Peng Chen

List of Publications by Year in descending order

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58	974	19	27
papers	citations	h-index	g-index
65	65	65	1137 citing authors
all docs	docs citations	times ranked	

#	Article	IF	CITATIONS
1	Anlotinib Induces a T Cell–Inflamed Tumor Microenvironment by Facilitating Vessel Normalization and Enhances the Efficacy of PD-1 Checkpoint Blockade in Neuroblastoma. Clinical Cancer Research, 2022, 28, 793-809.	7.0	43
2	Role of lncRNA FAM83H antisense RNA1 (FAM83H-AS1) in the progression of non-small cell lung cancer by regulating the miR-545-3p/heparan sulfate 6-O-sulfotransferase (HS6ST2) axis. Bioengineered, 2022, 13, 6476-6489.	3.2	4
3	Co-treatment with miR-21-5p inhibitor and Aurora kinase inhibitor reversine suppresses breast cancer progression by targeting sprouty RTK signaling antagonist 2. Bioengineered, 2022, 13, 455-468.	3.2	7
4	Inhibitors of PD-1 in Non-Small Cell Lung Cancer: A Meta-Analysis of Clinical and Molecular Features. Frontiers in Immunology, 2022, 13, 875093.	4.8	10
5	Effect of antibiotic use on the efficacy of nivolumab in the treatment of advanced/metastatic non-small cell lung cancer: A meta-analysis. Open Medicine (Poland), 2021, 16, 728-736.	1.3	6
6	PNO1 regulates autophagy and apoptosis of hepatocellular carcinoma via the MAPK signaling pathway. Cell Death and Disease, 2021, 12, 552.	6.3	28
7	A Pan-Cancer Analysis of the Oncogenic Role of Twinfilin Actin Binding Protein 1 in Human Tumors. Frontiers in Oncology, 2021, 11, 692136.	2.8	13
8	The incidence of gastrointestinal adverse events in patients with advanced non-small cell lung cancer (NSCLC) treated with PD-1 inhibitors: a meta-analysis. Translational Cancer Research, 2021, 10, 3389-3403.	1.0	1
9	The PSMD14 inhibitor Thiolutin as a novel therapeutic approach for esophageal squamous cell carcinoma through facilitating SNAIL degradation. Theranostics, 2021, 11, 5847-5862.	10.0	31
10	Durable efficacy of anlotinib in a patient with advanced thymic squamous cell carcinoma after multiline chemotherapy and apatinib: A case report and literature review. Thoracic Cancer, 2020, 11, 3383-3387.	1.9	8
11	Comparison of tumor mutational burden across eight types of human cancer Journal of Clinical Oncology, 2020, 38, e15170-e15170.	1.6	2
12	Identifying novel microhaplotypes for ancestry inference. International Journal of Legal Medicine, 2019, 133, 983-988.	2.2	47
13	The advances in DNA mixture interpretation. Forensic Science International, 2019, 301, 101-106.	2.2	18
14	Distribution of ALK Fusion Variants and Correlation with Clinical Outcomes in Chinese Patients with Non-Small Cell Lung Cancer Treated with Crizotinib. Targeted Oncology, 2019, 14, 159-168.	3.6	33
15	Evaluation of the microhaplotype markers in kinship analysis. Electrophoresis, 2019, 40, 1091-1095.	2.4	22
16	Forensic parameters and mutation analysis of 23 short tandem repeat (PowerPlex® Fusion System) loci in Fujian Han Chinese population. Legal Medicine, 2019, 37, 33-36.	1.3	9
17	Prevalence and clinical significance of pathogenic germline BRCA1/2 mutations in Chinese non-small cell lung cancer patients. Cancer Biology and Medicine, 2019, 16, 556-564.	3.0	36
18	A patient with classic biphasic pulmonary blastoma harboring CD74–ROS1 fusion responds to crizotinib. OncoTargets and Therapy, 2018, Volume 11, 157-161.	2.0	9

