

# Michael A Dyer

## List of Publications by Year in descending order

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135  
papers

11,548  
citations

31976

53  
h-index

30087

103  
g-index

147  
all docs

147  
docs citations

147  
times ranked

15355  
citing authors

#	ARTICLE	IF	CITATIONS
1	Neuroblastoma: developmental biology, cancer genomics and immunotherapy. <i>Nature Reviews Cancer</i> , 2013, 13, 397-411.	28.4	632
2	Recurrent Somatic Structural Variations Contribute to Tumorigenesis in Pediatric Osteosarcoma. <i>Cell Reports</i> , 2014, 7, 104-112.	6.4	583
3	Inactivation of the p53 pathway in retinoblastoma. <i>Nature</i> , 2006, 444, 61-66.	27.8	550
4	A novel retinoblastoma therapy from genomic and epigenetic analyses. <i>Nature</i> , 2012, 481, 329-334.	27.8	442
5	Genomic Landscape of Ewing Sarcoma Defines an Aggressive Subtype with Co-Association of <i>STAG2</i> and <i>TP53</i> Mutations. <i>Cancer Discovery</i> , 2014, 4, 1342-1353.	9.4	418
6	Control of Müller glial cell proliferation and activation following retinal injury. <i>Nature Neuroscience</i> , 2000, 3, 873-880.	14.8	400
7	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
8	Association of Age at Diagnosis and Genetic Mutations in Patients With Neuroblastoma. <i>JAMA - Journal of the American Medical Association</i> , 2012, 307, 1062.	7.4	379
9	Prox1 function controls progenitor cell proliferation and horizontal cell genesis in the mammalian retina. <i>Nature Genetics</i> , 2003, 34, 53-58.	21.4	364
10	Targeting Oxidative Stress in Embryonal Rhabdomyosarcoma. <i>Cancer Cell</i> , 2013, 24, 710-724.	16.8	252
11	TNF Counterbalances the Emergence of M2 Tumor Macrophages. <i>Cell Reports</i> , 2015, 12, 1902-1914.	6.4	232
12	Orthotopic patient-derived xenografts of paediatric solid tumours. <i>Nature</i> , 2017, 549, 96-100.	27.8	223
13	The Dynamic Epigenetic Landscape of the Retina During Development, Reprogramming, and Tumorigenesis. <i>Neuron</i> , 2017, 94, 550-568.e10.	8.1	222
14	Regulating proliferation during retinal development. <i>Nature Reviews Neuroscience</i> , 2001, 2, 333-342.	10.2	220
15	MDMX: from bench to bedside. <i>Journal of Cell Science</i> , 2007, 120, 371-378.	2.0	218
16	Cells previously identified as retinal stem cells are pigmented ciliary epithelial cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 6685-6690.	7.1	207
17	A Sleeping Beauty forward genetic screen identifies new genes and pathways driving osteosarcoma development and metastasis. <i>Nature Genetics</i> , 2015, 47, 615-624.	21.4	207
18	p27 <sup>Kip1</sup> and p57 <sup>Kip2</sup> Regulate Proliferation in Distinct Retinal Progenitor Cell Populations. <i>Journal of Neuroscience</i> , 2001, 21, 4259-4271.	3.6	206

