Renata bartesaghi

List of Publications by Year in descending order

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126907 144013 3,759 95 33 57 citations h-index g-index papers 95 95 95 3564 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Treatment with the flavonoid 7,8-Dihydroxyflavone: a promising strategy for a constellation of body and brain disorders. Critical Reviews in Food Science and Nutrition, 2022, 62, 13-50.	10.3	30
2	Early Appearance of Dendritic Alterations in Neocortical Pyramidal Neurons of the Ts65Dn Model of Down Syndrome. Developmental Neuroscience, 2022, 44, 23-38.	2.0	8
3	Prenatal and Postnatal Pharmacotherapy in Down Syndrome: The Search to Prevent or Ameliorate Neurodevelopmental and Neurodegenerative Disorders. Annual Review of Pharmacology and Toxicology, 2022, 62, 211-233.	9.4	7
4	Impaired Brain Mitochondrial Bioenergetics in the Ts65Dn Mouse Model of Down Syndrome Is Restored by Neonatal Treatment with the Polyphenol 7,8-Dihydroxyflavone. Antioxidants, 2022, 11, 62.	5.1	12
5	The Challenging Pathway of Treatment for Neurogenesis Impairment in Down Syndrome: Achievements and Perspectives. Frontiers in Cellular Neuroscience, 2022, 16 , .	3.7	6
6	Fatty Acids: A Safe Tool for Improving Neurodevelopmental Alterations in Down Syndrome?. Nutrients, 2022, 14, 2880.	4.1	3
7	The flavonoid 7,8-DHF fosters prenatal brain proliferation potency in a mouse model of Down syndrome. Scientific Reports, 2021, 11, 6300.	3.3	9
8	Obstructive sleep apneas naturally occur in mice during REM sleep and are highly prevalent in a mouse model of Down syndrome. Neurobiology of Disease, 2021, 159, 105508.	4.4	8
9	Building the Future Therapies for Down Syndrome: The Third International Conference of the T21 Research Society. Molecular Syndromology, 2021, 12, 202-218.	0.8	6
10	Epigallocatechin-3-gallate., 2021,, 619-630.		O
10	Epigallocatechin-3-gallate., 2021,, 619-630. Early appearance of developmental alterations in the dendritic tree of the hippocampal granule cells in the Ts65Dn model of Down syndrome. Hippocampus, 2021, 31, 435-447.	1.9	7
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11	Early appearance of developmental alterations in the dendritic tree of the hippocampal granule cells in the Ts65Dn model of Down syndrome. Hippocampus, 2021, 31, 435-447. Prenatal, but not Postnatal, Curcumin Administration Rescues Neuromorphological and Cognitive		7
11 12	Early appearance of developmental alterations in the dendritic tree of the hippocampal granule cells in the Ts65Dn model of Down syndrome. Hippocampus, 2021, 31, 435-447. Prenatal, but not Postnatal, Curcumin Administration Rescues Neuromorphological and Cognitive Alterations in Ts65Dn Down Syndrome Mice. Journal of Nutrition, 2020, 150, 2478-2489. Neuroanatomical alterations in higher-order thalamic nuclei of fetuses with Down syndrome.	2.9	7
11 12 13	Early appearance of developmental alterations in the dendritic tree of the hippocampal granule cells in the Ts65Dn model of Down syndrome. Hippocampus, 2021, 31, 435-447. Prenatal, but not Postnatal, Curcumin Administration Rescues Neuromorphological and Cognitive Alterations in Ts65Dn Down Syndrome Mice. Journal of Nutrition, 2020, 150, 2478-2489. Neuroanatomical alterations in higher-order thalamic nuclei of fetuses with Down syndrome. Clinical Neurology and Neurosurgery, 2020, 194, 105870. Neonatal therapy with clenbuterol and salmeterol restores spinogenesis and dendritic complexity in	2.9	7 7 11
