Josef Zamecnik

List of Publications by Year in descending order

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471509 289244 1,726 61 17 40 citations h-index g-index papers 61 61 61 3445 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	Atypical Teratoid/Rhabdoid Tumors Are Comprised of Three Epigenetic Subgroups with Distinct Enhancer Landscapes. Cancer Cell, 2016, 29, 379-393.	16.8	438
2	Alterations in ALK/ROS1/NTRK/MET drive a group of infantile hemispheric gliomas. Nature Communications, 2019, 10, 4343.	12.8	200
3	Value of thyroid transcription factor-1 and surfactant apoprotein A in the differential diagnosis of pulmonary carcinomas: a study of 109 cases. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2002, 440, 353-361.	2.8	140
4	The extracellular space and matrix of gliomas. Acta Neuropathologica, 2005, 110, 435-442.	7.7	121
5	Histone H3.3G34-Mutant Interneuron Progenitors Co-opt PDGFRA for Gliomagenesis. Cell, 2020, 183, 1617-1633.e22.	28.9	93
6	Pediatric Intracranial Ependymomas: Prognostic Relevance of Histological, Immunohistochemical, and Flow Cytometric Factors. Modern Pathology, 2003, 16, 980-991.	5.5	76
7	Novel Histopathological Patterns in Cortical Tubers of Epilepsy Surgery Patients with Tuberous Sclerosis Complex. PLoS ONE, 2016, 11, e0157396.	2.5	69
8	Outcomes of BRAF V600E Pediatric Gliomas Treated With Targeted BRAF Inhibition. JCO Precision Oncology, 2020, 4, 561-571.	3.0	62
9	Increasing incidence of immune-mediated necrotizing myopathy: single-centre experience. Rheumatology, 2015, 54, 2010-2014.	1.9	55
10	Functional aspects of early brain development are preserved in tuberous sclerosis complex (TSC) epileptogenic lesions. Neurobiology of Disease, 2016, 95, 93-101.	4.4	50
11	Thymidine kinase 2 and alanyl-tRNA synthetase 2 deficiencies cause lethal mitochondrial cardiomyopathy: case reports and review of the literature. Cardiology in the Young, 2017, 27, 936-944.	0.8	28
12	The coding and non-coding transcriptional landscape of subependymal giant cell astrocytomas. Brain, 2020, 143, 131-149.	7.6	24
13	Pattern of Relapse and Treatment Response in WNT-Activated Medulloblastoma. Cell Reports Medicine, 2020, 1, 100038.	6.5	24
14	Desminopathy: Novel Desmin Variants, a New Cardiac Phenotype, and Further Evidence for Secondary Mitochondrial Dysfunction. Journal of Clinical Medicine, 2020, 9, 937.	2.4	24
15	Radiation-induced gliomas represent H3-/IDH-wild type pediatric gliomas with recurrent PDGFRA amplification and loss of CDKN2A/B. Nature Communications, 2021, 12, 5530.	12.8	24
16	Specific pattern of maturation and differentiation in the formation of cortical tubers in tuberous sclerosis complex (TSC): evidence from layer-specific marker expression. Journal of Neurodevelopmental Disorders, 2016, 8, 9.	3.1	23
17	The extracellular matrix and diffusion barriers in focal cortical dysplasias. European Journal of Neuroscience, 2012, 36, 2017-2024.	2.6	22
18	Etiopathogenesis of adolescent idiopathic scoliosis: Expression of melatonin receptors 1A/1B, calmodulin and estrogen receptor 2 in deep paravertebral muscles revisited. Molecular Medicine Reports, 2016, 14, 5719-5724.	2.4	17

