Martin Schalling

List of Publications by Year in Descending Order

Source: https://exaly.com/author-pdf/5446/martin-schalling-publications-by-year.pdf

Version: 2024-04-28

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

69 25,412 155 307 h-index g-index citations papers 28,927 7.13 323 7.5 avg, IF L-index ext. citations ext. papers

#	Paper	IF	Citations
307	Using polygenic scores and clinical data for bipolar disorder patient stratification and lithium response prediction: machine learning approach <i>British Journal of Psychiatry</i> , 2022 , 1-10	5.4	1
306	Improving lithium dose prediction using population pharmacokinetics and pharmacogenomics: a cohort genome-wide association study in Sweden <i>Lancet Psychiatry,the</i> , 2022 , 9, 447-457	23.3	0
305	Association of polygenic score for major depression with response to lithium in patients with bipolar disorder. <i>Molecular Psychiatry</i> , 2021 , 26, 2457-2470	15.1	17
304	Genome-wide association study of panic disorder reveals genetic overlap with neuroticism and depression. <i>Molecular Psychiatry</i> , 2021 , 26, 4179-4190	15.1	8
303	The Anorectic Phenotype of the anx/anx Mouse Is Associated with Hypothalamic Dysfunction. <i>Neuromethods</i> , 2021 , 297-317	0.4	
302	Combining schizophrenia and depression polygenic risk scores improves the genetic prediction of lithium response in bipolar disorder patients. <i>Translational Psychiatry</i> , 2021 , 11, 606	8.6	1
301	Polymorphisms in the HTR2A and HTR3A Genes Contribute to Pain in TMD Myalgia <i>Frontiers in Oral Health</i> , 2021 , 2, 647924	0.8	1
300	Genome-wide association study of more than 40,000 bipolar disorder cases provides new insights into the underlying biology. <i>Nature Genetics</i> , 2021 , 53, 817-829	36.3	83
299	GRK3 deficiency elicits brain immune activation and psychosis. <i>Molecular Psychiatry</i> , 2021 ,	15.1	2
298	Serotonergic gene-to-gene interaction is associated with mood and GABA concentrations but not with pain-related cerebral processing in fibromyalgia subjects and healthy controls. <i>Molecular Brain</i> , 2021 , 14, 81	4.5	1
297	Polymorphisms of the Eppioid receptor gene influence cerebral pain processing in fibromyalgia. <i>European Journal of Pain</i> , 2021 , 25, 398-414	3.7	2
296	Germline genetic factors influence the outcome of interferon-Etherapy in polycythemia vera. <i>Blood</i> , 2021 , 137, 387-391	2.2	5
295	Biomimetics provides lessons from nature for contemporary ways to improve human health. <i>Journal of Clinical and Translational Science</i> , 2021 , 5, e128	0.4	O
294	Prediction of lithium response using genomic data. Scientific Reports, 2021, 11, 1155	4.9	1
293	HLA-DRB1 and HLA-DQB1 genetic diversity modulates response to lithium in bipolar affective disorders. <i>Scientific Reports</i> , 2021 , 11, 17823	4.9	1
292	Lithium and the Interplay Between Telomeres and Mitochondria in Bipolar Disorder. <i>Frontiers in Psychiatry</i> , 2020 , 11, 586083	5	6
291	Aberrant inflammatory profile in acute but not recovered anorexia nervosa. <i>Brain, Behavior, and Immunity</i> , 2020 , 88, 718-724	16.6	13

(2018-2020)

2 90	Effects of a synbiotic on symptoms, and daily functioning in attention deficit hyperactivity disorder - A double-blind randomized controlled trial. <i>Brain, Behavior, and Immunity</i> , 2020 , 89, 9-19	16.6	7
289	Expression of telomerase reverse transcriptase positively correlates with duration of lithium treatment in bipolar disorder. <i>Psychiatry Research</i> , 2020 , 286, 112865	9.9	7
288	EAAT2 as a Research Target in Bipolar Disorder and Unipolar Depression: A Systematic Review. <i>Molecular Neuropsychiatry</i> , 2020 , 5, 44-59	4.9	13
287	AKT1 and genetic vulnerability to bipolar disorder. <i>Psychiatry Research</i> , 2020 , 284, 112677	9.9	1
286	Single-nucleotide polymorphism in the human TIA1 gene interacts with stressful life events to predict the development of pathological anxiety symptoms in a Swedish population. <i>Journal of Affective Disorders</i> , 2020 , 260, 597-603	6.6	1
285	A genome-wide association meta-analysis of prognostic outcomes following cognitive behavioural therapy in individuals with anxiety and depressive disorders. <i>Translational Psychiatry</i> , 2019 , 9, 150	8.6	24
284	Genetic variant in SLC1A2 is associated with elevated anterior cingulate cortex glutamate and lifetime history of rapid cycling. <i>Translational Psychiatry</i> , 2019 , 9, 149	8.6	9
283	Mutational landscape of the transcriptome offers putative targets for immunotherapy of myeloproliferative neoplasms. <i>Blood</i> , 2019 , 134, 199-210	2.2	34
282	Genome-wide association study identifies 30 loci associated with bipolar disorder. <i>Nature Genetics</i> , 2019 , 51, 793-803	36.3	662
281	Investigating polygenic burden in age at disease onset in bipolar disorder: Findings from an international multicentric study. <i>Bipolar Disorders</i> , 2019 , 21, 68-75	3.8	15
2 80	Plasma neurofilament light chain concentration is increased in anorexia nervosa. <i>Translational Psychiatry</i> , 2019 , 9, 180	8.6	13
279	Early exposure to antibiotic drugs and risk for psychiatric disorders: a population-based study. <i>Translational Psychiatry</i> , 2019 , 9, 317	8.6	24
278	Contribution of Rare Copy Number Variants to Bipolar Disorder Risk Is Limited to Schizoaffective Cases. <i>Biological Psychiatry</i> , 2019 , 86, 110-119	7.9	20
277	Sex-specific effects of gain-of-function P2RX7 variation on bipolar disorder. <i>Journal of Affective Disorders</i> , 2019 , 245, 597-601	6.6	4
276	Impact of white blood cells on thrombotic risk in patients with optimized platelet count in essential thrombocythemia. <i>European Journal of Haematology</i> , 2018 , 101, 131	3.8	10
275	The WHO diagnostic criteria for polycythemia vera-role of red cell mass versus hemoglobin/hematocrit level and morphology. <i>Annals of Hematology</i> , 2018 , 97, 1581-1590	3	7
274	Analysis of the Influence of microRNAs in Lithium Response in Bipolar Disorder. <i>Frontiers in Psychiatry</i> , 2018 , 9, 207	5	15
273	Analysis of shared heritability in common disorders of the brain. <i>Science</i> , 2018 , 360,	33.3	666

