Suely K Marie

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9297702/publications.pdf

Version: 2024-02-01

289 papers	21,863 citations	31902 53 h-index	9839 141 g-index
299 all docs	299 docs citations	299 times ranked	32846 citing authors

#	Article	IF	Citations
1	An Integrated Genomic Analysis of Human Glioblastoma Multiforme. Science, 2008, 321, 1807-1812.	6.0	5,230
2	Detection of Circulating Tumor DNA in Early- and Late-Stage Human Malignancies. Science Translational Medicine, 2014, 6, 224ra24.	5.8	3,665
3	Altered Telomeres in Tumors with ATRX and DAXX Mutations. Science, 2011, 333, 425-425.	6.0	891
4	SOX2 is an amplified lineage-survival oncogene in lung and esophageal squamous cell carcinomas. Nature Genetics, 2009, 41, 1238-1242.	9.4	862
5	The Genetic Landscape of the Childhood Cancer Medulloblastoma. Science, 2011, 331, 435-439.	6.0	652
6	Transcriptomic analysis of purified human cortical microglia reveals age-associated changes. Nature Neuroscience, 2017, 20, 1162-1171.	7.1	575
7	Frequent ATRX, CIC, FUBP1 and IDH1 mutations refine the classification of malignant gliomas. Oncotarget, 2012, 3, 709-722.	0.8	532
8	Mutations in CIC and FUBP1 Contribute to Human Oligodendroglioma. Science, 2011, 333, 1453-1455.	6.0	485
9	Detection of tumor-derived DNA in cerebrospinal fluid of patients with primary tumors of the brain and spinal cord. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 9704-9709.	3.3	317
10	Evaluation of DNA from the Papanicolaou Test to Detect Ovarian and Endometrial Cancers. Science Translational Medicine, 2013, 5, 167ra4.	5.8	264
11	Increase in hand muscle strength of stroke patients after somatosensory stimulation. Annals of Neurology, 2002, 51, 122-125.	2.8	226
12	Title is missing!. Journal of Rehabilitation Research and Development, 2008, 45, 1215.	1.6	171
13	Congenital insensitivity to pain with anhidrosis (hereditary sensory and autonomic neuropathy type) Tj ETQq1 1 0.784314 rgBT /Over	1.0	170
14	The sarcoglycan complex in the six autosomal recessive limb-girdle muscular dystrophies. Human Molecular Genetics, 1996, 5, 1963-1969.	1.4	167
15	Therapeutic Impact of Cytoreductive Surgery and Irradiation of Posterior Fossa Ependymoma in the Molecular Era: A Retrospective Multicohort Analysis. Journal of Clinical Oncology, 2016, 34, 2468-2477.	0.8	160
16	Up-regulation of the inflammatory cytokines IFN-Î³ and IL-12 and down-regulation of IL-4 in cerebral cortex regions of APPSWE transgenic mice. Journal of Neuroimmunology, 2002, 126, 50-57.	1.1	150
17	New Molecular Mechanism for Ullrich Congenital Muscular Dystrophy: A Heterozygous In-Frame Deletion in the COL6A1 Gene Causes a Severe Phenotype. American Journal of Human Genetics, 2003, 73, 355-369.	2.6	150
18	PIK3CA Gene Mutations in Pediatric and Adult Glioblastoma Multiforme. Molecular Cancer Research, 2006, 4, 709-714.	1.5	148

#	Article	IF	Citations
19	Selection of suitable housekeeping genes for expression analysis in glioblastoma using quantitative RT-PCR. BMC Molecular Biology, 2009, 10, 17.	3.0	143
20	The Seventh Form of Autosomal Recessive Limb-Girdle Muscular Dystrophy Is Mapped to 17q11-12. American Journal of Human Genetics, 1997, 61, 151-159.	2.6	136
21	International consensus on a proposed score system for muscle biopsy evaluation in patients with juvenile dermatomyositis: A tool for potential use in clinical trials. Arthritis and Rheumatism, 2007, 57, 1192-1201.	6.7	132
22	Effects of somatosensory stimulation on motor function in chronic cortico-subcortical strokes. Journal of Neurology, 2007, 254, 333-339.	1.8	132
23	Effects of Somatosensory Stimulation on Motor Function After Subacute Stroke. Neurorehabilitation and Neural Repair, 2010, 24, 263-272.	1.4	130
24	Maternal embryonic leucine zipper kinase transcript abundance correlates with malignancy grade in human astrocytomas. International Journal of Cancer, 2008, 122, 807-815.	2.3	128
25	The Effects of School Crime Prevention on Studentsâ€™ Violent Victimization, Risk Perception, and Fear of Crime: A Multilevel Opportunity Perspective. Justice Quarterly, 2011, 28, 249-277.	1.1	126
26	The facioscapulohumeral muscular dystrophy (FSHD1) gene affects males more severely and more frequently than females. American Journal of Medical Genetics Part A, 1998, 77, 155-161.	2.4	123
27	The contribution of 700,000 ORF sequence tags to the definition of the human transcriptome. Proceedings of the National Academy of Sciences of the United States of America, 2001, 98, 12103-12108.	3.3	123
28	Automated genomic sequence analysis of the three collagen VI genes: applications to Ullrich congenital muscular dystrophy and Bethlem myopathy. Journal of Medical Genetics, 2005, 42, 108-120.	1.5	119
29	Bioinformatics construction of the human cell surfaceome. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 16752-16757.	3.3	119
30	Molecular basis of infantile reversible cytochrome c oxidase deficiency myopathy. Brain, 2009, 132, 3165-3174.	3.7	112
31	Genomic screening for beta-sarcoglycan gene mutations: missense mutations may cause severe limb-girdle muscular dystrophy type 2E (LGMD 2E). Human Molecular Genetics, 1996, 5, 1953-1961.	1.4	111
32	Gene expression profile analysis of primary glioblastomas and non-neoplastic brain tissue: identification of potential target genes by oligonucleotide microarray and real-time quantitative PCR. Journal of Neuro-Oncology, 2008, 88, 281-291.	1.4	109
33	Deconstructing Pompe Disease by Analyzing Single Muscle Fibers: â€œTo See a World in a Grain of Sandâ€¦â€• Autophagy, 2007, 3, 546-552.	4.3	102
34	Inhibition of Nuclear PTEN Tyrosine Phosphorylation Enhances Glioma Radiation Sensitivity through Attenuated DNA Repair. Cancer Cell, 2019, 35, 504-518.e7.	7.7	102
35	Resistance to EGF receptor inhibitors in glioblastoma mediated by phosphorylation of the PTEN tumor suppressor at tyrosine 240. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 14164-14169.	3.3	97
36	Metabolism and Brain Cancer. Clinics, 2011, 66, 33-43.	0.6	96

