

Jacqueline Crawley

List of Publications by Year in descending order

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119
papers

24,670
citations

9786

73
h-index

18130

120
g-index

122
all docs

122
docs citations

122
times ranked

19318
citing authors

#	ARTICLE	IF	CITATIONS
1	Behavioral phenotypes of inbred mouse strains: implications and recommendations for molecular studies. <i>Psychopharmacology</i> , 1997, 132, 107-124.	3.1	1,283
2	Behavioural phenotyping assays for mouse models of autism. <i>Nature Reviews Neuroscience</i> , 2010, 11, 490-502.	10.2	1,248
3	Sociability and preference for social novelty in five inbred strains: an approach to assess autistic-like behavior in mice. <i>Genes, Brain and Behavior</i> , 2004, 3, 287-302.	2.2	1,241
4	Preliminary report of a simple animal behavior model for the anxiolytic effects of benzodiazepines. <i>Pharmacology Biochemistry and Behavior</i> , 1980, 13, 167-170.	2.9	1,131
5	Autistic-like behaviour and cerebellar dysfunction in Purkinje cell Tsc1 mutant mice. <i>Nature</i> , 2012, 488, 647-651.	27.8	756
6	Mouse behavioral tasks relevant to autism: Phenotypes of 10 inbred strains. <i>Behavioural Brain Research</i> , 2007, 176, 4-20.	2.2	714
7	Autism-like behavioral phenotypes in BTBR T+tf/J mice. <i>Genes, Brain and Behavior</i> , 2008, 7, 152-163.	2.2	709
8	Automated apparatus for quantitation of social approach behaviors in mice. <i>Genes, Brain and Behavior</i> , 2004, 3, 303-314.	2.2	680
9	Exploratory behavior models of anxiety in mice. <i>Neuroscience and Biobehavioral Reviews</i> , 1985, 9, 37-44.	6.1	653
10	A Proposed Test Battery and Constellations of Specific Behavioral Paradigms to Investigate the Behavioral Phenotypes of Transgenic and Knockout Mice. <i>Hormones and Behavior</i> , 1997, 31, 197-211.	2.1	522
11	Haploinsufficiency of the autism-associated Shank3 gene leads to deficits in synaptic function, social interaction, and social communication. <i>Molecular Autism</i> , 2010, 1, 15.	4.9	521
12	Mouse Behavioral Assays Relevant to the Symptoms of Autism*. <i>Brain Pathology</i> , 2007, 17, 448-459.	4.1	511
13	Unusual Repertoire of Vocalizations in the BTBR T+tf/J Mouse Model of Autism. <i>PLoS ONE</i> , 2008, 3, e3067.	2.5	492
14	Pain responses, anxiety and aggression in mice deficient in pre-proenkephalin. <i>Nature</i> , 1996, 383, 535-538.	27.8	482
15	Social Interaction and Sensorimotor Gating Abnormalities in Mice Lacking Dvl1. <i>Cell</i> , 1997, 90, 895-905.	28.9	440
16	Designing mouse behavioral tasks relevant to autistic-like behaviors. <i>Mental Retardation and Developmental Disabilities Research Reviews</i> , 2004, 10, 248-258.	3.6	439
17	Automated Three-Chambered Social Approach Task for Mice. <i>Current Protocols in Neuroscience</i> , 2011, 56, Unit 8.26.	2.6	418
18	Ultrasonic vocalizations: A tool for behavioural phenotyping of mouse models of neurodevelopmental disorders. <i>Neuroscience and Biobehavioral Reviews</i> , 2009, 33, 508-515.	6.1	413

