Martin Schalling

List of Publications by Year in descending order

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315 papers 31,834 citations

72 h-index 167 g-index

323 all docs 323 docs citations

times ranked

323

36685 citing authors

#	Article	IF	CITATIONS
1	Identification of risk loci with shared effects on five major psychiatric disorders: a genome-wide analysis. Lancet, The, 2013, 381, 1371-1379.	13.7	2,643
2	Genetic relationship between five psychiatric disorders estimated from genome-wide SNPs. Nature Genetics, 2013, 45, 984-994.	21.4	2,067
3	Genetic variants in novel pathways influence blood pressure and cardiovascular disease risk. Nature, 2011, 478, 103-109.	27.8	1,855
4	Somatic Mutations of Calreticulin in Myeloproliferative Neoplasms. New England Journal of Medicine, 2013, 369, 2379-2390.	27.0	1,698
5	Large-scale genome-wide association analysis of bipolar disorder identifies a new susceptibility locus near ODZ4. Nature Genetics, 2011, 43, 977-983.	21.4	1,283
6	Genome-wide association study identifies 30 loci associated with bipolar disorder. Nature Genetics, 2019, 51, 793-803.	21.4	1,191
7	Analysis of shared heritability in common disorders of the brain. Science, 2018, 360, .	12.6	1,085
8	PDGF-A Signaling Is a Critical Event in Lung Alveolar Myofibroblast Development and Alveogenesis. Cell, 1996, 85, 863-873.	28.9	787
9	The neuropeptide Y/agouti gene-related protein (AGRP) brain circuitry in normal, anorectic, and monosodium glutamate-treated mice. Proceedings of the National Academy of Sciences of the United States of America, 1998, 95, 15043-15048.	7.1	713
10	Overexpression of the obese (ob) gene in adipose tissue of human obese subjects. Nature Medicine, 1995, 1, 950-953.	30.7	680
11	Genome-wide association study of more than 40,000 bipolar disorder cases provides new insights into the underlying biology. Nature Genetics, 2021, 53, 817-829.	21.4	629
12	Neuropeptide expression in rat dorsal root ganglion cells and spinal cord after peripheral nerve injury with special reference to galanin. Neuroscience, 1989, 33, 587-604.	2.3	449
13	Expression of the Wilms' tumor gene WT1 in the murine urogenital system Genes and Development, 1991, 5, 1345-1356.	5.9	400
14	ARSACS, a spastic ataxia common in northeastern Qu \tilde{A} Obec, is caused by mutations in a new gene encoding an 11.5-kb ORF. Nature Genetics, 2000, 24, 120-125.	21.4	395
15	Muscleblind localizes to nuclear foci of aberrant RNA in myotonic dystrophy types 1 and 2. Human Molecular Genetics, 2001, 10, 2165-2170.	2.9	381
16	Tissue specific expression of FMR–1 provides evidence for a functional role in fragile X syndrome. Nature Genetics, 1993, 3, 36-43.	21.4	358
17	Impact of inflammation on epigenetic DNA methylation – a novel risk factor for cardiovascular disease?. Journal of Internal Medicine, 2007, 261, 488-499.	6.0	344
18	The Swedish Twin Registry in the Third Millennium: An Update. Twin Research and Human Genetics, 2006, 9, 875-882.	0.6	323