#	Article	IF	Citations
37	Linkage analysis in autosomal recessive limb-girdle muscular dystrophy (AR LGMD) maps a sixth form to 5q33-34 (LGMD2F) and indicates that there is at least one more subtype of AR LGMD. Human Molecular Genetics, 1996, 5, 815-820.	1.4	92
38	Correlation of MGMT promoter methylation status with gene and protein expression levels in glioblastoma. Clinics, 2011, 66, 1747-1755.	0.6	84
39	Sarcoglycanopathies are responsible for 68% of severe autosomal recessive limb-girdle muscular dystrophy in the Brazilian population. Journal of the Neurological Sciences, 1999, 164, 44-49.	0.3	81
40	Analysis of the CTG repeat in skeletal muscle of young and adult myotonic dystrophy patients: when does the expansion occur?. Human Molecular Genetics, 1995, 4, 401-406.	1.4	79
41	A common missense mutation in the adhalin gene in three unrelated Brazilian families with a relatively mild form of autosomal recessive limb-girdle muscular dystrophy. Human Molecular Genetics, 1995, 4, 1163-1167.	1.4	75
42	Activation of Neural and Pluripotent Stem Cell Signatures Correlates with Increased Malignancy in Human Glioma. PLoS ONE, 2011, 6, e18454.	1.1	75
43	Uncovering the Role of N-Acetyl-Aspartyl-Glutamate as a Glutamate Reservoir in Cancer. Cell Reports, 2019, 27, 491-501.e6.	2.9	73
44	Identification of human chromosome 22 transcribed sequences with ORF expressed sequence tags. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 12690-12693.	3.3	70
45	14-3-3 protein in the CSF of patients with rapidly progressive dementia. Neurology, 2003, 61, 354-357.	1.5	69
46	Exomic Sequencing of Four Rare Central Nervous System Tumor Types. Oncotarget, 2013, 4, 572-583.	0.8	69
47	Angiogenesis and expression of <scp>PDGF</scp>â€<scp>C</scp>, <scp>VEGF</scp>, <scp>CD</scp>105 and <scp>HIF</scp>â€1Î± in human glioblastoma. Neuropathology, 2014, 34, 343-352.	0.7	68
48	Update of the pompe disease mutation database with 60 novel GAA sequence variants and additional studies on the functional effect of 34 previously reported variants. Human Mutation, 2012, 33, 1161-1165.	1.1	67
49	Decreased AKT1/mTOR pathway mRNA expression in short-term bipolar disorder. European Neuropsychopharmacology, 2015, 25, 468-473.	0.3	65
50	Melanocyte Transformation Associated with Substrate Adhesion Impediment. Neoplasia, 2006, 8, 231-241.	2.3	61
51	Effects of High Adherence to Mediterranean or Low-Fat Diets in Medicated Secondary Prevention Patients. American Journal of Cardiology, 2011, 108, 1523-1529.	0.7	60
52	Frequency of parafunctional oral habits in patients with cerebral palsy. Journal of Oral Rehabilitation, 2007, 34, 323-328.	1.3	57
53	Pompe disease in a Brazilian series: clinical and molecular analyses with identification of nine new mutations. Journal of Neurology, 2009, 256, 1881-1890.	1.8	57
54	Leukocyte mitochondrial DNA copy number in bipolar disorder. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2014, 48, 32-35.	2.5	57

#	Article	IF	Citations
55	Validation of a score tool for measurement of histological severity in juvenile dermatomyositis and association with clinical severity of disease. Annals of the Rheumatic Diseases, 2015, 74, 204-210.	0.5	56
56	Targeted Assessment of G0S2 Methylation Identifies a Rapidly Recurrent, Routinely Fatal Molecular Subtype of Adrenocortical Carcinoma. Clinical Cancer Research, 2019, 25, 3276-3288.	3.2	51
57	Genetic heterogeneity for Duchenne-like muscular dystrophy (DLMD) based on linkage and 50 DAG analysis. Human Molecular Genetics, 1993, 2, 1945-1947.	1.4	50
58	Fatores de risco associados Ã calcinose na dermatomiosite juvenil. Jornal De Pediatria, 2008, 84, 68-74.	0.9	50
59	Deficiency of Merosin (Laminin M or Î±2) in Congenital Muscular Dystrophy Associated with Cerebral White Matter Alterations. Neuropediatrics, 1995, 26, 293-297.	0.3	48
60	A first missense mutation in the delta sarcoglycan gene associated with a severe phenotype and frequency of limb-girdle muscular dystrophy type 2F (LGMD2F) in Brazilian sarcoglycanopathies.. Journal of Medical Genetics, 1998, 35, 951-953.	1.5	48
61	Knobloch syndrome in a large Brazilian consanguineous family: Confirmation of autosomal recessive inheritance. American Journal of Medical Genetics Part A, 1994, 52, 170-173.	2.4	47
62	Disruption of prion proteinâ€“HOP engagement impairs glioblastoma growth and cognitive decline and improves overall survival. Oncogene, 2015, 34, 3305-3314.	2.6	47
63	Main clinical features of the three mapped autosomal recessive limb-girdle muscular dystrophies and estimated proportion of each form in 13 Brazilian families.. Journal of Medical Genetics, 1996, 33, 97-102.	1.5	45
64	Juvenile dermatomyositis: clinical, laboratorial, histological, therapeutical and evolutive parameters of 35 patients. Arquivos De Neuro-Psiquiatria, 2002, 60, 889-899.	0.3	45
65	Identification of novel differentially expressed genes in human astrocytomas by cDNA representational difference analysis. Molecular Brain Research, 2005, 140, 25-33.	2.5	42
66	Galectinâ€3 as an Immunohistochemical Tool to Distinguish Pilocytic Astrocytomas from Diffuse Astrocytomas, and Glioblastomas from Anaplastic Oligodendrogliomas. Brain Pathology, 2004, 14, 399-405.	2.1	42
67	Confirmation of the 2p Locus for the Mild Autosomal Recessive Limb-Girdle Muscular Dystrophy Gene (LGMD2B) in Three Families Allows Refinement of the Candidate Region. Genomics, 1995, 27, 192-195.	1.3	41
68	Modulation of HJURP (Holliday Junction-Recognizing Protein) Levels Is Correlated with Glioblastoma Cells Survival. PLoS ONE, 2013, 8, e62200.	1.1	41
69	LOX Expression and Functional Analysis in Astrocytomas and Impact of IDH1 Mutation. PLoS ONE, 2015, 10, e0119781.	1.1	40
70	Natural history of intraventricular meningiomas: systematic review. Neurosurgical Review, 2020, 43, 513-523.	1.2	40
71	Neuroimaging Findings in Rasmussen's Syndrome. Journal of Neuroimaging, 1997, 7, 16-22.	1.0	39
72	Nebulin expression in patients with nemaline myopathy. Neuromuscular Disorders, 2001, 11, 154-162.	0.3	39

