# Pierandrea Muglia

List of Publications by Year
in descending order



| | | | |
|---|---|---|---|
| 135 papers | 23,094 citations | 23500 58 h-index | 12233 133 g-index |
| 139 all docs | 139 docs citations | 139 times ranked | 27639 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Use of a physiologically based pharmacokineticâ€“pharmacodynamic model for initial dose prediction and escalation during a paediatric clinical trial. British Journal of Clinical Pharmacology, 2021, 87, 1378-1389. | 1.1 | 12 |
| 2 | Clinical and dopamine transporter imaging characteristics of non-manifest LRRK2 and GBA mutation carriers in the Parkinson's Progression Markers Initiative (PPMI): a cross-sectional study. Lancet Neurology, The, 2020, 19, 71-80. | 4.9 | 94 |
| 3 | Longitudinal Measurements of Glucocerebrosidase activity in Parkinsonâ€™s patients. Annals of Clinical and Translational Neurology, 2020, 7, 1816-1830. | 1.7 | 23 |
| 4 | Radiprodil, a NR2B negative allosteric modulator, from bench to bedside in infantile spasm syndrome. Annals of Clinical and Translational Neurology, 2020, 7, 343-352. | 1.7 | 18 |
| 5 | Large-Scale Exome Sequencing Study Implicates Both Developmental and Functional Changes in the Neurobiology of Autism. Cell, 2020, 180, 568-584.e23. | 13.5 | 1,422 |
| 6 | Transcriptome signatures from discordant sibling pairs reveal changes in peripheral blood immune cell composition in Autism Spectrum Disorder. Translational Psychiatry, 2020, 10, 106. | 2.4 | 16 |
| 7 | Large-scale targeted sequencing identifies risk genes for neurodevelopmental disorders. Nature Communications, 2020, 11, 4932. | 5.8 | 105 |
| 8 | Padsevonil randomized Phase IIa trial in treatment-resistant focal epilepsy: a translational approach. Brain Communications, 2020, 2, fcaa183. | 1.5 | 11 |
| 9 | Feasibility and safety of lumbar puncture in the Parkinson's disease research participants: Parkinson's Progression Marker Initiative (PPMI). Parkinsonism and Related Disorders, 2019, 62, 201-209. | 1.1 | 15 |
| 10 | BLOOD-BASED AUTISM SPECTRUM DISORDER SIGNATURES FROM THE ITALIAN AUTISM NETWORK COLLECTION. European Neuropsychopharmacology, 2019, 29, S912. | 0.3 | 0 |
| 11 | A singleâ€center, openâ€label positron emission tomography study to evaluate brivaracetam and levetiracetam synaptic vesicle glycoprotein 2A binding in healthy volunteers. Epilepsia, 2019, 60, 958-967. | 2.6 | 45 |
| 12 | Drug Development for Rare Paediatric Epilepsies: Current State and Future Directions. Drugs, 2019, 79, 1917-1935. | 4.9 | 13 |
| 13 | Alzheimer's disease pathology explains association between dementia with Lewy bodies and APOEâ€ε4/TOMM40 long polyâ€T repeat allele variants. Alzheimer's and Dementia: Translational Research and Clinical Interventions, 2019, 5, 814-824. | 1.8 | 14 |
| 14 | Dopamine Transporter Neuroimaging as an Enrichment Biomarker in Early Parkinson's Disease Clinical Trials: A Disease Progression Modeling Analysis. Clinical and Translational Science, 2018, 11, 63-70. | 1.5 | 36 |
| 15 | Methylphenidate enhances implicit learning in healthy adults. Journal of Psychopharmacology, 2018, 32, 70-80. | 2.0 | 12 |
| 16 | Unravelling the GSK3Î²-related genotypic interaction network influencing hippocampal volume in recurrent major depressive disorder. Psychiatric Genetics, 2018, 28, 77-84. | 0.6 | 27 |
| 17 | The Italian autism network (ITAN): a resource for molecular genetics and biomarker investigations. BMC Psychiatry, 2018, 18, 369. | 1.1 | 6 |