11 12 13	Early appearance of developmental alterations in the dendritic tree of the hippocampal granule cells in the Ts65Dn model of Down syndrome. Hippocampus, 2021, 31, 435-447. Prenatal, but not Postnatal, Curcumin Administration Rescues Neuromorphological and Cognitive Alterations in Ts65Dn Down Syndrome Mice. Journal of Nutrition, 2020, 150, 2478-2489. Neuroanatomical alterations in higher-order thalamic nuclei of fetuses with Down syndrome. Clinical Neurology and Neurosurgery, 2020, 194, 105870. Neonatal therapy with clenbuterol and salmeterol restores spinogenesis and dendritic complexity in the dentate gyrus of the Ts65Dn model of Down syndrome. Neurobiology of Disease, 2020, 140, 104874. Prenatal Administration of Oleic Acid or Linolenic Acid Reduces Neuromorphological and Cognitive	2.9 1.4 4.4	7 7 11 12
11 12 13 14	Early appearance of developmental alterations in the dendritic tree of the hippocampal granule cells in the Ts65Dn model of Down syndrome. Hippocampus, 2021, 31, 435-447. Prenatal, but not Postnatal, Curcumin Administration Rescues Neuromorphological and Cognitive Alterations in Ts65Dn Down Syndrome Mice. Journal of Nutrition, 2020, 150, 2478-2489. Neuroanatomical alterations in higher-order thalamic nuclei of fetuses with Down syndrome. Clinical Neurology and Neurosurgery, 2020, 194, 105870. Neonatal therapy with clenbuterol and salmeterol restores spinogenesis and dendritic complexity in the dentate gyrus of the Ts65Dn model of Down syndrome. Neurobiology of Disease, 2020, 140, 104874. Prenatal Administration of Oleic Acid or Linolenic Acid Reduces Neuromorphological and Cognitive Alterations in Ts65dn Down Syndrome Mice. Journal of Nutrition, 2020, 150, 1631-1643. Early postnatal oleic acid administration enhances synaptic development and cognitive abilities in the	2.9 1.4 4.4 2.9	7 7 11 12 16

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19	Subicular hypotrophy in fetuses with Down syndrome and in the Ts65Dn model of Down syndrome. Brain Pathology, 2019, 29, 366-379.	4.1	9
20	Abnormal development of the inferior temporal region in fetuses with Down syndrome. Brain Pathology, 2018, 28, 986-998.	4.1	34
21	CDKL5 protein substitution therapy rescues neurological phenotypes of a mouse model of CDKL5 disorder. Human Molecular Genetics, 2018, 27, 1572-1592.	2.9	49
22	Selective inhibitors of GSK-3Î ² : a suitable therapy for Down syndrome?. European Neuropsychopharmacology, 2018, 28, S72-S73.	0.7	1
23	Neurogenesis impairment: An early developmental defect in Down syndrome. Free Radical Biology and Medicine, 2018, 114, 15-32.	2.9	75
24	Treatment with corn oil improves neurogenesis and cognitive performance in the Ts65Dn mouse model of Down syndrome. Brain Research Bulletin, 2018, 140, 378-391.	3.0	14
25	Translating molecular advances in Down syndrome and Fragile X syndrome into therapies. European Neuropsychopharmacology, 2018, 28, 675-690.	0.7	14
26	Epigallocatechin gallate: A useful therapy for cognitive disability in Down syndrome?. Neurogenesis (Austin, Tex), 2017, 4, e1270383.	1.5	13
27	Long-term effect of neonatal inhibition of APP gamma-secretase on hippocampal development in the Ts65Dn mouse model of Down syndrome. Neurobiology of Disease, 2017, 103, 11-23.	4.4	14
28	A flavonoid agonist of the TrkB receptor for BDNF improves hippocampal neurogenesis and hippocampus-dependent memory in the Ts65Dn mouse model of DS. Experimental Neurology, 2017, 298, 79-96.	4.1	50
29	Neuroanatomical alterations and synaptic plasticity impairment in the perirhinal cortex of the Ts65Dn mouse model of Down syndrome. Neurobiology of Disease, 2017, 106, 89-100.	4.4	19