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19	The search for the retinoblastoma cell of origin. <i>Nature Reviews Cancer</i> , 2005, 5, 91-101.	28.4	201
20	Rb regulates proliferation and rod photoreceptor development in the mouse retina. <i>Nature Genetics</i> , 2004, 36, 351-360.	21.4	191
21	Six3-mediated auto repression and eye development requires its interaction with members of the Groucho-related family of co-repressors. <i>Development (Cambridge)</i> , 2002, 129, 2835-2849.	2.5	177
22	Differentiated Horizontal Interneurons Clonally Expand to Form Metastatic Retinoblastoma in Mice. <i>Cell</i> , 2007, 131, 378-390.	28.9	174
23	Genomic landscape of paediatric adrenocortical tumours. <i>Nature Communications</i> , 2015, 6, 6302.	12.8	166
24	ATRX and DAXX: Mechanisms and Mutations. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2017, 7, a026567.	6.2	153
25	The Genomic Landscape of Childhood and Adolescent Melanoma. <i>Journal of Investigative Dermatology</i> , 2015, 135, 816-823.	0.7	148
26	Targeting the DNA Repair Pathway in Ewing Sarcoma. <i>Cell Reports</i> , 2014, 9, 829-840.	6.4	141
27	Coexpression of Normally Incompatible Developmental Pathways in Retinoblastoma Genesis. <i>Cancer Cell</i> , 2011, 20, 260-275.	16.8	123
28	Topotecan Combination Chemotherapy in Two New Rodent Models of Retinoblastoma. <i>Clinical Cancer Research</i> , 2005, 11, 7569-7578.	7.0	117
29	Retinoblastoma: One World, One Vision. <i>Pediatrics</i> , 2008, 122, e763-e770.	2.1	115
30	The First Knockout Mouse Model of Retinoblastoma. <i>Cell Cycle</i> , 2004, 3, 950-957.	2.6	113
31	Genetics and Epigenetics of Human Retinoblastoma. <i>Annual Review of Pathology: Mechanisms of Disease</i> , 2015, 10, 547-562.	22.4	109
32	St. Jude Cloud: A Pediatric Cancer Genomic Data-Sharing Ecosystem. <i>Cancer Discovery</i> , 2021, 11, 1082-1099.	9.4	109
33	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
34	RB1 gene inactivation by chromothripsis in human retinoblastoma. <i>Oncotarget</i> , 2014, 5, 438-450.	1.8	104
35	Compensation by tumor suppressor genes during retinal development in mice and humans. <i>BMC Biology</i> , 2006, 4, 14.	3.8	92
36	The first knockout mouse model of retinoblastoma. <i>Cell Cycle</i> , 2004, 3, 952-9.	2.6	90

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37	Targeting the p53 Pathway in Retinoblastoma with Subconjunctival Nutlin-3a. <i>Cancer Research</i> , 2011, 71, 4205-4213.	0.9	89
38	Quantification of Retinogenesis in 3D Cultures Reveals Epigenetic Memory and Higher Efficiency in iPSCs Derived from Rod Photoreceptors. <i>Cell Stem Cell</i> , 2015, 17, 101-115.	11.1	88
39	Pan-neuroblastoma analysis reveals age- and signature-associated driver alterations. <i>Nature Communications</i> , 2020, 11, 5183.	12.8	87
40	Six3-mediated auto repression and eye development requires its interaction with members of the Groucho-related family of co-repressors. <i>Development (Cambridge)</i> , 2002, 129, 2835-49.	2.5	87
41	Synthetic lethality between Rb, p53 and Dicer or miR-17-92 in retinal progenitors suppresses retinoblastoma formation. <i>Nature Cell Biology</i> , 2012, 14, 958-965.	10.3	79
42	The neoepitope landscape in pediatric cancers. <i>Genome Medicine</i> , 2017, 9, 78.	8.2	77
43	Preclinical Models for Neuroblastoma: Establishing a Baseline for Treatment. <i>PLoS ONE</i> , 2011, 6, e19133.	2.5	77
44	Developmental sources of conservation and variation in the evolution of the primate eye. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 8963-8968.	7.1	72
45	Nucleome Dynamics during Retinal Development. <i>Neuron</i> , 2019, 104, 512-528.e11.	8.1	70
46	Regulation of proliferation during central nervous system development. <i>Seminars in Cell and Developmental Biology</i> , 2005, 16, 407-421.	5.0	68
47	CONSORTING: integrating copy-number analysis with structural-variation detection. <i>Nature Methods</i> , 2015, 12, 527-530.	19.0	68
48	Analysis of MDM2 and MDM4 Single Nucleotide Polymorphisms, mRNA Splicing and Protein Expression in Retinoblastoma. <i>PLoS ONE</i> , 2012, 7, e42739.	2.5	68
49	MYCN amplification and ATRX mutations are incompatible in neuroblastoma. <i>Nature Communications</i> , 2020, 11, 913.	12.8	66
50	The Homeodomain Proteins Prox1, Six3 and Chx10 Regulate Proliferation, Cell Fate Specification and Differentiation in the Developing Retina. <i>Cell Cycle</i> , 2003, 2, 347-354.	2.6	65
51	The Childhood Solid Tumor Network: A new resource for the developmental biology and oncology research communities. <i>Developmental Biology</i> , 2016, 411, 287-293.	2.0	63
52	Retinal Cell Type DNA Methylation and Histone Modifications Predict Reprogramming Efficiency and Retinogenesis in 3D Organoid Cultures. <i>Cell Reports</i> , 2018, 22, 2601-2614.	6.4	63
53	ATRX binds to atypical chromatin domains at the 3' exons of zinc finger genes to preserve H3K9me3 enrichment. <i>Epigenetics</i> , 2016, 11, 398-414.	2.7	59
54	Preparation and square wave electroporation of retinal explant cultures. <i>Nature Protocols</i> , 2006, 1, 2710-2718.	12.0	58