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19	The genetic diversity and applicability assessment of autosomal STRs among Chinese populations by a novel Fixation Index and Nei's index. Legal Medicine, 2018, 31, 49-58.	1.3	2
20	Association of genetic variations in RTN4 3′-UTR with risk for clear cell renal cell carcinoma. Familial Cancer, 2018, 17, 129-134.	1.9	5
21	Role of Recursive Partitioning Analysis and Graded Prognostic Assessment on Identifying Non-Small Cell Lung Cancer Patients with Brain Metastases Who May Benefit from Postradiation Systemic Therapy. Chinese Medical Journal, 2018, 131, 1206-1213.	2.3	2
22	Association between a Single Nucleotide Polymorphism in the 3 <mml:math id="M1" xmlns:mml="http://www.w3.org/1998/Math/MathML"><mml:msup><mml:mrow></mml:mrow><mml:mrow><mml:mo>′</mml:mo></mml:mrow></mml:msup></mml:math> -UTR of <i>ARHGEF18</i> and the Risk of Nonidiopathic Pulmonary Arterial Hypertension in Chinese Population. Disease Markers, 2018, 2018, 1-6.	1.3	7
23	Genome-wide association study identifies loci and candidate genes for non-idiopathic pulmonary hypertensionÂin Eastern Chinese Han population. BMC Pulmonary Medicine, 2018, 18, 158.	2.0	12
24	Evaluation of the Microhaplotypes panel for DNA mixture analyses. Forensic Science International: Genetics, 2018, 35, 149-155.	3.1	64
25	CD31-labeled circulating endothelial cells as predictor in anlotinib-treated non-small-cell lung cancer: Analysis on ALTER-0303 study. Cancer Medicine, 2018, 7, 3011-3021.	2.8	29
26	Genetic Association of Interleukin-31 Gene Polymorphisms with Epithelial Ovarian Cancer in Chinese Population. Disease Markers, 2018, 2018, 1-7.	1.3	8
27	Forensic efficiency and genetic divergence of 30 autosomal InDels in Chinese Han population from Jiangsu province. Forensic Science International: Genetics, 2018, 37, e17-e19.	3.1	17
28	Association between polymorphisms in IL21 gene and risk for sepsis. Biomarkers, 2017, 22, 14-18.	1.9	2
29	Microhaplotype identified and performed in genetic investigation using PCR-SSCP. Forensic Science International: Genetics, 2017, 28, e1-e7.	3.1	15
30	Association between <i>IL-27</i> gene polymorphisms and risk of papillary thyroid carcinoma. Biomarkers in Medicine, 2017, 11, 141-149.	1.4	11
31	Associations of TIM-1 Genetic Polymorphisms with Asthma: A Meta-analysis. Lung, 2017, 195, 353-360.	3.3	5
32	Genotyping microhaplotype markers through massively parallel sequencing. Forensic Science International: Genetics Supplement Series, 2017, 6, e314-e316.	0.3	6
33	Microhaplotype: Ability of personal identification and being ancestry informative marker. Forensic Science International: Genetics Supplement Series, 2017, 6, e442-e444.	0.3	5
34	Estimate the heterozygote balance of microhaplotype marker with massively parallel sequencing. Forensic Science International: Genetics Supplement Series, 2017, 6, e375-e376.	0.3	6
35	Genetic association of polymorphisms in $\langle i \rangle AXIN1 \langle j \rangle$ gene with clear cell renal cell carcinoma in a Chinese population. Biomarkers in Medicine, 2017, 11, 947-955.	1.4	6
36	Level of activated circulating endothelial cells to predict progression-free survival of Anlotinib treatment in patients with NSCLC: Analysis on ALTER-0303 study Journal of Clinical Oncology, 2017, 35, e20530-e20530.	1.6	1