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19	A role for a new herpes virus (KSHV) in different forms of Kaposi's sarcoma. Nature Medicine, 1995, 1, 707-708.	30.7	320
20	Circadian Clock-Related Polymorphisms in Seasonal Affective Disorder and their Relevance to Diurnal Preference. Neuropsychopharmacology, 2003, 28, 734-739.	5.4	307
21	Genetic variants associated with response to lithium treatment in bipolar disorder: a genome-wide association study. Lancet, The, 2016, 387, 1085-1093.	13.7	306
22	Low fetuin-A levels are associated with cardiovascular death: Impact of variations in the gene encoding fetuin. Kidney International, 2005, 67, 2383-2392.	5.2	274
23	Direct detection of novel expanded trinucleotide repeats in the human genome. Nature Genetics, 1993, 4, 135-139.	21.4	272
24	All SNPs Are Not Created Equal: Genome-Wide Association Studies Reveal a Consistent Pattern of Enrichment among Functionally Annotated SNPs. PLoS Genetics, 2013, 9, e1003449.	3.5	268
25	LUMA (LUminometric Methylation Assay)—A high throughput method to the analysis of genomic DNA methylation. Experimental Cell Research, 2006, 312, 1989-1995.	2.6	261
26	Meta-analysis identifies multiple loci associated with kidney function–related traits in east Asian populations. Nature Genetics, 2012, 44, 904-909.	21.4	254
27	Genetic loci influencing kidney function and chronic kidney disease. Nature Genetics, 2010, 42, 373-375.	21.4	246
28	Three circadian clock genes Per2, Arntl, and Npas2 contribute to winter depression. Annals of Medicine, 2007, 39, 229-238.	3.8	234
29	Genetic Gating of Human Fear Learning and Extinction. Psychological Science, 2009, 20, 198-206.	3.3	228
30	Whole-exome sequencing identifies novel MPL and JAK2 mutations in triple-negative myeloproliferative neoplasms. Blood, 2016, 127, 325-332.	1.4	228
31	Differential effects of intracerebroventiricular colchicine administration on the expression of mrnas for neuropeptides and neurotransmitter enzymes, with specila emphasis on galanin: An in situ Hybridization Study. Synapse, 1990, 6, 369-391.	1.2	217
32	The Wilms tumour gene WT1 is expressed in murine mesoderm–derived tissues and mutated in a human mesothelioma. Nature Genetics, 1993, 4, 415-420.	21.4	199
33	Leptin secretion from adipose tissue in women. Relationship to plasma levels and gene expression Journal of Clinical Investigation, 1997, 99, 2398-2404.	8.2	184
34	Genetic and epigenetic associations of MAOA and NR3C1 with depression and childhood adversities. International Journal of Neuropsychopharmacology, 2013, 16, 1513-1528.	2.1	182
35	Genome-wide association study of 40,000 individuals identifies two novel loci associated with bipolar disorder. Human Molecular Genetics, 2016, 25, 3383-3394.	2.9	182
36	The Swedish Twin Registry in the Third Millennium: An Update. Twin Research and Human Genetics, 2006, 9, 875-882.	0.6	182