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55	Systematic Screening Identifies Dual PI3K and mTOR Inhibition as a Conserved Therapeutic Vulnerability in Osteosarcoma. <i>Clinical Cancer Research</i> , 2015, 21, 3216-3229.	7.0	58
56	The inherited blindness associated protein AIPL1 interacts with the cell cycle regulator protein NUB1. <i>Human Molecular Genetics</i> , 2002, 11, 2723-2733.	2.9	53
57	Whole-Body Physiologically Based Pharmacokinetic Model for Nutlin-3a in Mice after Intravenous and Oral Administration. <i>Drug Metabolism and Disposition</i> , 2011, 39, 15-21.	3.3	53
58	Use of Preclinical Models to Improve Treatment of Retinoblastoma. <i>PLoS Medicine</i> , 2005, 2, e332.	8.4	52
59	Retinoblastoma: From the Two-Hit Hypothesis to Targeted Chemotherapy. <i>Cancer Research</i> , 2007, 67, 7547-7550.	0.9	48
60	Retinoblastoma from human stem cell-derived retinal organoids. <i>Nature Communications</i> , 2021, 12, 4535.	12.8	48
61	Subconjunctival carboplatin and systemic topotecan treatment in preclinical models of retinoblastoma. <i>Cancer</i> , 2011, 117, 421-434.	4.1	46
62	Retinal degeneration in Aipl1-deficient mice: a new genetic model of Leber congenital amaurosis. <i>Molecular Brain Research</i> , 2004, 132, 208-220.	2.3	45
63	N-myc coordinates retinal growth with eye size during mouse development. <i>Genes and Development</i> , 2008, 22, 179-193.	5.9	45
64	The p57Kip2 cyclin kinase inhibitor is expressed by a restricted set of amacrine cells in the rodent retina. <i>Journal of Comparative Neurology</i> , 2001, 429, 601-614.	1.6	44
65	ATRX In-Frame Fusion Neuroblastoma Is Sensitive to EZH2 Inhibition via Modulation of Neuronal Gene Signatures. <i>Cancer Cell</i> , 2019, 36, 512-527.e9.	16.8	44
66	Progress in Small Molecule Therapeutics for the Treatment of Retinoblastoma. <i>Mini-Reviews in Medicinal Chemistry</i> , 2016, 16, 430-454.	2.4	42
67	Regulation of proliferation, cell fate specification and differentiation by the homeodomain proteins Prox1, Six3, and Chx10 in the developing retina. <i>Cell Cycle</i> , 2003, 2, 350-7.	2.6	41
68	Targeting MDM2 and MDMX in Retinoblastoma. <i>Current Cancer Drug Targets</i> , 2007, 7, 689-695.	1.6	40
69	The Clonal Evolution of Metastatic Osteosarcoma as Shaped by Cisplatin Treatment. <i>Molecular Cancer Research</i> , 2019, 17, 895-906.	3.4	40
70	Neuronal Differentiation and Synaptogenesis in Retinoblastoma. <i>Cancer Research</i> , 2007, 67, 2701-2711.	0.9	38
71	Cross-species genomic and epigenomic landscape of retinoblastoma. <i>Oncotarget</i> , 2013, 4, 844-859.	1.8	37
72	Genetic and Epigenetic Discoveries in Human Retinoblastoma. <i>Critical Reviews in Oncogenesis</i> , 2015, 20, 217-225.	0.4	36