272	Mitochondrial DNA copy number is associated with psychosis severity and anti-psychotic treatment. <i>Scientific Reports</i> , 2018 , 8, 12743	4.9	23
271	The Risk of Offspring Psychiatric Disorders in the Setting of Maternal Obesity and Diabetes. <i>Pediatrics</i> , 2018 , 142,	7.4	55
27 0	Association of Polygenic Score for Schizophrenia and HLA Antigen and Inflammation Genes With Response to Lithium in Bipolar Affective Disorder: A Genome-Wide Association Study. <i>JAMA Psychiatry</i> , 2018 , 75, 65-74	14.5	75
269	Long-Term Efficacy and Safety of Ropeginterferon Alfa-2b in Patients with Polycythemia Vera II Final Phase I/II Peginvera Study Results. <i>Blood</i> , 2018 , 132, 3030-3030	2.2	5
268	Effect of Ropeginterferon Alfa-2b in Prefibrotic Primary Myelofibrosis. <i>Blood</i> , 2018 , 132, 3029-3029	2.2	5
267	Reduced metabolism in the hypothalamus of the anorectic mouse. <i>Journal of Endocrinology</i> , 2017 , 233, 15-24	4.7	10
266	Gene-to-gene interactions regulate endogenous pain modulation in fibromyalgia patients and healthy controls-antagonistic effects between opioid and serotonin-related genes. <i>Pain</i> , 2017 , 158, 119	94 ⁸ 120	3 ³¹
265	Pre-fibrotic/early primary myelofibrosis vs. WHO-defined essential thrombocythemia: The impact of minor clinical diagnostic criteria on the outcome of the disease. <i>American Journal of Hematology</i> , 2017 , 92, 885-891	7.1	29
264	Troponin T levels associated with genetic variants in NOTCH2 and MTNR1B in women with psychosis. <i>Psychiatry Research</i> , 2017 , 250, 217-220	9.9	1
263	Plasma GDF15 level is elevated in psychosis and inversely correlated with severity. <i>Scientific Reports</i> , 2017 , 7, 7906	4.9	5
262	Genetic variants of increased waist circumference in psychosis. <i>Psychiatric Genetics</i> , 2017 , 27, 210-218	2.9	3
261	Essential thrombocythemia vs. pre-fibrotic/early primary myelofibrosis: discrimination by laboratory and clinical data. <i>Blood Cancer Journal</i> , 2017 , 7, 643	7	5
2 60	Impact of white blood cell counts at diagnosis and during follow-up in patients with essential thrombocythaemia and prefibrotic primary myelofibrosis. <i>British Journal of Haematology</i> , 2017 , 179, 166-169	4.5	15
259	expression is associated with vascular progeria in chronic kidney disease. <i>Aging</i> , 2017 , 9, 494-507	5.6	40
258	Increased Telomere Attrition After Renal Transplantation-Impact of Antimetabolite Therapy. <i>Transplantation Direct</i> , 2016 , 2, e116	2.3	12
257	Genome-wide association study of 40,000 individuals identifies two novel loci associated with bipolar disorder. <i>Human Molecular Genetics</i> , 2016 , 25, 3383-3394	5.6	125
256	Genetic variants associated with response to lithium treatment in bipolar disorder: a genome-wide association study. <i>Lancet, The</i> , 2016 , 387, 1085-1093	40	216
255	hTERT genetic variation in depression. <i>Journal of Affective Disorders</i> , 2016 , 189, 62-9	6.6	23

(2015-2016)

254	Negative Myeloproliferative Neoplasms - Data from the Austrian Reclassification Project. <i>Blood</i> , 2016 , 128, 4252-4252	2.2	2
253	Influence of Polymorphisms in the HTR3A and HTR3B Genes on Experimental Pain and the Effect of the 5-HT3 Antagonist Granisetron. <i>PLoS ONE</i> , 2016 , 11, e0168703	3.7	9
252	Survival in Essential Thrombocythemia and Prefibrotic Primary Myelofibrosis - Correlation of Clinical Phenotype with Histomorphological Diagnosis. <i>Blood</i> , 2016 , 128, 1932-1932	2.2	
251	Impact of High Molecular Risk Mutations on Overall Survival in WHO-Defined Essential Thrombocythemia and Prefibrotic Primary Myelofibrosis. <i>Blood</i> , 2016 , 128, 1931-1931	2.2	
250	Monitoring Molecular Response, Minimal Residual Disease and Clonal Structures in Polycythemia Vera Patients Treated with Interferon Alpha. <i>Blood</i> , 2016 , 128, 1944-1944	2.2	
249	Melatonin receptor 1B gene associated with hyperglycemia in bipolar disorder. <i>Psychiatric Genetics</i> , 2016 , 26, 136-9	2.9	2
248	Imaging of amyloid-In Alzheimer disease transgenic mouse brains with ToF-SIMS using immunoliposomes. <i>Biointerphases</i> , 2016 , 11, 02A312	1.8	11
247	Whole-exome sequencing identifies novel MPL and JAK2 mutations in triple-negative myeloproliferative neoplasms. <i>Blood</i> , 2016 , 127, 325-32	2.2	176
246	The translocator protein gene is associated with symptom severity and cerebral pain processing in fibromyalgia. <i>Brain, Behavior, and Immunity,</i> 2016 , 58, 218-227	16.6	26
245	Depression-associated ARNTL and PER2 genetic variants in psychotic disorders. <i>Chronobiology International</i> , 2015 , 32, 579-84	3.6	9
244	BDNFval66met affects neural activation pattern during fear conditioning and 24 h delayed fear recall. <i>Social Cognitive and Affective Neuroscience</i> , 2015 , 10, 664-71	4	32
243	A 16-year-old girl with anti-NMDA-receptor encephalitis and family history of psychotic disorders. <i>Acta Neuropsychiatrica</i> , 2015 , 27, 375-9	3.9	11
242	Ropeginterferon alfa-2b, a novel IFNDb, induces high response rates with low toxicity in patients with polycythemia vera. <i>Blood</i> , 2015 , 126, 1762-9	2.2	106
241	Mood Stabilizers and the Influence on Global Leukocyte DNA Methylation in Bipolar Disorder. <i>Molecular Neuropsychiatry</i> , 2015 , 1, 76-81	4.9	16
240	Association of brain-derived neurotrophic factor (BDNF) Val66Met polymorphism with early-onset bipolar disorder. <i>Bipolar Disorders</i> , 2015 , 17, 645-52	3.8	15
239	Diabetes and glucose disturbances in patients with psychosis in Sweden. <i>BMJ Open Diabetes Research and Care</i> , 2015 , 3, e000120	4.5	4
238	Molecular responses and chromosomal aberrations in patients with polycythemia vera treated with peg-proline-interferon alpha-2b. <i>American Journal of Hematology</i> , 2015 , 90, 288-94	7.1	38
237	Glucose intolerance and pancreatic Etell dysfunction in the anorectic anx/anx mouse. <i>American Journal of Physiology - Endocrinology and Metabolism</i> , 2015 , 309, E418-27	6	9