| 18 | The Parkinson's progression markers initiative (PPMI) â€“ establishing a PD biomarker cohort. Annals of Clinical and Translational Neurology, 2018, 5, 1460-1477. | 1.7 | 330 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Long-term seizure outcomes in patients with drug resistant epilepsy. Seizure: the Journal of the British Epilepsy Association, 2018, 62, 74-78. | 0.9 | 21 |
| 20 | A correction for sample overlap in genome-wide association studies in a polygenic pleiotropy-informed framework. BMC Genomics, 2018, 19, 494. | 1.2 | 37 |
| 21 | Genetics of schizophrenia: A consensus paper of the WFSBP Task Force on Genetics. World Journal of Biological Psychiatry, 2017, 18, 492-505. | 1.3 | 48 |
| 22 | Metoprololâ€pridopidine drugâ€“drug interaction and food effect assessments of pridopidine, a new drug for treatment of Huntington's disease. British Journal of Clinical Pharmacology, 2017, 83, 2214-2224. | 1.1 | 1 |
| 23 | GRIN2B gain of function mutations are sensitive to radiprodil, a negative allosteric modulator of GluN2B-containing NMDA receptors. Neuropharmacology, 2017, 123, 322-331. | 2.0 | 50 |
| 24 | Consensus paper of the WFSBP Task Force on Genetics: Genetics, epigenetics and gene expression markers of major depressive disorder and antidepressant response. World Journal of Biological Psychiatry, 2017, 18, 5-28. | 1.3 | 75 |
| 25 | Antiparkinsonian effects of the "Radiprodil and Tozadenant" combination in MPTP-treated marmosets. PLoS ONE, 2017, 12, e0182887. | 1.1 | 11 |
| 26 | NS11821, a partial subtype-selective GABA<sub>A</sub> agonist, elicits selective effects on the central nervous system in randomized controlled trial with healthy subjects. Journal of Psychopharmacology, 2016, 30, 253-262. | 2.0 | 25 |
| 27 | The association between lower educational attainment and depression owing to shared genetic effects? Results in ~25â€‰000 subjects. Molecular Psychiatry, 2015, 20, 735-743. | 4.1 | 59 |
| 28 | Genetic Differences in the Immediate Transcriptome Response to Stress Predict Risk-Related Brain Function and Psychiatric Disorders. Neuron, 2015, 86, 1189-1202. | 3.8 | 102 |
| 29 | Joint Analysis of Psychiatric Disorders Increases Accuracy of Risk Prediction for Schizophrenia, Bipolar Disorder, and Major Depressive Disorder. American Journal of Human Genetics, 2015, 96, 283-294. | 2.6 | 225 |
| 30 | Psychiatric genome-wide association study analyses implicate neuronal, immune and histone pathways. Nature Neuroscience, 2015, 18, 199-209. | 7.1 | 701 |
| 31 | Novel insights into the genetics of smoking behaviour, lung function, and chronic obstructive pulmonary disease (UK BiLEVE): a genetic association study in UK Biobank. Lancet Respiratory Medicine,the, 2015, 3, 769-781. | 5.2 | 346 |
| 32 | No Association Between NRG1 and ErbB4 Genes and Psychopathological Symptoms of Schizophrenia. NeuroMolecular Medicine, 2014, 16, 742-751. | 1.8 | 4 |
| 33 | Copy number variant study of bipolar disorder in Canadian and UK populations implicates synaptic genes. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2014, 165, 303-313. | 1.1 | 76 |
| 34 | Identification of Pathways for Bipolar Disorder. JAMA Psychiatry, 2014, 71, 657. | 6.0 | 204 |
| 35 | Genome-wide association study of bipolar disorder in Canadian and UK populations corroborates disease loci including SYNE1 and CSMD1. BMC Medical Genetics, 2014, 15, 2. | 2.1 | 106 |