30	Lithium Restores Age-related Olfactory Impairment in the Ts65Dn Mouse Model of Down Syndrome. CNS and Neurological Disorders - Drug Targets, 2017, 16, 812-819.	1.4	10
31	Targeting APP/AICD in Down syndrome. Oncotarget, 2017, 8, 50333-50334.	1.8	3
32	SNX27, a protein involved in down syndrome, regulates GPR17 trafficking and oligodendrocyte differentiation. Glia, 2016, 64, 1437-1460.	4.9	20
33	HDAC4: a key factor underlying brain developmental alterations in CDKL5 disorder. Human Molecular Genetics, 2016, 25, 3887-3907.	2.9	77
34	Short- and long-term effects of neonatal pharmacotherapy with epigallocatechin-3-gallate on hippocampal development in the Ts65Dn mouse model of Down syndrome. Neuroscience, 2016, 333, 277-301.	2.3	60
35	Timing of therapies for Down syndrome: the sooner, the better. Frontiers in Behavioral Neuroscience, 2015, 9, 265.	2.0	94
36	New Perspectives for the Rescue of Cognitive Disability in Down Syndrome. Journal of Neuroscience, 2015, 35, 13843-13852.	3.6	28

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37	Inhibition of GSK3 \hat{I}^2 rescues hippocampal development and learning in a mouse model of CDKL5 disorder. Neurobiology of Disease, 2015, 82, 298-310.	4.4	55
38	ISDN2014_0057: Inhibition of GSK3â€beta rescues hippocampal development in a knockout mouse model of CDKL5 encephalopathy. International Journal of Developmental Neuroscience, 2015, 47, 12-13.	1.6	0
39	Inhibition of APP gamma-secretase restores Sonic Hedgehog signaling and neurogenesis in the Ts65Dn mouse model of Down syndrome. Neurobiology of Disease, 2015, 82, 385-396.	4.4	37
40	Long-term effects of neonatal treatment with fluoxetine on cognitive performance in Ts65Dn mice. Neurobiology of Disease, 2015, 74, 204-218.	4.4	44
41	Prenatal pharmacotherapy rescues brain development in a Down's syndrome mouse model. Brain, 2014, 137, 380-401.	7.6	71
42	Loss of CDKL5 impairs survival and dendritic growth of newborn neurons by altering AKT/GSK-3Î ² signaling. Neurobiology of Disease, 2014, 70, 53-68.	4.4	105
43	APP-dependent alteration of GSK3 \hat{l}^2 activity impairs neurogenesis in the Ts65Dn mouse model of Down syndrome. Neurobiology of Disease, 2014, 67, 24-36.	4.4	33
44	Age-related impairment of olfactory bulb neurogenesis in the Ts65Dn mouse model of Down syndrome. Experimental Neurology, 2014, 251, 1-11.	4.1	18
45	The Amyloid Precursor Protein (APP) Triplicated Gene Impairs Neuronal Precursor Differentiation and Neurite Development through Two Different Domains in the Ts65Dn Mouse Model for Down Syndrome. Journal of Biological Chemistry, 2013, 288, 20817-20829.	3.4	46
46	Early Pharmacotherapy with Fluoxetine Rescues Dendritic Pathology in the <scp>Ts65Dn</scp> Mouse Model of <scp>D</scp> own Syndrome. Brain Pathology, 2013, 23, 129-143.	4.1	61
47	Pharmacotherapy with Fluoxetine Restores Functional Connectivity from the Dentate Gyrus to Field CA3 in the Ts65Dn Mouse Model of Down Syndrome. PLoS ONE, 2013, 8, e61689.	2.5	42
48	Early-occurring proliferation defects in peripheral tissues of the Ts65Dn mouse model of Down syndrome are associated with patched 1 over expression. Laboratory Investigation, 2012, 92, 1648-1660.	3.7	21
49	CDKL5, a novel MYCN-repressed gene, blocks cell cycle and promotes differentiation of neuronal cells. Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms, 2012, 1819, 1173-1185.	1.9	31
50	Postnatal neurogenesis in the hippocampal dentate gyrus and subventricular zone of the $G\tilde{A}^{\P}$ ttingen minipig. Brain Research Bulletin, 2011, 85, 169-179.	3.0	18
