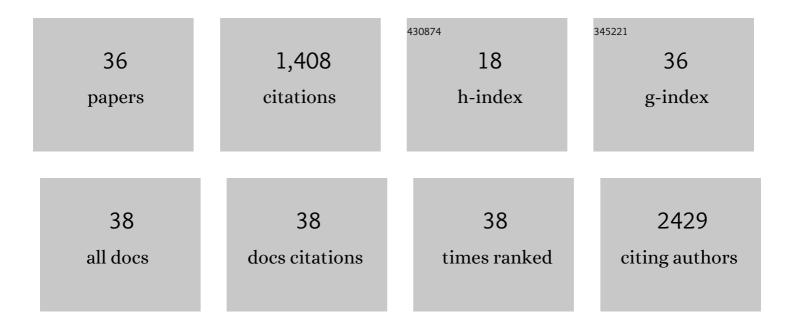
Corinne M Linardic

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	PAX3–FOXO1 fusion gene in rhabdomyosarcoma. Cancer Letters, 2008, 270, 10-18.	7.2	121
2	InÂVivo Imaging of Tumor-Propagating Cells, Regional Tumor Heterogeneity, and Dynamic Cell Movements in Embryonal Rhabdomyosarcoma. Cancer Cell, 2012, 21, 680-693.	16.8	110
3	Genomic Classification and Clinical Outcome in Rhabdomyosarcoma: A Report From an International Consortium. Journal of Clinical Oncology, 2021, 39, 2859-2871.	1.6	101
4	Alveolar rhabdomyosarcoma–associated PAX3-FOXO1 promotes tumorigenesis via Hippo pathway suppression. Journal of Clinical Investigation, 2014, 124, 285-296.	8.2	94
5	Modulation of cell growth and differentiation by ceramide. FEBS Letters, 1992, 307, 211-214.	2.8	92
6	Genetic Modeling of Human Rhabdomyosarcoma. Cancer Research, 2005, 65, 4490-4495.	0.9	79
7	FGFR4 Blockade Exerts Distinct Antitumorigenic Effects in Human Embryonal versus Alveolar Rhabdomyosarcoma. Clinical Cancer Research, 2012, 18, 3780-3790.	7.0	76
8	Defining the Cooperative Genetic Changes That Temporally Drive Alveolar Rhabdomyosarcoma. Cancer Research, 2008, 68, 9583-9588.	0.9	71
9	A Review: Molecular Aberrations within Hippo Signaling in Bone and Soft-Tissue Sarcomas. Frontiers in Oncology, 2015, 5, 190.	2.8	60
10	The <i>PAX3-FKHR</i> Fusion Gene of Rhabdomyosarcoma Cooperates with Loss of p16INK4A to Promote Bypass of Cellular Senescence. Cancer Research, 2007, 67, 6691-6699.	0.9	57
11	Insights into pediatric rhabdomyosarcoma research: Challenges and goals. Pediatric Blood and Cancer, 2019, 66, e27869.	1.5	57
12	Histone Deacetylase Inhibitors Inhibit Rhabdomyosarcoma by Reactive Oxygen Species–Dependent Targeting of Specificity Protein Transcription Factors. Molecular Cancer Therapeutics, 2015, 14, 2143-2153.	4.1	53
13	The NOTCH1/SNAIL1/MEF2C Pathway Regulates Growth and Self-Renewal in Embryonal Rhabdomyosarcoma. Cell Reports, 2017, 19, 2304-2318.	6.4	53
14	Inhibition of the Notch-Hey1 Axis Blocks Embryonal Rhabdomyosarcoma Tumorigenesis. Clinical Cancer Research, 2011, 17, 7324-7336.	7.0	51
15	A Novel Notch–YAP Circuit Drives Stemness and Tumorigenesis in Embryonal Rhabdomyosarcoma. Molecular Cancer Research, 2017, 15, 1777-1791.	3.4	49
16	Role of the YAP Oncoprotein in Priming Ras-Driven Rhabdomyosarcoma. PLoS ONE, 2015, 10, e0140781.	2.5	39
17	Pathology of childhood rhabdomyosarcoma: A consensus opinion document from the Children's Oncology Group, European Paediatric Soft Tissue Sarcoma Study Group, and the Cooperative Weichteilsarkom Studiengruppe. Pediatric Blood and Cancer, 2021, 68, e28798.	1.5	38
18	Soft Tissue Sarcoma Cancer Stem Cells: An Overview. Frontiers in Oncology, 2018, 8, 475.	2.8	37

