Yin Wang

List of Publications by Year
in descending order

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57 papers	1,160 citations	516215 16 h-index	414034 32 g-index
61 all docs	61 docs citations	61 times ranked	2147 citing authors

#	Article	IF	Citations
1	Associations of early nutrition with growth and body composition in very preterm infants: a prospective cohort study. European Journal of Clinical Nutrition, 2022, 76, 103-110.	1.3	7
2	Paraganglioma of the cauda equina: A clinicopathologic study of 12 cases with demonstration of cytokeratin positivity. Annals of Diagnostic Pathology, 2022, 57, 151887.	0.6	1
3	Maternal liver dysfunction in early pregnancy predisposes to gestational diabetes mellitus independent of preconception overweight: A prospective cohort study. BJOG: an International Journal of Obstetrics and Gynaecology, 2022, 129, 1695-1703.	1.1	4
4	Not Only in Sensorimotor Network: Local and Distant Cerebral Inherent Activity of Chronic Ankle Instabilityâ€"A Resting-State fMRI Study. Frontiers in Neuroscience, 2022, 16, 835538.	1.4	6
5	Mycoplasma pneumoniae and Adenovirus Coinfection Cause Pediatric Severe Community-Acquired Pneumonia. Microbiology Spectrum, 2022, 10, e0002622.	1.2	8
6	RBC Folate and Serum Folate, Vitamin B-12, and Homocysteine in Chinese Couples Prepregnancy in the Shanghai Preconception Cohort. Journal of Nutrition, 2022, 152, 1496-1506.	1.3	2
7	Association of Maternal Folate and Vitamin B12 in Early Pregnancy With Gestational Diabetes Mellitus: A Prospective Cohort Study. Diabetes Care, 2021, 44, 217-223.	4.3	54
8	Performance of waist-to-height ratio as a screening tool for identifying cardiometabolic risk in children: a meta-analysis. Diabetology and Metabolic Syndrome, 2021, 13, 66.	1.2	9
9	Etiology of Severe Pneumonia in Children in Alveolar Lavage Fluid Using a High-Throughput Gene Targeted Amplicon Sequencing Assay. Frontiers in Pediatrics, 2021, 9, 659164.	0.9	10
10	Higher Serum Bilirubin Levels in Response to Higher Carbohydrate Intake During Early Pregnancy and Lower Gestational Diabetes Mellitus Occurrence in Overweight and Obese Gravidae. Frontiers in Nutrition, 2021, 8, 701422.	1.6	2
11	Association Between Histopathology and Magnetic Resonance Imaging Texture in Grading Gliomas Based on Intraoperative Magnetic Resonance Navigated Stereotactic Biopsy. Journal of Computer Assisted Tomography, 2021, 45, 728-735.	0.5	1
12	The Hippo-TAZ axis mediates vascular endothelial growth factor C in glioblastoma-derived exosomes to promote angiogenesis. Cancer Letters, 2021, 513, 1-13.	3.2	18
13	Early skin contact combined with motherâ€™s breastfeeding to shorten the process of premature infants â‰¥ 30 weeks of gestation to achieve full oral feeding: the study protocol of a randomized controlled trial. Trials, 2021, 22, 637.	0.7	4
14	Microstructural Alteration of Lumbosacral Nerve Roots in Chronic Inflammatory Demyelinating Polyradiculoneuropathy: Insights From DTI and Correlations with Electrophysiological Parameters. Academic Radiology, 2021, , .	1.3	2
15	Effects of immunoglobulin plus prednisolone in reducing coronary artery lesions in patients with Kawasaki disease: study protocol for a phase III multicenter, open-label, blinded-endpoints randomized controlled trial. Trials, 2021, 22, 898.	0.7	2
16	Feasibility and effectiveness of prone position ventilation technique for postoperative acute lung injury in infants with congenital heart disease: study protocol for a prospective randomized study. Trials, 2021, 22, 929.	0.7	0
17	MR neurography of lumbosacral nerve roots: Diagnostic value in chronic inflammatory demyelinating polyradiculoneuropathy and correlation with electrophysiological parameters. European Journal of Radiology, 2020, 124, 108816.	1.2	7
18	OLIG2 Immunolabeling of Mesenchymal Chondrosarcoma: Report of 14 Cases. Journal of Neuropathology and Experimental Neurology, 2020, 79, 959-965.	0.9	6