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37	The chemotherapeutic drug boanmycin induces cell senescence and senescence-associated secretory phenotype factors, thus acquiring the potential to remodel the tumor microenvironment. Anti-Cancer Drugs, 2016, 27, 84-88.	1.4	3
38	Insertion/deletion polymorphism in <i>IL1A</i> 3′â€UTR is associated with susceptibility to endometrial cancer in Chinese Han women. Journal of Obstetrics and Gynaecology Research, 2016, 42, 983-989.	1.3	11
39	Fatal liver cyst rupture in polycystic liver disease complicated with autosomal dominant polycystic kidney disease: A case report. Forensic Science International, 2016, 262, e5-e8.	2.2	6
40	Combined analysis of pri-miR-34b/c rs4938723 and TP53 Arg72Pro with cervical cancer risk. Tumor Biology, 2016, 37, 6267-6273.	1.8	26
41	Association between polymorphisms in IL-27 gene and pre-eclampsia. Placenta, 2016, 37, 61-64.	1.5	16
42	A polymorphism rs4705341 in the flanking region of miR-143/145 predicts risk and prognosis of colorectal cancer. Oncotarget, 2016, 7, 62084-62090.	1.8	13
43	MicroRNA-10a silencing reverses cisplatin resistance in the A549/cisplatin human lung cancer cell line via the transforming growth factor-l²/Smad2/STAT3/STAT5 pathway. Molecular Medicine Reports, 2015, 11, 3854-3859.	2.4	44
44	Association Between Genetic Polymorphisms in the Promoter Regions of Let-7 and Risk of Papillary Thyroid Carcinoma. Medicine (United States), 2015, 94, e1879.	1.0	20
45	Pri-Mir-34b/C and Tp-53 Polymorphisms are Associated With The Susceptibility of Papillary Thyroid Carcinoma. Medicine (United States), 2015, 94, e1536.	1.0	37
46	The variations in the AXIN1 gene and susceptibility to cryptorchidism. Journal of Pediatric Urology, 2015, 11, 132.e1-132.e5.	1.1	13
47	A Functional Polymorphism in the Promoter of MiR-143/145 Is Associated With the Risk of Cervical Squamous Cell Carcinoma in Chinese Women. Medicine (United States), 2015, 94, e1289.	1.0	36
48	Polymorphisms and plasma levels of IL-27: impact on genetic susceptibility and clinical outcome of bladder cancer. BMC Cancer, 2015, 15, 433.	2.6	23
49	Association between polymorphisms in <i>IL27</i> gene and renal cell carcinoma. Biomarkers, 2015, 20, 202-205.	1.9	12
50	Prognostic value of serum nicotinamide phosphoribosyltransferase in patients with bladder cancer. Croatian Medical Journal, 2014, 55, 507-513.	0.7	18
51	Association between polymorphisms in AXIN1 gene and atrial septal defect. Biomarkers, 2014, 19, 674-678.	1.9	9
52	The variations in theIL1RL1gene and susceptibility to preeclampsia. Immunological Investigations, 2014, 43, 424-435.	2.0	6
53	An insertion–deletion polymorphism ininterleukin-1αgene associated with susceptibility to preeclampsia. Hypertension in Pregnancy, 2014, 33, 395-401.	1.1	5
54	Genetic variants in NAMPT predict bladder cancer risk and prognosis in individuals from southwest Chinese Han group. Tumor Biology, 2014, 35, 4031-4040.	1.8	22

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55	Association between a functional polymorphism rs712 within let-7-binding site and risk of papillary thyroid cancer. Medical Oncology, 2014, 31, 221.	2.5	20
56	Association between a functional insertion/deletion polymorphism in IL1A gene and risk of papillary thyroid carcinoma. Tumor Biology, 2014, 35, 3861-3865.	1.8	18
57	Association of an insertion/deletion polymorphism in IL1A 3′-UTR with risk for cervical carcinoma in Chinese Han Women. Human Immunology, 2014, 75, 740-744.	2.4	24
58	High NUAK1 expression correlates with poor prognosis and involved in NSCLC cells migration and invasion. Experimental Lung Research, 2013, 39, 9-17.	1.2	43