#	Article	IF	CITATIONS
19	Molecular testing of rhabdomyosarcoma in clinical trials to improve risk stratification and outcome: A consensus view from European paediatric Soft tissue sarcoma Study Group, Children's Oncology Group and Cooperative Weichteilsarkom-Studiengruppe. European Journal of Cancer, 2022, 172, 367-386.	2.8	19
20	RAS and ROS in Rhabdomyosarcoma. Cancer Cell, 2013, 24, 689-691.	16.8	18
21	Secreted Frizzled-Related Protein 3 (SFRP3) Is Required for Tumorigenesis of PAX3–FOXO1-Positive Alveolar Rhabdomyosarcoma. Clinical Cancer Research, 2015, 21, 4868-4880.	7.0	18
22	Clinical group and modified TNM stage for rhabdomyosarcoma: A review from the Children's Oncology Group. Pediatric Blood and Cancer, 2022, 69, e29644.	1.5	18
23	Loss of MST/Hippo Signaling in a Genetically Engineered Mouse Model of Fusion-Positive Rhabdomyosarcoma Accelerates Tumorigenesis. Cancer Research, 2018, 78, 5513-5520.	0.9	12
24	Epigenetic regulator BMI1 promotes alveolar rhabdomyosarcoma proliferation and constitutes a novel therapeutic target. Molecular Oncology, 2021, 15, 2156-2171.	4.6	11
25	Prioritization of Novel Agents for Patients with Rhabdomyosarcoma: A Report from the Children's Oncology Group (COG) New Agents for Rhabdomyosarcoma Task Force. Journal of Clinical Medicine, 2021, 10, 1416.	2.4	11
26	Relationship of DNA methylation to mutational changes and transcriptional organization in fusionâ€positive and fusionâ€negative rhabdomyosarcoma. International Journal of Cancer, 2019, 144, 2707-2717.	5.1	10
27	Detection of iron deficiency in children with Down syndrome. Genetics in Medicine, 2020, 22, 317-325.	2.4	10
28	Parenting a child with cancer: a couple-based approach. Translational Behavioral Medicine, 2019, 9, 504-513.	2.4	6
29	Recent Insights into Notch Signaling in Embryonal Rhabdomyosarcoma. Current Drug Targets, 2016, 17, 1235-1244.	2.1	6
30	Asparaginase-Induced Hypertriglyceridemia Presenting as Pseudohyponatremia during Leukemia Treatment. Case Reports in Pediatrics, 2014, 2014, 1-5.	0.4	4
31	RASSF4 is required for skeletal muscle differentiation. Cell Biology International, 2020, 44, 381-390.	3.0	4
32	Myelodysplasia as masquerader: A woman with hypereosinophilic syndrome and twelve years of ?chronic ITP?. Medical and Pediatric Oncology, 2002, 39, 137-138.	1.0	2
33	A method to culture human alveolar rhabdomyosarcoma cell lines as rhabdospheres demonstrates an enrichment in stemness and notch signaling. Biology Open, 2021, 10, .	1.2	2
34	Expression of oncogenic HRAS in human Rh28 and RMS-YM rhabdomyosarcoma cells leads to oncogene-induced senescence. Scientific Reports, 2021, 11, 16505.	3.3	1
35	Response to Zhang et al Genetics in Medicine, 2020, 22, 662-662.	2.4	0
36	Abstract LB213: Potent antitumor activity of a FGFR4 CAR-T in rhabdomyosarcoma. Cancer Research, 2022, 82, LB213-LB213.	0.9	0