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19	Mouse models of Tayâ€“Sachs and Sandhoff diseases differ in neurologic phenotype and ganglioside metabolism. <i>Nature Genetics</i> , 1995, 11, 170-176.	21.4	411
20	Simple Behavioral Assessment of Mouse Olfaction. <i>Current Protocols in Neuroscience</i> , 2009, 48, Unit 8.24.	2.6	401
21	Behavioral Phenotyping Strategies for Mutant Mice. <i>Neuron</i> , 2008, 57, 809-818.	8.1	393
22	Repetitive Self-Grooming Behavior in the BTBR Mouse Model of Autism is Blocked by the mGluR5 Antagonist MPEP. <i>Neuropsychopharmacology</i> , 2010, 35, 976-989.	5.4	374
23	Inbred strain differences in prepulse inhibition of the mouse startle response. <i>Psychopharmacology</i> , 1997, 132, 169-180.	3.1	359
24	Reduced Excitatory Neurotransmission and Mild Autism-Relevant Phenotypes in Adolescent<i>Shank3</i>Null Mutant Mice. <i>Journal of Neuroscience</i> , 2012, 32, 6525-6541.	3.6	342
25	Unusual repertoire of vocalizations in adult BTBR T+tf/j mice during three types of social encounters. <i>Genes, Brain and Behavior</i> , 2011, 10, 44-56.	2.2	316
26	Autism gene variant causes hyperserotonemia, serotonin receptor hypersensitivity, social impairment and repetitive behavior. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 5469-5474.	7.1	278
27	Minimal aberrant behavioral phenotypes of neuroliginâ€“3 R451C knockin mice. <i>Autism Research</i> , 2008, 1, 147-158.	3.8	263
28	Clustering autism: using neuroanatomical differences in 26 mouse models to gain insight into the heterogeneity. <i>Molecular Psychiatry</i> , 2015, 20, 118-125.	7.9	257
29	Evaluation of Antidepressant-related Behavioral Responses in Mice Lacking the Serotonin Transporter. <i>Neuropsychopharmacology</i> , 2002, 27, 914-923.	5.4	256
30	Drug development for neurodevelopmental disorders: lessons learned from fragile X syndrome. <i>Nature Reviews Drug Discovery</i> , 2018, 17, 280-299.	46.4	247
31	Negative Allosteric Modulation of the mGluR5 Receptor Reduces Repetitive Behaviors and Rescues Social Deficits in Mouse Models of Autism. <i>Science Translational Medicine</i> , 2012, 4, 131ra51.	12.4	238
32	Social approach in genetically engineered mouse lines relevant to autism. <i>Genes, Brain and Behavior</i> , 2009, 8, 129-142.	2.2	225
33	Social approach and repetitive behavior in eleven inbred mouse strains. <i>Behavioural Brain Research</i> , 2008, 191, 118-129.	2.2	215
34	Germline Chd8 haploinsufficiency alters brain development in mouse. <i>Nature Neuroscience</i> , 2017, 20, 1062-1073.	14.8	210
35	Behavioral Abnormalities and Circuit Defects in the Basal Ganglia of a Mouse Model of 16p11.2 Deletion Syndrome. <i>Cell Reports</i> , 2014, 7, 1077-1092.	6.4	208
36	Sociability and motor functions in Shank1 mutant mice. <i>Brain Research</i> , 2011, 1380, 120-137.	2.2	206