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37	Chapter 4 Coexistence of neuronal messengers â€" an overview. Progress in Brain Research, 1986, 68, 33-70.	1.4	180
38	Genetic pleiotropy between multiple sclerosis and schizophrenia but not bipolar disorder: differential involvement of immune-related gene loci. Molecular Psychiatry, 2015, 20, 207-214.	7.9	173
39	Epigenetic aberrations in leukocytes of patients with schizophrenia: association of global DNA methylation with antipsychotic drug treatment and disease onset. FASEB Journal, 2012, 26, 2712-2718.	0.5	170
40	Distinct Contributions of the Dorsolateral Prefrontal and Orbitofrontal Cortex during Emotion Regulation. PLoS ONE, 2012, 7, e48107.	2.5	169
41	Mapping of Spinocerebellar Ataxia 13 to Chromosome 19q13.3-q13.4 in a Family with Autosomal Dominant Cerebellar Ataxia and Mental Retardation. American Journal of Human Genetics, 2000, 67, 229-235.	6.2	166
42	Adiponectin in renal disease: Relationship to phenotype and genetic variation in the gene encoding adiponectin. Kidney International, 2004, 65, 274-281.	5.2	160
43	Delayed cytokine expression in rat brain following experimental contusion. Journal of Neurosurgery, 1997, 86, 493-504.	1.6	159
44	Assessment of Response to Lithium Maintenance Treatment in Bipolar Disorder: A Consortium on Lithium Genetics (ConLiGen) Report. PLoS ONE, 2013, 8, e65636.	2.5	156
45	Genome integrity of myeloproliferative neoplasms in chronic phase and during disease progression. Blood, 2011, 118, 167-176.	1.4	153
46	Detection of expanded CAG repeats in Bipolar Affective Disorder using the repeat expansion detection (RED) method. Neurobiology of Disease, 1995, 2, 55-62.	4.4	148
47	Low leptin gene expression and hyperleptinemia in chronic renal failure. Kidney International, 1998, 54, 1267-1275.	5. 2	148
48	Ropeginterferon alfa-2b, a novel IFN \hat{l} ±-2b, induces high response rates with low toxicity in patients with polycythemia vera. Blood, 2015, 126, 1762-1769.	1.4	142
49	Long-term lithium treatment in bipolar disorder is associated with longer leukocyte telomeres. Translational Psychiatry, 2013, 3, e261-e261.	4.8	134
50	A subpopulation of dopaminergic neurons in rat ventral mesencephalon contains both neurotensin and cholecystokinin. Brain Research, 1988, 455, 88-98.	2.2	133
51	CRY2 Is Associated with Depression. PLoS ONE, 2010, 5, e9407.	2.5	132
52	Localization of neuropeptide receptor mRNA in rat brain: Initial observations using probes for neurotensin and substance P receptors. Neuroscience Letters, 1990, 120, 134-138.	2.1	124
53	<i>PER2</i> variantion is associated with depression vulnerability. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2010, 153B, 570-581.	1.7	118
54	A gene encoding a fibroblast growth factor receptor isolated from the Huntington disease gene region of human chromosome 4. Genomics, 1991, 11, 1133-1142.	2.9	115

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55	Calcitonin Gene-Related Peptide in the Brain, Spinal Cord, and Some Peripheral Systems. Annals of the New York Academy of Sciences, 1992, 657, 119-134.	3.8	113
56	Circulating Levels of Visfatin/Pre–B-Cell Colony–Enhancing Factor 1 in Relation to Genotype, GFR, Body Composition, and Survival in Patients With CKD. American Journal of Kidney Diseases, 2007, 49, 237-244.	1.9	109
57	No evidence for heritability of Parkinson disease in Swedish twins. Neurology, 2004, 63, 305-311.	1.1	108
58	A functional variant of the myeloperoxidase gene is associated with cardiovascular disease in end-stage renal disease patients. Kidney International, 2003, 63, S172-S176.	5. 2	105
59	Association of Polygenic Score for Schizophrenia and HLA Antigen and Inflammation Genes With Response to Lithium in Bipolar Affective Disorder. JAMA Psychiatry, 2018, 75, 65-74.	11.0	102
60	Increased Sensitivity to Thermal Pain Following a Single Opiate Dose Is Influenced by the COMT val158met Polymorphism. PLoS ONE, 2009, 4, e6016.	2.5	97
61	Clinical and genetic outcome determinants of Internetâ€and groupâ€based cognitive behavior therapy for social anxiety disorder. Acta Psychiatrica Scandinavica, 2012, 126, 126-136.	4.5	96
62	An expanded CAG repeat sequence in spinocerebellar ataxia type 7 Genome Research, 1996, 6, 965-971.	5.5	94
63	The KMO allele encoding Arg452 is associated with psychotic features in bipolar disorder type 1, and with increased CSF KYNA level and reduced KMO expression. Molecular Psychiatry, 2014, 19, 334-341.	7.9	91
64	Increase of insulin-like growth factor (IGF)-1, IGF binding protein-2 and â^4 mRNAs following cerebral contusion. Molecular Brain Research, 1996, 38, 285-293.	2.3	87
65	Conditioned Pain Modulation Is Associated with Common Polymorphisms in the Serotonin Transporter Gene. PLoS ONE, 2011, 6, e18252.	2.5	87
66	The Risk of Offspring Psychiatric Disorders in the Setting of Maternal Obesity and Diabetes. Pediatrics, 2018, 142, .	2.1	84
67	Neuropeptide tyrosine in the rat adrenal gland—immunohistochemical and in situ hybridization studies. Neuroscience, 1988, 24, 337-349.	2.3	82
68	Multigenic Control of Disease Severity after Virulent Mycobacterium tuberculosis Infection in Mice. Infection and Immunity, 2003, 71, 126-131.	2.2	81
69	The COMTval158met polymorphism is associated with symptom relief during exposure-based cognitive-behavioral treatment in panic disorder. BMC Psychiatry, 2010, 10, 99.	2.6	81
70	Severity of Tuberculosis in Mice is Linked to Distal Chromosome 3 and Proximal Chromosome 9. Journal of Infectious Diseases, 1999, 180, 150-155.	4.0	78
71	Spinocerebellar ataxia with sensory neuropathy (SCA25) maps to chromosome 2p. Annals of Neurology, 2004, 55, 97-104.	5.3	78
72	CLOCK is suggested to associate with comorbid alcohol use and depressive disorders. Journal of Circadian Rhythms, 2014, 8, 1.	1.3	78

