Xiong Wang

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/4592014/publications.pdf

Version: 2024-02-01

81	6,239	448610 19	84171 75
papers	citations	h-index	g-index

83	83	83	14673
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Prognostic Value and Immune Infiltration Analysis of Nuclear Factor Erythroid-2 Family Members in Ovarian Cancer. BioMed Research International, 2022, 2022, 1-9.	0.9	5
2	Effective Diagnosis of Prostate Cancer Based on mRNAs From Urinary Exosomes. Frontiers in Medicine, 2022, 9, 736110.	1.2	9
3	VHL Ser65 mutations enhance HIF2Î± signaling and promote epithelial-mesenchymal transition of renal cancer cells. Cell and Bioscience, 2022, 12, 52.	2.1	4
4	ceRNA Network Analysis Reveals AP-1 Transcription Factor Components as Potential Biomarkers for Alzheimerâ€™s Disease. Current Alzheimer Research, 2022, 19, 387-406.	0.7	3
5	Trioâ€WES reveals a novel de novo missense mutation of KMT2A in a Chinese patient with Wiedemannâ€Steiner syndrome: A case report. Molecular Genetics & Genomic Medicine, 2021, 9, e1533.	0.6	4
6	The emerging role of non-coding RNAs from extracellular vesicles in Alzheimerâ€™s disease. Journal of Integrative Neuroscience, 2021, 20, 239.	0.8	6
7	TWIN BROTHERS WITH VARIABLE PHENOTYPES OF HAEMOPHILIA A RESULTING FROM MATERNAL MOSAICISM OF ARG550CYS OF F8. Journal of Paediatrics and Child Health, 2021, 57, 307-309.	0.4	0
8	Risk Factors for Mortality of COVID-19 Patient Based on Clinical Course: A Single Center Retrospective Case-Control Study. Frontiers in Immunology, 2021, 12, 581469.	2.2	14
9	Harnessing Big Data to Optimize an Algorithm for Rapid Diagnosis of Pulmonary Tuberculosis in a Real-World Setting. Frontiers in Cellular and Infection Microbiology, 2021, 11, 650163.	1.8	6
10	Apolipoprotein E genotype predicts subarachnoid extension in spontaneous intracerebral haemorrhage. European Journal of Neurology, 2021, 28, 1992-1999.	1.7	3
11	Toxicities and Associated Factors in Patients Receiving Temozolomide-Containing Regimens: A 12-Year Analysis of Hospital Data. Drug Design, Development and Therapy, 2021, Volume 15, 2151-2159.	2.0	4
12	A novel nonsense mutation in ARMC5 causes primary bilateral macronodular adrenocortical hyperplasia. BMC Medical Genomics, 2021, 14, 126.	0.7	6
13	Identification of a recurrent missense mutation in the FGA gene likely causing a congenital fibrinogen disorder. Blood Coagulation and Fibrinolysis, 2021, 32, 424-426.	0.5	0
14	Prognostic and Immunological Role of THBS2 in Colorectal cancer. BioMed Research International, 2021, 2021, 1-11.	0.9	7
15	A Case of Parturient with Hereditary Thrombotic Thrombocytopenic Purpura: Case Report of a Novel Variant. Seminars in Thrombosis and Hemostasis, 2021, , .	1.5	2
16	An aberrant F8 intron 1 inversion with concomitant large duplication and deletion in a Chinese severe hemophilia A patient. Hematology, 2021, 26, 53-57.	0.7	1
17	Expression of Immune Related Genes and Possible Regulatory Mechanisms in Alzheimerâ€™s Disease. Frontiers in Immunology, 2021, 12, 768966.	2.2	33
18	Successful treatment of a kidney transplant patient with COVID-19 and late-onset Pneumocystis jirovecii pneumonia. Annals of Clinical Microbiology and Antimicrobials, 2021, 20, 83.	1.7	8