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73	Changes in Retinoblastoma Cell Adhesion Associated with Optic Nerve Invasion. <i>Molecular and Cellular Biology</i> , 2009, 29, 6268-6282.	2.3	35
74	Chromatin remodelers HELLS and UHRF1 mediate the epigenetic deregulation of genes that drive retinoblastoma tumor progression. <i>Oncotarget</i> , 2014, 5, 9594-9608.	1.8	35
75	The chemotherapeutic CX-5461 primarily targets TOP2B and exhibits selective activity in high-risk neuroblastoma. <i>Nature Communications</i> , 2021, 12, 6468.	12.8	35
76	Glutamate regulates retinal progenitors cells proliferation during development. <i>European Journal of Neuroscience</i> , 2006, 24, 969-980.	2.6	34
77	An E2F Binding-Deficient Rb1 Protein Partially Rescues Developmental Defects Associated with Rb1 Nullizygosity. <i>Molecular and Cellular Biology</i> , 2006, 26, 1527-1537.	2.3	34
78	Lessons from Retinoblastoma: Implications for Cancer, Development, Evolution, and Regenerative Medicine. <i>Trends in Molecular Medicine</i> , 2016, 22, 863-876.	6.7	34
79	Regenerative and restorative medicine for eye disease. <i>Nature Medicine</i> , 2022, 28, 1149-1156.	30.7	34
80	Interleukin-15 Enhances Anti-GD2 Antibody-Mediated Cytotoxicity in an Orthotopic PDX Model of Neuroblastoma. <i>Clinical Cancer Research</i> , 2019, 25, 7554-7564.	7.0	33
81	Developmental defects in Rb-deficient retinæ. <i>Vision Research</i> , 2004, 44, 3323-3333.	1.4	31
82	Superselective Intraocular Artery Chemotherapy. <i>JAMA Ophthalmology</i> , 2011, 129, 1490.	2.4	31
83	Brg1 coordinates multiple processes during retinogenesis and is a tumor suppressor in retinoblastoma. <i>Development (Cambridge)</i> , 2015, 142, 4092-4106.	2.5	30
84	Development and characterization of a human orthotopic neuroblastoma xenograft. <i>Developmental Biology</i> , 2015, 407, 344-355.	2.0	30
85	Monitoring Ligand-Induced Protein Ordering in Drug Discovery. <i>Journal of Molecular Biology</i> , 2016, 428, 1290-1303.	4.2	29
86	AAV-mediated Local Delivery of Interferon- $\beta$ for the Treatment of Retinoblastoma in Preclinical Models. <i>NeuroMolecular Medicine</i> , 2009, 11, 43-52.	3.4	28
87	Posttranscriptional gene expression control by NANOS is upregulated and functionally important in pR <sup>-</sup> deficient cells. <i>EMBO Journal</i> , 2014, 33, 2201-2215.	7.8	25
88	Pharmacokinetics and Efficacy of the Spleen Tyrosine Kinase Inhibitor R406 after Ocular Delivery for Retinoblastoma. <i>Pharmaceutical Research</i> , 2014, 31, 3060-3072.	3.5	24
89	Small-Molecule and CRISPR Screening Converge to Reveal Receptor Tyrosine Kinase Dependencies in Pediatric Rhabdoid Tumors. <i>Cell Reports</i> , 2019, 28, 2331-2344.e8.	6.4	24
90	The myogenesis program drives clonal selection and drug resistance in rhabdomyosarcoma. <i>Developmental Cell</i> , 2022, 57, 1226-1240.e8.	7.0	24