236	Genetic pleiotropy between multiple sclerosis and schizophrenia but not bipolar disorder: differential involvement of immune-related gene loci. <i>Molecular Psychiatry</i> , 2015 , 20, 207-14	15.1	144
235	Four Weeks Administration Schedule of Ropeginterferon Alfa-2b (AOP2014/P1101) in Polycythemia Very Patients Allows Maintaining of Efficacy with Favorable Toxicity Profile in the Phase I/II Peginvera Stud. <i>Blood</i> , 2015 , 126, 1603-1603	2.2	1
234	Fusion Gene Detection Using Whole Transcriptome Analysis in Patients with Chronic Myeloproliferative Neoplasms and Secondary Acute Myeloid Leukemia. <i>Blood</i> , 2015 , 126, 4093-4093	2.2	1
233	Whole Exome Sequencing Identifies Novel MPL and JAK2 M utations in Triple Negative Myeloproliferative Neoplasms. <i>Blood</i> , 2015 , 126, 606-606	2.2	1
232	Long Term Efficacy and Safety Results and Analysis of Dose Correlations from the Phase I/II Peginvera Study of Ropeginterferon Alfa-2b, a Novel IFNa-2b, in Polycythemia Vera Patient. <i>Blood</i> , 2015 , 126, 4056-4056	2.2	
231	Influence of Platelet and White Blood Cell Counts until Major Thrombosis: Analysis from a Patient Registry in Essential Thrombocythemia. <i>Blood</i> , 2015 , 126, 2814-2814	2.2	
230	Clinical Impact of Bone Marrow Morphology for the Diagnosis of Essential Thrombocythemia: Comparison Between the British Standards (BCSH) and the WHO Criteria. <i>Blood</i> , 2015 , 126, 2816-2816	2.2	
229	Common Variation at 6q25.3 (TULP4) Influences Risk for Arterial Thrombosis in Myeloproliferative Neoplasms. <i>Blood</i> , 2015 , 126, 4088-4088	2.2	
228	Long-Term Follow-up in Interferon-Alpha Treated Polycythemia Vera Patients - a Single Center Analysis. <i>Blood</i> , 2015 , 126, 1608-1608	2.2	1
227	Simultaneous imaging of amyloid-land lipids in brain tissue using antibody-coupled liposomes and time-of-flight secondary ion mass spectrometry. <i>Journal of the American Chemical Society</i> , 2014 , 136, 9973-81	16.4	31
226	Selection of genetic and phenotypic features associated with inflammatory status of patients on dialysis using relaxed linear separability method. <i>PLoS ONE</i> , 2014 , 9, e86630	3.7	4
225	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. <i>Molecular Psychiatry</i> , 2014 , 19, 334-47	l ^{15.1}	73
224	How can genetics and epigenetics help the nephrologist improve the diagnosis and treatment of chronic kidney disease patients?. <i>Nephrology Dialysis Transplantation</i> , 2014 , 29, 972-80	4.3	12
223	Novel insights from genetic and epigenetic studies in understanding the complex uraemic phenotype. <i>Nephrology Dialysis Transplantation</i> , 2014 , 29, 964-71	4.3	9
222	Human adenovirus-36 is uncommon in type 2 diabetes and is associated with increased insulin sensitivity in adults in Sweden. <i>Annals of Medicine</i> , 2014 , 46, 539-46	1.5	15
221	Restoration of response to ruxolitinib upon brief withdrawal in two patients with myelofibrosis. <i>American Journal of Hematology</i> , 2014 , 89, 344-6	7.1	14
220	Decanucleotide insertion polymorphism of F7 significantly influences the risk of thrombosis in patients with essential thrombocythemia. <i>European Journal of Haematology</i> , 2014 , 93, 103-11	3.8	11
219	Attention biases and habituation of attention biases are associated with 5-HTTLPR and COMTval158met. <i>Cognitive, Affective and Behavioral Neuroscience</i> , 2014 , 14, 354-63	3.5	11

218	Calreticulin Mutation Status Predicts Improved Disease Outcome in Prefibrotic Primary Myelofibrosis but Not in WHO-Defined Essential Thrombocythemia. <i>Blood</i> , 2014 , 124, 3167-3167	2.2	
217	Anorexia and hypothalamic degeneration. <i>Vitamins and Hormones</i> , 2013 , 92, 27-60	2.5	11
216	Somatic mutations of calreticulin in myeloproliferative neoplasms. <i>New England Journal of Medicine</i> , 2013 , 369, 2379-90	59.2	1367
215	Genetic relationship between five psychiatric disorders estimated from genome-wide SNPs. <i>Nature Genetics</i> , 2013 , 45, 984-94	36.3	1628
214	Localization of cholesterol, amyloid and glia in AlzheimerQ disease transgenic mouse brain tissue using time-of-flight secondary ion mass spectrometry (ToF-SIMS) and immunofluorescence imaging. <i>Acta Neuropathologica</i> , 2013 , 125, 145-57	14.3	42
213	Genetic and epigenetic associations of MAOA and NR3C1 with depression and childhood adversities. <i>International Journal of Neuropsychopharmacology</i> , 2013 , 16, 1513-28	5.8	153
212	Identification of risk loci with shared effects on five major psychiatric disorders: a genome-wide analysis. <i>Lancet, The</i> , 2013 , 381, 1371-1379	40	2112
211	All SNPs are not created equal: genome-wide association studies reveal a consistent pattern of enrichment among functionally annotated SNPs. <i>PLoS Genetics</i> , 2013 , 9, e1003449	6	209
210	Genetic polymorphisms in monoamine systems and outcome of cognitive behavior therapy for social anxiety disorder. <i>PLoS ONE</i> , 2013 , 8, e79015	3.7	27
209	Long-term lithium treatment in bipolar disorder is associated with longer leukocyte telomeres. <i>Translational Psychiatry</i> , 2013 , 3, e261	8.6	106
208	Cognitive manic symptoms in bipolar disorder associated with polymorphisms in the DAOA and COMT genes. <i>PLoS ONE</i> , 2013 , 8, e67450	3.7	7
207	Correlation Of Longitudinal Blood Counts With Thromboembolic Complications In WHO 2008 Defined Prefibrotic Primary Myelofibrosis. <i>Blood</i> , 2013 , 122, 2834-2834	2.2	1
206	Frequent Mutations in the Calreticulin Gene CALR in Myeloproliferative Neoplasms. <i>Blood</i> , 2013 , 122, LBA-1-LBA-1	2.2	2
205	Assessment of Response to Lithium Maintenance Treatment in Bipolar Disorder: A Consortium on Lithium Genetics (ConLiGen) Report. <i>PLoS ONE</i> , 2013 , 8, e65636	3.7	113
204	The Anorectic Phenotype of the anx/anx Mouse Is Related to Hypothalamic Dysfunction. <i>Neuromethods</i> , 2013 , 333-350	0.4	
203	Clinical and genetic outcome determinants of Internet- and group-based cognitive behavior therapy for social anxiety disorder. <i>Acta Psychiatrica Scandinavica</i> , 2012 , 126, 126-36	6.5	77
202	Polymorphisms in AKR1C4 and HSD3B2 and differences in serum DHEAS and progesterone are associated with paranoid ideation during mania or hypomania in bipolar disorder. <i>European Neuropsychopharmacology</i> , 2012 , 22, 632-40	1.2	19
201	Kynurenine 3-monooxygenase polymorphisms: relevance for kynurenic acid synthesis in patients with schizophrenia and healthy controls. <i>Journal of Psychiatry and Neuroscience</i> , 2012 , 37, 53-7	4.5	51