#	Article	IF	Citations
19	Concurrent hematoma and venous thrombosis in a patient with autoimmune acquired factor XIII deficiency. International Journal of Laboratory Hematology, 2020, 42, e4-e6.	0.7	5
20	Association of APOE Îµ4 with progressive hemorrhagic injury in patients with traumatic intracerebral hemorrhage. Journal of Neurosurgery, 2020, 133, 496-503.	0.9	5
21	Alpha and beta-Thalassemia mutations in Hubei area of China. BMC Medical Genetics, 2020, 21, 6.	2.1	12
22	Vitamin D receptor rs2228570 polymorphism and Parkinsonâ€™s disease risk in a Chinese population. Neuroscience Letters, 2020, 717, 134722.	1.0	11
23	Two cases of von Willebrand disease type 3 in consanguineous Chinese families. Molecular Genetics & Genomic Medicine, 2020, 8, e1075.	0.6	4
24	Deep intronic F8 c.5999-27A>G variant causes exon 19 skipping and leads to moderate hemophilia A. Blood Coagulation and Fibrinolysis, 2020, 31, 476-480.	0.5	3
25	APOE rs405509 polymorphism and Parkinsonâ€™s disease risk in the Chinese population. Neuroscience Letters, 2020, 736, 135256.	1.0	6
26	Author Response to â€œGeneralizability of COVID-19 Mortality Risk Score Modelâ€• American Journal of Preventive Medicine, 2020, 59, e251.	1.6	3
27	Genetic and Clinical Characteristics of Patients With Hereditary Spherocytosis in Hubei Province of China. Frontiers in Genetics, 2020, 11, 953.	1.1	24
28	Epidemiological and clinical characteristics of 1663 hospitalized patients infected with COVID-19 in Wuhan, China: a single-center experience. Journal of Infection and Public Health, 2020, 13, 1202-1209.	1.9	42
29	Normal activated partial thromboplastin time in Chinese patients with mild hemophilia B. Hematology, 2020, 25, 484-488.	0.7	0
30	Clinical Characteristics, Associated Factors, and Predicting COVID-19 Mortality Risk: A Retrospective Study in Wuhan, China. American Journal of Preventive Medicine, 2020, 59, 168-175.	1.6	115
31	The association between the C-reactive protein gene +1444C/T polymorphism and Parkinson's disease susceptibility in a Chinese population. Gene, 2020, 753, 144808.	1.0	3
32	Abnormal coagulation parameters are associated with poor prognosis in patients with novel coronavirus pneumonia. Journal of Thrombosis and Haemostasis, 2020, 18, 844-847.	1.9	4,615
33	Rapid genotyping of F8 intron 22 inversion by nested PCR based on long-distance PCR. Journal of Thrombosis and Thrombolysis, 2020, 49, 591-601.	1.0	2
34	Correcting abnormalities in miRâ€124/PTPN1 signaling rescues tau pathology in Alzheimerâ€™s disease. Journal of Neurochemistry, 2020, 154, 441-457.	2.1	43
35	Congenital fibrinogen disorder caused by digenic mutations of the FGA and FGB genes. Hematology, 2020, 25, 145-148.	0.7	2
36	Clinical characteristics of 80 hospitalized frontline medical workers infected with COVID-19 in Wuhan, China. Journal of Hospital Infection, 2020, 105, 399-403.	1.4	64