51	Widespread Proliferation Impairment and Hypocellularity in the Cerebellum of Fetuses with Down Syndrome. Brain Pathology, 2011, 21, 361-373.	4.1	137
52	Impact of environmental enrichment on neurogenesis in the dentate gyrus during the early postnatal period. Brain Research, 2011, 1415, 23-33.	2.2	30
53	Is it possible to improve neurodevelopmental abnormalities in Down syndrome?. Reviews in the Neurosciences, 2011, 22, 419-455.	2.9	66
54	APP-dependent up-regulation of Ptch1 underlies proliferation impairment of neural precursors in Down syndrome. Human Molecular Genetics, 2011, 20, 1560-1573.	2.9	106

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55	Lithium Restores Neurogenesis in the Subventricular Zone of the Ts65Dn Mouse, a Model for Down Syndrome. Brain Pathology, 2010, 20, 106-118.	4.1	75
56	Early Pharmacotherapy Restores Neurogenesis and Cognitive Performance in the Ts65Dn Mouse Model for Down Syndrome. Journal of Neuroscience, 2010, 30, 8769-8779.	3.6	164
57	CB1 Cannabinoid Receptors Increase Neuronal Precursor Proliferation through AKT/Glycogen Synthase Kinase-3 \hat{l}^2/\hat{l}^2 -Catenin Signaling. Journal of Biological Chemistry, 2010, 285, 10098-10109.	3.4	73
58	Lot1 Is a Key Element of the Pituitary Adenylate Cyclase-activating Polypeptide (PACAP)/Cyclic AMP Pathway That Negatively Regulates Neuronal Precursor Proliferation. Journal of Biological Chemistry, 2009, 284, 15325-15338.	3.4	18
59	Changes in hippocampal morphology and neuroplasticity induced by adolescent THC treatment are associated with cognitive impairment in adulthood. Hippocampus, 2009, 19, 763-772.	1.9	244
60	Widespread impairment of cell proliferation in the neonate Ts65Dn mouse, a model for Down syndrome. Cell Proliferation, 2009, 42, 171-181.	5.3	35
61	Cell Cycle Elongation Impairs Proliferation of Cerebellar Granule Cell Precursors in the Ts65Dn Mouse, an Animal Model for Down Syndrome. Brain Pathology, 2009, 19, 224-237.	4.1	60
62	RESEARCH ARTICLE: Neurogenesis Impairment and Increased Cell Death Reduce Total Neuron Number in the Hippocampal Region of Fetuses with Down Syndrome. Brain Pathology, 2008, 18, 180-197.	4.1	230
63	Neonatal isolation impairs neurogenesis in thedentate gyrus of the guinea pig. Hippocampus, 2007, 17, 78-91.	1.9	23
64	Cell cycle alteration and decreased cell proliferation in the hippocampal dentate gyrus and in the neocortical germinal matrix of fetuses with down syndrome and in Ts65Dn mice. Hippocampus, 2007, 17, 665-678.	1.9	234
65	Effect of early isolation on signal transfer in the entorhinal cortex–dentate–hippocampal system. Neuroscience, 2006, 137, 875-890.	2.3	17
66	Sex differences in the hilar mossy cells of the guinea-pig before puberty. Neuroscience, 2006, 139, 565-576.	2.3	7
67	Input–output relations in the entorhinal cortex–dentate–hippocampal system: Evidence for a non-linear transfer of signals. Neuroscience, 2006, 142, 247-265.	2.3	45
68	Choline acetyltransferase activity at different ages in brain of Ts65Dn mice, an animal model for Down's syndrome and related neurodegenerative diseases. Journal of Neurochemistry, 2006, 97, 515-526.	3.9	63
69	Proliferation of cerebellar precursor cells is negatively regulated by nitric oxide in newborn rat. Journal of Cell Science, 2006, 119, 3161-3170.	2.0	35
70	Postnatal neurogenesis in the dentate gyrus of the guinea pig. Hippocampus, 2005, 15, 285-301.	1.9	52
71	Topographic activation of the medial entorhinal cortex by presubicular commissural projections. Journal of Comparative Neurology, 2005, 487, 283-299.	1.6	9