| 36 | Variability in Working Memory Performance Explained by Epistasis vs Polygenic Scores in theZNF804APathway. JAMA Psychiatry, 2014, 71, 778. | 6.0 | 28 |

| # | Article | IF | Citations |
|---|---|---|---|
| 37 | Investigating the genetic variation underlying episodicity in major depressive disorder: Suggestive evidence for a bipolar contribution. Journal of Affective Disorders, 2014, 155, 81-89. | 2.0 | 15 |
| 38 | Analysis of RBFOX1 gene expression in lymphoblastoid cell lines of Italian discordant autism spectrum disorders sib-pairs. Molecular and Cellular Probes, 2014, 28, 242-245. | 0.9 | 6 |
| 39 | Genome-wide association analysis of copy number variation in recurrent depressive disorder. Molecular Psychiatry, 2013, 18, 183-189. | 4.1 | 45 |
| 40 | Genetic relationship between five psychiatric disorders estimated from genome-wide SNPs. Nature Genetics, 2013, 45, 984-994. | 9.4 | 2,067 |
| 41 | Identification of altered dipeptidyl-peptidase activities as potential biomarkers for unipolar depression. Journal of Affective Disorders, 2013, 151, 667-672. | 2.0 | 16 |
| 42 | Genomeâ€wide association analysis accounting for environmental factors through propensityâ€score matching: Application to stressful live events in major depressive disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2013, 162, 521-529. | 1.1 | 16 |
| 43 | Neuropsychological effects of the <scp>CSMD1</scp> genomeâ€wide associated schizophrenia risk variant rs10503253. Genes, Brain and Behavior, 2013, 12, 203-209. | 1.1 | 48 |
| 44 | A mega-analysis of genome-wide association studies for major depressive disorder. Molecular Psychiatry, 2013, 18, 497-511. | 4.1 | 1,002 |
| 45 | Genetic Schizophrenia Risk Variants Jointly Modulate Total Brain and White Matter Volume. Biological Psychiatry, 2013, 73, 525-531. | 0.7 | 119 |
| 46 | Genome-wide association study meta-analysis of European and Asian-ancestry samples identifies three novel loci associated with bipolar disorder. Molecular Psychiatry, 2013, 18, 195-205. | 4.1 | 180 |
| 47 | A common biological basis of obesity and nicotine addiction. Translational Psychiatry, 2013, 3, e308-e308. | 2.4 | 51 |
| 48 | Estimating the heritability of reporting stressful life events captured by common genetic variants. Psychological Medicine, 2013, 43, 1965-1971. | 2.7 | 46 |
| 49 | Schizophrenia genetic variants are not associated with intelligence. Psychological Medicine, 2013, 43, 2563-2570. | 2.7 | 40 |
| 50 | Runs of Homozygosity Implicate Autozygosity as a Schizophrenia Risk Factor. PLoS Genetics, 2012, 8, e1002656. | 1.5 | 109 |
| 51 | Investigation of the Genetic Association between Quantitative Measures of Psychosis and Schizophrenia: A Polygenic Risk Score Analysis. PLoS ONE, 2012, 7, e37852. | 1.1 | 60 |
| 52 | The association of rs4307059 and rs35678 markers with autism spectrum disorders is replicated in Italian families. Psychiatric Genetics, 2012, 22, 177-181. | 0.6 | 39 |
| 53 | Genetic variation in GOLM1 and prefrontal cortical volume in Alzheimer's disease. Neurobiology of Aging, 2012, 33, 457-465. | 1.5 | 14 |
| 54 | Dissecting the Genetic Heterogeneity of Depression Through Age at Onset. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 859-868. | 1.1 | 31 |

| # | Article | IF | Citations |
|---|---|---|---|
| 55 | Don't give up on GWAS. Molecular Psychiatry, 2012, 17, 2-3. | 4.1 | 54 |