200	Meta-analysis identifies multiple loci associated with kidney function-related traits in east Asian populations. <i>Nature Genetics</i> , 2012 , 44, 904-9	36.3	201
199	Serotonin-1A receptor polymorphism (rs6295) associated with thermal pain perception. <i>PLoS ONE</i> , 2012 , 7, e43221	3.7	27
198	Distinct contributions of the dorsolateral prefrontal and orbitofrontal cortex during emotion regulation. <i>PLoS ONE</i> , 2012 , 7, e48107	3.7	119
197	Epigenetic aberrations in leukocytes of patients with schizophrenia: association of global DNA methylation with antipsychotic drug treatment and disease onset. <i>FASEB Journal</i> , 2012 , 26, 2712-8	0.9	144
196	Adenovirus-36 is associated with obesity in children and adults in Sweden as determined by rapid ELISA. <i>PLoS ONE</i> , 2012 , 7, e41652	3.7	57
195	P2RX7: expression responds to sleep deprivation and associates with rapid cycling in bipolar disorder type 1. <i>PLoS ONE</i> , 2012 , 7, e43057	3.7	26
194	Genetic studies in chronic kidney disease: basic concepts. <i>Journal of Nephrology</i> , 2012 , 25, 141-9	4.8	3
193	Genetic studies in chronic kidney disease: interpretation and clinical applicability. <i>Journal of Nephrology</i> , 2012 , 25, 851-64	4.8	6
192	Decanucleotide Insertion Polymorphism of F7 Influences Significantly the Risk of Thrombosis in Patients with Essential Thrombocythemia. <i>Blood</i> , 2012 , 120, 1730-1730	2.2	
191	Kynurenine 3-monooxygenase (KMO) polymorphisms in schizophrenia: an association study. <i>Schizophrenia Research</i> , 2011 , 127, 270-2	3.6	19
190	Genetic variants in novel pathways influence blood pressure and cardiovascular disease risk. <i>Nature</i> , 2011 , 478, 103-9	50.4	1564
189	5-HTTLPR and COMTval158met genotype gate amygdala reactivity and habituation. <i>Biological Psychology</i> , 2011 , 87, 106-12	3.2	55
188	Perception of thermal pain and the thermal grill illusion is associated with polymorphisms in the serotonin transporter gene. <i>PLoS ONE</i> , 2011 , 6, e17752	3.7	52
187	Conditioned pain modulation is associated with common polymorphisms in the serotonin transporter gene. <i>PLoS ONE</i> , 2011 , 6, e18252	3.7	75
186	Cognitive manic symptoms associated with the P2RX7 gene in bipolar disorder. <i>Bipolar Disorders</i> , 2011 , 13, 500-8	3.8	35
185	Expression of osteoprotegerin in human fat tissue; implications for chronic kidney disease. <i>European Journal of Clinical Investigation</i> , 2011 , 41, 498-506	4.6	12
184	AKR1C4 gene variant associated with low euthymic serum progesterone and a history of mood irritability in males with bipolar disorder. <i>Journal of Affective Disorders</i> , 2011 , 133, 346-51	6.6	20
183	Genome integrity of myeloproliferative neoplasms in chronic phase and during disease progression. <i>Blood</i> , 2011 , 118, 167-76	2.2	129

(2009-2011)

182	Large-scale genome-wide association analysis of bipolar disorder identifies a new susceptibility locus near ODZ4. <i>Nature Genetics</i> , 2011 , 43, 977-83	36.3	1094
181	Hypothalamic mitochondrial dysfunction associated with anorexia in the anx/anx mouse. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 18108-13	11.5	33
180	CLOCK is suggested to associate with comorbid alcohol use and depressive disorders. <i>Journal of Circadian Rhythms</i> , 2010 , 8, 1	2.5	67
179	Genetic loci influencing kidney function and chronic kidney disease. <i>Nature Genetics</i> , 2010 , 42, 373-5	36.3	205
178	CRY2 is associated with depression. <i>PLoS ONE</i> , 2010 , 5, e9407	3.7	102
177	CRY2 is associated with rapid cycling in bipolar disorder patients. <i>PLoS ONE</i> , 2010 , 5, e12632	3.7	58
176	Multidrug resistance 1 (MDR1) gene polymorphisms in childhood drug-resistant epilepsy. <i>Journal of Child Neurology</i> , 2010 , 25, 1485-90	2.5	24
175	Visfatin is increased in chronic kidney disease patients with poor appetite and correlates negatively with fasting serum amino acids and triglyceride levels. <i>Nephrology Dialysis Transplantation</i> , 2010 , 25, 901-6	4.3	44
174	Expression of inflammatory and insulin signaling genes in adipose tissue in response to elective surgery. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010 , 95, 3460-9	5.6	26
173	Analysis of opioid and amyloid peptides using time-of-flight secondary ion mass spectrometry. <i>Analytical Chemistry</i> , 2010 , 82, 1964-74	7.8	15
172	Amygdala-dependent fear conditioning in humans is modulated by the BDNFval66met polymorphism. <i>Behavioral Neuroscience</i> , 2010 , 124, 9-15	2.1	52
171	PER2 variantion is associated with depression vulnerability. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2010 , 153B, 570-581	3.5	88
170	The COMTval158met polymorphism is associated with symptom relief during exposure-based cognitive-behavioral treatment in panic disorder. <i>BMC Psychiatry</i> , 2010 , 10, 99	4.2	69
169	Increased sensitivity to thermal pain following a single opiate dose is influenced by the COMT val(158)met polymorphism. <i>PLoS ONE</i> , 2009 , 4, e6016	3.7	81
168	Genetic gating of human fear learning and extinction: possible implications for gene-environment interaction in anxiety disorder. <i>Psychological Science</i> , 2009 , 20, 198-206	7.9	211
167	Azithromycin increases chloride efflux from cystic fibrosis airway epithelial cells. <i>Experimental Lung Research</i> , 2009 , 35, 210-21	2.3	12
166	CCR5 deletion protects against inflammation-associated mortality in dialysis patients. <i>Journal of the American Society of Nephrology: JASN</i> , 2009 , 20, 1641-9	12.7	59
165	Increased expression of inflammatory pathway genes in skeletal muscle during surgery. <i>Clinical Nutrition</i> , 2009 , 28, 291-8	5.9	21

