Michel Simonneau

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7591839/publications.pdf

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47 papers

2,459 citations

304602 22 h-index 214721 47 g-index

48 all docs 48 docs citations

48 times ranked

 $\begin{array}{c} 3429 \\ \text{citing authors} \end{array}$

#	Article	IF	CITATIONS
1	Increased expression of BDNF mRNA in the frontal cortex of autistic patients. Behavioural Brain Research, 2019, 359, 903-909.	1.2	11
2	Translational research identifies a metabolism pathway involved in first-episode of schizophrenia: Towards precision medicine. EBioMedicine, 2019, 46, 19-20.	2.7	0
3	Looking beyond the usual suspects: sulfide stress in schizophrenia pathophysiology. EMBO Molecular Medicine, 2019, 11, e10983.	3.3	2
4	Netrin G1: its downregulation in the nucleus accumbens of cocaineâ€conditioned mice and genetic association in human cocaine dependence. Addiction Biology, 2018, 23, 448-460.	1.4	3
5	Single particle tracking of fluorescent nanodiamonds in cells and organisms. Current Opinion in Solid State and Materials Science, 2017, 21, 35-42.	5.6	56
6	Fluorescent nanodiamond tracking reveals intraneuronal transport abnormalities induced by brain-disease-related genetic risk factors. Nature Nanotechnology, 2017, 12, 322-328.	15.6	111
7	Somatostatin-IRES-Cre Mice: Between Knockout and Wild-Type?. Frontiers in Endocrinology, 2017, 8, 131.	1.5	26
8	Single KTP nanocrystals as second-harmonic generation biolabels in cortical neurons. Nanoscale, 2013, 5, 8466.	2.8	37
9	Genetics of dopamine receptors and drug addiction. Human Genetics, 2012, 131, 803-822.	1.8	93
10	Altered axonal targeting and short-term plasticity in the hippocampus of Disc1 mutant mice. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, E1349-58.	3.3	100
11	Association of DISC1 gene with schizophrenia in families from two distinct French and Algerian populations. Psychiatric Genetics, 2010, 20, 298-303.	0.6	16
12	SMARCA2 and other genome-wide supported schizophrenia-associated genes: regulation by REST/NRSF, network organization and primate-specific evolution. Human Molecular Genetics, 2010, 19, 2841-2857.	1.4	78
13	DYRK1A interacts with the REST/NRSF-SWI/SNF chromatin remodelling complex to deregulate gene clusters involved in the neuronal phenotypic traits of Down syndrome. Human Molecular Genetics, 2009, 18, 1405-1414.	1.4	128
14	Convergent evidence identifying MAP/microtubule affinity-regulating kinase 1 (MARK1) as a susceptibility gene for autism. Human Molecular Genetics, 2008, 17, 2541-2551.	1.4	78
15	Nrxn3 upregulation in the globus pallidus of mice developing cocaine addiction. NeuroReport, 2008, 19, 751-755.	0.6	30
16	Polymorphisms of coding trinucleotide repeats of homeogenes in neurodevelopmental psychiatric disorders. Psychiatric Genetics, 2008, 18, 295-301.	0.6	19
17	The fragile X mental retardation protein is a molecular adaptor between the neurospecific KIF3C kinesin and dendritic RNA granules. Human Molecular Genetics, 2007, 16, 3047-3058.	1.4	119
18	Nrsf silencing induces molecular and subcellular changes linked to neuronal plasticity. NeuroReport, 2007, 18, 441-446.	0.6	17