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73	Neuropeptide gene expression in hypothalamic magnocellular neurons of normal and hypophysectomized rats: A combined immunohistochemical and in situ hybridization study. Neuroscience, 1990, 36, 181-199.	2.3	74
74	Neuropeptide Y and catecholamine synthesizing enzymes and their mRNAs in rat sympathetic neurons and adrenal glands: Studies on expression, synthesis and axonal transport after pharmacological and experimental manipulations using hybridization techniques and radioimmunoassay. Neuroscience, 1991, 41, 753-766.	2.3	71
75	Gender differences in heritability of depressive symptoms in the elderly. Psychological Medicine, 2004, 34, 471-479.	4.5	71
76	CRY2 Is Associated with Rapid Cycling in Bipolar Disorder Patients. PLoS ONE, 2010, 5, e12632.	2.5	71
77	Maturation of the hypothalamic arcuate agouti-related protein system during postnatal development in the mouse. Developmental Brain Research, 2005, 155, 147-154.	1.7	70
78	Analysis of expression of cholecystokinin in dopamine cells in the ventral mesencephalon of several species and in humans with schizophrenia Proceedings of the National Academy of Sciences of the United States of America, 1990, 87, 8427-8431.	7.1	69
79	Leptin gene is expressed in rat brown adipose tissue at birth. FASEB Journal, 1997, 11, 382-387.	0.5	68
80	Fat tissue accumulation during peritoneal dialysis is associated with a polymorphism in uncoupling protein 2. Kidney International, 2000, 57, 1713-1719.	5.2	68
81	Chemical analysis of osmium tetroxide staining in adipose tissue using imaging ToF-SIMS. Histochemistry and Cell Biology, 2009, 132, 105-115.	1.7	68
82	Large-scale genotyping of single nucleotide polymorphisms by Pyrosequencing? and validation against the 5?nuclease (Taqmanı̈ $^1_2/^2$) assay. Human Mutation, 2002, 19, 395-401.	2.5	66
83	CCR5 Deletion Protects Against Inflammation-Associated Mortality in Dialysis Patients. Journal of the American Society of Nephrology: JASN, 2009, 20, 1641-1649.	6.1	66
84	Adenovirus-36 Is Associated with Obesity in Children and Adults in Sweden as Determined by Rapid ELISA. PLoS ONE, 2012, 7, e41652.	2.5	66
85	Genetic Variation in the Serotonin Transporter Gene (5-HTTLPR, Rs25531) Influences the Analgesic Response to the Short Acting Opioid Remifentanil in Humans. Molecular Pain, 2009, 5, 1744-8069-5-37.	2.1	65
86	Kynurenine 3-monooxygenase polymorphisms: relevance for kynurenic acid synthesis in patients with schizophrenia and healthy controls. Journal of Psychiatry and Neuroscience, 2012, 37, 53-57.	2.4	65
87	Androgen receptor trinucleotide repeat polymorphism and personality traits. Psychiatric Genetics, 2001, 11, 19-23.	1.1	63
88	Perception of Thermal Pain and the Thermal Grill Illusion Is Associated with Polymorphisms in the Serotonin Transporter Gene. PLoS ONE, 2011, 6, e17752.	2.5	61
89	Basal cell carcinoma and variants in genes coding for immune response, DNA repair, folate and iron metabolism. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2005, 574, 105-111.	1.0	60
90	Early exposure to antibiotic drugs and risk for psychiatric disorders: a population-based study. Translational Psychiatry, 2019, 9, 317.	4.8	60