| 56 | Common variants at VRK2 and TCF4 conferring risk of schizophrenia. Human Molecular Genetics, 2011, 20, 4076-4081. | 1.4 | 193 |
| 57 | Candidate Gene Analysis of the Human Natural Killer-1 Carbohydrate Pathway and Perineuronal Nets in Schizophrenia: B3GAT2 Is Associated with Disease Risk and Cortical Surface Area. Biological Psychiatry, 2011, 69, 90-96. | 0.7 | 42 |
| 58 | The AVPR1A Gene and Substance Use Disorders: Association, Replication, and Functional Evidence. Biological Psychiatry, 2011, 70, 519-527. | 0.7 | 45 |
| 59 | Genome-wide association study identifies five new schizophrenia loci. Nature Genetics, 2011, 43, 969-976. | 9.4 | 1,758 |
| 60 | A neuropsychological investigation of the genome wide associated schizophrenia risk variant NRGN rs12807809. Schizophrenia Research, 2011, 125, 304-306. | 1.1 | 23 |
| 61 | ADAMTSL3 as a candidate gene for schizophrenia: Gene sequencing and ultra-high density association analysis by imputation. Schizophrenia Research, 2011, 127, 28-34. | 1.1 | 42 |
| 62 | From genes to therapeutic targets for psychiatric disorders â€“ what to expect?. Current Opinion in Pharmacology, 2011, 11, 563-571. | 1.7 | 22 |
| 63 | Admixture analysis of age at onset in bipolar disorder. Psychiatry Research, 2011, 185, 27-32. | 1.7 | 51 |
| 64 | Genomewide Association Scan of Suicidal Thoughts and Behaviour in Major Depression. PLoS ONE, 2011, 6, e20690. | 1.1 | 98 |
| 65 | Structural Brain Changes in Patients with Recurrent Major Depressive Disorder Presenting with Anxiety Symptoms. , 2011, 21, 375-382. | | 44 |
| 66 | Copy number variations of chromosome 16p13.1 region associated with schizophrenia. Molecular Psychiatry, 2011, 16, 17-25. | 4.1 | 227 |
| 67 | Thyroid hormone transporter genes and grey matter changes in patients with major depressive disorder and healthy controls. Psychoneuroendocrinology, 2011, 36, 929-934. | 1.3 | 6 |
| 68 | No association between a common single nucleotide polymorphism, rs4141463, in the MACROD2 gene and autism spectrum disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 633-639. | 1.1 | 30 |
| 69 | A followâ€up caseâ€“control association study of tractable (druggable) genes in recurrent major depression. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 640-650. | 1.1 | 17 |
| 70 | Maternally Derived Microduplications at 15q11-q13: Implication of Imprinted Genes in Psychotic Illness. American Journal of Psychiatry, 2011, 168, 408-417. | 4.0 | 95 |
| 71 | A Genome-Wide Significant Linkage for Severe Depression on Chromosome 3: The Depression Network Study. American Journal of Psychiatry, 2011, 168, 840-847. | 4.0 | 51 |
| 72 | Large-scale genome-wide association analysis of bipolar disorder identifies a new susceptibility locus near ODZ4. Nature Genetics, 2011, 43, 977-983. | 9.4 | 1,283 |

| # | Article | IF | Citations |
|---|---|---|---|
| 73 | Evidence of statistical epistasis between DISC1, CIT and NDEL1 impacting risk for schizophrenia: biological validation with functional neuroimaging. Human Genetics, 2010, 127, 441-452. | 1.8 | 93 |
| 74 | Gene variants associated with schizophrenia in a Norwegian genome-wide study are replicated in a large European cohort. Journal of Psychiatric Research, 2010, 44, 748-753. | 1.5 | 183 |
| 75 | Stressful life events and the brain-derived neurotrophic factor gene in bipolar disorder. Journal of Affective Disorders, 2010, 125, 345-349. | 2.0 | 68 |