72	Cyclic AMP-mediated Regulation of Transcription Factor Lot1 Expression in Cerebellar Granule Cells. Journal of Biological Chemistry, 2005, 280, 33541-33551.	3.4	17

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73	Neurochemical Correlates of Nicotine Neurotoxicity on Rat Habenulo-Interpeduncular Cholinergic Neurons. NeuroToxicology, 2005, 26, 467-474.	3.0	22
74	Sex differences in the stereological parameters of the hippocampal dentate gyrus of the guinea-pig before puberty. Neuroscience, 2005, 132, 375-387.	2.3	16
75	Nitric oxide negatively regulates proliferation and promotes neuronal differentiation through N-Myc downregulation. Journal of Cell Science, 2004, 117, 4727-4737.	2.0	69
76	Nitric oxide regulates cGMP-dependent cAMP-responsive element binding protein phosphorylation and Bcl-2 expression in cerebellar neurons: implication for a survival role of nitric oxide. Journal of Neurochemistry, 2004, 82, 1282-1289.	3.9	128
77	Effect of early isolation on the synaptic function in the dentate gyrus and field CA1 of the guinea pig. Hippocampus, 2004, 14, 482-498.	1.9	26
78	Parallel activation of field CA2 and dentate gyrus by synaptically elicited perforant path volleys. Hippocampus, 2004, 14, 948-963.	1.9	46
79	Effects of early environment on field CA2 pyramidal neurons in the guinea-pig. Neuroscience, 2004, 123, 703-714.	2.3	11
80	Activation of perforant path neurons to field CA1 by hippocampal projections. Hippocampus, 2003, 13, 235-249.	1.9	25
81	> effects of early environment on pyramidal neuron morphology in field CA1 of the guinea-pig. Neuroscience, 2003, 116, 715-732.	2.3	19
82	Effects of early isolation on layer ii neurons in the entorhinal cortex of the guinea pig. Neuroscience, 2003, 120, 721-732.	2.3	10
83	Sex differences in the hippocampal dentate gyrus of the guinea-pig before puberty. Neuroscience, 2003, 121, 327-339.	2.3	13
84	Nitric Oxide Protects Neuroblastoma Cells from Apoptosis Induced by Serum Deprivation through cAMP-response Element-binding Protein (CREB) Activation. Journal of Biological Chemistry, 2002, 277, 49896-49902.	3.4	76
85	Effects of early environment on field CA3a pyramidal neuron morphology in the guinea-pig. Neuroscience, 2002, 110, 475-488.	2.3	18
86	Effects of early environment on granule cell morphology in the dentate gyrus of the guinea-pig. Neuroscience, 2001, 102, 87-100.	2.3	29
87	Pyramidal neuron types in field CA2 of the guinea pig. Brain Research Bulletin, 1999, 50, 263-273.	3.0	24
88	Input-output relations in the entorhinal-hippocampal-entorhinal loop: Entorhinal cortex and dentate gyrus. Hippocampus, 1995, 5, 440-451.	1.9	16
89	Hippocampal-entorhinal relationships: Electrophysiological analysis of the ventral hippocampal projections to the ventral entorhinal cortex. Neuroscience, 1994, 61, 457-466.	2.3	9
90	Electrophysiological analysis of the hippocampal output to the presubiculum. Neuroscience, 1990, 37, 335-345.	2.3	6

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91	Electrophysiological analysis of the hippocampal projections to the entorhinal area. Neuroscience, 1989, 30, 51-62.	2.3	40
92	Electrophysiological analysis of the dorsal hippocampal commissure projections to the entorhinal area. Neuroscience, 1988, 26, 55-67.	2.3	25
93	Hippocampal output to the subicular cortex: An electrophysiological study. Experimental Neurology, 1986, 92, 114-133.	4.1	21
94	Fiber groups in the dorsal psalterium of the guinea pig. Experimental Neurology, 1985, 88, 500-514.	4.1	6
95	Interlamellar transfer of impulses in the hippocampal formation. Experimental Neurology, 1983, 82, 550-567.	4.1	27