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91	Mosaic deletion of Rb arrests rod differentiation and stimulates ectopic synaptogenesis in the mouse retina. <i>Journal of Comparative Neurology</i> , 2006, 498, 112-128.	1.6	23
92	The role of interleukin-2, all-trans retinoic acid, and natural killer cells: surveillance mechanisms in anti-GD2 antibody therapy in neuroblastoma. <i>Cancer Immunology, Immunotherapy</i> , 2018, 67, 615-626.	4.2	23
93	Combinatorial screening using orthotopic patient derived xenograft-expanded early phase cultures of osteosarcoma identify novel therapeutic drug combinations. <i>Cancer Letters</i> , 2019, 442, 262-270.	7.2	23
94	Childhood Cancer and Developmental Biology. <i>Current Topics in Developmental Biology</i> , 2011, 94, 1-13.	2.2	20
95	Next-generation humanized patient-derived xenograft mouse model for pre-clinical antibody studies in neuroblastoma. <i>Cancer Immunology, Immunotherapy</i> , 2021, 70, 721-732.	4.2	17
96	Distinct Developmental Mechanisms Act Independently to Shape Biased Synaptic Divergence from an Inhibitory Neuron. <i>Current Biology</i> , 2020, 30, 1258-1268.e2.	3.9	15
97	A new model of tumor susceptibility following tumor suppressor gene inactivation. <i>Cell Cycle</i> , 2008, 7, 735-740.	2.6	14
98	Preclinical Models Provide Scientific Justification and Translational Relevance for Moving Novel Therapeutics into Clinical Trials for Pediatric Cancer. <i>Cancer Research</i> , 2015, 75, 5176-5186.	0.9	14
99	Longitudinal NK cell kinetics and cytotoxicity in children with neuroblastoma enrolled in a clinical phase II trial. , 2020, 8, e000176.		14
100	Performance of a docking/molecular dynamics protocol for virtual screening of nutlin-class inhibitors of Mdmx. <i>Journal of Molecular Graphics and Modelling</i> , 2017, 74, 54-60.	2.4	12
101	Identification of a modular super-enhancer in murine retinal development. <i>Nature Communications</i> , 2022, 13, 253.	12.8	12
102	Stem Cells Expand Insights into Human Brain Evolution. <i>Cell Stem Cell</i> , 2016, 18, 425-426.	11.1	11
103	Reprogramming of mouse retinal neurons and standardized quantification of their differentiation in 3D retinal cultures. <i>Nature Protocols</i> , 2016, 11, 1955-1976.	12.0	11
104	Perspective: New genetic tools for studying retinal development and disease. <i>Visual Neuroscience</i> , 2005, 22, 553-560.	1.0	10
105	Statistical analysis of data from retroviral clonal experiments in the developing retina. <i>Brain Research</i> , 2008, 1192, 178-185.	2.2	8
106	Methylation profiling reveals novel molecular classes of rhabdomyosarcoma. <i>Scientific Reports</i> , 2021, 11, 22213.	3.3	8
107	Mutations and cancer: one or two historical perspectives?. <i>Lancet Oncology</i> , The, 2009, 10, 834.	10.7	7
108	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

