Jean-FranÃ\sois Brunet

List of Publications by Year in descending order

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49 papers

5,911 citations

34 h-index 206112 48 g-index

53 all docs 53 docs citations

times ranked

53

4506 citing authors

#	Article	lF	CITATIONS
1	The homeobox gene Phox2b is essential for the development of autonomic neural crest derivatives. Nature, 1999, 399, 366-370.	27.8	775
2	Spatiotemporal structure of cell fate decisions in murine neural crest. Science, 2019, 364, .	12.6	345
3	Defects in Sensory and Autonomic Ganglia and Absence of Locus Coeruleus in Mice Deficient for the Homeobox Gene Phox2a. Neuron, 1997, 18, 411-423.	8.1	315
4	Phox2b controls the development of peripheral chemoreceptors and afferent visceral pathways. Development (Cambridge), 2003, 130, 6635-6642.	2.5	279
5	A human mutation in Phox2b causes lack of CO ₂ chemosensitivity, fatal central apnea, and specific loss of parafacial neurons. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 1067-1072.	7.1	271
6	The Expression Pattern of the Transcription Factor Phox2 Delineates Synaptic Pathways of the Autonomic Nervous System. Journal of Neuroscience, 1996, 16, 7649-7660.	3.6	266
7	Phox2 genes â€" from patterning to connectivity. Current Opinion in Genetics and Development, 2002, 12, 435-440.	3.3	222
8	Specification of the Central Noradrenergic Phenotype by the Homeobox Gene Phox2b. Molecular and Cellular Neurosciences, 2000, 15, 235-243.	2.2	198
9	Coordinated temporal and spatial control of motor neuron and serotonergic neuron generation from a common pool of CNS progenitors. Genes and Development, 2003, 17, 729-737.	5.9	196
10	The inducible cytotoxic T-lymphocyte-associated gene transcript CTLA-1 sequence and gene localization to mouse chromosome 14. Nature, 1986, 322, 268-271.	27.8	194
11	Essential role of Gata transcription factors in sympathetic neuron development. Development (Cambridge), 2004, 131, 4775-4786.	2.5	184
12	Ascl1/Mash1 is required for the development of central serotonergic neurons. Nature Neuroscience, 2004, 7, 589-595.	14.8	172
13	Specification of Neurotransmitter Identity by Phox2 Proteins in Neural Crest Stem Cells. Neuron, 1999, 22, 693-705.	8.1	161
14	Breathing without CO ₂ Chemosensitivity in Conditional <i>Phox2b</i> Mutants. Journal of Neuroscience, 2011, 31, 12880-12888.	3.6	149
15	Precraniate origin of cranial motoneurons. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 8727-8732.	7.1	144
16	Molecular consequences of PHOX2B missense, frameshift and alanine expansion mutations leading to autonomic dysfunction. Human Molecular Genetics, 2005, 14, 3697-3708.	2.9	135
17	Task2 potassium channels set central respiratory CO ₂ and O ₂ sensitivity. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 2325-2330.	7.1	132
18	Deconstructing cell determination: proneural genes and neuronal identity. BioEssays, 1999, 21, 313-318.	2.5	131