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19	Pro-inflammatory S100A11 is elevated in inflammatory myopathies and reflects disease activity and extramuscular manifestations in myositis. Cytokine, 2019, 116, 13-20.	3.2	17
20	Muscle lymphocytic infiltrates in thymoma-associated myasthenia gravis are phenotypically different from those in polymyositis. Neuromuscular Disorders, 2007, 17, 935-942.	0.6	16
21	Novel variant in the KCNK9 gene in a girl with Birk Barel syndrome. European Journal of Medical Genetics, 2020, 63, 103619.	1.3	16
22	GATOR1-related focal cortical dysplasia in epilepsy surgery patients and their families: A possible gradient in severity?. European Journal of Paediatric Neurology, 2021, 30, 88-96.	1.6	16
23	Evolution of pediatric epilepsy surgery program over 2000–2017: Improvement of care?. European Journal of Paediatric Neurology, 2019, 23, 456-465.	1.6	15
24	Three New PLP1 Splicing Mutations Demonstrate Pathogenic and Phenotypic Diversity of Pelizaeus-Merzbacher Disease. Journal of Child Neurology, 2014, 29, 924-931.	1.4	14
25	Brain gliomas, hydrocephalus and idiopathic aqueduct stenosis in children with neurofibromatosis type 1. Brain and Development, 2019, 41, 678-690.	1.1	12
26	Homozygous missense mutation in UQCRC2 associated with severe encephalomyopathy, mitochondrial complex III assembly defect and activation of mitochondrial protein quality control. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2021, 1867, 166147.	3.8	11
27	Unique presentation of LHON/MELAS overlap syndrome caused by m.13046T>C in <i>MTND5</i> Ophthalmic Genetics, 2016, 37, 419-423.	1.2	10
28	Limited clinical significance of tissue calprotectin levels in bowel mucosa for the prediction of complicated course of the disease in children with ulcerative colitis. Pathology Research and Practice, 2019, 215, 152689.	2.3	10
29	CO2-related vasoconstriction superimposed on ischemic medullary brain autonomic nuclei may contribute to sudden death. Cardiovascular Pathology, 2019, 38, 42-45.	1.6	8
30	The Phenotypic Spectrum of 47 Czech Patients with Single, Large-Scale Mitochondrial DNA Deletions. Brain Sciences, 2020, 10, 766.	2.3	8
31	Alemtuzumab and intrathecal methotrexate failed in the therapy of Rasmussen encephalitis. Neurology: Neuroimmunology and NeuroInflammation, 2017, 4, e354.	6.0	7
32	Low predictive value of histopathological scoring system for complications development in children with Crohn's disease. Pathology Research and Practice, 2017, 213, 353-358.	2.3	7
33	Rare IDH1 variants are common in pediatric hemispheric diffuse astrocytomas and frequently associated with Li-Fraumeni syndrome. Acta Neuropathologica, 2020, 139, 795-797.	7.7	7
34	The clinical utility of intraoperative electrocorticography in pediatric epilepsy surgical strategy and planning. Journal of Neurosurgery: Pediatrics, 2020, 26, 533-542.	1.3	7
35	Familial temporal lobe epilepsy due to focal cortical dysplasia type IIIa. Seizure: the Journal of the British Epilepsy Association, 2015, 31, 120-123.	2.0	6
36	Atrophy of type II fibres in myasthenia gravis muscle in thymectomized patients: steroidâ€induced change with prognostic impact. Journal of Cellular and Molecular Medicine, 2009, 13, 2008-2018.	3.6	5