#	Article	IF	CITATIONS
19	Expression pattern of NOGO and NgR genes during human development. Gene Expression Patterns, 2005, 5, 561-568.	0.3	25
20	Molecular cloning and expression pattern of the Fkbp25 gene during cerebral cortical neurogenesis. Gene Expression Patterns, 2005, 5, 577-585.	0.3	6
21	Genetics and Early Disturbances of Breathing Control: The Genetics of Childhood Disease and Development: A Series of Review Articles. Pediatric Research, 2004, 55, 729-733.	1.1	64
22	Identification of a novel brain-specific and reelin-regulated gene that encodes a protein colocalized with synapsin. European Journal of Neuroscience, 2004, 20, 603-610.	1.2	17
23	Survival motor neuron SMN1 and SMN2 gene promoters: identical sequences and differential expression in neurons and non-neuronal cells. European Journal of Human Genetics, 2004, 12, 729-737.	1.4	37
24	Expression of the RSK2 gene during early human development. Gene Expression Patterns, 2004, 4, 111-114.	0.3	6
25	Modulation of the respiratory rhythm generator by the pontine noradrenergic A5 and A6 groups in rodents. Respiratory Physiology and Neurobiology, 2004, 143, 187-197.	0.7	145
26	Polyalanine expansion and frameshift mutations of the paired-like homeobox gene PHOX2B in congenital central hypoventilation syndrome. Nature Genetics, 2003, 33, 459-461.	9.4	771
27	Genes modulating chemical breathing control: lessons from mutant animals. Respiratory Physiology and Neurobiology, 2003, 136, 105-114.	0.7	16
28	Noradrenergic neuronal development is impaired by mutation of the proneural HASH-1 gene in congenital central hypoventilation syndrome (Ondine's curse). Human Molecular Genetics, 2003, 12, 3173-3180.	1.4	72
29	Early neuronal and glial determination from mouse E10.5 telencephalon embryonic stem cells: an in vitro study. NeuroReport, 2002, 13, 1209-1214.	0.6	5
30	Murine peripherin gene sequences direct Cre recombinase expression to peripheral neurons in transgenic mice. FEBS Letters, 2002, 523, 68-72.	1.3	24
31	Ventilatory responses to hypercapnia and hypoxia in heterozygous c-ret newborn mice. Respiratory Physiology and Neurobiology, 2002, 131, 213-222.	0.7	16
32	Heart transplantation changes the expression of distinct gene families. Physiological Genomics, 2001, 7, 115-126.	1.0	7
33	MASH-1/RET pathway involvement in development of brain stem control of respiratory frequency in newborn mice. Physiological Genomics, 2001, 7, 149-157.	1.0	35
34	Impaired Ventilatory Responses to Hypoxia in Mice Deficient in Endothelin-Converting-Enzyme-1. Pediatric Research, 2001, 49, 705-712.	1.1	41
35	Cloning and Expression Analysis of a Novel Gene, RP42, Mapping to an Autism Susceptibility Locus on 6q16. Genomics, 2000, 65, 70-74.	1.3	10
36	Ventilatory Responses to Hypercapnia and Hypoxia in Mash-1 Heterozygous Newborn and Adult Mice. Pediatric Research, 1999, 46, 535-535.	1.1	49

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37	CRE and TRE sequences of the rat tyrosine hydroxylase promoter are required for TH basal expression in adult mice but not in the embryo. European Journal of Neuroscience, 1998, 10, 508-521.	1.2	54
38	Cell type-specific expression of the mouse peripherin gene requires both upstream and intragenic sequences in transgenic mouse embryos. Developmental Brain Research, 1996, 92, 1-9.	2.1	11
39	Neuronal defects in genotyped dominant megacolon (Dom) mouse embryos, a model for Hirschsprung disease. NeuroReport, 1996, 7, 489-492.	0.6	6
40	Both upstream and intragenic sequences of the human neurofilament light gene direct expression of of of transgenic mouse embryos. Journal of Molecular Neuroscience, 1994, 5, 273-295.	1.1	14
41	Calcium alginate immobilization of mammalian neurons: A potential tool to purify ligands of neuronal membrane proteins. Biotechnology Letters, 1991, 5, 289-294.	0.5	2
42	Storage and growth of neuroblastoma cells immobilized in calcium-alginate beads. Applied Microbiology and Biotechnology, 1990, 33, 442-7.	1.7	20
43	Large unit conductance voltage chloride channels are expressed in mouse neural crest cells and embryonic dorsal root ganglion cells. Developmental Brain Research, 1990, 51, 283-286.	2.1	4
44	Expression of voltage-dependent sodium and transient potassium currents in an identified sub-population of dorsal root ganglion cells acutely isolated from 12-day-old mouse embryos. Pflugers Archiv European Journal of Physiology, 1989, 414, 360-368.	1.3	24
45	Potassium channels in mouse neonate dorsal root ganglion cells: a patch-clamp study. Brain Research, 1987, 412, 224-232.	1.1	27
46	Scorpion venom inhibits selectively Ca2+ -activated K+ channels in situ. FEBS Letters, 1986, 209, 165-168.	1.3	9
47	Single channel currents in mouse embryonal multipotential carcinoma cells. Cell Differentiation, 1985, 17, 21-28.	1.3	5