#	Article	IF	Citations
73	Quantitative proteomic analysis shows differentially expressed HSPB1 in glioblastoma as a discriminating short from long survival factor and NOVA1 as a differentiation factor between low-grade astrocytoma and oligodendroglioma. BMC Cancer, 2015, 15, 481.	1.1	39
74	Difference in adhesion molecule expression (ICAM-1 and VCAM-1) in juvenile and adult dermatomyositis, polymyositis and inclusion body myositis. Autoimmunity Reviews, 2006, 5, 93-100.	2.5	38
75	Inhibition of phospholipase A2 reduces neurite outgrowth and neuronal viability. Prostaglandins Leukotrienes and Essential Fatty Acids, 2007, 76, 47-55.	1.0	38
76	Mitochondrial DNA depletion and its correlation with TFAM, TFB1M, TFB2M and POLG in human diffusely infiltrating astrocytomas. Mitochondrion, 2011, 11, 48-53.	1.6	38
77	Limited Ca2+ and PKA-pathway dependent neurogenic differentiation of human adult mesenchymal stem cells as compared to fetal neuronal stem cells. Experimental Cell Research, 2010, 316, 216-231.	1.2	37
78	Half the dystrophin gene is apparently enough for a mild clinical course: confirmation of its potential use for gene therapy. Human Molecular Genetics, 1994, 3, 919-922.	1.4	36
79	Expression of HOXC9 and E2F2 are up-regulated in CD133+ cells isolated from human astrocytomas and associate with transformation of human astrocytes. Biochimica Et Biophysica Acta Gene Regulatory Mechanisms, 2007, 1769, 437-442.	2.4	36
80	Proteomic analysis of lowâ€•to highâ€grade astrocytomas reveals an alteration of the expression level of raf kinase inhibitor protein and nucleophosmin. Proteomics, 2010, 10, 2812-2821.	1.3	36
81	Double Pathology in Rasmussen's Encephalitis: Etiologic Considerations. Epilepsia, 1996, 37, 495-500.	2.6	35
82	Merosin-deficient congenital muscular dystrophy (CMD): a study of 25 Brazilian patients using MRI. Pediatric Radiology, 2005, 35, 572-579.	1.1	33
83	Clinical Outcome, Tumor Recurrence, and Causes of Death: A Long-Term Follow-Up of Surgically Treated Meningiomas. World Neurosurgery, 2017, 102, 139-143.	0.7	33
84	Liver-specific Enhancer of the Glucokinase Gene. Journal of Biological Chemistry, 1996, 271, 29113-29120.	1.6	31
85	Diffusion-weighted MRI in two cases of familial Creutzfeldtâ€“Jakob disease. Journal of the Neurological Sciences, 2001, 184, 163-167.	0.3	31
86	Prognostic significance of co-overexpression of the EGFR/IGFBP-2/HIF-2A genes in astrocytomas. Journal of Neuro-Oncology, 2007, 83, 233-239.	1.4	31
87	Evidence of genetic heterogeneity in the autosomal recessive adult forms of limb-girdle muscular dystrophy following linkage analysis with 15q probes in Brazilian families.. Journal of Medical Genetics, 1993, 30, 385-387.	1.5	30
88	Mitochondria Transcription Factor A: A Putative Target for the Effect of Melatonin on U87MG Malignant Glioma Cell Line. Molecules, 2018, 23, 1129.	1.7	30
89	Risk factors associated with calcinosis of juvenile dermatomyositis. Jornal De Pediatria, 2007, 84, 68-74.	0.9	30
90	Pleiotrophin expression in astrocytic and oligodendroglial tumors and itâ€™s correlation with histological diagnosis, microvascular density, cellular proliferation and overall survival. Journal of Neuro-Oncology, 2007, 84, 255-261.	1.4	29

#	Article	IF	Citations
91	Changes in the expression of proteins associated with aerobic glycolysis and cell migration are involved in tumorigenic ability of two glioma cell lines. Proteome Science, 2012, 10, 53.	0.7	29
92	Intraoperative assistive technologies and extent of resection in glioma surgery: a systematic review of prospective controlled studies. Neurosurgical Review, 2015, 38, 217-227.	1.2	29
93	Immunohistological analysis of CD59 and membrane attack complex of complement in muscle in juvenile dermatomyositis. Journal of Rheumatology, 2002, 29, 1301-7.	1.0	29
94	Merosin-positive congenital muscular dystrophy in two siblings with cataract and slight mental retardation. Brain and Development, 1999, 21, 274-278.	0.6	28
95	Factors of morbidity in hemispherectomies: Surgical techniqueÃ—pathology. Brain and Development, 2006, 28, 215-222.	0.6	28
96	Molecular alterations in meningiomas: Literature review. Clinical Neurology and Neurosurgery, 2019, 176, 89-96.	0.6	28
97	Identiication of COL6A1 as a differentially expressed gene in human astrocytomas. Genetics and Molecular Research, 2008, 7, 371-378.	0.3	28
98	Myotonic dystrophy: genetic, clinical, and molecular analysis of patients from 41 Brazilian families.. Journal of Medical Genetics, 1995, 32, 14-18.	1.5	27
99	Frequency of temporomandibular disorder signs in individuals with cerebral palsy. Journal of Oral Rehabilitation, 2008, 35, 191-195.	1.3	27
100	Differential expression of E-cadherin gene in human neuroepithelial tumors. Genetics and Molecular Research, 2008, 7, 295-304.	0.3	27
101	Immunohistochemical analysis of adhesion molecule expression on muscle biopsy specimens from patients with juvenile dermatomyositis. Journal of Rheumatology, 2004, 31, 801-7.	1.0	27
102	Bite force and handgrip force in patients with molecular diagnosis of myotonic dystrophy. Journal of Oral Rehabilitation, 2007, 34, 195-200.	1.3	26
103	The Brazilian Consensus on the Management of Pompe Disease. Journal of Pediatrics, 2009, 155, S47-S56.	0.9	26
104	IDH1 mutations in a Brazilian series of Glioblastoma. Clinics, 2011, 66, 163-165.	0.6	26
105	Clinical characteristics and surgical outcome of patients with temporal lobe tumors and epilepsy. Arquivos De Neuro-Psiquiatria, 2000, 58, 1002-1008.	0.3	25
106	Volumetric evidence of a left laterality effect in epileptic psychosis. Epilepsy and Behavior, 2003, 4, 234-240.	0.9	25
107	Survival and Neuronal Differentiation of Mesenchymal Stem Cells Transplanted into the Rodent Brain Are Dependent upon Microenvironment. Tissue Engineering - Part A, 2010, 16, 2769-2782.	1.6	25
108	Adult stem cells in neural repair: Current options, limitations and perspectives. World Journal of Stem Cells, 2015, 7, 477.	1.3	25