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109	An eye on retinal recovery. <i>Nature</i> , 2016, 540, 350-351.	27.8	5
110	Phase I study of talazoparib and irinotecan in children and young adults with recurrent/refractory solid tumors.. <i>Journal of Clinical Oncology</i> , 2017, 35, 10542-10542.	1.6	5
111	Antigen cross-presentation in young tumor-bearing hosts promotes CD8 <sup>+</sup> T cell terminal differentiation. <i>Science Immunology</i> , 2022, 7, eabf6136.	11.9	5
112	Retinoblastoma (Rb) regulates laminar dendritic arbor reorganization in retinal horizontal neurons. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 21111-21116.	7.1	4
113	The ATRX cDNA is prone to bacterial IS10 element insertions that alter its structure. <i>SpringerPlus</i> , 2014, 3, 222.	1.2	3
114	First pointwise encoding time reduction with radial acquisition (PETRA) implementation for catheter detection in interstitial high-dose-rate (HDR) brachytherapy. <i>Brachytherapy</i> , 2022, 21, 501-510.	0.5	3
115	Automated Tracing of Horizontal Neuron Processes During Retinal Development. <i>Neurochemical Research</i> , 2011, 36, 583-593.	3.3	2
116	Genetically Engineered Mouse and Orthotopic Human Tumor Xenograft Models of Retinoblastoma. <i>Methods in Molecular Biology</i> , 2015, 1267, 307-317.	0.9	2
117	Association of age at diagnosis and stage of disease with <i>ATRX</i> mutations in neuroblastoma.. <i>Journal of Clinical Oncology</i> , 2016, 34, 10525-10525.	1.6	2
118	Targeting retinoblastoma: therapeutic inhibition using catalytic antioxidant cerium oxide nanoparticles. <i>FASEB Journal</i> , 2013, 27, 1088.16.	0.5	2
119	Abstract 1543: Mining cancer-specific isoforms as CAR T-cell therapy targets for pediatric solid and brain tumors. , 2021, , .		1
120	Biology of Retinoblastoma. <i>Pediatric Oncology</i> , 2010, , 1-9.	0.5	1
121	Targeting the cell cycle for cancer therapy in rhabdomyosarcoma.. <i>Journal of Clinical Oncology</i> , 2017, 35, 10535-10535.	1.6	1
122	Comprehensive preclinical testing for neuroblastoma using orthotopic xenografts of a patient tumor.. <i>Journal of Clinical Oncology</i> , 2012, 30, e13584-e13584.	1.6	1
123	SYK expression in metastatic retinoblastoma.. <i>Journal of Clinical Oncology</i> , 2012, 30, e13544-e13544.	1.6	1
124	Special issue on retinal development. <i>Brain Research</i> , 2008, 1192, 1-2.	2.2	0
125	Current and emerging therapy for improving outcomes in patients with intraocular retinoblastoma. <i>Expert Opinion on Orphan Drugs</i> , 2015, 3, 1155-1166.	0.8	0
126	Retinoblastoma Tumorigenesis. , 2019, , 67-77.		0



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127	Abstract 2289: Empowering point-and-click genomic analysis with large pediatric genomic reference data on St. Jude Cloud. , 2021, , .		0
128	Cellular and genetic events in retinoblastoma tumorigenesis. , 2007, , 405-409.		0
129	Incorporating PARP inhibitors in translational studies for Ewing's sarcoma.. Journal of Clinical Oncology, 2014, 32, e21025-e21025.	1.6	0
130	Preclinical evaluation of PARP inhibitors in combination with DNA-damaging agents in a Ewing sarcoma orthotopic xenograft model.. Journal of Clinical Oncology, 2014, 32, 10073-10073.	1.6	0
131	Molecular analysis of solid tumors (MAST): A protocol for comprehensive preclinical evaluation of pediatric solid tumors.. Journal of Clinical Oncology, 2014, 32, 10036-10036.	1.6	0
132	Retinoblastoma Tumorigenesis. , 2015, , 61-68.		0
133	The Childhood Solid Tumor Network: A St. Jude Initiative.. Journal of Clinical Oncology, 2015, 33, e21022-e21022.	1.6	0
134	Loss of STAG2 expression and prognosis in Ewing sarcoma family of tumors.. Journal of Clinical Oncology, 2015, 33, 10024-10024.	1.6	0
135	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. Journal of the Endocrine Society, 2019, 3, .	0.2	0