| 76 | The Bipolar Association Caseâ€“Control Study (BACCS) and metaâ€analysis: No association with the 5,10â€Methylenetetrahydrofolate reductase gene and bipolar disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2010, 153B, 1298-1304. | 1.1 | 26 |
| 77 | Population-based linkage analysis of schizophrenia and bipolar caseâ€“control cohorts identifies a potential susceptibility locus on 19q13. Molecular Psychiatry, 2010, 15, 319-325. | 4.1 | 38 |
| 78 | Genome-wide association study of recurrent major depressive disorder in two European caseâ€“control cohorts. Molecular Psychiatry, 2010, 15, 589-601. | 4.1 | 215 |
| 79 | Association of DISC1 and TSNAX genes and affective disorders in the depression caseâ€“control (DeCC) and bipolar affective caseâ€“control (BACCS) studies. Molecular Psychiatry, 2010, 15, 844-849. | 4.1 | 59 |
| 80 | Meta-analysis of genome-wide association data identifies a risk locus for major mood disorders on 3p21.1. Nature Genetics, 2010, 42, 128-131. | 9.4 | 152 |
| 81 | Meta-analysis and imputation refines the association of 15q25 with smoking quantity. Nature Genetics, 2010, 42, 436-440. | 9.4 | 581 |
| 82 | Genome-wide association study of migraine implicates a common susceptibility variant on 8q22.1. Nature Genetics, 2010, 42, 869-873. | 9.4 | 332 |
| 83 | Association analysis of DAOA and DAO in bipolar disorder: results from two independent caseâ€control studies. Bipolar Disorders, 2010, 12, 579-581. | 1.1 | 9 |
| 84 | A Genome-Wide Association Study of Neuroticism in a Population-Based Sample. PLoS ONE, 2010, 5, e11504. | 1.1 | 71 |
| 85 | Genome-Wide Association Study of Major Recurrent Depression in the U.K. Population. American Journal of Psychiatry, 2010, 167, 949-957. | 4.0 | 221 |
| 86 | A large replication study and meta-analysis in European samples provides further support for association of AHI1 markers with schizophrenia. Human Molecular Genetics, 2010, 19, 1379-1386. | 1.4 | 51 |
| 87 | Psychosis Susceptibility Gene ZNF804A and Cognitive Performance in Schizophrenia. Archives of General Psychiatry, 2010, 67, 692. | 13.8 | 129 |
| 88 | Genome-Wide Pharmacogenetics of Antidepressant Response in the GENDEP Project. American Journal of Psychiatry, 2010, 167, 555-564. | 4.0 | 314 |
| 89 | Biological Validation of Increased Schizophrenia Risk With NRG1, ERBB4, and AKT1 Epistasis via Functional Neuroimaging in Healthy Controls. Archives of General Psychiatry, 2010, 67, 991. | 13.8 | 113 |
| 90 | Plasma Protein Biomarkers for Depression and Schizophrenia by Multi Analyte Profiling of Case-Control Collections. PLoS ONE, 2010, 5, e9166. | 1.1 | 294 |

| # | Article | IF | Citations |
|---|---|---|---|
| 91 | Pathway-based approaches to imaging genetics association studies: Wnt signaling, GSK3beta substrates and major depression. NeuroImage, 2010, 53, 908-917. | 2.1 | 86 |
| 92 | Discovering genetic polymorphism associated with gene expression levels across the whole genome. , 2009, 2009, 5466-9. | | 1 |
| 93 | Genome-wide association and meta-analysis of bipolar disorder in individuals of European ancestry. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 7501-7506. | 3.3 | 274 |
| 94 | Association of GSK3Î² Polymorphisms With Brain Structural Changes in Major Depressive Disorder. Archives of General Psychiatry, 2009, 66, 721. | 13.8 | 121 |
