

D Wade Clapp

List of Publications by Year in descending order

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Version: 2024-02-01

51
papers

2,452
citations

430874

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233421

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docs citations

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4304
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#	ARTICLE	IF	CITATIONS
1	SIK2 kinase synthetic lethality is driven by spindle assembly defects in <i>FANCA</i> -deficient cells. <i>Molecular Oncology</i> , 2022, 16, 860-884.	4.6	2
2	Irradiation of Nf1 mutant mouse models of spinal plexiform neurofibromas drives pathologic progression and decreases survival. <i>Neuro-Oncology Advances</i> , 2021, 3, vdab063.	0.7	4
3	Cabozantinib for neurofibromatosis type 1-related plexiform neurofibromas: a phase 2 trial. <i>Nature Medicine</i> , 2021, 27, 165-173.	30.7	46
4	Exploring transcriptional regulators Ref-1 and STAT3 as therapeutic targets in malignant peripheral nerve sheath tumours. <i>British Journal of Cancer</i> , 2021, 124, 1566-1580.	6.4	12
5	PAK1 inhibition reduces tumor size and extends the lifespan of mice in a genetically engineered mouse model of Neurofibromatosis Type 2 (NF2). <i>Human Molecular Genetics</i> , 2021, 30, 1607-1617.	2.9	12
6	Brigatinib causes tumor shrinkage in both NF2-deficient meningioma and schwannoma through inhibition of multiple tyrosine kinases but not ALK. <i>PLoS ONE</i> , 2021, 16, e0252048.	2.5	19
7	Mitotic Errors Promote Genomic Instability and Leukemia in a Novel Mouse Model of Fanconi Anemia. <i>Frontiers in Oncology</i> , 2021, 11, 752933.	2.8	4
8	<i>Nf1</i> -Mutant Tumors Undergo Transcriptome and Kinome Remodeling after Inhibition of either mTOR or MEK. <i>Molecular Cancer Therapeutics</i> , 2020, 19, 2382-2395.	4.1	3
9	Genetic disruption of the small GTPase RAC1 prevents plexiform neurofibroma formation in mice with neurofibromatosis type 17. <i>Journal of Biological Chemistry</i> , 2020, 295, 9948-9958.	3.4	7
10	Early administration of imatinib mesylate reduces plexiform neurofibroma tumor burden with durable results after drug discontinuation in a mouse model of neurofibromatosis type 1. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28372.	1.5	3
11	Addressing Gaps in Pediatric Scientist Development: The Department Chair View of 2 AMSPDC-Sponsored Programs. <i>Journal of Pediatrics</i> , 2020, 222, 7-12.e4.	1.8	6
12	Selumetinib in Children with Inoperable Plexiform Neurofibromas. <i>New England Journal of Medicine</i> , 2020, 382, 1430-1442.	27.0	360
13	A molecular basis for neurofibroma-associated skeletal manifestations in NF1. <i>Genetics in Medicine</i> , 2020, 22, 1786-1793.	2.4	12
14	Schwannoma development is mediated by Hippo pathway dysregulation and modified by RAS/MAPK signaling. <i>JCI Insight</i> , 2020, 5, .	5.0	14
15	Feasibility of using NF1-GRD and AAV for gene replacement therapy in NF1-associated tumors. <i>Gene Therapy</i> , 2019, 26, 277-286.	4.5	21
16	Cdkn2a (Arf) loss drives NF1-associated atypical neurofibroma and malignant transformation. <i>Human Molecular Genetics</i> , 2019, 28, 2752-2762.	2.9	54
17	A proteasome-resistant fragment of NIK mediates oncogenic NF- κ B signaling in schwannomas. <i>Human Molecular Genetics</i> , 2019, 28, 572-583.	2.9	5
18	Hospitalist Medicine's Chairs' Perspective of Specialty Status and Training Requirements. <i>Journal of Pediatrics</i> , 2018, 193, 4-8.e1.	1.8	3