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37	Social approach behaviors in oxytocin knockout mice: Comparison of two independent lines tested in different laboratory environments. <i>Neuropeptides</i> , 2007, 41, 145-163.	2.2	204
38	Communication Impairments in Mice Lacking Shank1: Reduced Levels of Ultrasonic Vocalizations and Scent Marking Behavior. <i>PLoS ONE</i> , 2011, 6, e20631.	2.5	196
39	Translational animal models of autism and neurodevelopmental disorders. <i>Dialogues in Clinical Neuroscience</i> , 2012, 14, 293-305.	3.7	195
40	Mice lacking both subunits of lysosomal β -hexosaminidase display gangliosidosis and mucopolysaccharidosis. <i>Nature Genetics</i> , 1996, 14, 348-352.	21.4	194
41	Development of a mouse test for repetitive, restricted behaviors: Relevance to autism. <i>Behavioural Brain Research</i> , 2008, 188, 178-194.	2.2	192
42	GABAB Receptor Agonist R-Baclofen Reverses Social Deficits and Reduces Repetitive Behavior in Two Mouse Models of Autism. <i>Neuropsychopharmacology</i> , 2015, 40, 2228-2239.	5.4	187
43	Preclinical research in Rett syndrome: setting the foundation for translational success. <i>DMM Disease Models and Mechanisms</i> , 2012, 5, 733-745.	2.4	183
44	Modeling fragile X syndrome in the <i>Fmr1</i> knockout mouse. <i>Intractable and Rare Diseases Research</i> , 2014, 3, 118-133.	0.9	183
45	The Female Urine Sniffing Test: A Novel Approach for Assessing Reward-Seeking Behavior in Rodents. <i>Biological Psychiatry</i> , 2010, 67, 864-871.	1.3	174
46	Reduced scent marking and ultrasonic vocalizations in the BTBR T+tf/J mouse model of autism. <i>Genes, Brain and Behavior</i> , 2011, 10, 35-43.	2.2	166
47	Neurogranin null mutant mice display performance deficits on spatial learning tasks with anxiety related components. <i>Hippocampus</i> , 2001, 11, 763-775.	1.9	159
48	Autism-Relevant Social Abnormalities and Cognitive Deficits in Engrailed-2 Knockout Mice. <i>PLoS ONE</i> , 2012, 7, e40914.	2.5	143
49	The role of galanin in feeding behavior. <i>Neuropeptides</i> , 1999, 33, 369-375.	2.2	140
50	Behavioral phenotypes of genetic mouse models of autism. <i>Genes, Brain and Behavior</i> , 2016, 15, 7-26.	2.2	137
51	Replicable in vivo physiological and behavioral phenotypes of the Shank3B null mutant mouse model of autism. <i>Molecular Autism</i> , 2017, 8, 26.	4.9	135
52	Dysbindin-1 modulates prefrontal cortical activity and schizophrenia-like behaviors via dopamine/D2 pathways. <i>Molecular Psychiatry</i> , 2012, 17, 85-98.	7.9	128
53	Low stress reactivity and neuroendocrine factors in the BTBR T+tf/J mouse model of autism. <i>Neuroscience</i> , 2010, 171, 1197-1208.	2.3	125
54	Social deficits in BTBR T+tf/J mice are unchanged by cross-fostering with C57BL/6J mothers. <i>International Journal of Developmental Neuroscience</i> , 2007, 25, 515-521.	1.6	124