#	Article	IF	Citations
37	Comparison of nasopharyngeal and oropharyngeal swabs for SARS-CoV-2 detection in 353 patients received tests with both specimens simultaneously. International Journal of Infectious Diseases, 2020, 94, 107-109.	1.5	219
38	Regulatory Effects of Circular RNAs on Host Genes in Human Cancer. Frontiers in Oncology, 2020, 10, 586163.	1.3	22
39	Circular RNAs in Human Cancer. Frontiers in Oncology, 2020, 10, 577118.	1.3	20
40	Abnormal coagulation parameters are associated with poor prognosis in patients with novel coronavirus pneumonia. , 2020, 18, 844.		1
41	C-Reactive Protein +1444C/T Polymorphism Is Associated with the Susceptibility to Pulmonary Tuberculosis. BioMed Research International, 2020, 2020, 1-4.	0.9	2
42	Mutation of Factor IX Cys178 is intolerant and may cause severe hemophilia B. Thrombosis Research, 2019, 183, 108-110.	0.8	2
43	Hereditary elliptocytosis with variable expression and incomplete penetrance in a Chinese family. British Journal of Haematology, 2019, 186, e159-e162.	1.2	3
44	Circular HDAC9/microRNA-138/Sirtuin-1 Pathway Mediates Synaptic and Amyloid Precursor Protein Processing Deficits in Alzheimerâ€™s Disease. Neuroscience Bulletin, 2019, 35, 877-888.	1.5	107
45	Novel compound heterozygous mutations in the SPTA1 gene, causing hereditary spherocytosis in a neonate with Coombsâ€negative hemolytic jaundice. Molecular Medicine Reports, 2019, 19, 2801-2807.	1.1	7
46	Vitamin D receptor polymorphisms and the susceptibility of Parkinsonâ€™s disease. Neuroscience Letters, 2019, 699, 206-211.	1.0	13
47	F8 IVS9+5G>A mutation causes moderate haemophilia A. Haemophilia, 2019, 25, e132-e135.	1.0	1
48	PROS1 IVS10+5G>A mutation causes hereditary protein S deficiency in a Chinese patient with pulmonary embolism and venous thromboembolism. Thrombosis Research, 2019, 174, 1-4.	0.8	6
49	Association of RAGE rs1800625 Polymorphism and Cancer Risk: A Meta-Analysis of 18 Case-Control Studies. Medical Science Monitor, 2019, 25, 7026-7034.	0.5	6
50	Novel hereditary spherocytosis-associated splice site mutation in the ANK1 gene caused by parental gonosomal mosaicism. Haematologica, 2018, 103, e219-e222.	1.7	9
51	Tau-Induced Ca2+/Calmodulin-Dependent Protein Kinase-IV Activation Aggravates Nuclear Tau Hyperphosphorylation. Neuroscience Bulletin, 2018, 34, 261-269.	1.5	20
52	A Novel MicroRNA-124/PTPN1 Signal Pathway Mediates Synaptic and Memory Deficits in Alzheimerâ€™s Disease. Biological Psychiatry, 2018, 83, 395-405.	0.7	153
53	Hemophagocytic lymphohistiocytosis and congenital factor VII deficiency: a case report. BMC Medical Genetics, 2018, 19, 163.	2.1	1
54	Identification of a Homozygous Missense Mutation in the TYR Gene in a Chinese Family with OCA1. Current Medical Science, 2018, 38, 932-936.	0.7	1