164	Chemical analysis of osmium tetroxide staining in adipose tissue using imaging ToF-SIMS. Histochemistry and Cell Biology, 2009 , 132, 105-15	2.4	53
163	282 INCREASED SENSITIVITY TO PAIN FOLLOWING A SINGLE OPIATE DOSE IS INFLUENCED BY THE COMT val158met POLYMORPHISM. <i>European Journal of Pain</i> , 2009 , 13, 588	3.7	
162	Genetics/Genomics in chronic kidney diseasetowards personalized medicine?. <i>Seminars in Dialysis</i> , 2009 , 22, 417-22	2.5	20
161	The symptomatic profile of panic disorder is shaped by the 5-HTTLPR polymorphism. <i>Progress in Neuro-Psychopharmacology and Biological Psychiatry</i> , 2009 , 33, 1479-83	5.5	34
160	Genetic variation in the serotonin transporter gene (5-HTTLPR, rs25531) influences the analgesic response to the short acting opioid Remifentanil in humans. <i>Molecular Pain</i> , 2009 , 5, 37	3.4	55
159	Idiopathic megalencephaly-possible cause and treatment opportunities: from patient to lab. <i>European Journal of Paediatric Neurology</i> , 2008 , 12, 438-45	3.8	10
158	CREB-regulated diurnal activity patterns are not indicative for depression-like symptoms in mice and men. <i>Medical Hypotheses</i> , 2008 , 70, 117-21	3.8	0
157	TOF-SIMS analysis of adipose tissue from patients with chronic kidney disease. <i>Applied Surface Science</i> , 2008 , 255, 1177-1180	6.7	25
156	Complex aetiology of an apparently Mendelian form of mental retardation. <i>BMC Medical Genetics</i> , 2008 , 9, 6	2.1	4
155	Three circadian clock genes Per2, Arntl, and Npas2 contribute to winter depression. <i>Annals of Medicine</i> , 2007 , 39, 229-38	1.5	193
154	Lack of potassium channel induces proliferation and survival causing increased neurogenesis and two-fold hippocampus enlargement. <i>Hippocampus</i> , 2007 , 17, 292-304	3.5	16
153	Impact of inflammation on epigenetic DNA methylation - a novel risk factor for cardiovascular disease?. <i>Journal of Internal Medicine</i> , 2007 , 261, 488-99	10.8	304
152	Circulating levels of visfatin/pre-B-cell colony-enhancing factor 1 in relation to genotype, GFR, body composition, and survival in patients with CKD. <i>American Journal of Kidney Diseases</i> , 2007 , 49, 237-44	7.4	92
151	Changes in fat mass after initiation of maintenance dialysis is influenced by the uncoupling protein 2 exon 8 insertion/deletion polymorphism. <i>Nephrology Dialysis Transplantation</i> , 2007 , 22, 196-202	4.3	28
150	Evidence for hypothalamic dysregulation in mouse models of anorexia as well as in humans. <i>Physiology and Behavior</i> , 2007 , 92, 278-82	3.5	13
149	Carbamazepine protects against megencephaly and abnormal expression of BDNF and Nogo signaling components in the mceph/mceph mouse. <i>Neurobiology of Disease</i> , 2006 , 24, 374-83	7.5	23
148	The Swedish Twin Registry in the Third Millennium: An Update. <i>Twin Research and Human Genetics</i> , 2006 , 9, 875-882	2.2	290
147	LUMA (LUminometric Methylation Assay)a high throughput method to the analysis of genomic DNA methylation. <i>Experimental Cell Research</i> , 2006 , 312, 1989-95	4.2	231

(2004-2006)

146	The Swedish Twin Registry in the third millennium: an update. <i>Twin Research and Human Genetics</i> , 2006 , 9, 875-82	2.2	145
145	Interleukin-1 gene cluster polymorphisms are associated with nutritional status and inflammation in patients with end-stage renal disease. <i>Blood Purification</i> , 2005 , 23, 384-93	3.1	21
144	Peroxisome proliferator-activated receptor gamma polymorphisms affect systemic inflammation and survival in end-stage renal disease patients starting renal replacement therapy. <i>Atherosclerosis</i> , 2005 , 182, 105-11	3.1	28
143	Low fetuin-A levels are associated with cardiovascular death: Impact of variations in the gene encoding fetuin. <i>Kidney International</i> , 2005 , 67, 2383-92	9.9	232
142	Alterations of arcuate nucleus neuropeptidergic development in contactin-deficient mice: comparison with anorexia and food-deprived mice. <i>European Journal of Neuroscience</i> , 2005 , 22, 3217-28	3.5	34
141	Association study between chromosome 10q26.11 and obesity among Swedish men. <i>International Journal of Obesity</i> , 2005 , 29, 1422-8	5.5	
140	Maturation of the hypothalamic arcuate agouti-related protein system during postnatal development in the mouse. <i>Developmental Brain Research</i> , 2005 , 155, 147-54		60
139	Basal cell carcinoma and variants in genes coding for immune response, DNA repair, folate and iron metabolism. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2005 , 574, 105	;3 ₁ 3	54
138	A truncated Kv1.1 protein in the brain of the megencephaly mouse: expression and interaction. <i>BMC Neuroscience</i> , 2005 , 6, 65	3.2	22
137	MAOA haplotypes associated with thrombocyte-MAO activity. <i>BMC Genetics</i> , 2005 , 6, 46	2.6	29
136	Cosegregation of bipolar disorder and autosomal-dominant medullary cystic kidney disease in a large family. <i>American Journal of Psychiatry</i> , 2005 , 162, 1972-4	11.9	8
135	Detection and isolation of trinucleotide repeat expansions using the RED method. <i>Methods in Molecular Biology</i> , 2004 , 277, 47-59	1.4	3
134	No evidence for heritability of Parkinson disease in Swedish twins. <i>Neurology</i> , 2004 , 63, 305-11	6.5	94
133	Adiponectin in renal disease: relationship to phenotype and genetic variation in the gene encoding adiponectin. <i>Kidney International</i> , 2004 , 65, 274-81	9.9	125
132	Reduced gene expression of adiponectin in fat tissue from patients with end-stage renal disease. <i>Kidney International</i> , 2004 , 66, 46-50	9.9	45
131	Selective noradrenaline reuptake inhibition enhances serotonergic neuronal activity and transmitter release in the rat forebrain. <i>Journal of Neural Transmission</i> , 2004 , 111, 127-39	4.3	32
130	Expanded RED products and loci containing CAG/CTG repeats on chromosome 17 (ERDA1) and chromosome 18 (CTG18.1) in trans-generational pairs with bipolar affective disorder 2004 , 128B, 71-5		3
129	Spinocerebellar ataxia with sensory neuropathy (SCA25) maps to chromosome 2p. <i>Annals of Neurology</i> , 2004 , 55, 97-104	9.4	64

