

Angela Bentivegna

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1214168/publications.pdf>

Version: 2024-02-01

50
papers

1,342
citations

394421

19
h-index

345221

36
g-index

51
all docs

51
docs citations

51
times ranked

2627
citing authors

#	ARTICLE	IF	CITATIONS
1	MV1035 Overcomes Temozolomide Resistance in Patient-Derived Glioblastoma Stem Cell Lines. <i>Biology</i> , 2022, 11, 70.	2.8	5
2	Characterization of Chromosomal Breakpoints in 12 Cases with 8p Rearrangements Defines a Continuum of Fragility of the Region. <i>International Journal of Molecular Sciences</i> , 2022, 23, 3347.	4.1	6
3	Analysis of copy number alterations in bladder cancer stem cells revealed a prognostic role of LRP1B. <i>World Journal of Urology</i> , 2022, 40, 2267-2273.	2.2	5
4	<i>UGT1A1</i> mutations and psychoses: towards understanding the relationship with unconjugated bilirubin. <i>CNS Spectrums</i> , 2021, 26, 188-190.	1.2	2
5	Genomic and Epigenomic Profile of Uterine Smooth Muscle Tumors of Uncertain Malignant Potential (STUMPs) Revealed Similarities and Differences with Leiomyomas and Leiomyosarcomas. <i>International Journal of Molecular Sciences</i> , 2021, 22, 1580.	4.1	10
6	Human Chromosome 18 and Acrocentrics: A Dangerous Liaison. <i>International Journal of Molecular Sciences</i> , 2021, 22, 5637.	4.1	0
7	Characterizing the Genomic Profile in High-Grade Gliomas: From Tumor Core to Peritumoral Brain Zone, Passing through Glioma-Derived Tumorspheres. <i>Biology</i> , 2021, 10, 1157.	2.8	9
8	Notch Signaling and MicroRNA: The Dynamic Duo Steering Between Neurogenesis and Glioblastomas. <i>Cellular and Molecular Biology</i> , 2021, 67, 33-43.	0.9	4
9	Instability of Short Arm of Acrocentric Chromosomes: Lesson from Non-Acrocentric Satellited Chromosomes. Report of 24 Unrelated Cases. <i>International Journal of Molecular Sciences</i> , 2020, 21, 3431.	4.1	3
10	Unbalanced X;Autosome Translocations May Lead to Mild Phenotypes and Are Associated with Autoimmune Diseases. <i>Cytogenetic and Genome Research</i> , 2020, 160, 80-84.	1.1	4
11	A Ploidy Increase Promotes Sensitivity of Glioma Stem Cells to Aurora Kinases Inhibition. <i>Journal of Oncology</i> , 2019, 2019, 1-15.	1.3	1
12	Role of Notch Signaling Pathway in Glioblastoma Multiforme Pathogenesis. <i>Cancers</i> , 2019, 11, 292.	3.7	113
13	Refining the Phenotype of Recurrent Rearrangements of Chromosome 16. <i>International Journal of Molecular Sciences</i> , 2019, 20, 1095.	4.1	34
14	Specific Expression of a New Bruton Tyrosine Kinase Isoform (p65BTK) in the Glioblastoma Gemistocytic Histotype. <i>Frontiers in Molecular Neuroscience</i> , 2019, 12, 2.	2.9	16
15	Familial glioma. <i>Atlas of Genetics and Cytogenetics in Oncology and Haematology</i> , 2019, , .	0.1	0
16	Analysis of Chromosomal Alterations in Urothelial Carcinoma. <i>Methods in Molecular Biology</i> , 2018, 1655, 3-17.	0.9	2
17	Valproic Acid Inhibits Proliferation and Reduces Invasiveness in Glioma Stem Cells Through Wnt/ β 2 Catenin Signalling Activation. <i>Genes</i> , 2018, 9, 522.	2.4	24
18	Epigenetic and transcriptional modulation of WDR5, a chromatin remodeling protein, in Huntington's disease human induced pluripotent stem cell (hiPSC) model. <i>Molecular and Cellular Neurosciences</i> , 2017, 82, 46-57.	2.2	8

