

Csaba Barta

List of Publications by Year in descending order

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Version: 2024-02-01

60
papers

2,527
citations

185998

28
h-index

214527

47
g-index

74
all docs

74
docs citations

74
times ranked

3886
citing authors

#	ARTICLE	IF	CITATIONS
1	Development and validation of the Reward Deficiency Syndrome Questionnaire (RDSQ-29). <i>Journal of Psychopharmacology</i> , 2022, 36, 409-422.	2.0	14
2	FOXP3 and GDNF Polymorphisms as Common Genetic Factors of Substance Use and Addictive Behaviors. <i>Journal of Personalized Medicine</i> , 2022, 12, 690.	1.1	10
3	Synaptic processes and immune-related pathways implicated in Tourette syndrome. <i>Translational Psychiatry</i> , 2021, 11, 56.	2.4	31
4	Riluzole Administration to Rats with Levodopa-Induced Dyskinesia Leads to Loss of DNA Methylation in Neuronal Genes. <i>Cells</i> , 2021, 10, 1442.	1.8	0
5	Investigating Shared Genetic Basis Across Tourette Syndrome and Comorbid Neurodevelopmental Disorders Along the Impulsivity-Compulsivity Spectrum. <i>Biological Psychiatry</i> , 2021, 90, 317-327.	0.7	49
6	Elevated common variant genetic risk for tourette syndrome in a densely-affected pedigree. <i>Molecular Psychiatry</i> , 2021, 26, 7522-7529.	4.1	8
7	Genetics of obsessive-compulsive disorder and Tourette disorder. , 2020, , 239-252.		1
8	International Consensus Statement for the Screening, Diagnosis, and Treatment of Adolescents with Concurrent Attention-Deficit/Hyperactivity Disorder and Substance Use Disorder. <i>European Addiction Research</i> , 2020, 26, 223-232.	1.3	28
9	Association of Genetic Variation in the 3'UTR of LHX6, IMMP2L, and AADAC With Tourette Syndrome. <i>Frontiers in Neurology</i> , 2020, 11, 803.	1.1	13
10	Validity of the ADHD module of the Mini International Neuropsychiatric Interview PLUS for screening of adult ADHD in treatment seeking substance use disorder patients: ADHD screening with MINI-Plus. <i>Revista De Psiquiatria Y Salud Mental</i> , 2020, , .	1.0	6
11	Co-occurrences of substance use and other potentially addictive behaviors: Epidemiological results from the Psychological and Genetic Factors of the Addictive Behaviors (PGA) Study. <i>Journal of Behavioral Addictions</i> , 2020, 9, 272-288.	1.9	56
12	Association of GDNF and CNTNAP2 gene variants with gambling. <i>Journal of Behavioral Addictions</i> , 2019, 8, 471-478.	1.9	1
13	Interrogating the Genetic Determinants of Tourette's Syndrome and Other Tic Disorders Through Genome-Wide Association Studies. <i>American Journal of Psychiatry</i> , 2019, 176, 217-227.	4.0	242
14	T12THE EFFECT OF ETHANOL EXPOSURE ON LONGEVITY AND LOCOMOTION IN A CAENORHABDITIS ELEGANS ANIMAL MODEL. <i>European Neuropsychopharmacology</i> , 2019, 29, S225-S226.	0.3	0
15	New insights and perspectives on the genetics of obsessive-compulsive disorder. <i>Psychiatric Genetics</i> , 2019, 29, 142-151.	0.6	16
16	The psychological and genetic factors of the addictive behaviors (PGA) study. <i>International Journal of Methods in Psychiatric Research</i> , 2019, 28, e1748.	1.1	16
17	Riluzole Attenuates L-DOPA-Induced Abnormal Involuntary Movements Through Decreasing CREB1 Activity: Insights from a Rat Model. <i>Molecular Neurobiology</i> , 2019, 56, 5111-5121.	1.9	3
18	Persistence and Subtype Stability of ADHD Among Substance Use Disorder Treatment Seekers. <i>Journal of Attention Disorders</i> , 2019, 23, 1438-1453.	1.5	34