128	Pyrosequencing-based SNP allele frequency estimation in DNA pools. <i>Human Mutation</i> , 2004 , 23, 92-7	4.7	48
127	Seasonal affective disorder and the G-protein beta-3-subunit C825T polymorphism. <i>Biological Psychiatry</i> , 2004 , 55, 317-9	7.9	22
126	Gender differences in heritability of depressive symptoms in the elderly. <i>Psychological Medicine</i> , 2004 , 34, 471-9	6.9	63
125	Gender diagnosticity and androgen receptor gene CAG repeat sequence. <i>Twin Research and Human Genetics</i> , 2004 , 7, 456-61		7
124	Possible pathologic involvement of receptor for advanced glycation end products (RAGE) for development of encapsulating peritoneal sclerosis in Japanese CAPD patients. <i>Clinical Nephrology</i> , 2004 , 62, 455-60	2.1	26
123	Circadian clock-related polymorphisms in seasonal affective disorder and their relevance to diurnal preference. <i>Neuropsychopharmacology</i> , 2003 , 28, 734-9	8.7	276
122	The serotonin transporter promoter repeat length polymorphism, seasonal affective disorder and seasonality. <i>Psychological Medicine</i> , 2003 , 33, 785-92	6.9	33
121	No linkage to obesity in candidate regions of chromosome 2 and 10 in a selected sample of Swedish twins. <i>Twin Research and Human Genetics</i> , 2003 , 6, 162-9		3
120	A linkage study of candidate loci in familial Parkinson@ Disease. BMC Neurology, 2003, 3, 6	3.1	14
119	Approaches to anorexia in rodents: focus on the anx/anx mouse. <i>European Journal of Pharmacology</i> , 2003 , 480, 171-6	5.3	18
118	A functional variant of the myeloperoxidase gene is associated with cardiovascular disease in end-stage renal disease patients. <i>Kidney International</i> , 2003 , S172-6	9.9	93
117	Genetic approaches in the clinical investigation of complex disorders: malnutrition, inflammation, and atherosclerosis (MIA) as a prototype. <i>Kidney International</i> , 2003 , S162-7	9.9	24
116	MRI and in situ hybridization reveal early disturbances in brain size and gene expression in the megencephalic (mceph/mceph) mouse. <i>European Journal of Neuroscience</i> , 2003 , 18, 3218-30	3.5	27
115	Truncation of the Shaker-like voltage-gated potassium channel, Kv1.1, causes megencephaly. <i>European Journal of Neuroscience</i> , 2003 , 18, 3231-40	3.5	49
114	Multigenic control of disease severity after virulent Mycobacterium tuberculosis infection in mice. <i>Infection and Immunity</i> , 2003 , 71, 126-31	3.7	74
113	Repeat expansion detection (RED) and the RED cloning strategy. <i>Methods in Molecular Biology</i> , 2003 , 217, 51-60	1.4	
112	The androgen receptor gene and psychological traits: are results consistent in Sweden and Australia?. <i>Twin Research and Human Genetics</i> , 2003 , 6, 201-8		5
111	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, 691-8	8.7	32

(2001-2002)

110	Large-scale genotyping of single nucleotide polymorphisms by Pyrosequencingtrade mark and validation against the 5@uclease (Taqman((R))) assay. <i>Human Mutation</i> , 2002 , 19, 395-401	4.7	63
109	The hormone-sensitive lipase i6 gene polymorphism and body fat accumulation. <i>European Journal of Clinical Investigation</i> , 2002 , 32, 938-42	4.6	20
108	Physical map of the chromosome 6q22 region containing the oculodentodigital dysplasia locus: analysis of thirteen candidate genes and identification of novel ESTs and DNA polymorphisms. <i>Cytogenetic and Genome Research</i> , 2002 , 98, 29-37	1.9	6
107	European combined analysis of the CTG18.1 and the ERDA1 CAG/CTG repeats in bipolar disorder. <i>European Journal of Human Genetics</i> , 2002 , 10, 276-80	5.3	20
106	Genetic survival analysis of age-at-onset of bipolar disorder: evidence for anticipation or cohort effect in families. <i>Psychiatric Genetics</i> , 2001 , 11, 129-37	2.9	27
105	A common hormone-sensitive lipase i6 gene polymorphism is associated with decreased human adipocyte lipolytic function. <i>Diabetes</i> , 2001 , 50, 2410-3	0.9	31
104	Insulin-like growth factor-II in endocrine pancreatic tumours. Immunohistochemical, biochemical and in situ hybridization findings. <i>Apmis</i> , 2001 , 109, 127-40	3.4	15
103	The effect of the beta(2) adrenoceptor gene Thr164lle polymorphism on human adipose tissue lipolytic function. <i>British Journal of Pharmacology</i> , 2001 , 133, 708-12	8.6	28
102	Cytotoxic T lymphocyte-associated antigen 4 (CTLA-4) blockade enhances incidence and severity of experimental autoimmune neuritis in resistant mice. <i>Journal of Neuroimmunology</i> , 2001 , 115, 111-7	3.5	21
101	Dominantly inherited familial myasthenia gravis as a separate genetic entity without involvement of defined candidate gene loci. <i>International Journal of Molecular Medicine</i> , 2001 , 7, 289-94	4.4	5
100	A cloning strategy for identification of genes containing trinucleotide repeat expansions. <i>International Journal of Molecular Medicine</i> , 2001 , 8, 427-31	4.4	1
99	Muscleblind localizes to nuclear foci of aberrant RNA in myotonic dystrophy types 1 and 2. <i>Human Molecular Genetics</i> , 2001 , 10, 2165-70	5.6	330
98	Genetics of affective disorders. European Neuropsychopharmacology, 2001, 11, 385-94	1.2	44
97	Seasonal affective disorder and serotonin-related polymorphisms. <i>Neurobiology of Disease</i> , 2001 , 8, 351	1 -7 .5	42
96	Expression of alpha-synuclein in the human brain: relation to Lewy body disease. <i>Molecular Brain Research</i> , 2001 , 92, 58-65		22
95	Androgen receptor trinucleotide repeat polymorphism and personality traits. <i>Psychiatric Genetics</i> , 2001 , 11, 19-23	2.9	54
94	Altered dopaminergic transmission in the anorexic anx/anx mouse striatum. <i>NeuroReport</i> , 2001 , 12, 273	37 <u>r.</u> ∳1	15
93	The Anorexia Mouse 2001 , 193-203		

92	Long repeat tracts at SCA8 in major psychosis. <i>American Journal of Medical Genetics Part A</i> , 2000 , 96, 873-6		48
91	P0 protein peptide 180-199 together with pertussis toxin induces experimental autoimmune neuritis in resistant C57BL/6 mice. <i>Journal of Neuroscience Research</i> , 2000 , 62, 717-21	4.4	36
90	Fat tissue accumulation during peritoneal dialysis is associated with a polymorphism in uncoupling protein 2. <i>Kidney International</i> , 2000 , 57, 1713-9	9.9	50
89	ARSACS, a spastic ataxia common in northeastern QuBec, is caused by mutations in a new gene encoding an 11.5-kb ORF. <i>Nature Genetics</i> , 2000 , 24, 120-5	36.3	332
88	No evidence for long CAG/CTG repeats in families with spastic paraplegia linked to chromosome 2p21-24. <i>Neuroscience Letters</i> , 2000 , 279, 41-4	3.3	6
87	Expression of cholecystokinin, enkephalin, galanin and neuropeptide Y is markedly changed in the brain of the megencephaly mouse. <i>Neuroscience</i> , 2000 , 100, 297-317	3.9	17
86	Hypothalamic CART and serum leptin levels are reduced in the anorectic (anx/anx) mouse. <i>Molecular Brain Research</i> , 2000 , 84, 97-105		57
85	Mapping of spinocerebellar ataxia 13 to chromosome 19q13.3-q13.4 in a family with autosomal dominant cerebellar ataxia and mental retardation. <i>American Journal of Human Genetics</i> , 2000 , 67, 229-	3 ¹ 5 ¹	151
84	Changes in neuropeptide Y receptors and pro-opiomelanocortin in the anorexia (anx/anx) mouse hypothalamus. <i>Journal of Neuroscience</i> , 1999 , 19, 7130-9	6.6	53
83	Expanded repeat sequences and disease. <i>Seminars in Neurology</i> , 1999 , 19, 289-99	3.2	21
82	Severity of tuberculosis in mice is linked to distal chromosome 3 and proximal chromosome 9. Journal of Infectious Diseases, 1999 , 180, 150-5	7	67
81	Genes involved in animal models of obesity and anorexia. <i>Journal of Internal Medicine</i> , 1999 , 245, 613-9	10.8	9
80	Leptin and its potential role in human obesity. <i>Journal of Internal Medicine</i> , 1999 , 245, 643-52	10.8	41
79	An integrated map of chromosome 18 CAG trinucleotide repeat loci. <i>European Journal of Human Genetics</i> , 1999 , 7, 12-9	5.3	5
78	Exclusion of dominant mutations within the FTDP-17 locus on chromosome 17 for Parkinson@ disease. <i>Neuroscience Letters</i> , 1999 , 272, 140-2	3.3	2
77	The megencephaly mouse has disturbances in the insulin-like growth factor (IGF) system. <i>Molecular Brain Research</i> , 1999 , 72, 80-8		13
76	Molecular interpretation of expanded RED products in bipolar disorder by CAG/CTG repeats located at chromosomes 17q and 18q. <i>Neurobiology of Disease</i> , 1999 , 6, 424-32	7·5	25
75	Chapter 5.3 Trinucleotide repeat disorders. <i>Handbook of Behavioral Neuroscience</i> , 1999 , 13, 783-804		