#	Article	IF	Citations
55	Validation of the PLASMIC score, a clinical prediction tool for thrombotic thrombocytopenic purpura diagnosis, in Chinese patients. Thrombosis Research, 2018, 172, 9-13.	0.8	23
56	Exacerbation of ichthyosis vulgaris phenotype by co-inheritance of STS and FLG mutations in a Chinese family with ichthyosis: a case report. BMC Medical Genetics, 2018, 19, 120.	2.1	7
57	A de novo deletion mutation in SOX10 in a Chinese family with Waardenburg syndrome type 4. Scientific Reports, 2017, 7, 41513.	1.6	5
58	Meta-analysis Reveals the Prognostic Value of Circulating Tumour Cells Detected in the Peripheral Blood in Patients with Non-Metastatic Colorectal Cancer. Scientific Reports, 2017, 7, 905.	1.6	30
59	Melatonin ameliorates amygdala-dependent emotional memory deficits in Tg2576 mice by up-regulating the CREB/c-Fos pathway. Neuroscience Letters, 2017, 638, 76-82.	1.0	12
60	[P4â€“112]: ROLE AND MECHANISMS OF MICRORNAâ€124 IN THE PATHOGENESIS OF ALZHEIMER'S DISEASE. Alzheimer's and Dementia, 2017, 13, P1300.	0.4	0
61	Identification of a novel de novo ANK1 R1426* nonsense mutation in a Chinese family with hereditary spherocytosis by NGS. Oncotarget, 2017, 8, 96791-96797.	0.8	9
62	An ANK1 IVS3-2A>C mutation causes exon 4 skipping in two patients from a Chinese family with hereditary spherocytosis. Oncotarget, 2017, 8, 113282-113286.	0.8	6
63	Association between rs2853669 in TERT gene and the risk and prognosis of human cancer: a systematic review and meta-analysis. Oncotarget, 2017, 8, 50864-50872.	0.8	24
64	Association between the p.V37I variant of GJB2 and hearing loss: a pedigree and meta-analysis. Oncotarget, 2017, 8, 46681-46690.	0.8	21
65	Mutational analysis of a Chinese family with oculocutaneous albinism type 2. Oncotarget, 2017, 8, 70345-70355.	0.8	3
66	Mutation analysis of a Chinese family with oculocutaneous albinism. Oncotarget, 2016, 7, 84981-84988.	0.8	8
67	The significant prognostic value of circulating tumor cells in triple-negative breast cancer: a meta-analysis. Oncotarget, 2016, 7, 37361-37369.	0.8	27
68	Genetic association of CALHM1 rs2986017 polymorphism with risk of Alzheimerâ€™s disease: a meta-analysis. Neurological Sciences, 2016, 37, 525-532.	0.9	1
69	Possible association of CCDC62 rs12817488 polymorphism and Parkinsonâ€™s disease risk in Chinese population: a meta-analysis. Scientific Reports, 2016, 6, 23991.	1.6	5
70	ATP1B3: a virus-induced host factor against EV71 replication by up-regulating the production of type-I interferons. Virology, 2016, 496, 28-34.	1.1	15
71	Association of autophagy-related IRGM polymorphisms with latent versus active tuberculosis infection in a Chinese population. Tuberculosis, 2016, 97, 47-51.	0.8	19
72	FOXO3 rs12212067: T > G Association with Active Tuberculosis in Han Chinese Population. Inflammation, 2016, 39, 10-15.	1.7	15

#	Article	IF	Citations
73	Association of lncRNA H19 rs217727 polymorphism and cancer risk in the Chinese population: a meta-analysis. Oncotarget, 2016, 7, 59580-59588.	0.8	17
74	Genetic association of RIT2 rs12456492 polymorphism and Parkinsonâ€™s disease susceptibility in Asian populations: a meta-analysis. Scientific Reports, 2015, 5, 13805.	1.6	21
75	Intraperitoneal Administration of a Novel TAT-BDNF Peptide Ameliorates Cognitive Impairments via Modulating Multiple Pathways in Two Alzheimerâ€™s Rodent Models. Scientific Reports, 2015, 5, 15032.	1.6	43
76	Genotype Distribution and Molecular Epidemiology of Hepatitis C Virus in Hubei, Central China. PLoS ONE, 2015, 10, e0137059.	1.1	20
77	TREM2 variants and risk of Alzheimerâ€™s disease: a meta-analysis. Neurological Sciences, 2015, 36, 1881-1888.	0.9	30
78	MicroRNAâ€138 promotes tau phosphorylation by targeting retinoic acid receptor alpha. FEBS Letters, 2015, 589, 726-729.	1.3	96
79	Activation of Glycogen Synthase Kinase-3 Mediates the Olfactory Deficit-Induced Hippocampal Impairments. Molecular Neurobiology, 2015, 52, 1601-1617.	1.9	22
80	Lack of association between FOXO1 polymorphisms and bacteremia. International Journal of Clinical and Experimental Medicine, 2015, 8, 16384-8.	1.3	0
81	Melatonin Attenuates Scopolamine-Induced Memory/Synaptic Disorder by Rescuing EPACs/miR-124/Egr1 Pathway. Molecular Neurobiology, 2013, 47, 373-381.	1.9	84