Roberta Morosetti

List of Publications by Year in descending order

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49 papers	2,632 citations	236833 25 h-index	243529 44 g-index
49 all docs	49 docs citations	49 times ranked	3759 citing authors

#	Article	IF	Citations
1	Role of Favorable Perfusion Imaging in Predicting the Outcome of Patients with Acute Ischemic Stroke due to Large Vessel Occlusion Undergoing Effective Thrombectomy: A Single-Center Study. Cerebrovascular Diseases Extra, 2021, 11, 1-8.	0.5	2
2	Teaching NeuroImages: A cutaneous vascular malformation hides giant cerebral aneurysms. Neurology, 2018, 90, e1362-e1363.	1.5	0
3	Dysphagia and Obstructive Sleep Apnea in Acute, First-Ever, Ischemic Stroke. Journal of Stroke and Cerebrovascular Diseases, 2018, 27, 539-546.	0.7	10
4	Thrombus in Transit. Neurologist, 2017, 22, 21-23.	0.4	0
5	Novel SEC61Gâ€“EGFR Fusion Gene in Pediatric Ependymomas Discovered by Clonal Expansion of Stem Cells in Absence of Exogenous Mitogens. Cancer Research, 2017, 77, 5860-5872.	0.4	21
6	Estrogens enhance myoblast differentiation in facioscapulohumeral muscular dystrophy by antagonizing DUX4 activity. Journal of Clinical Investigation, 2017, 127, 1531-1545.	3.9	46
7	Thalamic Amnesia Mimicking Transient Global Amnesia. Neurologist, 2015, 19, 149-152.	0.4	5
8	Neurofibromatosis Type 1 Associated with Vertebrobasilar Dolichoectasia and Pontine Ischemic Stroke. Journal of Neuroimaging, 2015, 25, 505-506.	1.0	13
9	Risk Factor and Etiology Analysis of Ischemic Stroke in Young Adult Patients. Journal of Stroke and Cerebrovascular Diseases, 2014, 23, e221-e227.	0.7	69
10	Spontaneous sternocleidomastoid muscle hematoma following thrombolysis for acute ischemic stroke. Journal of the Neurological Sciences, 2014, 341, 189-190.	0.3	2
11	Sleep disordered breathing in a cohort of patients with sporadic inclusion body myositis. Clinical Neurophysiology, 2013, 124, 1615-1621.	0.7	13
12	Sleep Modifications in Acute Transient Global Amnesia. Journal of Clinical Sleep Medicine, 2013, 09, 921-927.	1.4	3
13	TWEAK in Inclusion-Body Myositis Muscle. American Journal of Pathology, 2012, 180, 1603-1613.	1.9	30
14	The recovery of platelet cyclooxygenase activity explains interindividual variability in responsiveness to lowâ€dose aspirin in patients with and without diabetes. Journal of Thrombosis and Haemostasis, 2012, 10, 1220-1230.	1.9	211
15	Bilateral thalamic stroke transiently reduces arousals and NREM sleep instability. Journal of the Neurological Sciences, 2011, 300, 151-154.	0.3	33
16	Mesoangioblasts from Facioscapulohumeral Muscular Dystrophy Display in Vivo a Variable Myogenic Ability Predictable by their in Vitro Behavior. Cell Transplantation, 2011, 20, 1299-1313.	1.2	26
17	Vertebral Artery Dissection Presenting With Isolated Occipital Headache. Headache, 2010, 50, 1378-1380.	1.8	4
18	Pseudoperipheral tongue weakness. Journal of Neurology, Neurosurgery and Psychiatry, 2010, 81, 1024-1025.	0.9	0