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55	Autism and Cancer Share Risk Genes, Pathways, and Drug Targets. <i>Trends in Genetics</i> , 2016, 32, 139-146.	6.7	123
56	Mouse models of autism spectrum disorders: The challenge for behavioral genetics. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2006, 142C, 40-51.	1.6	116
57	Long-term exposure to intranasal oxytocin in a mouse autism model. <i>Translational Psychiatry</i> , 2014, 4, e480-e480.	4.8	112
58	Assessing behavioural and cognitive domains of autism spectrum disorders in rodents: current status and future perspectives. <i>Psychopharmacology</i> , 2014, 231, 1125-1146.	3.1	111
59	Developmental delays and reduced pup ultrasonic vocalizations but normal sociability in mice lacking the postsynaptic cell adhesion protein <i>neuroligin2</i> . <i>Behavioural Brain Research</i> , 2013, 251, 50-64.	2.2	110
60	Social approach behaviors are similar on conventional versus reverse lighting cycles, and in replications across cohorts, in BTBR T+ tf/J, C57BL/6J, and vasopressin receptor 1B mutant mice. <i>Frontiers in Behavioral Neuroscience</i> , 2007, 1, 1.	2.0	109
61	Behavioral and Neuroanatomical Phenotypes in Mouse Models of Autism. <i>Neurotherapeutics</i> , 2015, 12, 521-533.	4.4	108
62	Social deficits, stereotypy and early emergence of repetitive behavior in the C58/J inbred mouse strain. <i>Behavioural Brain Research</i> , 2010, 208, 178-188.	2.2	107
63	Postnatal lesion evidence against a primary role for the corpus callosum in mouse sociability. <i>European Journal of Neuroscience</i> , 2009, 29, 1663-1677.	2.6	104
64	Galanin receptor subtype 2 (<i>GalR2</i>) null mutant mice display an anxiogenic-like phenotype specific to the elevated plus-maze. <i>Pharmacology Biochemistry and Behavior</i> , 2007, 86, 8-20.	2.9	100
65	Low sociability in BTBR T+tf/J mice is independent of partner strain. <i>Physiology and Behavior</i> , 2012, 107, 649-662.	2.1	100
66	Translational Mouse Models of Autism: Advancing Toward Pharmacological Therapeutics. <i>Current Topics in Behavioral Neurosciences</i> , 2015, 28, 1-52.	1.7	100
67	Social transmission of food preference in mice: Methodology and application to galanin-overexpressing transgenic mice. <i>Behavioral Neuroscience</i> , 2003, 117, 21-31.	1.2	99
68	AMPAKINE enhancement of social interaction in the BTBR mouse model of autism. <i>Neuropharmacology</i> , 2013, 64, 268-282.	4.1	98
69	Absence of preference for social novelty and increased grooming in integrin $\beta 3$ knockout mice: Initial studies and future directions. <i>Autism Research</i> , 2011, 4, 57-67.	3.8	97
70	Subtype-selective cholecystokinin receptor antagonists block cholecystokinin modulation of dopamine-mediated behaviors in the rat mesolimbic pathway. <i>Journal of Neuroscience</i> , 1992, 12, 3380-3391.	3.6	94
71	Genetic analysis of anxiety-related behaviors and responses to benzodiazepine-related drugs in AXB and BXA recombinant inbred mouse strains. <i>Behavior Genetics</i> , 1995, 25, 557-568.	2.1	90
72	Female urine-induced male mice ultrasonic vocalizations, but not scent-marking, is modulated by social experience. <i>Behavioural Brain Research</i> , 2011, 216, 19-28.	2.2	85

