

Adriana S Beltran

List of Publications by Year in descending order

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Version: 2024-02-01

19
papers

1,182
citations

567281

15
h-index

839539

18
g-index

20
all docs

20
docs citations

20
times ranked

2232
citing authors

#	ARTICLE	IF	CITATIONS
1	Abstract P3-15-01: Patients and Researchers Together (PART); a patient-centered tumor tissue collection PARTnership between patients and researchers to increase tissue donations for breast cancer research. <i>Cancer Research</i> , 2022, 82, P3-15-01-P3-15-01.	0.9	0
2	Generation of an induced pluripotent stem cell line (UNCCI002-A) from a healthy donor using a non-integration system to study Cerebral Cavernous Malformation (CCM). <i>Stem Cell Research</i> , 2021, 54, 102421.	0.7	11
3	Pathogenic SPTBN1 variants cause an autosomal dominant neurodevelopmental syndrome. <i>Nature Genetics</i> , 2021, 53, 1006-1021.	21.4	44
4	Generation of an integration-free induced pluripotent stem cell line (UNC001-A) from blood of a healthy individual. <i>Stem Cell Research</i> , 2020, 49, 102015.	0.7	10
5	GSK2801, a BAZ2/BRD9 Bromodomain Inhibitor, Synergizes with BET Inhibitors to Induce Apoptosis in Triple-Negative Breast Cancer. <i>Molecular Cancer Research</i> , 2019, 17, 1503-1518.	3.4	39
6	Inheritance of OCT4 predetermines fate choice in human embryonic stem cells. <i>Molecular Systems Biology</i> , 2018, 14, e8140.	7.2	27
7	Enhancer Remodeling during Adaptive Bypass to MEK Inhibition Is Attenuated by Pharmacologic Targeting of the P-TEFb Complex. <i>Cancer Discovery</i> , 2017, 7, 302-321.	9.4	128
8	Inhibition of Lapatinib-Induced Kinome Reprogramming in ERBB2-Positive Breast Cancer by Targeting BET Family Bromodomains. <i>Cell Reports</i> , 2015, 11, 390-404.	6.4	254
9	Breaking through an epigenetic wall. <i>Epigenetics</i> , 2013, 8, 164-176.	2.7	20
10	Expression of the Pluripotency Transcription Factor OCT4 in the Normal and Aberrant Mammary Gland. <i>Frontiers in Oncology</i> , 2013, 3, 79.	2.8	28
11	Targeting Serous Epithelial Ovarian Cancer with Designer Zinc Finger Transcription Factors. <i>Journal of Biological Chemistry</i> , 2012, 287, 29873-29886.	3.4	38
12	Targeted silencing of the oncogenic transcription factor SOX2 in breast cancer. <i>Nucleic Acids Research</i> , 2012, 40, 6725-6740.	14.5	138
13	Epigenetic reprogramming of cancer cells via targeted DNA methylation. <i>Epigenetics</i> , 2012, 7, 350-360.	2.7	189
14	Generation of tumor-initiating cells by exogenous delivery of OCT4 transcription factor. <i>Breast Cancer Research</i> , 2011, 13, R94.	5.0	81
15	Suppression of Breast Tumor Growth and Metastasis by an Engineered Transcription Factor. <i>PLoS ONE</i> , 2011, 6, e24595.	2.5	45
16	Reactivation of <i>MASPIN</i> in non-small cell lung carcinoma (NSCLC) cells by artificial transcription factors (ATFs). <i>Epigenetics</i> , 2011, 6, 224-235.	2.7	42
17	Remodeling Genomes with Artificial Transcription Factors (ATFs). <i>Methods in Molecular Biology</i> , 2010, 649, 163-182.	0.9	9
18	Rational Design, Selection and Specificity of Artificial Transcription Factors (ATFs): The Influence of Chromatin in Target Gene Regulation. <i>Combinatorial Chemistry and High Throughput Screening</i> , 2008, 11, 146-158.	1.1	17

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19	Reprogramming epigenetic silencing: artificial transcription factors synergize with chromatin remodeling drugs to reactivate the tumor suppressor <i>p16</i> in mammary serine protease inhibitor. <i>Molecular Cancer Therapeutics</i> , 2008, 7, 1080-1090.	4.1	58