74	Linkage analysis of candidate loci in families with recurrent major depression. <i>Molecular Psychiatry</i> , 1998 , 3, 162-8	15.1	12
73	Large CAG/CTG repeats are associated with childhood-onset schizophrenia. <i>Molecular Psychiatry</i> , 1998 , 3, 321-7	15.1	29
72	Two commonly expanded CAG/CTG repeat loci: involvement in affective disorders?. <i>Molecular Psychiatry</i> , 1998 , 3, 405-10	15.1	52
71	Isolation and characterization of trinucleotide repeat containing partial transcripts in human spinal cord. <i>Neurogenetics</i> , 1998 , 1, 239-47	3	6
70	Long CAG/CTG repeats in mice. <i>Mammalian Genome</i> , 1998 , 9, 392-3	3.2	2
69	Reduced gene expression of UCP2 but not UCP3 in skeletal muscle of human obese subjects. <i>Diabetologia</i> , 1998 , 41, 935-9	10.3	51
68	Low leptin gene expression and hyperleptinemia in chronic renal failure. <i>Kidney International</i> , 1998 , 54, 1267-75	9.9	122
67	A triplet repeat on 17q accounts for most expansions detected by the repeat-expansion-detection technique. <i>American Journal of Human Genetics</i> , 1998 , 62, 1548-51	11	34
66	Differential expression of IGF-I and IGF-binding protein-1 and -2 in periportal and perivenous zones of rat liver. <i>Journal of Endocrinology</i> , 1998 , 157, 285-94	4.7	14
65	The neuropeptide Y/agouti gene-related protein (AGRP) brain circuitry in normal, anorectic, and monosodium glutamate-treated mice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1998 , 95, 15043-8	11.5	653
64	Multivariate analysis of factors influencing repeat expansion detection. <i>Genome Research</i> , 1998 , 8, 1085	5 -9. 4	9
63	Effect of in vitro promoter methylation and CGG repeat expansion on FMR-1 expression. <i>Nucleic Acids Research</i> , 1997 , 25, 2883-7	20.1	26
62	Delayed cytokine expression in rat brain following experimental contusion. <i>Journal of Neurosurgery</i> , 1997 , 86, 493-504	3.2	137
61	Cystic fibrosis mRNA expression in rat brain: cerebral cortex and medial preoptic area. <i>NeuroReport</i> , 1997 , 8, 535-9	1.7	36
60	MK-801 inhibits the cortical increase in IGF-1, IGFBP-2 and IGFBP-4 expression following trauma. <i>NeuroReport</i> , 1997 , 8, 455-60	1.7	18
59	Polymorphic expression of multidrug resistance mRNA in lung parenchyma of nonpregnant and pregnant rats: a comparison to cystic fibrosis mRNA expression. <i>Biochemical and Biophysical Research Communications</i> , 1997 , 239, 606-11	3.4	9
58	Cerebral autosomal dominant arteriopathy with subcortical infarcts and leucoencephalopathy. <i>Journal of Clinical Neuroscience</i> , 1997 , 4, 176-80	2.2	2
57	Expanded trinucleotide CAG repeats in families with bipolar affective disorder. <i>Biological Psychiatry</i> , 1997 , 42, 1115-22	7.9	50

56	Leptin gene is expressed in rat brown adipose tissue at birth. FASEB Journal, 1997, 11, 382-7	0.9	61
55	Protein kinase C in the developing kidney: isoform expression and effects of ceramide and PKC inhibitors. <i>Kidney International</i> , 1997 , 52, 901-10	9.9	25
54	Schizophrenia and neurotrophin-3 alleles. <i>Acta Psychiatrica Scandinavica</i> , 1997 , 95, 414-9	6.5	43
53	Hypothalamic neurohistochemistry of the murine anorexia (anx/anx) mutation: altered processing of neuropeptide Y in the arcuate nucleus. <i>Journal of Comparative Neurology</i> , 1997 , 387, 124-35	3.4	50
52	Oculopharyngeal muscular dystrophy (OPMD)report and genetic studies of an Australian kindred. <i>Clinical Genetics</i> , 1997 , 51, 52-5	4	14
51	Leptin secretion from adipose tissue in women. Relationship to plasma levels and gene expression. Journal of Clinical Investigation, 1997 , 99, 2398-404	15.9	138
50	Increased leptin messenger RNA and serum leptin levels in children with Prader-Willi syndrome and nonsyndromal obesity. <i>Pediatric Research</i> , 1997 , 42, 593-6	3.2	19
49	Lack of expansion of cytosine adenine guanine trinucleotide repeat in the dentatorubral and pallidoluysian atrophy gene in Swedish schizophrenic patients. <i>Biological Psychiatry</i> , 1996 , 40, 228-9	7.9	1
48	Increase of insulin-like growth factor (IGF)-1, IGF binding protein-2 and -4 mRNAs following cerebral contusion. <i>Molecular Brain Research</i> , 1996 , 38, 285-93		80
47	PDGF-A signaling is a critical event in lung alveolar myofibroblast development and alveogenesis. <i>Cell</i> , 1996 , 85, 863-73	56.2	704
46	Mutation Detection in Machado-Joseph Disease Using Repeat Expansion Detection. <i>Molecular Medicine</i> , 1996 , 2, 77-85	6.2	17
45	An expanded CAG repeat sequence in spinocerebellar ataxia type 7. <i>Genome Research</i> , 1996 , 6, 965-71	9.7	72
44	A role for a new herpes virus (KSHV) in different forms of KaposiQ sarcoma. <i>Nature Medicine</i> , 1995 , 1, 707-8	50.5	278
43	Overexpression of the obese (ob) gene in adipose tissue of human obese subjects. <i>Nature Medicine</i> , 1995 , 1, 950-3	50.5	606
42	Detection of expanded CAG repeats in bipolar affective disorder using the repeat expansion detection (RED) method. <i>Neurobiology of Disease</i> , 1995 , 2, 55-62	7.5	135
41	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. <i>Mammalian Genome</i> , 1994 , 5, 597-607	3.2	26
40	Growing triplet repeats. <i>Nature Genetics</i> , 1994 , 7, 124	36.3	25
39	Plasticity in expression of neuropeptides. <i>European Neuropsychopharmacology</i> , 1993 , 3, 162-163		

