Roberta Morosetti

List of Publications by Year in descending order

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Version: 2024-02-01

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	Pericytes of human skeletal muscle are myogenic precursors distinct from satellite cells. Nature Cell Biology, 2007, 9, 255-267.	4.6	899
2	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
3	A Novel, Myeloid Transcription Factor, C/EBPÎµ, Is Upregulated During Granulocytic, But Not Monocytic, Differentiation. Blood, 1997, 90, 2591-2600.	0.6	177
4	Therapy Related Leukemias: Susceptibility, Prevention and Treatment. Leukemia and Lymphoma, 2001, 41, 255-276.	0.6	115
5	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
6	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
7	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
8	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
9	NCAM is hyposialylated in hereditary inclusion body myopathy due to GNE mutations. Neurology, 2006, 66, 755-758.	1.5	66
10	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
11	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
12	Estrogens enhance myoblast differentiation in facioscapulohumeral muscular dystrophy by antagonizing DUX4 activity. Journal of Clinical Investigation, 2017, 127, 1531-1545.	3.9	46
13	Allelotype Analysis in the Evolution of Chronic Myelocytic Leukemia. Blood, 1997, 90, 2010-2014.	0.6	45
14	Is Kaposi's Sarcomaâ€“Associated Herpesvirus Ubiquitous in Urogenital and Prostate Tissues?. Blood, 1997, 89, 1686-1689.	0.6	39
15	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
16	Expression of cyclin-dependent kinase inhibitor p15INK4B during normal and leukemic myeloid differentiation. Experimental Hematology, 2000, 28, 519-526.	0.2	37
17	Isolation and Characterization of Mesoangioblasts from Facioscapulohumeral Muscular Dystrophy Muscle Biopsies. Stem Cells, 2007, 25, 3173-3182.	1.4	37
18	Progression of myelodysplastic syndrome: allelic loss on chromosomal arm 1p. British Journal of Haematology, 2003, 122, 226-230.	1.2	36

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19	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
20	Increased aging in primary muscle cultures of sporadic inclusion-body myositis. Neurobiology of Aging, 2010, 31, 1205-1214.	1.5	35
21	Bilateral thalamic stroke transiently reduces arousals and NREM sleep instability. Journal of the Neurological Sciences, 2011, 300, 151-154.	0.3	33
22	Granulocyte colony-stimulating factor perturbs lymphocyte mitochondrial function and inhibits cell cycle progression. Experimental Hematology, 2000, 28, 612-625.	0.2	32
23	TWEAK in Inclusion-Body Myositis Muscle. American Journal of Pathology, 2012, 180, 1603-1613.	1.9	30
24	Integrity of the 1,25-dihydroxyvitamin D3 receptor in bone, lung, and other cancers. , 1997, 19, 254-257.		29
25	Analysis of NCAM helps identify unusual phenotypes of hereditary inclusion-body myopathy. Neurology, 2010, 75, 265-272.	1.5	28
26	Hereditary inclusionâ€body myopathy: Clues on pathogenesis and possible therapy. Muscle and Nerve, 2009, 40, 340-349.	1.0	26
27	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
28	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
29	Analysis of p18INK4C in adult T-cell leukaemia and non-Hodgkin's lymphoma. British Journal of Haematology, 1997, 99, 668-670.	1.2	21
30	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
31	The Cellular Response to PPARÎ³ Ligands Is Related to the Phenotype of Neuroblastoma Cell Lines. Oncology Research, 2004, 14, 345-354.	0.6	20
32	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
33	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
34	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
35	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
36	Sleep disordered breathing in a cohort of patients with sporadic inclusion body myositis. Clinical Neurophysiology, 2013, 124, 1615-1621.	0.7	13

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37	Neurofibromatosis Type 1 Associated with Vertebrobasilar Dolichoectasia and Pontine Ischemic Stroke. Journal of Neuroimaging, 2015, 25, 505-506.	1.0	13
38	Dysphagia and Obstructive Sleep Apnea in Acute, First-Ever, Ischemic Stroke. Journal of Stroke and Cerebrovascular Diseases, 2018, 27, 539-546.	0.7	10
39	Thalamic Amnesia Mimicking Transient Global Amnesia. Neurologist, 2015, 19, 149-152.	0.4	5
40	Vertebral Artery Dissection Presenting With Isolated Occipital Headache. Headache, 2010, 50, 1378-1380.	1.8	4
41	Vitamin D and Hematological Malignancy. , 2005, , 1727-1740.		4
42	Sleep Modifications in Acute Transient Global Amnesia. Journal of Clinical Sleep Medicine, 2013, 09, 921-927.	1.4	3
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	Spontaneous sternocleidomastoid muscle hematoma following thrombolysis for acute ischemic stroke. Journal of the Neurological Sciences, 2014, 341, 189-190.	0.3	2
45	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
46	The Stolen Memory: A Case of Transient Global Amnesia. Biological Psychiatry, 2010, 67, e31-e32.	0.7	1
47	Pseudoperipheral tongue weakness. Journal of Neurology, Neurosurgery and Psychiatry, 2010, 81, 1024-1025.	0.9	0
48	Thrombus in Transit. Neurologist, 2017, 22, 21-23.	0.4	0
49	Teaching NeuroImages: A cutaneous vascular malformation hides giant cerebral aneurysms. Neurology, 2018, 90, e1362-e1363.	1.5	0