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19	ETMR-22. TITLE: DEFINING THE CLINICAL AND PROGNOSTIC LANDSCAPE OF EMBRYONAL TUMORS WITH MULTI-LAYERED ROSETTES (ETMRs), A RARE BRAIN TUMOR REGISTRY (RBTC) STUDY. Neuro-Oncology, 2020, 22, iii327-iii328.	0.6	0
20	Waist-to-height ratio as a screening tool for cardiometabolic risk in children and adolescents: a nationwide cross-sectional study in China. BMJ Open, 2020, 10, e037040.	0.8	20
21	Acupuncture treatment on the motor area of the scalp for motor dysfunction in children with cerebral palsy: study protocol for a multicenter randomized controlled trial. Trials, 2020, 21, 29.	0.7	9
22	Central nervous system impairment detected by somatosensory evoked potentials in patients with Charcot-Marie-Tooth disease type 1A. Journal of Clinical Neuroscience, 2020, 79, 191-196.	0.8	1
23	A C19MC-LIN28A-MYCN Oncogenic Circuit Driven by Hijacked Super-enhancers Is a Distinct Therapeutic Vulnerability in ETMRs: A Lethal Brain Tumor. Cancer Cell, 2019, 36, 51-67.e7.	7.7	69
24	Loss-of-function mutations in Lysyl-tRNA synthetase cause various leukoencephalopathy phenotypes. Neurology: Genetics, 2019, 5, e565.	0.9	9
25	A case of solitary fibrous tumor/hemangiopericytoma in the central nervous system with papillary morphology. Neuropathology, 2019, 39, 141-146.	0.7	2
26	Efficacy of adjuvant radiotherapy for atypical and anaplastic meningioma. Cancer Medicine, 2019, 8, 13-20.	1.3	55
27	MR textural analysis on contrast enhanced 3D-SPACE images in assessment of consistency of pituitary macroadenoma. European Journal of Radiology, 2019, 110, 219-224.	1.2	21
28	Decreased Distance between Representation Sites of Distinct Facial Movements in Facial Synkinesisâ€"A Task fMRI Study. Neuroscience, 2019, 397, 12-17.	1.1	14
29	Detection of H3K27M mutation in cases of brain stem subependymoma. Human Pathology, 2019, 84, 262-269.	1.1	16
30	Preliminary Exploration of the Diagnosis and Treatment of Skull-Based Chondromyxoid Fibromas. Operative Neurosurgery, 2018, 15, 270-277.	0.4	3
31	Hospitalization Rate and Population-based Incidence of Hospitalization for Community-acquired Pneumonia Among Children in Suzhou, China. Pediatric Infectious Disease Journal, 2018, 37, 1242-1247.	1.1	6
32	MR textural analysis on T₂ FLAIR images for the prediction of true oligodendroglioma by the 2016 WHO genetic classification. Journal of Magnetic Resonance Imaging, 2018, 48, 74-83.	1.9	18
33	Statin-naÃ¯ve anti-HMGCR antibody-mediated necrotizing myopathy in China. Journal of Clinical Neuroscience, 2018, 57, 13-19.	0.8	18
34	Patterns of cortical reorganization in facial synkinesis: a task functional magnetic resonance imaging study. Neural Regeneration Research, 2018, 13, 1637.	1.6	14
35	Glioma groups classified by IDH and TERT promoter mutations remain stable among primary and recurrent gliomas. Neuro-Oncology, 2017, 19, 1008-1010.	0.6	12
36	Differences in the Prognostic Value of Tumor Extent of Resection among the Molecular Subgroups of Medulloblastoma: A Single Centre Study of 113 Cases. Translational Neuroscience and Clinics, 2017, 3, 66-73.	0.1	0