#	Article	IF	Citations
109	CoGA: An R Package to Identify Differentially Co-Expressed Gene Sets by Analyzing the Graph Spectra. PLoS ONE, 2015, 10, e0135831.	1.1	25
110	Spontaneous cervical artery dissection: an update on clinical and diagnostic aspects. Arquivos De Neuro-Psiquiatria, 2008, 66, 922-927.	0.3	24
111	Antiâ€C1q Antibodies in Juvenileâ€Onset Systemic Lupus Erythematosus. Annals of the New York Academy of Sciences, 2009, 1173, 235-238.	1.8	24
112	Myositis in mixed connective tissue disease: a unique syndrome characterized by immunohistopathologic elements of both polymyositis and dermatomyositis. Arquivos De Neuro-Psiquiatria, 2004, 62, 923-934.	0.3	24
113	No evidence of genetic heterogeneity in Brazilian facioscapulohumeral muscular dystrophy familes (FSHD) with 4q markers. Human Molecular Genetics, 1993, 2, 557-562.	1.4	23
114	Congenital muscular dystrophy with cerebral white matter hypodensity. Correlation of clinical features and merosin deficiency. Brain and Development, 1996, 18, 53-58.	0.6	23
115	Deficiency of Î±-Actinin-3 (ACTN3) Occurs in Different Forms of Muscular Dystrophy. Neuropediatrics, 1997, 28, 223-228.	0.3	23
116	Congenital Muscular Dystrophy with Merosin Deficiency:1H MR Spectroscopy and Diffusion-weighted MR Imaging. Radiology, 2005, 235, 190-196.	3.6	23
117	Identification of FAM46D as a novel cancer/testis antigen using EST data and serological analysis. Genomics, 2009, 94, 153-160.	1.3	23
118	Comparison of motor strength and function in patients with Duchenne muscular dystrophy with or without steroid therapy. Arquivos De Neuro-Psiquiatria, 2010, 68, 683-688.	0.3	23
119	Expression of tissue factor signaling pathway elements correlates with the production of vascular endothelial growth factor and interleukin-8 in human astrocytoma patients. Oncology Reports, 2014, 31, 679-686.	1.2	23
120	Serum amyloid A1 is upregulated in human glioblastoma. Journal of Neuro-Oncology, 2017, 132, 383-391.	1.4	23
121	Impact of radiotherapy in atypical meningioma recurrence: literature review. Neurosurgical Review, 2019, 42, 631-637.	1.2	23
122	A Caucasian Family with the 3271 Mutation in Mitochondrial DNA. Biochemical Medicine and Metabolic Biology, 1994, 52, 136-139.	0.7	22
123	Further evidence for the organisation of the four sarcoglycans proteins within the dystrophinâ€“glycoprotein complex. European Journal of Human Genetics, 1999, 7, 251-254.	1.4	22
124	A Transcript Finishing Initiative for Closing Gaps in the Human Transcriptome. Genome Research, 2004, 14, 1413-1423.	2.4	22
125	Recessive COL6A2 C-globular Missense Mutations in Ullrich Congenital Muscular Dystrophy. Journal of Biological Chemistry, 2010, 285, 10005-10015.	1.6	22
126	Quantification of muscle strength and motor ability in patients with Duchenne muscular dystrophy on steroid therapy. Arquivos De Neuro-Psiquiatria, 2007, 65, 245-250.	0.3	22

#	Article	IF	Citations
127	Polymorphisms of APOE and LRP Genes in Brazilian Individuals With Alzheimer Disease. Alzheimer Disease and Associated Disorders, 2008, 22, 61-65.	0.6	21
128	Transcriptional response to GAA deficiency (Pompe disease) in infantile-onset patients. Molecular Genetics and Metabolism, 2012, 106, 287-300.	0.5	20
129	CTNNB1, AXIN1 and APC expression analysis of different medulloblastoma variants. Clinics, 2013, 68, 167-172.	0.6	20
130	Selection of suitable housekeeping genes for expression analysis in glioblastoma using quantitative RT-PCR. Annals of Neurosciences, 2014, 21, 62-3.	0.9	20
131	Stem cells in neurology - current perspectives. Arquivos De Neuro-Psiquiatria, 2014, 72, 457-465.	0.3	20
132	Melatonergic systemâ€based twoâ€gene index is prognostic in human gliomas. Journal of Pineal Research, 2016, 60, 84-94.	3.4	20
133	LOXL3 Function Beyond Amino Oxidase and Role in Pathologies, Including Cancer. International Journal of Molecular Sciences, 2019, 20, 3587.	1.8	20
134	Familial Creutzfeldt-Jakob disease associated with a point mutation at codon 210 of the prion protein gene. Arquivos De Neuro-Psiquiatria, 2001, 59, 932-935.	0.3	19
135	Helicobacter pylori Seropositivity among 963 Japanese Brazilians According to Sex, Age, Generation, and Lifestyle Factors. Japanese Journal of Cancer Research, 2001, 92, 1150-1156.	1.7	19
136	Lifestyle factors associated with atrophic gastritis among Helicobacter pylori-seropositive Japanese-Brazilians in Sï¿½o Paulo. International Journal of Clinical Oncology, 2003, 8, 362-368.	1.0	19
137	Expression of cytochrome P-450 isozymes in the liver of hypophysectomized rats. Evidence for different regulation mechanisms concerning P450IIB and P450IIIA subfamilies. FEBS Journal, 1988, 177, 597-604.	0.2	18
138	A comparison of the prevalence of the metabolic syndrome and its components among native Japanese and Japanese Brazilians residing in Japan and Brazil. European Journal of Cardiovascular Prevention and Rehabilitation, 2007, 14, 508-514.	3.1	18
139	ICAM-1 (Lys469Glu) and PECAM-1 (Leu125Val) polymorphisms in diffuse astrocytomas. Clinical and Experimental Medicine, 2009, 9, 157-163.	1.9	18
140	Differential Expression of ID4 and Its Association with TP53 Mutation, SOX2, SOX4 and OCT-4 Expression Levels. PLoS ONE, 2013, 8, e61605.	1.1	18
141	Adult Neurogenesis and Glial Oncogenesis: When the Process Fails. BioMed Research International, 2014, 2014, 1-10.	0.9	18
142	A simplified approach using Taqman low-density array for medulloblastoma subgrouping. Acta Neuropathologica Communications, 2019, 7, 33.	2.4	18
143	Assessment of the 50-kDa dystrophin-associated glycoprotein in Brazilian patients with severe childhood autosomal recessive muscular dystrophy. Journal of the Neurological Sciences, 1994, 123, 122-128.	0.3	17
144	Lack of the C-terminal domain of nebulin in a patient with nemaline myopathy. Muscle and Nerve, 2002, 25, 747-752.	1.0	17