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37	An immunotherapy effect analysis in Rasmussen encephalitis. BMC Neurology, 2020, 20, 359.	1.8	5
38	Survival and functional outcomes in paediatric thalamic and thalamopeduncular low grade gliomas. Acta Neurochirurgica, 2022, 164, 1459-1472.	1.7	5
39	Immunohistochemical Assessment of CD30+ Lymphocytes in the Intestinal Mucosa Facilitates Diagnosis of Pediatric Ulcerative Colitis. Digestive Diseases and Sciences, 2018, 63, 1811-1818.	2.3	3
40	MicroRNA expression in pediatric intracranial ependymomas and their potential value for tumor grading. Oncology Letters, 2019, 17, 1379-1383.	1.8	3
41	Cytometric analysis of cell suspension generated by cavitron ultrasonic surgical aspirator in pediatric brain tumors. Journal of Neuro-Oncology, 2019, 143, 15-25.	2.9	3
42	A novel PSMA/GCPIIâ€deficient mouse model shows enlarged seminal vesicles upon aging. Prostate, 2019, 79, 126-139.	2.3	3
43	Interleukin-35 in idiopathic inflammatory myopathies. Cytokine, 2021, 137, 155350.	3.2	3
44	Expression of cancer stem cells markers in urinary bladder urothelial carcinoma and its precursor lesions. Biomedical Papers of the Medical Faculty of the University Palacký, Olomouc, Czechoslovakia, 2021, 165, 316-321.	0.6	3
45	Expression of molecules of the Wnt pathway and of E‑cadherin in the etiopathogenesis of human thymomas. Oncology Letters, 2020, 19, 2413-2421.	1.8	3
46	Complex Immunometabolic Profiling Reveals the Activation of Cellular Immunity and Biliary Lesions in Patients with Severe COVID-19. Journal of Clinical Medicine, 2020, 9, 3000.	2.4	2
47	Cognitive performance in distinct groups of children undergoing epilepsy surgery—a single-centre experience. PeerJ, 2019, 7, e7790.	2.0	2
48	Hsp90 Levels in Idiopathic Inflammatory Myopathies and Their Association With Muscle Involvement and Disease Activity: A Cross-Sectional and Longitudinal Study. Frontiers in Immunology, 2022, 13, 811045.	4.8	2
49	Increased visfatin levels are associated with higher disease activity in anti-Jo-1-positive myositis patients. Clinical and Experimental Rheumatology, 2016, 34, 222-9.	0.8	2
50	GENE-14. UNIQUE MOLECULAR AND CLINICAL FEATURES OF LI-FRAUMENI SYNDROME ASSOCIATED BRAIN TUMOURS. Neuro-Oncology, 2019, 21, ii84-ii84.	1.2	1
51	Primary angiosarcoma of the femoral artery in patient with kidney and liver polycystosis and multiple arterial aneurysms: report of the case and review of the literature. Cardiovascular Pathology, 2019, 39, 8-11.	1.6	1
52	Histopathological examination of the ectocervical biopsy in nonâ€transplanted uteri: A study contributing to the provisional scoring system of subclinical graft rejection after uterus transplantation. Acta Obstetricia Et Gynecologica Scandinavica, 2021, 101, 37.	2.8	1
53	HGG-31. UNIQUE BIOLOGICAL CHARACTERISTICS OF RADIATION-INDUCED GLIOMAS. Neuro-Oncology, 2020, 22, iii349-iii349.	1.2	1
54	Clusterin is upregulated in serum and muscle tissue in idiopathic inflammatory myopathies and associates with clinical disease activity and cytokine profile. Clinical and Experimental Rheumatology, 2021, 39, 1021-1032.	0.8	1

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55	Inflammatory myopathy associated with statins: report of three cases. Modern Rheumatology, 2012, , 1.	1.8	0
56	$08.01 \hat{a} \in$ Heat shock protein 90 is increased in muscle tissue and plasma in idiopathic inflammatory myopathies. , $2017, , .$		0
57	ATRT-11. MOLECULAR BACKGROUND AND SURVIVAL OF PATIENTS WITH ATRT AND RHABDOID TUMOURS; SINGLE CENTRE EXPERIENCE. Neuro-Oncology, 2019, 21, ii65-ii65.	1.2	O
58	LGG-46. MOLECULAR CHARACTERIZATION OF HEMISPHERIC LOW-GRADE GLIOMAS IN CHILDREN. Neuro-Oncology, 2020, 22, iii374-iii375.	1.2	0
59	HGG-14. Molecular characterization of unique biological subgroups among H3 wild type high-grade gliomas. Neuro-Oncology, 2022, 24, i63-i63.	1.2	O
60	SURG-05. Survival and functional outcomes in pediatric thalamic and thalamopeduncular low grade gliomas. Neuro-Oncology, 2022, 24, i142-i143.	1.2	0
61	HGG-49. Gliomatosis cerebri in children: A collaborative report from the European Society for Pediatric Oncology (SIOPE). Neuro-Oncology, 2022, 24, i72-i73.	1.2	0