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73	Impaired Learning and Motor Behavior in Heterozygous <i>En1</i> Mutant Mice. <i>Learning and Memory</i> , 1999, 6, 521-537.	1.3	84
74	16p11.2 Deletion Syndrome Mice Display Sensory and Ultrasonic Vocalization Deficits During Social Interactions. <i>Autism Research</i> , 2015, 8, 507-521.	3.8	80
75	Olfactory cues are sufficient to elicit social approach behaviors but not social transmission of food preference in C57BL/6J mice. <i>Behavioural Brain Research</i> , 2008, 193, 235-242.	2.2	76
76	R-Baclofen Reverses Cognitive Deficits and Improves Social Interactions in Two Lines of 16p11.2 Deletion Mice. <i>Neuropsychopharmacology</i> , 2018, 43, 513-524.	5.4	75
77	Absence of deficits in social behaviors and ultrasonic vocalizations in later generations of mice lacking <i>neuroligin4</i> . <i>Genes, Brain and Behavior</i> , 2012, 11, 928-941.	2.2	71
78	Hippocampal Transcriptomic and Proteomic Alterations in the BTBR Mouse Model of Autism Spectrum Disorder. <i>Frontiers in Physiology</i> , 2015, 6, 324.	2.8	70
79	Rigor and reproducibility in rodent behavioral research. <i>Neurobiology of Learning and Memory</i> , 2019, 165, 106780.	1.9	65
80	Male mice emit distinct ultrasonic vocalizations when the female leaves the social interaction arena. <i>Frontiers in Behavioral Neuroscience</i> , 2013, 7, 159.	2.0	56
81	Quantitative Trait Loci for Interhemispheric Commissure Development and Social Behaviors in the BTBR T+ <i>tf/J</i> Mouse Model of Autism. <i>PLoS ONE</i> , 2013, 8, e61829.	2.5	53
82	16p11.2 Deletion mice display cognitive deficits in touchscreen learning and novelty recognition tasks. <i>Learning and Memory</i> , 2015, 22, 622-632.	1.3	53
83	Social transmission of food preference in mice: methodology and application to galanin-overexpressing transgenic mice. <i>Behavioral Neuroscience</i> , 2003, 117, 21-31.	1.2	53
84	Mouse Models of Autism: Testing Hypotheses About Molecular Mechanisms. <i>Current Topics in Behavioral Neurosciences</i> , 2011, 7, 187-212.	1.7	51
85	Centrally administered cholecystokinin suppresses feeding through a peripheral-type receptor mechanism. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 1991, 257, 1076-80.	2.5	48
86	Behavioral Phenotyping Assays for Genetic Mouse Models of Neurodevelopmental, Neurodegenerative, and Psychiatric Disorders. <i>Annual Review of Animal Biosciences</i> , 2017, 5, 371-389.	7.4	46
87	Galanin: Neurobiologic Mechanisms and Therapeutic Potential for Alzheimer's Disease. <i>CNS Neuroscience & Therapeutics</i> , 2001, 7, 445-470.	4.0	45
88	<i>Engrailed-2</i> (<i>En2</i>) deletion produces multiple neurodevelopmental defects in monoamine systems, forebrain structures and neurogenesis and behavior. <i>Human Molecular Genetics</i> , 2015, 24, 5805-5827.	2.9	45
89	Neuregulin-2 ablation results in dopamine dysregulation and severe behavioral phenotypes relevant to psychiatric disorders. <i>Molecular Psychiatry</i> , 2018, 23, 1233-1243.	7.9	45
90	Evaluation of the neuroactive steroid ganaxolone on social and repetitive behaviors in the BTBR mouse model of autism. <i>Psychopharmacology</i> , 2016, 233, 309-323.	3.1	43

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91	Hypothesis-driven investigations of diverse pharmacological targets in two mouse models of autism. <i>Autism Research</i> , 2019, 12, 401-421.	3.8	42
92	Autism-specific maternal autoantibodies produce behavioral abnormalities in an endogenous antigen-driven mouse model of autism. <i>Molecular Psychiatry</i> , 2020, 25, 2994-3009.	7.9	42
93	SynDIG4/Prrt1 Is Required for Excitatory Synapse Development and Plasticity Underlying Cognitive Function. <i>Cell Reports</i> , 2018, 22, 2246-2253.	6.4	41
94	Cognitive Abilities on Transitive Inference Using a Novel Touchscreen Technology for Mice. <i>Cerebral Cortex</i> , 2015, 25, 1133-1142.	2.9	39
95	Behavioral assessment of NIH Swiss mice acutely intoxicated with tetramethylenedisulfotetramine. <i>Neurotoxicology and Teratology</i> , 2015, 47, 36-45.	2.4	38
96	Genetic background modulates phenotypes of serotonin transporter Ala56 knock-in mice. <i>Molecular Autism</i> , 2013, 4, 35.	4.9	35
97	Galanin – 25 years with a multitalented neuropeptide. <i>Cellular and Molecular Life Sciences</i> , 2008, 65, 1836-1841.	5.4	34
98	Early motor phenotype detection in a female mouse model of Rett syndrome is improved by cross-fostering. <i>Human Molecular Genetics</i> , 2017, 26, 1839-1854.	2.9	32
99	Coexistence of Neuropeptides and "Classical" Neurotransmitters.. <i>Annals of the New York Academy of Sciences</i> , 1990, 579, 233-241.	3.8	29
100	The promising trajectory of autism therapeutics discovery. <i>Drug Discovery Today</i> , 2014, 19, 838-844.	6.4	29
101	Galanin peptide levels in hippocampus and cortex of galanin-overexpressing transgenic mice evaluated for cognitive performance. <i>Neuropeptides</i> , 2002, 36, 413-426.	2.2	26
102	Chronic desipramine treatment rescues depression-related, social and cognitive deficits in <i>Engrailed2</i> knockout mice. <i>Genes, Brain and Behavior</i> , 2014, 13, 286-298.	2.2	24
103	In tribute to Bob Blanchard: Divergent behavioral phenotypes of 16p11.2 deletion mice reared in same-genotype versus mixed-genotype cages. <i>Physiology and Behavior</i> , 2015, 146, 16-27.	2.1	24
104	Touchscreen learning deficits in <i>Ube3a</i> , <i>Ts65Dn</i> and <i>Mecp2</i> mouse models of neurodevelopmental disorders with intellectual disabilities. <i>Genes, Brain and Behavior</i> , 2018, 17, e12452.	2.2	24
105	Normal Performance of <i>Fmr1</i> Mice on a Touchscreen Delayed Nonmatching to Position Working Memory Task. <i>ENeuro</i> , 2016, 3, ENEURO.0143-15.2016.	1.9	21
106	Modulation of Mesolimbic Dopaminergic Behaviors by Cholecystokinin. <i>Annals of the New York Academy of Sciences</i> , 1988, 537, 380-396.	3.8	19
107	Lack of effect of chronic morphine treatment and naloxone-precipitated withdrawal on tyrosine hydroxylase, galanin, and neuropeptide Y mRNA levels in the rat locus coeruleus. <i>Synapse</i> , 1995, 19, 197-205.	1.2	18
108	3D visualization of the regional differences. <i>Molecular Psychiatry</i> , 2015, 20, 1-1.	7.9	16