#	Article	IF	Citations
145	Bilateral olivary hypertrophy after unilateral cerebellar infarction: case report. Arquivos De Neuro-Psiquiatria, 2005, 63, 321-323.	0.3	17
146	ADAM23 methylation and expression analysis in brain tumors. Neuroscience Letters, 2005, 380, 260-264.	1.0	17
147	Intracranial and spinal ependymoma: series at Faculdade de Medicina, Universidade de SÃ£o Paulo. Arquivos De Neuro-Psiquiatria, 2009, 67, 626-632.	0.3	17
148	ASPM gene expression in medulloblastoma. Child's Nervous System, 2011, 27, 71-74.	0.6	17
149	Effects of yoga breathing exercises on pulmonary function in patients with Duchenne muscular dystrophy: an exploratory analysis. Jornal Brasileiro De Pneumologia, 2014, 40, 128-133.	0.4	17
150	Stathmin involvement in the maternal embryonic leucine zipper kinase pathway in glioblastoma. Proteome Science, 2016, 14, 6.	0.7	17
151	Detection of somatic TP53 splice site mutations in diffuse astrocytomas. Cancer Letters, 2005, 224, 321-327.	3.2	16
152	Mapping of direction and muscle representation in the human primary motor cortex controlling thumb movements. Journal of Physiology, 2009, 587, 1977-1987.	1.3	16
153	CD99 Expression in Glioblastoma Molecular Subtypes and Role in Migration and Invasion. International Journal of Molecular Sciences, 2019, 20, 1137.	1.8	16
154	Intracranial meningiomas: magnetic resonance imaging findings in 78 cases. Arquivos De Neuro-Psiquiatria, 2007, 65, 610-614.	0.3	15
155	Effects of somatosensory stimulation on the excitability of the unaffected hemisphere in chronic stroke patients. Clinics, 2008, 63, 735-740.	0.6	15
156	The association of post-stroke anhedonia with salivary cortisol levels and stroke lesion in hippocampal/parahippocampal region. Neuropsychiatric Disease and Treatment, 2015, 11, 233.	1.0	15
157	Multiple Intracranial Meningiomas: A Case Series and Review of the Literature. World Neurosurgery, 2019, 122, e1536-e1541.	0.7	15
158	Methylenetetrahydrofolate reductase gene polymorphism is not related to the risk of ischemic cerebrovascular disease in a Brazilian population. Clinics, 2007, 62, 295-300.	0.6	15
159	SELAdb: A database of exonic variants in a Brazilian population referred to a quaternary medical center in SÃ£o Paulo. Clinics, 2020, 75, e1913.	0.6	15
160	Transplacental induction of cytochromes P-450IA1 and P-450IA2 by polycyclic aromatic carcinogens: TCDD-binding protein level as the rate-limiting step. Carcinogenesis, 1988, 9, 2059-2063.	1.3	14
161	Brazilian family with pure autosomal dominant spastic paraplegia maps to 8q: Analysis of muscle beta 1 syntrophin. , 2000, 92, 122-127.		14
162	Glutaminolysis dynamics during astrocytoma progression correlates with tumor aggressiveness. Cancer & Metabolism, 2021, 9, 18.	2.4	14

#	Article	IF	Citations
163	Different behavior in the paternally vs. maternally inherited mutated allele in Brazilian Machado-Joseph (MJD1) Families. , 1998, 77, 246-248.		13
164	Rod Distribution and Muscle Fiber Type Modification in the Progression of Nemaline Myopathy. Journal of Child Neurology, 2003, 18, 235-240.	0.7	13
165	Associations of TNF-A-1031TT and -857TT genotypes with Helicobacter pylori seropositivity and gastric atrophy among Japanese Brazilians. International Journal of Clinical Oncology, 2006, 11, 140-145.	1.0	13
166	Xenograft Transplantation of Human Malignant Astrocytoma Cells Into Immunodeficient Rats: An Experimental Model of Glioblastoma. Clinics, 2010, 65, 305-309.	0.6	13
167	CXCR7 and CXCR4 Expressions in Infiltrative Astrocytomas and Their Interactions with HIF1Î± Expression and IDH1 Mutation. Pathology and Oncology Research, 2015, 21, 229-240.	0.9	13
168	N-acetylcysteine Counteracts Adipose Tissue Macrophage Infiltration and Insulin Resistance Elicited by Advanced Glycated Albumin in Healthy Rats. Frontiers in Physiology, 2017, 8, 723.	1.3	13
169	Kearns-Sayre syndrome "plus": classical clinical findings and dystonia. Arquivos De Neuro-Psiquiatria, 1999, 57, 1017-1023.	0.3	13
170	Age- and tissue-dependent expression of CYP2C23 in the rat. Biochimica Et Biophysica Acta Gene Regulatory Mechanisms, 1993, 1172, 124-130.	2.4	12
171	Ullrich congenital muscular dystrophy and bethlem myopathy: clinical and genetic heterogeneity. Arquivos De Neuro-Psiquiatria, 2005, 63, 785-790.	0.3	12
172	Community-based familial study of Helicobacter pylori infection among healthy Japanese Brazilians. Gastric Cancer, 2006, 9, 208-216.	2.7	12
173	Significant association between PTPN11 polymorphism and gastric atrophy among Japanese Brazilians. Gastric Cancer, 2006, 9, 277-283.	2.7	12
174	Translational Neurorehabilitation Research in the Third World. Stroke, 2014, 45, 1495-1497.	1.0	12
175	A coordinated approach for the assessment of molecular subgroups in pediatric ependymomas using low-cost methods. Journal of Molecular Medicine, 2021, 99, 1101-1113.	1.7	12
176	Extracellular Matrix Proteome Remodeling in Human Glioblastoma and Medulloblastoma. Journal of Proteome Research, 2021, 20, 4693-4707.	1.8	12
177	Skeletal muscle major histocompatibility complex class I and II expression differences in adult and juvenile dermatomyositis. Clinics, 2012, 67, 885-890.	0.6	12
178	Helicobacter pylori seropositivity and IL-1B C-31T polymorphism among Japanese Brazilians. International Journal of Molecular Medicine, 2002, 10, 321.	1.8	11
179	Association of Lewis and Secretor gene polymorphisms and Helicobacter pylori seropositivity among Japanese-Brazilians. Journal of Gastroenterology, 2004, 39, 717-723.	2.3	11
180	Pituitary volume and the effects of phototherapy in patients with seasonal winter depression: a controlled study. Revista Brasileira De Psiquiatria, 2008, 30, 50-54.	0.9	11