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19	De Novo Sequence and Copy Number Variants Are Strongly Associated with Tourette Disorder and Implicate Cell Polarity in Pathogenesis. <i>Cell Reports</i> , 2018, 24, 3441-3454.e12.	2.9	91
20	Rare Copy Number Variants in NRXN1 and CNTN6 Increase Risk for Tourette Syndrome. <i>Neuron</i> , 2017, 94, 1101-1111.e7.	3.8	137
21	Effect of Chronic Treatment With Psychiatric Medications Aripiprazole And Riluzole on Dna Methylation Profiles In The Rat Striatum And Prefrontal Cortex. <i>European Neuropsychopharmacology</i> , 2017, 27, S394.	0.3	0
22	Childhood trauma exposure in substance use disorder patients with and without ADHD. <i>Addictive Behaviors</i> , 2017, 65, 118-124.	1.7	26
23	From Genetics to Epigenetics: New Perspectives in Tourette Syndrome Research. <i>Frontiers in Neuroscience</i> , 2016, 10, 277.	1.4	40
24	Targeted Re-Sequencing Approach of Candidate Genes Implicates Rare Potentially Functional Variants in Tourette Syndrome Etiology. <i>Frontiers in Neuroscience</i> , 2016, 10, 428.	1.4	29
25	TS-EUROTRAIN: A European-Wide Investigation and Training Network on the Etiology and Pathophysiology of Gilles de la Tourette Syndrome. <i>Frontiers in Neuroscience</i> , 2016, 10, 384.	1.4	21
26	Association of AADAC Deletion and Gilles de la Tourette Syndrome in a Large European Cohort. <i>Biological Psychiatry</i> , 2016, 79, 383-391.	0.7	41
27	Epigenome-Wide Association Study of Tic Disorders. <i>Twin Research and Human Genetics</i> , 2015, 18, 699-709.	0.3	31
28	Risk Factors for Borderline Personality Disorder in Treatment Seeking Patients with a Substance Use Disorder: An International Multicenter Study. <i>European Addiction Research</i> , 2015, 21, 188-194.	1.3	11
29	Early developmental, temperamental and educational problems in "substance use disorder"™ patients with and without ADHD. Does ADHD make a difference?. <i>Addictive Behaviors Reports</i> , 2015, 2, 13-18.	1.0	4
30	Psychiatric comorbidity in treatment-seeking substance use disorder patients with and without attention deficit hyperactivity disorder: results of the <sc>IASP</sc> study. <i>Addiction</i> , 2014, 109, 262-272.	1.7	129
31	Genetic association signal near <sc><i>NTN4</i></sc> in <sc>T</sc>ourette syndrome. <i>Annals of Neurology</i> , 2014, 76, 310-315.	2.8	53
32	Variability in the prevalence of adult ADHD in treatment seeking substance use disorder patients: Results from an international multi-center study exploring DSM-IV and DSM-5 criteria. <i>Drug and Alcohol Dependence</i> , 2014, 134, 158-166.	1.6	150
33	The Genetics of Problem and Pathological Gambling: A Systematic Review. <i>Current Pharmaceutical Design</i> , 2014, 20, 3993-3999.	0.9	57
34	Support of the histaminergic hypothesis in Tourette Syndrome: association of the histamine decarboxylase gene in a large sample of families. <i>Journal of Medical Genetics</i> , 2013, 50, 760-764.	1.5	92
35	Validity of the Adult ADHD Self-Report Scale (ASRS) as a screener for adult ADHD in treatment seeking substance use disorder patients. <i>Drug and Alcohol Dependence</i> , 2013, 132, 587-596.	1.6	135
36	The International ADHD in Substance Use Disorders Prevalence (IASP) study: background, methods and study population. <i>International Journal of Methods in Psychiatric Research</i> , 2013, 22, 232-244.	1.1	53