| 95 | Influence of NOS1 on Verbal Intelligence and Working Memory in Both Patients With Schizophrenia and Healthy Control Subjects. Archives of General Psychiatry, 2009, 66, 1045. | 13.8 | 45 |
| 96 | A Genome-Wide Investigation of SNPs and CNVs in Schizophrenia. PLoS Genetics, 2009, 5, e1000373. | 1.5 | 383 |
| 97 | Genomewide Association Studies: History, Rationale, and Prospects for Psychiatric Disorders. American Journal of Psychiatry, 2009, 166, 540-556. | 4.0 | 391 |
| 98 | Association of the dystrobrevin binding protein 1 gene (DTNBP1) in a bipolar caseâ€“control study (BACCS). American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2009, 150B, 836-844. | 1.1 | 33 |
| 99 | The PsyCoLaus study: methodology and characteristics of the sample of a population-based survey on psychiatric disorders and their association with genetic and cardiovascular risk factors. BMC Psychiatry, 2009, 9, 9. | 1.1 | 182 |
| 100 | Common variants conferring risk of schizophrenia. Nature, 2009, 460, 744-747. | 13.7 | 1,572 |
| 101 | Genome-wide association study identifies 19p13.3 (UNC13A) and 9p21.2 as susceptibility loci for sporadic amyotrophic lateral sclerosis. Nature Genetics, 2009, 41, 1083-1087. | 9.4 | 344 |
| 102 | Failure to replicate effect of kibra on human memory in two large cohorts of European origin. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 667-668. | 1.1 | 62 |
| 103 | Serotonin transporter gene and adverse life events in adult ADHD. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 1461-1469. | 1.1 | 41 |
| 104 | Association study of brainâ€derived neurotrophic factor (BDNF) and LINâ€7 homolog (LIN‐7) genes with adult attentionâ€deficit/hyperactivity disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 945-951. | 1.1 | 45 |
| 105 | Large recurrent microdeletions associated with schizophrenia. Nature, 2008, 455, 232-236. | 13.7 | 1,619 |
| 106 | Î±-5/Î±-3 nicotinic receptor subunit alleles increase risk for heavy smoking. Molecular Psychiatry, 2008, 13, 368-373. | 4.1 | 437 |
| 107 | Investigation of the dopamine D5 receptor gene (DRD5) in adult attention deficit hyperactivity disorder. Neuroscience Letters, 2008, 432, 50-53. | 1.0 | 15 |
| 108 | Family history of depression is associated with younger age of onset in patients with recurrent depression. Psychological Medicine, 2008, 38, 641-649. | 2.7 | 53 |

| # | Article | IF | Citations |
|---|---|---|---|
| 109 | Hunting for Peripheral Biomarkers to Support Drug Development in Psychiatry. , 2008, , 405-426. | | 2 |
| 110 | The search for peripheral disease markers in psychiatry by genomic and proteomic approaches. Expert Opinion on Medical Diagnostics, 2007, 1, 235-251. | 1.6 | 23 |
| 111 | LRRTM1 on chromosome 2p12 is a maternally suppressed gene that is associated paternally with handedness and schizophrenia. Molecular Psychiatry, 2007, 12, 1129-1139. | 4.1 | 300 |
| 112 | LRRTM1 protein is located in the endoplasmic reticulum (ER) in mammalian cells. Molecular Psychiatry, 2007, 12, 1057-1057. | 4.1 | 8 |
| 113 | Factor structure and external validity of the PANSS revisited. Schizophrenia Research, 2006, 82, 213-223. | 1.1 | 124 |
| 114 | Decisionâ€Making Deficits and Overeating: A Risk Model for Obesity. Obesity, 2004, 12, 929-935. | 4.0 | 166 |
| 115 | No evidence of linkage or association between the norepinephrine transporter (NET) geneMnlI polymorphism and adult ADHD. American Journal of Medical Genetics Part A, 2004, 124B, 38-40. | 2.4 | 31 |