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91	Expression and localization of gastrin messenger RNA and peptide in spermatogenic cells Journal of Clinical Investigation, 1990, 86, 660-669.	8.2	60
92	Reduced gene expression of UCP2 but not UCP3 in skeletal muscle of human obese subjects. Diabetologia, 1998, 41, 935-939.	6.3	58
93	Hypothalamic CART and serum leptin levels are reduced in the anorectic (anx/anx) mouse. Molecular Brain Research, 2000, 84, 97-105.	2.3	58
94	Truncation of the Shaker-like voltage-gated potassium channel, Kv1.1, causes megencephaly. European Journal of Neuroscience, 2003, 18, 3231-3240.	2.6	58
95	5-HTTLPR and COMTval158met genotype gate amygdala reactivity and habituation. Biological Psychology, 2011, 87, 106-112.	2.2	58
96	Genome-wide association study of panic disorder reveals genetic overlap with neuroticism and depression. Molecular Psychiatry, 2021, 26, 4179-4190.	7.9	58
97	Reduced gene expression of adiponectin in fat tissue from patients with end-stage renal disease. Kidney International, 2004, 66, 46-50.	5.2	57
98	Amygdala-dependent fear conditioning in humans is modulated by the BDNFval66met polymorphism Behavioral Neuroscience, 2010, 124, 9-15.	1.2	57
99	Analysis of a murine male germ cell-specific transcript that encodes a putative zinc finger protein. Molecular Reproduction and Development, 1991, 30, 173-181.	2.0	56
100	Hypothalamic neurohistochemistry of the murineanorexia (anx/anx) mutation: Altered processing of neuropeptide Y in the arcuate nucleus., 1997, 387, 124-135.		56
101	Changes in Neuropeptide Y Receptors and Pro-Opiomelanocortin in the Anorexia (anx/anx) Mouse Hypothalamus. Journal of Neuroscience, 1999, 19, 7130-7139.	3.6	56
102	Long repeat tracts atSCA8 in major psychosis. American Journal of Medical Genetics Part A, 2000, 96, 873-876.	2.4	55
103	Two commonly expanded CAG/CTG repeat loci: involvement in affective disorders?. Molecular Psychiatry, 1998, 3, 405-410.	7.9	54
104	Genetics of affective disorders. European Neuropsychopharmacology, 2001, 11, 385-394.	0.7	54
105	Gene-to-gene interactions regulate endogenous pain modulation in fibromyalgia patients and healthy controls—antagonistic effects between opioid and serotonin-related genes. Pain, 2017, 158, 1194-1203.	4.2	54
106	Mutational landscape of the transcriptome offers putative targets for immunotherapy of myeloproliferative neoplasms. Blood, 2019, 134, 199-210.	1.4	54
107	Distribution and cellular localization of DARPP-32 mRNA in rat brain. Molecular Brain Research, 1990, 7, 139-149.	2.3	53
108	Expanded trinucleotide CAG repeats in families with bipolar affective disorder. Biological Psychiatry, 1997, 42, 1115-1122.	1.3	53