#	Article	IF	Citations
181	Low-grade astrocytoma: surgical outcomes in eloquent versus non-eloquent brain areas. Arquivos De Neuro-Psiquiatria, 2013, 71, 31-34.	0.3	11
182	CD99 is upregulated in placenta and astrocytomas with a differential subcellular distribution according to the malignancy stage. Journal of Neuro-Oncology, 2014, 119, 59-70.	1.4	11
183	Morphometric analyses of normal pediatric brachial biceps and quadriceps muscle tissue. Histology and Histopathology, 2013, 28, 525-30.	0.5	11
184	Isolation and characterization of novel RECK tumor suppressor gene splice variants. Oncotarget, 2015, 6, 33120-33133.	0.8	11
185	Lower HDL-cholesterol among healthy middle-aged Japanese-Brazilians in SÃ£o Paulo compared to Natives and Japanese-Brazilians in Japan. European Journal of Epidemiology, 2007, 22, 33-42.	2.5	10
186	Alcohol Use Disorder is Associated with Upregulation of MicroRNAâ€34a and MicroRNAâ€34c in Hippocampal Postmortem Tissue. Alcoholism: Clinical and Experimental Research, 2021, 45, 64-68.	1.4	10
187	The effects of postmortem delay on mouse and human microglia gene expression. Glia, 2021, 69, 1053-1060.	2.5	10
188	Late p65 nuclear translocation in glioblastoma cells indicates non-canonical TLR4 signaling and activation of DNA repair genes. Scientific Reports, 2021, 11, 1333.	1.6	10
189	Dystrophin-glycoproteins associated in congenital muscular dystrophy: immunohistochemical analysis of 59 Brazilian cases. Arquivos De Neuro-Psiquiatria, 2005, 63, 791-800.	0.3	10
190	Extraneural metastases in medulloblastoma. Arquivos De Neuro-Psiquiatria, 2011, 69, 328-331.	0.3	10
191	Gene Structure and Promoter Analysis of the Rat ConstitutiveCYP2C23Gene. DNA and Cell Biology, 1995, 14, 777-788.	0.9	9
192	Different origins of mutations at the Machado-Joseph locus (MJD1). Journal of Medical Genetics, 1996, 33, 439-439.	1.5	9
193	Heterogeneity of Classic Congenital Muscular Dystrophy With Involvement of the Central Nervous System: Report of Five Atypical Cases. Journal of Child Neurology, 2000, 15, 172-178.	0.7	9
194	Confiabilidade de medidas volumÃ©tricas de estruturas temporais mesiais. Arquivos De Neuro-Psiquiatria, 2002, 60, 420-428.	0.3	9
195	Activation of EGFR Signaling from Pilocytic Astrocytomas to Glioblastomas. International Journal of Biological Markers, 2014, 29, 120-128.	0.7	9
196	Immunohistochemical expression of cyclin <scp>D1</scp> is higher in supratentorial ependymomas and predicts relapses in gross total resection cases. Neuropathology, 2015, 35, 312-323.	0.7	9
197	Treatment of unilateral spatial neglect after stroke using transcranial direct current stimulation (ELETRON trial): study protocol for a randomized controlled trial. Trials, 2016, 17, 479.	0.7	9
198	Schwartz-jampel syndrome: report of five cases. Arquivos De Neuro-Psiquiatria, 2002, 60, 734-738.	0.3	9

#	Article	IF	Citations
199	Phenobarbital-inducible gene expression in developing rat liver: relationship to hepatocyte function. Biochimica Et Biophysica Acta Gene Regulatory Mechanisms, 1989, 1009, 221-228.	2.4	8
200	King-Denborough Syndrome: report of two Brazilian cases. Arquivos De Neuro-Psiquiatria, 2002, 60, 739-741.	0.3	8
201	Basilar artery occlusive disease in stroke survivors in a multiethnic population. Clinical Neurology and Neurosurgery, 2010, 112, 233-236.	0.6	8
202	A Brazilian family with inclusion body myopathy associated with Pagetâ€™s disease of bone and frontotemporal dementia linked to the VCP pGly97Glu mutation. Clinical Rheumatology, 2018, 37, 1129-1136.	1.0	8
203	Plasmatic membrane toll-like receptor expressions in human astrocytomas. PLoS ONE, 2018, 13, e0199211.	1.1	8
204	Exercise Training Attenuates Ubiquitin-Proteasome Pathway and Increases the Genes Related to Autophagy on the Skeletal Muscle of Patients With Inflammatory Myopathies. Journal of Clinical Rheumatology, 2021, 27, S224-S231.	0.5	8
205	Histopathological findings in skeletal muscle used in human dynamic cardiomyoplasty. Journal of Pathology, 2001, 194, 116-121.	2.1	7
206	Rasmussen Encephalitis Associated With Segmental Vitiligo of the Scalp: Clinicopathologic Report. Journal of Child Neurology, 2001, 16, 374-377.	0.7	7
207	Treatment of subclavian steal syndrome with percutaneous transluminal angioplasty and stenting: case report. Arquivos De Neuro-Psiquiatria, 2003, 61, 95-99.	0.3	7
208	Clinical presentation, treatment and outcome of patients with cerebral metastases: the University of SÃ£o Paulo series. Arquivos De Neuro-Psiquiatria, 2004, 62, 808-814.	0.3	7
209	â€œSalt and Pepperâ€•in the Eye and Face: A Prelude to Brainstem Ischemia. American Journal of Ophthalmology, 2007, 144, 322-325.	1.7	7
210	Experimental nodel of C6 brain tumors in athymic rats. Arquivos De Neuro-Psiquiatria, 2008, 66, 238-241.	0.3	7
211	Current perspectives in stem cell therapy for spinal cord repair in humans: a review of work from the past 10 years. Arquivos De Neuro-Psiquiatria, 2014, 72, 451-456.	0.3	7
212	Urinary Sediment Transcriptomic and Longitudinal Data to Investigate Renal Function Decline in Type 1 Diabetes. Frontiers in Endocrinology, 2020, 11, 238.	1.5	7
213	LOXL3 Silencing Affected Cell Adhesion and Invasion in U87MG Glioma Cells. International Journal of Molecular Sciences, 2021, 22, 8072.	1.8	7
214	Exclusion of the 15q locus as a candidate gene for severe childhood autosomal recessive Duchenne-like muscular dystrophy in Brazilian families. Human Molecular Genetics, 1993, 2, 201-202.	1.4	6
215	Migraine and motion sickness independently contribute to visual discomfort. Cephalalgia, 2010, 30, 161-169.	1.8	6
216	Quantitative proteomic analysis and functional studies reveal that nucleophosmin is involved in cell death in glioblastoma cell line transfected with siRNA. Proteomics, 2012, 12, 2632-2640.	1.3	6