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37	Multivariate Analysis of Dopaminergic Gene Variants as Risk Factors of Heroin Dependence. PLoS ONE, 2013, 8, e66592.	1.1	67
38	Replication of association between a SLITRK1 haplotype and Tourette Syndrome in a large sample of families. Molecular Psychiatry, 2012, 17, 665-668.	4.1	69
39	A global view of the OCA2-HERC2 region and pigmentation. Human Genetics, 2012, 131, 683-696.	1.8	113
40	Evaluation of the LIM homeobox genes <i>LHX6</i> and <i>LHX8</i> as candidates for Tourette syndrome. Genes, Brain and Behavior, 2012, 11, 444-451.	1.1	23
41	The Distribution and Most Recent Common Ancestor of the 17q21 Inversion in Humans. American Journal of Human Genetics, 2010, 86, 161-171.	2.6	59
42	The complex global pattern of genetic variation and linkage disequilibrium at catechol-O-methyltransferase. Molecular Psychiatry, 2010, 15, 216-225.	4.1	48
43	Association between Novelty Seeking of opiate-dependent patients and the catechol-O-methyltransferase Val158Met polymorphism. Comprehensive Psychiatry, 2010, 51, 510-515.	1.5	32
44	Haplotype evolution of SLITRK1, a candidate gene for Gilles de la Tourette Syndrome. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 463-466.	1.1	16
45	Geographically Separate Increases in the Frequency of the Derived ADH1B*47His Allele in Eastern and Western Asia. American Journal of Human Genetics, 2007, 81, 842-846.	2.6	90
46	No direct effect of the -521 C/T polymorphism in the human dopamine D4 receptor gene promoter on transcriptional activity. BMC Molecular Biology, 2006, 7, 18.	3.0	39
47	Combined effect of promoter polymorphisms in the dopamine D4 receptor and the serotonin transporter genes in heroin dependence. Neuropsychopharmacologia Hungarica, 2005, 7, 28-33.	0.1	29
48	Mutational Analysis of Hungarian Patients with Androgen Insensitivity Syndrome. Journal of Pediatric Endocrinology and Metabolism, 2003, 16, 367-73.	0.4	5
49	Hormonal evaluation and mutation screening for steroid 21-hydroxylase deficiency in patients with unilateral and bilateral adrenal incidentalomas. European Journal of Endocrinology, 2002, 147, 349-355.	1.9	33
50	Membrane-mediated ultrafast restriction digestion and subsequent rapid gel microchip electrophoresis of DNA. Electrophoresis, 2002, 23, 1524.	1.3	15
51	Analysis of Mutations in the Plasma Cholinesterase Gene of Patients with a History of Prolonged Neuromuscular Block during Anesthesia. Molecular Genetics and Metabolism, 2001, 74, 484-488.	0.5	20
52	Prenatal Diagnosis of Steroid 21-Hydroxylase Deficiency by Allele-Specific Amplification. Fetal Diagnosis and Therapy, 2001, 16, 237-240.	0.6	3
53	Genotyping the -521C/T functional polymorphism in the promoter region of dopamine D4 receptor (DRD4) gene. Electrophoresis, 2001, 22, 1102-1105.	1.3	30
54	DNA analysis on electrophoretic microchips: Effect of operational variables. Electrophoresis, 2001, 22, 294-299.	1.3	45

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55	Rapid single nucleotide polymorphism analysis by primer extension and capillary electrophoresis using polyvinyl pyrrolidone matrix. <i>Electrophoresis</i> , 2001, 22, 779-782.	1.3	30
56	Analysis of dopamine D4 receptor gene polymorphism using microchip electrophoresis. <i>Journal of Chromatography A</i> , 2001, 924, 285-290.	1.8	23
57	Screening for Mutations of 21-Hydroxylase Gene in Hungarian Patients with Congenital Adrenal Hyperplasia1. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1999, 84, 2369-2372.	1.8	43
58	Human dopamine D4 receptor allele genotyping by ultrathin agarose gel electrophoresis with To-Pro-3 complexation. <i>Electrophoresis</i> , 1999, 20, 497-501.	1.3	10
59	Simultaneous analysis of various mutations on the 21-hydroxylase gene by multi-allele specific amplification and capillary gel electrophoresis. <i>Journal of Chromatography A</i> , 1998, 817, 281-286.	1.8	17
60	Real-time detection of allele-specific polymerase chain reaction products by automated ultra-thin-layer agarose gel electrophoresis. <i>Journal of Chromatography A</i> , 1998, 828, 481-487.	1.8	25