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19	Analysis of NCAM helps identify unusual phenotypes of hereditary inclusion-body myopathy. Neurology, 2010, 75, 265-272.	1.5	28
20	The Stolen Memory: A Case of Transient Global Amnesia. Biological Psychiatry, 2010, 67, e31-e32.	0.7	1
21	Increased aging in primary muscle cultures of sporadic inclusion-body myositis. Neurobiology of Aging, 2010, 31, 1205-1214.	1.5	35
22	Vessel-associated stem cells from skeletal muscle: From biology to future uses in cell therapy. World Journal of Stem Cells, 2010, 2, 39.	1.3	15
23	Hereditary inclusionâ€body myopathy: Clues on pathogenesis and possible therapy. Muscle and Nerve, 2009, 40, 340-349.	1.0	26
24	Hyposialylation of neprilysin possibly affects its expression and enzymatic activity in hereditary inclusionâ€body myopathy muscle. Journal of Neurochemistry, 2008, 105, 971-981.	2.1	53
25	The ER-Bound RING Finger Protein 5 (RNF5/RMA1) Causes Degenerative Myopathy in Transgenic Mice and Is Deregulated in Inclusion Body Myositis. PLoS ONE, 2008, 3, e1609.	1.1	57
26	Isolation and Characterization of Mesoangioblasts from Facioscapulohumeral Muscular Dystrophy Muscle Biopsies. Stem Cells, 2007, 25, 3173-3182.	1.4	37
27	Pericytes of human skeletal muscle are myogenic precursors distinct from satellite cells. Nature Cell Biology, 2007, 9, 255-267.	4.6	899
28	The effect of disease activity on leptin, leptin receptor and suppressor of cytokine signalling-3 expression in relapsingâ€“remitting multiple sclerosis. Journal of Neuroimmunology, 2007, 192, 174-183.	1.1	74
29	Neprilysin participates in skeletal muscle regeneration and is accumulated in abnormal muscle fibres of inclusion body myositis. Journal of Neurochemistry, 2006, 96, 777-789.	2.1	35
30	NCAM is hyposialylated in hereditary inclusion body myopathy due to GNE mutations. Neurology, 2006, 66, 755-758.	1.5	66
31	MyoD expression restores defective myogenic differentiation of human mesoangioblasts from inclusion-body myositis muscle. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 16995-17000.	3.3	75
32	Î±-Dystroglycan does not play a major pathogenic role in autosomal recessive hereditary inclusion-body myopathy. Neuromuscular Disorders, 2005, 15, 177-184.	0.3	39
33	Vitamin D and Hematological Malignancy. , 2005, , 1727-1740.		4
34	The Cellular Response to PPARÎ³ Ligands Is Related to the Phenotype of Neuroblastoma Cell Lines. Oncology Research, 2004, 14, 345-354.	0.6	20
35	Cyclooxygenase-1, but not -2, is upregulated in NB4 leukemic cells and human primary promyelocytic blasts during differentiation. Leukemia, 2004, 18, 1373-1379.	3.3	20
36	The PPARgamma ligands PGJ2 and rosiglitazone show a differential ability to inhibit proliferation and to induce apoptosis and differentiation of human glioblastoma cell lines. International Journal of Oncology, 2004, 25, 493-502.	1.4	25

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37	Progression of myelodysplastic syndrome: allelic loss on chromosomal arm 1p. British Journal of Haematology, 2003, 122, 226-230.	1.2	36
38	Therapy Related Leukemias: Susceptibility, Prevention and Treatment. Leukemia and Lymphoma, 2001, 41, 255-276.	0.6	115
39	High cyclin-dependent kinase inhibitors in Bcl-2 and Bcl-xL -expressing CD34+ -proliferating haematopoietic progenitors. British Journal of Haematology, 2000, 110, 654-662.	1.2	16
40	Expression of cyclin-dependent kinase inhibitor p15INK4B during normal and leukemic myeloid differentiation. Experimental Hematology, 2000, 28, 519-526.	0.2	37
41	Granulocyte colony-stimulating factor perturbs lymphocyte mitochondrial function and inhibits cell cycle progression. Experimental Hematology, 2000, 28, 612-625.	0.2	32
42	Chromosome Band 1p36 Contains a Putative Tumor Suppressor Gene Important in the Evolution of Chronic Myelocytic Leukemia. Blood, 1998, 92, 3405-3409.	0.6	70
43	Chromosome Band 1p36 Contains a Putative Tumor Suppressor Gene Important in the Evolution of Chronic Myelocytic Leukemia. Blood, 1998, 92, 3405-3409.	0.6	3
44	Is Kaposi's Sarcomaâ€“Associated Herpesvirus Ubiquitous in Urogenital and Prostate Tissues?. Blood, 1997, 89, 1686-1689.	0.6	39
45	Allelotype Analysis in the Evolution of Chronic Myelocytic Leukemia. Blood, 1997, 90, 2010-2014.	0.6	45
46	A Novel, Myeloid Transcription Factor, C/EBPÎµ, Is Upregulated During Granulocytic, But Not Monocytic, Differentiation. Blood, 1997, 90, 2591-2600.	0.6	177
47	Analysis of p18INK4C in adult T-cell leukaemia and non-Hodgkin's lymphoma. British Journal of Haematology, 1997, 99, 668-670.	1.2	21
48	Integrity of the 1,25-dihydroxyvitamin D3 receptor in bone, lung, and other cancers. , 1997, 19, 254-257.		29
49	Effect of Sodium on the Energetics of Thrombin â€“ Thrombomodulin Interaction and its Relevance for Protein C Hydrolysis. Journal of Molecular Biology, 1996, 258, 190-200.	2.0	15