38	Tissue specific expression of FMR-1 provides evidence for a functional role in fragile X syndrome. <i>Nature Genetics</i> , 1993 , 3, 36-43	36.3	320
37	Direct detection of novel expanded trinucleotide repeats in the human genome. <i>Nature Genetics</i> , 1993 , 4, 135-9	36.3	251
36	The Wilms tumour gene WT1 is expressed in murine mesoderm-derived tissues and mutated in a human mesothelioma. <i>Nature Genetics</i> , 1993 , 4, 415-20	36.3	185
35	Calcitonin gene-related peptide in the brain, spinal cord, and some peripheral systems. <i>Annals of the New York Academy of Sciences</i> , 1992 , 657, 119-34	6.5	100
34	In Situ Hybridization and Immunohistochemical Methods in Study of Regulatory Molecules. <i>Methods in Neurosciences</i> , 1992 , 231-255		7
33	Analysis of a murine male germ cell-specific transcript that encodes a putative zinc finger protein. <i>Molecular Reproduction and Development</i> , 1991 , 30, 173-81	2.6	49
32	Calcitonin gene-related peptide and its mRNA in pulmonary neuroendocrine cells and ganglia. <i>Histochemistry</i> , 1991 , 96, 311-5		44
31	Expression of the Wilms Qumor gene WT1 in the murine urogenital system. <i>Genes and Development</i> , 1991 , 5, 1345-56	12.6	350
30	Colocalization of neurotransmitters analyzed by in situ hybridization. <i>European Neuropsychopharmacology</i> , 1991 , 1, 173-6	1.2	8
29	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	3.9	67
28	A gene encoding a fibroblast growth factor receptor isolated from the Huntington disease gene region of human chromosome 4. <i>Genomics</i> , 1991 , 11, 1133-42	4.3	98
27	Comparison of gene expression of the dopamine D-2 receptor and DARPP-32 in rat brain, pituitary and adrenal gland. <i>European Journal of Pharmacology</i> , 1990 , 188, 277-81		23
26	Differential effects of intracerebroventricular colchicine administration on the expression of mRNAs for neuropeptides and neurotransmitter enzymes, with special emphasis on galanin: an in situ hybridization study. <i>Synapse</i> , 1990 , 6, 369-91	2.4	207
25	Analysis of expression of cholecystokinin in dopamine cells in the ventral mesencephalon of several species and in humans with schizophrenia. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1990 , 87, 8427-31	11.5	59
24	Distribution and cellular localization of DARPP-32 mRNA in rat brain. <i>Molecular Brain Research</i> , 1990 , 7, 139-49		47
23	Effects of reserpine on phenylethanolamine N-methyltransferase mRNA levels in rat adrenal gland: Role of steroids. <i>Neurochemistry International</i> , 1990 , 17, 343-56	4.4	23
22	Localization of neuropeptide receptor mRNA in rat brain: initial observations using probes for neurotensin and substance P receptors. <i>Neuroscience Letters</i> , 1990 , 120, 134-8	3.3	119
21	Neuropeptide gene expression in hypothalamic magnocellular neurons of normal and hypophysectomized rats: a combined immunohistochemical and in situ hybridization study. <i>Neuroscience</i> , 1990 , 36, 181-99	3.9	7 ²

20	Evidence for coexistence between calcitonin gene-related peptide and serotonin in the bulbospinal pathway in the monkey. <i>Brain Research</i> , 1990 , 532, 47-57	3.7	33
19	Expression and localization of gastrin messenger RNA and peptide in spermatogenic cells. <i>Journal of Clinical Investigation</i> , 1990 , 86, 660-9	15.9	49
18	Demonstration of DARPP-32 in pituicytes of the neurohypophysis-decreased expression after administration of hyperosmotic stimuli. <i>Acta Physiologica Scandinavica</i> , 1989 , 137, 461-2		9
17	Immunocytochemical demonstration of cholecystokinin-like immunoreactivity in spermatozoa in monkey testis and epididymis. <i>Acta Physiologica Scandinavica</i> , 1989 , 137, 465-6		13
16	Presence of cholecystokinin mRNA in dopamine cells in the ventral mesencephalon of a human with schizophrenia. <i>Acta Physiologica Scandinavica</i> , 1989 , 137, 467-8		11
15	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. <i>Neuroscience Letters</i> , 1989 , 107, 1-5	3.3	30
14	Neuropeptide expression in rat dorsal root ganglion cells and spinal cord after peripheral nerve injury with special reference to galanin. <i>Neuroscience</i> , 1989 , 33, 587-604	3.9	433
13	Decrease in gene expression of substance P and thyrotropin releasing hormone (TRH) in rat medulla oblongata following treatment with an antidepressive drug. <i>Acta Physiologica Scandinavica</i> , 1988 , 134, 455-6		7
12	Increased neuropeptide Y messenger RNA and peptide in sympathetic ganglia after reserpine pretreatment. <i>European Journal of Pharmacology</i> , 1988 , 156, 419-20	5.3	22
11	A subpopulation of dopaminergic neurons in rat ventral mesencephalon contains both neurotensin and cholecystokinin. <i>Brain Research</i> , 1988 , 455, 88-98	3.7	117
10	Tachykinins in the central nervous system. <i>Regulatory Peptides</i> , 1988 , 22, 6-8		1
9	Neuropeptide tyrosine in the rat adrenal glandimmunohistochemical and in situ hybridization studies. <i>Neuroscience</i> , 1988 , 24, 337-49	3.9	76
8	Detection of cholecystokinin in spermatogenic cells. Acta Physiologica Scandinavica, 1988, 134, 565-566		7
7	Chemical Neuroanatomy of the Hypothalamo-Pituitary Axis: Focus on Multimessenger Systems 1987, 1-34		1
6	Tyrosine 3-hydroxylase in rat brain and adrenal medulla: hybridization histochemistry and immunohistochemistry combined with retrograde tracing. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1986 , 83, 6208-12	11.5	44
5	Leukotriene C4 binding sites in the rat central nervous system. <i>European Journal of Pharmacology</i> , 1986 , 122, 251-7	5.3	39
4	Coexistence of neuronal messengersan overview. <i>Progress in Brain Research</i> , 1986 , 68, 33-70	2.9	142
3	Neurons with multiple messengers with special reference in neuroendocrine systems. <i>Endocrine Reviews</i> , 1986 , 42, 1-70		14

2 Animal Models of Eating Disorders1115-1125

Polygenic scores for major depressive disorder and depressive symptoms predict response to lithium in patients with bipolar disorder

1