#	ARTICLE	IF	CITATIONS
19	Thyrospheres from B-CPAP Cell Line with <i>BRAF</i> and <i>TERT</i> Promoter Mutations have Different Functional and Molecular Features than Parental Cells. <i>Journal of Cancer</i> , 2017, 8, 1629-1639.	2.5	9
20	Resveratrol Impairs Glioma Stem Cells Proliferation and Motility by Modulating the Wnt Signaling Pathway. <i>PLoS ONE</i> , 2017, 12, e0169854.	2.5	103
21	Pioglitazone Effect on Glioma Stem Cell Lines: Really a Promising Drug Therapy for Glioblastoma?. <i>PPAR Research</i> , 2016, 2016, 1-8.	2.4	10
22	The Effect of Culture on Human Bone Marrow Mesenchymal Stem Cells: Focus on DNA Methylation Profiles. <i>Stem Cells International</i> , 2016, 2016, 1-12.	2.5	18
23	Using Copy Number Alterations to Identify New Therapeutic Targets for Bladder Carcinoma. <i>International Journal of Molecular Sciences</i> , 2016, 17, 271.	4.1	9
24	Epigenetic targeting of glioma stem cells: Short-term and long-term treatments with valproic acid modulate DNA methylation and differentiation behavior, but not temozolomide sensitivity. <i>Oncology Reports</i> , 2016, 35, 2811-2824.	2.6	22
25	Chromosomal imbalances in human bladder urothelial carcinoma: similarities and differences between biopsy samples and cancer stem-like cells. <i>BMC Cancer</i> , 2014, 14, 646.	2.6	6
26	In vitro anticancer drug test: A new method emerges from the model of glioma stem cells. <i>Toxicology Reports</i> , 2014, 1, 188-199.	3.3	21
27	Synchrotron-based photon activation therapy effect on cisplatin pre-treated human glioma stem cells. <i>Anticancer Research</i> , 2014, 34, 5351-5.	1.1	7
28	DNA Methylation Changes during In Vitro Propagation of Human Mesenchymal Stem Cells: Implications for Their Genomic Stability?. <i>Stem Cells International</i> , 2013, 2013, 1-9.	2.5	45
29	Delineating the Cytogenomic and Epigenomic Landscapes of Glioma Stem Cell Lines. <i>PLoS ONE</i> , 2013, 8, e57462.	2.5	31
30	Investigating the role of X chromosome breakpoints in premature ovarian failure. <i>Molecular Cytogenetics</i> , 2012, 5, 32.	0.9	33
31	DNA copy number alterations and PPARG amplification in a patient with multifocal bladder urothelial carcinoma. <i>BMC Research Notes</i> , 2012, 5, 607.	1.4	5
32	From cytogenomic to epigenomic profiles: monitoring the biologic behavior of in vitro cultured human bone marrow mesenchymal stem cells. <i>Stem Cell Research and Therapy</i> , 2012, 3, 47.	5.5	93
33	A de novo supernumerary genomic discontinuous ring chromosome 21 in a child with mild intellectual disability. <i>American Journal of Medical Genetics, Part A</i> , 2011, 155, 1425-1431.	1.2	1
34	Cytogenetics of Premature Ovarian Failure: An Investigation on 269 Affected Women. <i>Journal of Biomedicine and Biotechnology</i> , 2011, 2011, 1-9.	3.0	42
35	Chromosomal Aberrations in Bladder Cancer: Fresh versus Formalin Fixed Paraffin Embedded Tissue and Targeted FISH versus Wide Microarray-Based CGH Analysis. <i>PLoS ONE</i> , 2011, 6, e24237.	2.5	21
36	UroVysion Multiprobe FISH on transitional cell carcinoma of the urinary bladder: comparative analysis on fresh isolated interphasic nuclei and paraffin-embedded tissue. <i>Cancer Genetics and Cytogenetics</i> , 2010, 203, 79.	1.0	0

#	ARTICLE	IF	CITATIONS
37	Biological heterogeneity of putative bladder cancer stem-like cell populations from human bladder transitional cell carcinoma samples. <i>Cancer Science</i> , 2010, 101, 416-424.	3.9	60
38	Monitoring the genomic stability of in vitro cultured rat bone-marrow-derived mesenchymal stem cells. <i>Chromosome Research</i> , 2009, 17, 1025-39.	2.2	76
39	Distinct pools of cancer stem-like cells coexist within human glioblastomas and display different tumorigenicity and independent genomic evolution. <i>Oncogene</i> , 2009, 28, 1807-1811.	5.9	177
40	Prenatal/neonatal pathology in two cases of Cornelia de Lange syndrome harboring novel mutations of NIPBL. <i>Genetics in Medicine</i> , 2007, 9, 188-194.	2.4	18
41	High frequency of mosaic CREBBP deletions in Rubinstein-Taybi syndrome patients and mapping of somatic and germ-line breakpoints. <i>Genomics</i> , 2007, 90, 567-573.	2.9	42
42	Clinical score of 62 Italian patients with Cornelia de Lange syndrome and correlations with the presence and type of NIPBL mutation. <i>Clinical Genetics</i> , 2007, 72, 98-108.	2.0	93
43	Rubinstein-Taybi Syndrome: spectrum of CREBBP mutations in Italian patients. <i>BMC Medical Genetics</i> , 2006, 7, 77.	2.1	60
44	Evidence by Expression Analysis of Candidate Genes for Congenital Heart Defects in the NF1 Microdeletion Interval. <i>Annals of Human Genetics</i> , 2005, 69, 508-516.	0.8	22
45	Evidence for non-homologous end joining and non-allelic homologous recombination in atypical NF1 microdeletions. <i>Human Genetics</i> , 2004, 115, 69-80.	3.8	41
46	Tandem duplication of the NF1 gene detected by high-resolution FISH in the 17q11.2 region. <i>Human Genetics</i> , 2002, 110, 314-321.	3.8	17
47	FISH with locus-specific probes on stretched chromosomes: a useful tool for genome organization studies. <i>Chromosome Research</i> , 2001, 9, 167-170.	2.2	5
48	Title is missing!. <i>Chromosome Research</i> , 2001, 9, 520-520.	2.2	0
49	Identification of duplicated genes in 17q11.2 using FISH on stretched chromosomes and DNA fibers. <i>Human Genetics</i> , 2001, 109, 48-54.	3.8	6
50	Mapping of genes and ESTs assigned to 17q11.2 to a YAC contig centred on the NF1 gene. <i>GeneScreen</i> , 2000, 1, 21-27.	0.6	3