 38, 1494-1500.	3.7	1
38	MAST2 and NOTCH1 translocations in breast carcinoma and associated pre-invasive lesions. Human Pathology, 2013, 44, 2837-2844.	2.0	14
39	Clinicopathologic Characteristics of HER2 FISH-ambiguous Breast Cancer at a Single Institution. American Journal of Surgical Pathology, 2013, 37, 120-127.	3.7	6
40	No Evidence of Human Clonally Transmissible Anogenital Cancer. International Journal of Surgical Pathology, 2010, 18, 304-304.	0.8	0