Josep Roma

List of Publications by Year in descending order

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LOSED ROMA

#	Article	IF	CITATIONS
1	Urokinase-dependent plasminogen activation is required for efficient skeletal muscle regeneration in vivo. Blood, 2001, 97, 1703-1711.	1.4	114
2	uPA deficiency exacerbates muscular dystrophy in <i>MDX</i> mice. Journal of Cell Biology, 2007, 178, 1039-1051.	5.2	66
3	BRG1/SMARCA4 is essential for neuroblastoma cell viability through modulation of cell death and survival pathways. Oncogene, 2016, 35, 5179-5190.	5.9	65
4	microRNAs as pharmacological targets in cancer. Pharmacological Research, 2013, 75, 3-14.	7.1	56
5	Notch Pathway Inhibition Significantly Reduces Rhabdomyosarcoma Invasiveness and Mobility <i>In Vitro</i> . Clinical Cancer Research, 2011, 17, 505-513.	7.0	49
6	Clonal dynamics in osteosarcoma defined by RGB marking. Nature Communications, 2018, 9, 3994.	12.8	40
7	Detection of bone marrow micrometastasis and microcirculating disease in rhabdomyosarcoma by a real-time RT-PCR assay. Journal of Cancer Research and Clinical Oncology, 2006, 132, 356-362.	2.5	36
8	<i>Notch, Wnt,</i> and <i>Hedgehog</i> Pathways in Rhabdomyosarcoma: From Single Pathways to an Integrated Network. Sarcoma, 2012, 2012, 1-7.	1.3	34
9	MicroRNA-497 impairs the growth of chemoresistant neuroblastoma cells by targeting cell cycle, survival and vascular permeability genes. Oncotarget, 2016, 7, 9271-9287.	1.8	31
10	Longitudinal pathologic study of the gastrocnemius muscle group in mdx mice. Acta Neuropathologica, 2004, 107, 27-34.	7.7	29
11	DNA methylation profiling identifies PTRF/Cavin-1 as a novel tumor suppressor in Ewing sarcoma when co-expressed with caveolin-1. Cancer Letters, 2017, 386, 196-207.	7.2	25
12	Evolution of pathological changes in the gastrocnemius of the mdx mice correlate with utrophin and ?-dystroglycan expression. Acta Neuropathologica, 2004, 108, 443-452.	7.7	23
13	Notch-mediated induction of N-cadherin and α9-integrin confers higher invasive phenotype on rhabdomyosarcoma cells. British Journal of Cancer, 2012, 107, 1374-1383.	6.4	23
14	Origins of DNA methylation defects in Wilms tumors. Cancer Letters, 2019, 457, 119-128.	7.2	23
15	Sequential combinations of chemotherapeutic agents with BH3 mimetics to treat rhabdomyosarcoma and avoid resistance. Cell Death and Disease, 2020, 11, 634.	6.3	17
16	Patient-derived xenografts for childhood solid tumors: a valuable tool to test new drugs and personalize treatments. Clinical and Translational Oncology, 2017, 19, 44-50.	2.4	15
17	Engineering pH‣ensitive Stable Nanovesicles for Delivery of MicroRNA Therapeutics. Small, 2022, 18, e2101959.	10.0	13
18	Hedgehog Pathway Inhibition Hampers Sphere and Holoclone Formation in Rhabdomyosarcoma. Stem Cells International, 2017, 2017, 1-14.	2.5	10

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19	CN133, a Novel Brain-Penetrating Histone Deacetylase Inhibitor, Hampers Tumor Growth in Patient-Derived Pediatric Posterior Fossa Ependymoma Models. Cancers, 2020, 12, 1922.	3.7	7
20	The antitumour drug ABTL0812 impairs neuroblastoma growth through endoplasmic reticulum stress-mediated autophagy and apoptosis. Cell Death and Disease, 2020, 11, 773.	6.3	7
21	Optimization of rhabdomyosarcoma disseminated disease assessment by flow cytometry. Cytometry Part A: the Journal of the International Society for Analytical Cytology, 2014, 85, 1020-1029.	1.5	4
22	MEK and MCL-1 sequential inhibition synergize to enhance rhabdomyosarcoma treatment. Cell Death Discovery, 2022, 8, 172.	4.7	4
23	DAX-1 Expression in Pediatric Rhabdomyosarcomas: Another Immunohistochemical Marker Useful in the Diagnosis of Translocation Positive Alveolar Rhabdomyosarcoma. PLoS ONE, 2015, 10, e0133019.	2.5	0