#	Article	IF	CITATIONS
19	<i>Paired</i> àêLike Homeodomain Proteins, Phox2a and Phox2b, Are Responsible for Noradrenergic Cellâ€Specific Transcription of the Dopamine βâ€Hydroxylase Gene. Journal of Neurochemistry, 1998, 71, 1813-1826.	3.9	123
20	Specific and integrated roles of Lmx1a, Lmx1b and Phox2a in ventral midbrain development. Development (Cambridge), 2011, 138, 3399-3408.	2.5	119
21	Defective Respiratory Rhythmogenesis and Loss of Central Chemosensitivity in Phox2b Mutants Targeting Retrotrapezoid Nucleus Neurons. Journal of Neuroscience, 2009, 29, 14836-14846.	3.6	115
22	Noradrenergic-Specific Transcription of the Dopamine Î ² -Hydroxylase Gene Requires Synergy of MultipleCis-Acting Elements Including at Least Two Phox2a-Binding Sites. Journal of Neuroscience, 1998, 18, 8247-8260.	3.6	112
23	Dual origin of enteric neurons in vagal Schwann cell precursors and the sympathetic neural crest. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 11980-11985.	7.1	108
24	Homeoprotein Phox2b commands a somatic-to-visceral switch in cranial sensory pathways. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 20018-20023.	7.1	101
25	Autonomic neurocristopathy-associated mutations in PHOX2B dysregulate Sox10 expression. Journal of Clinical Investigation, 2012, 122, 3145-3158.	8.2	89
26	The retrotrapezoid nucleus neurons expressing Atoh1 and Phox2b are essential for the respiratory response to CO2. ELife, 2015, 4, .	6.0	83
27	Delays in neuronal differentiation in Mash1/Ascl1 mutants. Developmental Biology, 2006, 295, 67-75.	2.0	78
28	Genetic identification of a hindbrain nucleus essential for innate vocalization. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 8095-8100.	7.1	74
29	PHOX2B in respiratory control: Lessons from congenital central hypoventilation syndrome and its mouse models. Respiratory Physiology and Neurobiology, 2009, 168, 125-132.	1.6	63
30	Epibranchial ganglia orchestrate the development of the cranial neurogenic crest. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 2066-2071.	7.1	51
31	Dynamic Expression of <i>RGS4</i> in the Developing Nervous System and Regulation by the Neural Type-Specific Transcription Factor Phox2b. Journal of Neuroscience, 2003, 23, 10613-10621.	3.6	50
32	Breathing with <i>Phox2b </i> . Philosophical Transactions of the Royal Society B: Biological Sciences, 2009, 364, 2477-2483.	4.0	43
33	Ongoing roles of Phox2 homeodomain transcription factors during neuronal differentiation. Development (Cambridge), 2010, 137, 4211-4220.	2.5	43
34	A <i>Phox2b::FLPo</i> transgenic mouse line suitable for intersectional genetics. Genesis, 2013, 51, 506-514.	1.6	38
35	Ancient origin of somatic and visceral neurons. BMC Biology, 2013, 11, 53.	3.8	38
36	Phox2b, congenital central hypoventilation syndrome and the control of respiration. Seminars in Cell and Developmental Biology, 2010, 21, 814-822.	5.0	37

#	Article	IF	Citations
37	Forced expression of Phox2 homeodomain transcription factors induces a branchio-visceromotor axonal phenotype. Developmental Biology, 2007, 303, 687-702.	2.0	34
38	<i>In Vitro</i> studies of non poly alanine PHOX2B mutations argue against a loss-of-function mechanism for congenital central hypoventilation. Human Mutation, 2009, 30, E421-E431.	2.5	34
39	Haploinsufficiency for Phox2b in mice causes dilated pupils and atrophy of the ciliary ganglion: mechanistic insights into human congenital central hypoventilation syndrome. Human Molecular Genetics, 2004, 13, 1433-1439.	2.9	31
40	Central chemoreception: Lessons from mouse and human genetics. Respiratory Physiology and Neurobiology, 2010, 173, 312-321.	1.6	28
41	Mutation in $\langle i \rangle$ LBX1/Lbx1 $\langle i \rangle$ precludes transcription factor cooperativity and causes congenital hypoventilation in humans and mice. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, 13021-13026.	7.1	27
42	The "sacral parasympathetic― ontogeny and anatomy of a myth. Clinical Autonomic Research, 2018, 28, 13-21.	2.5	25
43	Role of the Target in the Pathfinding of Facial Visceral Motor Axons. Molecular and Cellular Neurosciences, 2000, 16, 14-26.	2.2	23
44	A medullary centre for lapping in mice. Nature Communications, 2021, 12, 6307.	12.8	19
45	Taste bud formation depends on taste nerves. ELife, 2019, 8, .	6.0	13
46	Phox2b and the homeostatic brain. , 2008, , 25-44.		11
47	<i>Phox2b</i> expression in the taste centers of fish. Journal of Comparative Neurology, 2012, 520, 3633-3649.	1.6	9
48	Tridimensional mapping of Phox2b expressing neurons in the brainstem of adult <i>Macaca fascicularis</i> and identification of the retrotrapezoid nucleus. Journal of Comparative Neurology, 2019, 527, 2875-2884.	1.6	8
49	Molecular Cloning of an Inducible Cytotoxic T-Lymphocyte-Associated Gene (Hu-CTLA 1) and Gene Localization to Human Chromosome 14. , 1989, , 574-577.		O