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109	CDKN2A/p16INK4a expression is associated with vascular progeria in chronic kidney disease. Aging, 2017, 9, 494-507.	3.1	52
110	Calcitonin gene-related peptide and its mRNA in pulmonary neuroendocrine cells and ganglia. Histochemistry, 1991, 96, 311-315.	1.9	51
111	Pyrosequencing?-based SNP allele frequency estimation in DNA pools. Human Mutation, 2004, 23, 92-97.	2.5	51
112	Visfatin is increased in chronic kidney disease patients with poor appetite and correlates negatively with fasting serum amino acids and triglyceride levels. Nephrology Dialysis Transplantation, 2010, 25, 901-906.	0.7	50
113	Tyrosine 3-hydroxylase in rat brain and adrenal medulla: hybridization histochemistry and immunohistochemistry combined with retrograde tracing Proceedings of the National Academy of Sciences of the United States of America, 1986, 83, 6208-6212.	7.1	49
114	Localization of cholesterol, amyloid and glia in Alzheimer's disease transgenic mouse brain tissue using time-of-flight secondary ion mass spectrometry (ToF-SIMS) and immunofluorescence imaging. Acta Neuropathologica, 2013, 125, 145-157.	7.7	48
115	Schizophrenia and neurotrophin-3 alleles. Acta Psychiatrica Scandinavica, 1997, 95, 414-419.	4.5	47
116	Leptin and its potential role in human obesity. Journal of Internal Medicine, 1999, 245, 643-652.	6.0	47
117	Seasonal Affective Disorder and Serotonin-Related Polymorphisms. Neurobiology of Disease, 2001, 8, 351-357.	4.4	47
118	Preâ€fibrotic/early primary myelofibrosis vs. WHOâ€defined essential thrombocythemia: The impact of minor clinical diagnostic criteria on the outcome of the disease. American Journal of Hematology, 2017, 92, 885-891.	4.1	47
119	Hypothalamic mitochondrial dysfunction associated with anorexia in the <i>anx/anx</i> mouse. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 18108-18113.	7.1	46
120	Contribution of Rare Copy Number Variants toÂBipolar Disorder Risk Is Limited to Schizoaffective Cases. Biological Psychiatry, 2019, 86, 110-119.	1.3	45
121	Molecular responses and chromosomal aberrations in patients with polycythemia vera treated with pegâ€prolineâ€interferon alphaâ€2b. American Journal of Hematology, 2015, 90, 288-294.	4.1	44
122	Association of polygenic score for major depression with response to lithium in patients with bipolar disorder. Molecular Psychiatry, 2021, 26, 2457-2470.	7.9	44
123	Leukotriene C4 binding sites in the rat central nervous system. European Journal of Pharmacology, 1986, 122, 251-257.	3.5	43
124	Cognitive manic symptoms associated with the <i>P2RX7</i> gene in bipolar disorder. Bipolar Disorders, 2011, 13, 500-508.	1.9	43
125	The symptomatic profile of panic disorder is shaped by the 5-HTTLPR polymorphism. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2009, 33, 1479-1483.	4.8	42
126	Cystic fibrosis mRNA expression in rat brain. NeuroReport, 1997, 8, 535-539.	1.2	41

