Brigitte Wolf

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8196174/publications.pdf

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		1170033	1427216	
12	242	9	11	
papers	citations	h-index	g-index	
12	12	12	595	
all docs	docs citations	times ranked	citing authors	

#	Article	IF	CITATIONS
1	Long interspersed element-1 ribonucleoprotein particles protect telomeric ends in alternative lengthening of telomeres dependent cells. Neoplasia, 2020, 22, 61-75.	2.3	13
2	Short-course radiotherapy promotes pro-inflammatory macrophages via extracellular vesicles in human rectal cancer., 2020, 8, e000667.		24
3	•ÂPancho trial (p53-adapted neoadjuvant chemotherapy for resectable esophageal cancer) completed—mutation rate of the marker higher than expected. European Surgery - Acta Chirurgica Austriaca, 2018, 50, 160-166.	0.3	5
4	DNA damage predicts prognosis and treatment response in colorectal liver metastases superior to immunogenic cell death and T cells. Theranostics, 2018, 8, 3198-3213.	4.6	18
5	Careful neuropsychological testing reveals a novel genetic marker, <i>GSTO1*C </i> , linked to the pre-stage of Alzheimer's disease. Oncotarget, 2016, 7, 39108-39117.	0.8	10
6	Potential of DNA methylation in rectal cancer as diagnostic and prognostic biomarkers. British Journal of Cancer, 2015, 113, 1035-1045.	2.9	25
7	TP53 Mutational Status and Prediction of Benefit from Adjuvant 5-Fluorouracil in Stage III Colon Cancer Patients. EBioMedicine, 2015, 2, 825-830.	2.7	37
8	The biomarker TP53 divides patients with neoadjuvantly treated esophageal cancer into 2 subgroups with markedly different outcomes. A p53 Research Group study. Journal of Thoracic and Cardiovascular Surgery, 2014, 148, 2280-2286.	0.4	48
9	Adaptive mutation in nuclear export protein allows stable transgene expression in a chimaeric influenza A virus vector. Journal of General Virology, 2014, 95, 337-349.	1.3	11
10	Multiplex SNaPshot Genotyping for Detecting Loss of Heterozygosity in the Mismatch-Repair Genes MLH1 and MSH2 in Microsatellite-Unstable Tumors. Clinical Chemistry, 2008, 54, 1844-1854.	1.5	18
11	Efficiency of the revised Bethesda guidelines (2003) for the detection of mutations in mismatch repair genes in Austrian HNPCC patients. International Journal of Cancer, 2006, 118, 1465-1470.	2.3	22
12	Spectrum of germ-line MLH1 and MSH2 mutations in Austrian patients with hereditary nonpolyposis colorectal cancer. Wiener Klinische Wochenschrift, 2005, 117, 269-277.	1.0	11