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19	Fanconi anaemia and cancer: an intricate relationship. <i>Nature Reviews Cancer</i> , 2018, 18, 168-185.	28.4	275
20	From bedside to bench and back: Translating ASD models. <i>Progress in Brain Research</i> , 2018, 241, 113-158.	1.4	2
21	NFM-09. PRELIMINARY REPORT OF A MULTICENTER, PHASE 2 STUDY OF BEVACIZUMAB IN CHILDREN AND ADULTS WITH NEUROFIBROMATOSIS 2 AND PROGRESSIVE VESTIBULAR SCHWANNOMAS: AN NF CLINICAL TRIALS CONSORTIUM STUDY. <i>Neuro-Oncology</i> , 2018, 20, i144-i144.	1.2	0
22	Traditional and systems biology based drug discovery for the rare tumor syndrome neurofibromatosis type 2. <i>PLoS ONE</i> , 2018, 13, e0197350.	2.5	17
23	Contributions of inflammation and tumor microenvironment to neurofibroma tumorigenesis. <i>Journal of Clinical Investigation</i> , 2018, 128, 2848-2861.	8.2	101
24	Chemopreventative celecoxib fails to prevent schwannoma formation or sensorineural hearing loss in genetically engineered murine model of neurofibromatosis type 2. <i>Oncotarget</i> , 2018, 9, 718-725.	1.8	6
25	A Collaborative Model for Accelerating the Discovery and Translation of Cancer Therapies. <i>Cancer Research</i> , 2017, 77, 5706-5711.	0.9	22
26	Preclinical Evidence for the Use of Sunitinib Malate in the Treatment of Plexiform Neurofibromas. <i>Pediatric Blood and Cancer</i> , 2016, 63, 206-213.	1.5	20
27	Fanconi Anemia Proteins Function in Mitophagy and Immunity. <i>Cell</i> , 2016, 165, 867-881.	28.9	205
28	The importance of nerve microenvironment for schwannoma development. <i>Acta Neuropathologica</i> , 2016, 132, 289-307.	7.7	62
29	<i>Nf1</i> ^{+/Δ} monocytes/macrophages induce neointima formation via CCR2 activation. <i>Human Molecular Genetics</i> , 2016, 25, 1129-1139.	2.9	13
30	Spatially- and temporally-controlled postnatal p53 knockdown cooperates with embryonic Schwann cell precursor <i>Nf1</i> gene loss to promote malignant peripheral nerve sheath tumor formation. <i>Oncotarget</i> , 2016, 7, 7403-7414.	1.8	30
31	A murine model of neurofibromatosis type 2 that accurately phenocopies human schwannoma formation. <i>Human Molecular Genetics</i> , 2015, 24, 1-8.	2.9	76
32	Social learning and amygdala disruptions in <i>Nf1</i> mice are rescued by blocking p21-activated kinase. <i>Nature Neuroscience</i> , 2014, 17, 1583-1590.	14.8	106
33	Fanconi anemia and the cell cycle: new perspectives on aneuploidy. <i>F1000prime Reports</i> , 2014, 6, 23.	5.9	23
34	FANCA Controls Mitotic Phosphosignaling Networks To Ensure Genome Stability During Cell Division. <i>Blood</i> , 2013, 122, 801-801.	1.4	0
35	Generation Of FANCA ^{-/-} Human CD34 ⁺ Hematopoietic Stem Cells By shRNA Knockdown. <i>Blood</i> , 2013, 122, 2903-2903.	1.4	2
36	Normal Hematopoiesis and Neurofibromin-Deficient Myeloproliferative Disease Require Erk. <i>Blood</i> , 2012, 120, 704-704.	1.4	0

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37	PAK1 Regulates Eotaxin-Mediated Murine Eosinophil Migration in Vitro and In Vivo. <i>Blood</i> , 2011, 118, 18-18.	1.4	1
38	Ezrin Regulates Hematopoietic Stem/Progenitor Cell Motility. <i>Blood</i> , 2011, 118, 1282-1282.	1.4	0
39	A Modified Foamy Viral Envelope Enhances Gene Transfer Efficiency and Reduces Toxicity of Lentiviral FANCA Vectors in Fanca ^{-/-} HSCs. <i>Blood</i> , 2009, 114, 696-696.	1.4	2
40	Mast Cells and Tumor Progression. <i>Blood</i> , 2009, 114, SCI-33-SCI-33.	1.4	0
41	Kinase Suppressor of Ras Plays a Critical Role in Modulating Inflammatory Mast Cell Functions. <i>Blood</i> , 2007, 110, 2407-2407.	1.4	0
42	Loss of Pak1 Corrects Multiple Gain of Function Phenotypes in Nf1 ^{+/-} Mast Cells. <i>Blood</i> , 2007, 110, 236-236.	1.4	1
43	Developmental Regulation of the Immune System. <i>Seminars in Perinatology</i> , 2006, 30, 69-72.	2.5	88
44	Suprasynergistic Peripheral Blood Stem Cell Mobilization in Normal and Fanconi Anemia Knockout Mice by the Combination of G-CSF Plus the CXCR4 Antagonist AMD3100 and the CXCR2 Agonist GRO β . <i>Blood</i> , 2006, 108, 3185-3185.	1.4	2
45	Comparative Functional Genomic Analysis of Myelodysplasia (MDS) in Fanconi Anemia (FA). <i>Blood</i> , 2006, 108, 2636-2636.	1.4	0
46	Foamy Viral Vectors Efficiently Transduce Quiescent Hematopoietic Stem/Progenitor Cells (HSC) and Restore the Long Term Repopulating Activity of FancC Δ Stem Cells. <i>Blood</i> , 2005, 106, 182-182.	1.4	1
47	Murine and Human NF1 Haploinsufficient Mast Cells Promote Alterations in Fibroblast Function and Organization of the Extracellular Matrix in Three-Dimensional Collagen Lattices and this Gain in Function Is Abrogated by the Addition of STI-571. <i>Blood</i> , 2004, 104, 1465-1465.	1.4	3
48	Functional Analysis of Leukemia-Associated PTPN11 Mutations in Primary Hematopoietic Cells. <i>Blood</i> , 2004, 104, 2423-2423.	1.4	0
49	Loss of FancC Function Results in Decreased Hematopoietic Stem Cell Repopulating Ability. <i>Blood</i> , 1999, 94, 1-8.	1.4	185
50	Loss of NF1 results in activation of the Ras signaling pathway and leads to aberrant growth in haematopoietic cells. <i>Nature Genetics</i> , 1996, 12, 144-148.	21.4	555
51	The Highest Concentration of Primitive Hematopoietic Progenitor Cells in Cord Blood Is Found in Extremely Premature Infants. <i>Pediatric Research</i> , 1996, 39, 820-825.	2.3	49