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109	Mesolimbic dopaminergic mechanisms underlying individual differences in sugar consumption and amphetamine hyperlocomotion in Wistar rats. <i>European Journal of Neuroscience</i> , 1998, 10, 1895-1902.	2.6	15
110	Behavioral Evaluation of Angelman Syndrome Mice at Older Ages. <i>Neuroscience</i> , 2020, 445, 163-171.	2.3	15
111	Transcription Factor 2I Regulates Neuronal Development via TRPC3 in 7q11.23 Disorder Models. <i>Molecular Neurobiology</i> , 2019, 56, 3313-3325.	4.0	13
112	Sexually dimorphic neuroanatomical differences relate to ASD-relevant behavioral outcomes in a maternal autoantibody mouse model. <i>Molecular Psychiatry</i> , 2021, 26, 7530-7537.	7.9	12
113	Galanin Impairs Cognitive Abilities in Rodents: Relevance to Alzheimer's Disease. <i>Exs</i> , 2010, 102, 133-141.	1.4	11
114	Spaced training improves learning in Ts65Dn and Ube3a mouse models of intellectual disabilities. <i>Translational Psychiatry</i> , 2019, 9, 166.	4.8	8
115	The CCK-B Antagonist CI-988 Increases Dopamine Levels in Microdialysate from the Rat Nucleus Accumbens via a Tetrodotoxin- and Calcium-Independent Mechanism. <i>Journal of Neurochemistry</i> , 2002, 65, 208-217.	3.9	6
116	Curiosity as an approach to ethoexperimental analysis: Behavioral neuroscience as seen by students and colleagues of Bob Blanchard. <i>Neuroscience and Biobehavioral Reviews</i> , 2017, 76, 415-422.	6.1	5
117	Rigor in science and science reporting: updated guidelines for submissions to <i>Molecular Autism</i> . <i>Molecular Autism</i> , 2019, 10, 6.	4.9	4
118	Evaluation of a TrkB agonist on spatial and motor learning in the Ube3a mouse model of Angelman syndrome. <i>Learning and Memory</i> , 2020, 27, 346-354.	1.3	4
119	Behavioral analyses of animal models of intellectual and developmental disabilities. <i>Neurobiology of Learning and Memory</i> , 2019, 165, 107087.	1.9	1