| 116 | Adrenergic alpha 2C receptor genomic organization: Association study in adult ADHD. American Journal of Medical Genetics Part A, 2004, 127B, 65-67. | 2.4 | 22 |
| 117 | The dopamine-4 receptor gene associated with binge eating and weight gain in women with seasonal affective disorder: An evolutionary perspective. Biological Psychiatry, 2004, 56, 665-669. | 0.7 | 94 |
| 118 | Pharmacogenetics of antipsychotic-induced weight gain. Pharmacological Research, 2004, 49, 309-329. | 3.1 | 69 |
| 119 | Childhood Inattention and Dysphoria and Adult Obesity Associated with the Dopamine D4 receptor Gene in Overeating Women with Seasonal Affective Disorder. Neuropsychopharmacology, 2004, 29, 179-186. | 2.8 | 90 |
| 120 | Linkage disequilibrium analysis of the dopamine beta-hydroxylase gene in persistent attention deficit hyperactivity disorder. Psychiatric Genetics, 2004, 14, 117-120. | 0.6 | 22 |
| 121 | Polymorphisms in glutamate decarboxylase genes: analysis in schizophrenia. Psychiatric Genetics, 2004, 14, 39-42. | 0.6 | 21 |
| 122 | Association between the BDNF gene and schizophrenia. Molecular Psychiatry, 2003, 8, 147-148. | 4.1 | 77 |
| 123 | Discovery of a null mutation in a human trace amine receptor gene. Genomics, 2003, 82, 531-536. | 1.3 | 28 |
| 124 | The Brain-Derived Neurotrophic Factor Gene Confers Susceptibility to Bipolar Disorder: Evidence from a Family-Based Association Study. American Journal of Human Genetics, 2002, 71, 651-655. | 2.6 | 544 |
| 125 | Psychiatric pharmacogenetics: personalizing psychostimulant therapy in attention-deficit/hyperactivity disorder. Behavioural Brain Research, 2002, 130, 85-90. | 1.2 | 27 |
| 126 | A transmission disequilibrium test of the Ser9/Gly dopamine D3 receptor gene polymorphism in adult attention-deficit hyperactivity disorder. Behavioural Brain Research, 2002, 130, 91-95. | 1.2 | 38 |

| # | Article | IF | Citations |
|---|---|---|---|
| 127 | 99A Quantitative Trait Locus Analysis of the Dopamine Transporter Gene in Adults with ADHD. Neuropsychopharmacology, 2002, 27, 655-62. | 2.8 | 32 |
| 128 | Dopamine D4 receptor and tyrosine hydroxylase genes in bipolar disorder: evidence for a role of DRD4. Molecular Psychiatry, 2002, 7, 860-866. | 4.1 | 77 |
| 129 | A Drosophila Model for Attention Deficit Hyperactivity Disorder (ADHD) : No Evidence of Association with PRKG1 Gene. NeuroMolecular Medicine, 2002, 2, 281-288. | 1.8 | 12 |
| 130 | Adult attention deficit hyperactivity disorder and the dopamine D4 receptor gene. American Journal of Medical Genetics Part A, 2000, 96, 273-277. | 2.4 | 127 |
| 131 | Long-Term Risperidone for Pervasive Developmental Disorder: Efficacy, Tolerability, and Discontinuation. Journal of Child and Adolescent Psychopharmacology, 2000, 10, 79-90. | 0.7 | 93 |
| 132 | Attention-deficit/hyperactivity disorder: a neuropsychiatric disorder with childhood onset. European Journal of Paediatric Neurology, 2000, 4, 53-62. | 0.7 | 29 |
| 133 | The Neurodevelopmental Hypothesis of Schizophrenia: Genetic Investigations. CNS Spectrums, 1999, 4, 78-84. | 0.7 | 3 |
| 134 | Effects of the calcium antagonist isradipine on cocaine intravenous self-administration in rats. Psychopharmacology, 1994, 113, 378-380. | 1.5 | 41 |
| 135 | Calcium antagonists antagonize cocaine-induced place-preference and self-administration in rats. Pharmacological Research, 1992, 26, 78. | 3.1 | 1 |