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127	Noradrenaline Reuptake Inhibition Enhances the Antipsychotic-like Effect of Raclopride and Potentiates D2-blockage–induced Dopamine Release in the Medial Prefrontal Cortex of the Rat. Neuropsychopharmacology, 2002, 27, 691-698.	5.4	41
128	The translocator protein gene is associated with symptom severity and cerebral pain processing in fibromyalgia. Brain, Behavior, and Immunity, 2016, 58, 218-227.	4.1	39
129	PO protein peptide 180-199 together with pertussis toxin induces experimental autoimmune neuritis in resistant C57BL/6 mice. Journal of Neuroscience Research, 2000, 62, 717-721.	2.9	38
130	A Common Hormone-Sensitive Lipase i6 Gene Polymorphism Is Associated With Decreased Human Adipocyte Lipolytic Function. Diabetes, 2001, 50, 2410-2413.	0.6	37
131	The serotonin transporter promoter repeat length polymorphism, seasonal affective disorder and seasonality. Psychological Medicine, 2003, 33, 785-792.	4.5	37
132	MAOA haplotypes associated with thrombocyte-MAO activity. BMC Genetics, 2005, 6, 46.	2.7	37
133	Simultaneous Imaging of Amyloid- \hat{l}^2 and Lipids in Brain Tissue Using Antibody-Coupled Liposomes and Time-of-Flight Secondary Ion Mass Spectrometry. Journal of the American Chemical Society, 2014, 136, 9973-9981.	13.7	37
134	A Triplet Repeat on 17q Accounts for Most Expansions Detected by the Repeat-Expansion–Detection Technique. American Journal of Human Genetics, 1998, 62, 1548-1551.	6.2	36
135	Genetic Polymorphisms in Monoamine Systems and Outcome of Cognitive Behavior Therapy for Social Anxiety Disorder. PLoS ONE, 2013, 8, e79015.	2.5	35
136	<i>BDNF</i> val66met affects neural activation pattern during fear conditioning and 24 h delayed fear recall. Social Cognitive and Affective Neuroscience, 2015, 10, 664-671.	3.0	35
137	A genome-wide association meta-analysis of prognostic outcomes following cognitive behavioural therapy in individuals with anxiety and depressive disorders. Translational Psychiatry, 2019, 9, 150.	4.8	35
138	P2RX7: Expression Responds to Sleep Deprivation and Associates with Rapid Cycling in Bipolar Disorder Type 1. PLoS ONE, 2012, 7, e43057.	2.5	35
139	Substance P mRNA is present in a population of CGRP-immunoreactive cholinergic postganglionic sympathetic neurons of the cat: evidence from combined in situ hybridization and immunohistochemistry. Neuroscience Letters, 1989, 107, 1-5.	2.1	34
140	Selective noradrenaline reuptake inhibition enhances serotonergic neuronal activity and transmitter release in the rat forebrain. Journal of Neural Transmission, 2004, 111, 127-139.	2.8	34
141	Alterations of arcuate nucleus neuropeptidergic development in contactin-deficient mice: comparison with anorexia and food-deprived mice. European Journal of Neuroscience, 2005, 22, 3217-3228.	2.6	34
142	Mitochondrial DNA copy number is associated with psychosis severity and anti-psychotic treatment. Scientific Reports, 2018, 8, 12743.	3.3	34
143	Evidence for coexistence between calcitonin gene-related peptide and serotonin in the bulbospinal pathway in the monkey. Brain Research, 1990, 532, 47-57.	2.2	33
144	Effect of in vitro promoter methylation and CGG repeat expansion on FMR- 1 expression. Nucleic Acids Research, 1997, 25, 2883-2887.	14.5	33