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37	Detecting isocitrate dehydrogenase gene mutations in oligodendroglial tumors using diffusion tensor imaging metrics and their correlations with proliferation and microvascular density. Journal of Magnetic Resonance Imaging, 2016, 43, 45-54.	1.9	28
38	Clinicopathologic and neuroradiologic studies of papillary glioneuronal tumors. Acta Neurochirurgica, 2016, 158, 695-702.	0.9	20
39	Clinicopathological analysis of UHRF1 expression in medulloblastoma tissues and its regulation on tumor cell proliferation. Medical Oncology, 2016, 33, 99.	1.2	10
40	Extraventricular neurocytoma of the sellar region: case report and literature review. SpringerPlus, 2016, 5, 987.	1.2	17
41	Prognostic Factors in Patients with Primary Hemangiopericytomas of the Central Nervous System: A Series of 103 Cases at a Single Institution. World Neurosurgery, 2016, 90, 414-419.	0.7	14
42	Pilomyxoid astrocytomas with rare rosenthal fibers. Brain Tumor Pathology, 2016, 33, 35-39.	1.1	3
43	Combination of diffusion tensor imaging and conventional MRI correlates with isocitrate dehydrogenase 1/2 mutations but not 1p/19q genotyping in oligodendroglial tumours. European Radiology, 2016, 26, 1705-1715.	2.3	54
44	Common variants at 10p12.31, 10q21.1 and 13q12.13 are associated with sporadic pituitary adenoma. Nature Genetics, 2015, 47, 793-797.	9.4	43
45	Recurrent gain-of-function USP8 mutations in Cushing's disease. Cell Research, 2015, 25, 306-317.	5.7	263
46	New disease allele and de novo mutation indicate mutational vulnerability of titin exon 343 in hereditary myopathy with early respiratory failure. Neuromuscular Disorders, 2015, 25, 172-176.	0.3	16
47	Analysis of prognostic factors and treatment of anaplastic meningioma in China. Journal of Clinical Neuroscience, 2015, 22, 690-695.	0.8	24
48	Muscle pathology and whole-body MRI in a polyglucosan myopathy associated with a novel glycogenin-1 mutation. Neuromuscular Disorders, 2015, 25, 780-785.	0.3	28
49	TERTpromoter mutations contribute toIDHmutations in predicting differential responses to adjuvant therapies in WHO grade II and III diffuse gliomas. Oncotarget, 2015, 6, 24871-24883.	0.8	34
50	Gene mutation profiling of primary glioblastoma through multiple tumor biopsy guided by 1H-magnetic resonance spectroscopy. International Journal of Clinical and Experimental Pathology, 2015, 8, 5327-35.	0.5	13
51	Primary testicular lymphoma with subcutaneous masses as the sole manifestation of the first relapse and central nervous system lymphoma as the second relapse: A case report and literature review. Oncology Letters, 2014, 7, 1881-1884.	0.8	1
52	Giant cell polymyositis associated with myasthenia gravis and thymoma. Journal of Clinical Neuroscience, 2014, 21, 2252-2254.	0.8	11
53	Diagnosis and Surgical Treatment of Cavernous Sinus Angioleiomyoma: A Report of Four Cases. Japanese Journal of Clinical Oncology, 2014, 44, 1052-1057.	0.6	7
54	Central nervous system tumors: a single center pathology review of 34,140 cases over 60 years. BMC Clinical Pathology, 2013, 13, 14.	1.8	42

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55	Rosette-forming glioneuronal tumor: report of an unusual case with intraventricular dissemination. Acta Neuropathologica, 2009, 118, 813-819.	3.9	54
56	Histopathological study of five cases with sporadic meningioangiomatosis. Neuropathology, 2006, 26, 249-256.	0.7	36
57	Report on the first Chinese family with Gerstmann-StrÃussler-Scheinker disease manifesting the codon 102 mutation in the prion protein gene. Neuropathology, 2006, 26, 429-432.	0.7	11