#	Article	IF	Citations
217	A game of hide and seek: Is it possible to recruit more patients for NIBS studies in stroke?. Journal of the Neurological Sciences, 2015, 358, 472-474.	0.3	6
218	Atypical and Malignant Meningiomas: Neurooncologic Management in a BrazilianÂcohort. World Neurosurgery, 2018, 110, e20-e23.	0.7	6
219	Identification ofÂITPR1Âas a Hub Gene of Group 3 Medulloblastoma and Coregulated Genes with Potential Prognostic Values. Journal of Molecular Neuroscience, 2022, 72, 633-641.	1.1	6
220	Serum interleukin-17A level is associated with disease activity of adult patients with dermatomyositis and polymyositis. Clinical and Experimental Rheumatology, 2019, 37, 656-662.	0.4	6
221	Clinical and molecular analysis of spinal muscular atrophy in Brazilian patients. Genetics and Molecular Biology, 1999, 22, 487-492.	0.6	5
222	Neuroinflammatory responses to traumatic brain injury. Neuropsychiatric Disease and Treatment, 2015, 11, 773.	1.0	5
223	CXCR7, CXCR4, and Their Ligand Expression Profile in Traumatic Brain Injury. World Neurosurgery, 2021, 147, e16-e24.	0.7	5
224	Transcriptional profiling of macaque microglia reveals an evolutionary preserved gene expression program. Brain, Behavior, & Immunity - Health, 2021, 15, 100265.	1.3	5
225	Phenotypic and immunohistochemical characterization of sarcoglycanopathies. Clinics, 2011, 66, 1713-9.	0.6	5
226	Does previous corticosteroid treatment affect the inflammatory infiltrate found in polymyositis muscle biopsies?. Clinical and Experimental Rheumatology, 2015, 33, 310-4.	0.4	5
227	Cerebral microbleeds and intravenous thrombolysis: case report. Arquivos De Neuro-Psiquiatria, 2006, 64, 855-857.	0.3	4
228	Screening for MELAS mutations in young patients with stroke of undetermined origin. Arquivos De Neuro-Psiquiatria, 2007, 65, 371-376.	0.3	4
229	Muscle Fiber Type Composition, Fiber Diameter, Capillary Density in Temporalis and Masseter Muscles and Correlation with Bite Force. International Journal of Morphology, 2013, 31, 747-753.	0.1	4
230	Angiotensin-converting enzyme insertion/deletion gene polymorphism is associated with dermatomyositis. JRAAS - Journal of the Renin-Angiotensin-Aldosterone System, 2015, 16, 666-671.	1.0	4
231	Correlation between molecular features and genetic subtypes of Glioblastoma: critical analysis in 109 cases. Medical Express, 2017, 4, .	0.2	4
232	Distinct response to GDF15 knockdown in pediatric and adult glioblastoma cell lines. Journal of Neuro-Oncology, 2018, 139, 51-60.	1.4	4
233	The expression of the aminoacid transporters ASCT2 (SLC1A5) and LAT1 (SLC7A5) in astrocytomas. Medical Express, 2016, 3, .	0.2	4
234	The chromatin remodeler complex ATRX-DAXX-H3.3 and telomere length in meningiomas. Clinical Neurology and Neurosurgery, 2021, 210, 106962.	0.6	4

#	Article	IF	Citations
235	Facial sensory symptoms in medullary infarcts. Arquivos De Neuro-Psiquiatria, 2005, 63, 947-950.	0.3	4
236	Distribution of leptin receptors in the brain stem: possible route in the pathophysiology of neuromuscular control of airway resistance during sleep. Sleep Medicine, 2022, 93, 56-62.	0.8	4
237	Activation of Hedgehog signaling by the oncogenic RELA fusion reveals a primary cilia-dependent vulnerability in supratentorial ependymoma. Neuro-Oncology, 2023, 25, 185-198.	0.6	4
238	Homozygotic intronic GAA mutation in three siblings with late-onset Pompe's disease. Arquivos De Neuro-Psiquiatria, 2010, 68, 194-197.	0.3	3
239	Comparison between treatment naive juvenile and adult dermatomyositis muscle biopsies: difference of inflammatory cells phenotyping. Advances in Rheumatology, 2018, 58, 37.	0.8	3
240	A novel type of C11orf95-LOC-RELA fusion in a grade II supratentorial ependymoma: report of a case with literature review. Child's Nervous System, 2019, 35, 689-694.	0.6	3
241	Cyclin E1 expression and malignancy in meningiomas. Clinical Neurology and Neurosurgery, 2020, 190, 105647.	0.6	3
242	Report of the first Brazilian infantile Pompe disease patient to be treated with recombinant human acid alpha-glucosidase. Jornal De Pediatria, 2008, 84, 272-275.	0.9	3
243	Relevant coexpression of STMN1, MELK and FOXM1 in glioblastoma and review of the impact of STMN1 in cancer biology. Medical Express, 2017, 4, .	0.2	3
244	Magnetic Resonance Imaging of Wallerian Degeneration in Stroke. Archives of Neurology, 2003, 60, 1466.	4.9	2
245	Possible association of interleukin 1B C-31T polymorphism among Helicobacter pylori seropositive Japanese Brazilians with susceptibility to atrophic gastritis. International Journal of Molecular Medicine, 2004, 14, 421.	1.8	2
246	Comparing Methods for Determining Motor-Hand Lateralization Based on fTCD Signals. Journal of Medical Systems, 2015, 39, 4.	2.2	2
247	Cyclooxygenase-2 gene polymorphisms and susceptibility to colorectal cancer in a Brazilian population. Journal of Gastrointestinal Oncology, 2017, 8, 629-635.	0.6	2
248	The impact of interleukin-13 receptor expressions in cell migration of astrocytomas. Medical Express, 2015, 2, .	0.2	2
249	The effect of prior corticosteroid use in muscle biopsies from patients with dermatomyositis. Clinical and Experimental Rheumatology, 2015, 33, 336-40.	0.4	2
250	Cellular Model of Malignant Transformation of Primary Human Astrocytes Induced by Deadhesion/Readhesion Cycles. International Journal of Molecular Sciences, 2022, 23, 4471.	1.8	2
251	Factors associated with serum CA19-9 levels among healthy children: a cross-sectional study. BMC Clinical Pathology, 2012, 12, 23.	1.8	1
252	Interventions to Enhance Adaptive Plasticity after Stroke: From Mechanisms to Therapeutic Perspectives. Neural Plasticity, 2016, 2016, 1-2.	1.0	1