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145	The effect of the beta2 adrenoceptor gene Thr164lle polymorphism on human adipose tissue lipolytic function. British Journal of Pharmacology, 2001, 133, 708-712.	5.4	33
146	Serotonin-1A Receptor Polymorphism (rs6295) Associated with Thermal Pain Perception. PLoS ONE, 2012, 7, e43221.	2.5	33
147	Possible pathologic involvement of receptor for advanced glycation end products (RAGE) for development of encapsulating peritoneal sclerosis in Japanese CAPD patients. Clinical Nephrology, 2004, 62, 455-460.	0.7	32
148	Rapid and efficient construction of yeast artificial chromosome contigs in the mouse genome with interspersed repetitive sequence PCR (IRS-PCR): Generation of a 5-cM, >5 megabase contig on mouse Chromosome 1. Mammalian Genome, 1994, 5, 597-607.	2.2	31
149	Growing triplet repeats. Nature Genetics, 1994, 7, 124-124.	21.4	31
150	MRI and in situ hybridization reveal early disturbances in brain size and gene expression in the megencephalic (mceph/mceph) mouse. European Journal of Neuroscience, 2003, 18, 3218-3230.	2.6	31
151	Carbamazepine protects against megencephaly and abnormal expression of BDNF and Nogo signaling components in the mceph/mceph mouse. Neurobiology of Disease, 2006, 24, 374-383.	4.4	31
152	Aberrant inflammatory profile in acute but not recovered anorexia nervosa. Brain, Behavior, and Immunity, 2020, 88, 718-724.	4.1	31
153	Protein kinase C in the developing kidney: Isoform expression and effects of ceramide and PKC inhibitors. Kidney International, 1997, 52, 901-910.	5.2	30
154	Large CAG/CTG repeats are associated with childhood-onset schizophrenia. Molecular Psychiatry, 1998, 3, 321-327.	7.9	30
155	Genetic survival analysis of age-at-onset of bipolar disorder: evidence for anticipation or cohort effect in families. Psychiatric Genetics, 2001, 11, 129-137.	1.1	30
156	Changes in fat mass after initiation of maintenance dialysis is influenced by the uncoupling protein 2 exon 8 insertion/deletion polymorphism. Nephrology Dialysis Transplantation, 2006, 22, 196-202.	0.7	30
157	Peroxisome proliferator-activated receptor \hat{l}^3 polymorphisms affect systemic inflammation and survival in end-stage renal disease patients starting renal replacement therapy. Atherosclerosis, 2005, 182, 105-111.	0.8	29
158	Effects of a synbiotic on symptoms, and daily functioning in attention deficit hyperactivity disorder – A double-blind randomized controlled trial. Brain, Behavior, and Immunity, 2020, 89, 9-19.	4.1	29
159	Analysis of the Influence of microRNAs in Lithium Response in Bipolar Disorder. Frontiers in Psychiatry, 2018, 9, 207.	2.6	28
160	Molecular Interpretation of Expanded RED Products in Bipolar Disorder by CAG/CTG Repeats Located at Chromosomes 17q and 18q. Neurobiology of Disease, 1999, 6, 424-432.	4.4	27
161	Multidrug Resistance 1 (MDR1) Gene Polymorphisms in Childhood Drug-Resistant Epilepsy. Journal of Child Neurology, 2010, 25, 1485-1490.	1.4	27
162	Expression of Inflammatory and Insulin Signaling Genes in Adipose Tissue in Response to Elective Surgery. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 3460-3469.	3.6	27

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163	Expression of $\hat{l}\pm$ -synuclein in the human brain: relation to Lewy body disease. Molecular Brain Research, 2001, 92, 58-65.	2.3	26
164	Genetic approaches in the clinical investigation of complex disorders: Malnutrition, inflammation, and atherosclerosis (MIA) as a prototype. Kidney International, 2003, 63, S162-S167.	5.2	26
165	Seasonal affective disorder and the G-protein \hat{I}^2 -3-subunit C825T polymorphism. Biological Psychiatry, 2004, 55, 317-319.	1.3	26
166	TOF-SIMS analysis of adipose tissue from patients with chronic kidney disease. Applied Surface Science, 2008, 255, 1177-1180.	6.1	26
167	Plasma neurofilament light chain concentration is increased in anorexia nervosa. Translational Psychiatry, 2019, 9, 180.	4.8	26
168	Genetics of response to cognitive behavior therapy in adults with major depression: a preliminary report. Molecular Psychiatry, 2019, 24, 484-490.	7.9	26
169	Increased Leptin Messenger RNA and Serum Leptin Levels in Children with Prader-Willi Syndrome and Nonsyndromal Obesity. Pediatric Research, 1997, 42, 593-596.	2.3	26
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