#	Article	IF	Citations
253	ATRX-DAXX Complex Expression Levels and Telomere Length in Normal Young and Elder Autopsy Human Brains. DNA and Cell Biology, 2019, 38, 955-961.	0.9	1
254	The facioscapulohumeral muscular dystrophy (FSHD1) gene affects males more severely and more frequently than females. , 1998, 77, 155.		1
255	The prevalence of mitochondrial DNA mutations in Leigh syndrome in a Brazilian series. Medical Express, 2014, 1, .	0.2	1
256	Stroke: an ongoing revolution. Arquivos De Neuro-Psiquiatria, 2015, 73, 892-893.	0.3	1
257	Pituitary size in seasonal affective disorder: A controlled study. Biological Psychiatry, 1994, 35, 669.	0.7	0
258	Sensitive and quantitative chimerism analysis in all patients with real-time PCR. Biology of Blood and Marrow Transplantation, 2005, 11, 42.	2.0	0
259	T.P.1 06 Improvement of muscle mass after injection of AAV vectors expressing either myostatin shRNA or activin receptor IIb shRNA. Neuromuscular Disorders, 2006, 16, 686.	0.3	0
260	G.P.7.08 FRG1 over-expression induces severe dystrophic phenotype in mouse. Neuromuscular Disorders, 2007, 17, 808.	0.3	0
261	Dermatomiosite juvenil e linfoma de Hodgkin: uma rara associaÃ§Ã£o. Revista Brasileira De Reumatologia, 2007, 47, 458-462.	0.8	0
262	Immunoglobulins Profile in Juvenile-onset Systemic Lupus Erythematosus (JSLE). Journal of Allergy and Clinical Immunology, 2008, 121, S226-S226.	1.5	0
263	G.P.8.07 Increased oxidative metabolism in mdx muscle treated by a combination of exon skipping and myostatin blockade. Neuromuscular Disorders, 2008, 18, 783-784.	0.3	0
264	P4.52 Transcriptional response to GAA deficiency in mice and humans. Neuromuscular Disorders, 2010, 20, 674.	0.3	0
265	Transcriptional Response to GAA Deficiency in Mice and Humans. Clinical Therapeutics, 2011, 33, S25.	1.1	0
266	66 Proffered Paper: PTEN Phosphorylation by Fibroblast Growth Factor Receptors and SRC Mediates Resistance to Epidermal Growth Factor Receptor Inhibitors in Glioblastoma. European Journal of Cancer, 2012, 48, S16.	1.3	0
267	Temporomandibular Joint Magnetic Resonance Imaging Analysis in Adults with Steinert's Myotonic Dystrophy. International Journal of Morphology, 2013, 31, 301-306.	0.1	0
268	MP66-19 TARGETED RESEQUENCING FOUND POLYMORPHISMS IN COMMON METABOLIC PATHWAYS INVOLVED IN MALE INFERTILITY. Journal of Urology, 2014, 191, .	0.2	0
269	The study of central nervous system involvement in mithochondrial disorders with mitochondrial DNA mutation. Arquivos De Neuro-Psiquiatria, 2002, 60, 173-173.	0.3	0
270	Abstract 2354: Characterization of three novel splice variants of the RECK tumor and metastasis suppressor gene. , 2011, , .		0

#	Article	IF	Citations
271	Referred Dental Pain, an Analysis of their Prevalence and Clinical Implication. International Journal of Odontostomatology, 2012, 6, 169-173.	0.0	0
272	Abstract 4845: VEGF-A and Cox-2 gene polymorphisms and susceptibility to colorectal cancer in Brazilian population.. , 2013, , .		0
273	Pompe disease: further challenges to pursue. Arquivos De Neuro-Psiquiatria, 2013, 71, 273-274.	0.3	0
274	Abstract B134: Stathmin is involved in the maternal embryonic leucine zipper kinase pathway in human astrocytomas.. , 2013, , .		0
275	Abstract 4607: Stathmin is involved in the maternal embryonic leucine zipper kinase pathway and impacts in the outcome of glioblastoma. , 2014, , .		0
276	Abstract 5594: Cyclin D1 expression correlates with supratentorial location of ependymomas. , 2014, , .		0
277	Abstract 3047: Mitochondrial DNA copy variation and TFAM expression in astrocytoma. , 2015, , .		0
278	Abstract 66: CD99 functional analysis in glioblastoma by RNAseq. , 2015, , .		0
279	Challenges to understand the mitochondrial respiratory diseases caused by mitochondrial DNA mutations. Arquivos De Neuro-Psiquiatria, 2015, 73, 897-898.	0.3	0
280	Do cyclin e levels correlate with recurrence in meningioma? Results from an observational study.. Journal of Clinical Oncology, 2016, 34, e23123-e23123.	0.8	0
281	Abstract 3229: Microglia/macrophages activation status in diffuse gliomas. , 2016, , .		0
282	Abstract 1458: Toll-like receptors 1, 2, 4 and 6 expression levels in diffusely infiltrative astrocytomas. , 2016, , .		0
283	The expression of gene ANKRD1 correlates with hypoxia status in muscle biopsies of treatment-nÃive adult dermatomyositis. Medical Express, 2017, 4, .	0.2	0
284	Cerebrospinal Fluid. , 2017, , .		0
285	Abstract 899: CD99 plays an important role in glioblastoma cell migration. , 2017, , .		0
286	Abstract 2958: Transcriptome analysis of astrocytomaversusnon-neoplastic human microglia. , 2017, , .		0
287	Abstract 1498: Melatonin inhibits mitochondrial transcription factor A expression in glioblastoma U87MG cell culture inducing an anti-tumorigenic effect. , 2017, , .		0
288	Mitochondriogenesis and brain aging. , 2018, 97, 604-606.	0.0	0

#	Article	IF	Citations
289	Analysis of the use of photosensitization in human glioblastoma multiforme to induce cell death. , 2019, 98, 25-25.	0.0	0