

# Nagahide Takahashi

## List of Publications by Year in descending order

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Version: 2024-02-01

57  
papers

4,082  
citations

201385

27  
h-index

161609

54  
g-index

58  
all docs

58  
docs citations

58  
times ranked

7277  
citing authors

#	ARTICLE	IF	CITATIONS
1	Autism spectrum disorder comorbid with obsessive compulsive disorder and eating disorder in a woman with <i>NBEA</i> deletion. <i>Psychiatry and Clinical Neurosciences</i> , 2022, 76, 36-38.	1.0	0
2	Early temperament as a predictor of language skills at 40 months. <i>BMC Pediatrics</i> , 2022, 22, 56.	0.7	1
3	Trajectories of Adaptive Behaviors During Childhood in Females and Males in the General Population. <i>Frontiers in Psychiatry</i> , 2022, 13, 817383.	1.3	5
4	Identification of neurodevelopmental transition patterns from infancy to early childhood and risk factors predicting descending transition. <i>Scientific Reports</i> , 2022, 12, 4822.	1.6	2
5	Long-term effect of persistent postpartum depression on children's psychological problems in childhood. <i>Journal of Affective Disorders</i> , 2022, 305, 71-76.	2.0	9
6	Peripheral biomarkers of attention-deficit hyperactivity disorder: Current status and future perspective. <i>Journal of Psychiatric Research</i> , 2021, 137, 465-470.	1.5	12
7	Elevated risk of attention deficit hyperactivity disorder (ADHD) in Japanese children with higher genetic susceptibility to ADHD with a birth weight under 2000 g. <i>BMC Medicine</i> , 2021, 19, 229.	2.3	10
8	Associations Among Maternal Metabolic Conditions, Cord Serum Leptin Levels, and Autistic Symptoms in Children. <i>Frontiers in Psychiatry</i> , 2021, 12, 816196.	1.3	4
9	Polygenic risk score analysis revealed shared genetic background in attention deficit hyperactivity disorder and narcolepsy. <i>Translational Psychiatry</i> , 2020, 10, 284.	2.4	17
10	Association of Genetic Risks With Autism Spectrum Disorder and Early Neurodevelopmental Delays Among Children Without Intellectual Disability. <i>JAMA Network Open</i> , 2020, 3, e1921644.	2.8	21
11	A randomized, double-blind, placebo-controlled, parallel-group study to evaluate the efficacy and safety of osmotic-controlled release oral delivery system methylphenidate HCl in adults with attention-deficit/hyperactivity disorder in Japan. <i>World Journal of Biological Psychiatry</i> , 2014, 15, 488-498.	1.3	13
12	Introduction of the human <i>AVPR1A</i> gene significantly alters brain receptor expression patterns, and may enhance aspects of social behavior in transgenic mice. <i>DMM Disease Models and Mechanisms</i> , 2014, 7, 1013-22.	1.2	17
13	Roles of glial cells in schizophrenia: Possible targets for therapeutic approaches. <i>Neurobiology of Disease</i> , 2013, 53, 49-60.	2.1	59
14	Randomized, placebo-controlled, double-blind study assessing the efficacy and safety of paliperidone palmitate in Asian patients with schizophrenia. <i>Neuropsychiatric Disease and Treatment</i> , 2013, 9, 1889.	1.0	25
15	Copy Number Variations in Alternative Splicing Gene Networks Impact Lifespan. <i>PLoS ONE</i> , 2013, 8, e53846.	1.1	13
16	HDAC2 regulates atypical antipsychotic responses through the modulation of mGlu2 promoter activity. <i>Nature Neuroscience</i> , 2012, 15, 1245-1254.	7.1	247
17	Genome-wide copy number variation study associates metabotropic glutamate receptor gene networks with attention deficit hyperactivity disorder. <i>Nature Genetics</i> , 2012, 44, 78-84.	9.4	334
18	Haploinsufficiency of <i>Cyfp1</i> Produces Fragile X-Like Phenotypes in Mice. <i>PLoS ONE</i> , 2012, 7, e42422.	1.1	95

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19	Rare structural variation of synapse and neurotransmission genes in autism. <i>Molecular Psychiatry</i> , 2012, 17, 402-411.	4.1	151
20	Loss of Function Studies in Mice and Genetic Association Link Receptor Protein Tyrosine Phosphatase $\hat{\pm}$ to Schizophrenia. <i>Biological Psychiatry</i> , 2011, 70, 626-635.	0.7	22
21	Linking oligodendrocyte and myelin dysfunction to neurocircuitry abnormalities in schizophrenia. <i>Progress in Neurobiology</i> , 2011, 93, 13-24.	2.8	263
22	Association analysis of the GDNF gene with methamphetamine use disorder in a Japanese population. <i>Progress in Neuro-Psychopharmacology and Biological Psychiatry</i> , 2011, 35, 1268-1272.	2.5	13
23	Analysis of a purported SHANK3 mutation in a boy with autism: Clinical impact of rare variant research in neurodevelopmental disabilities. <i>Brain Research</i> , 2011, 1380, 98-105.	1.1	28
24	Haploinsufficiency of <i>Gtf2i</i> , a gene deleted in Williams Syndrome, leads to increases in social interactions. <i>Autism Research</i> , 2011, 4, 28-39.	2.1	109
25	Increased expression of receptor phosphotyrosine phosphatase- $\hat{2}/\hat{1}\uparrow$ is associated with molecular, cellular, behavioral and cognitive schizophrenia phenotypes. <i>Translational Psychiatry</i> , 2011, 1, e8-e8.	2.4	37
26	A Genome-wide Study Reveals Copy Number Variants Exclusive to Childhood Obesity Cases. <i>American Journal of Human Genetics</i> , 2010, 87, 661-666.	2.6	91
27	Extensive proteomic screening identifies the obesity-related NYGGF4 protein as a novel LRP1-interactor, showing reduced expression in early Alzheimer's disease. <i>Molecular Neurodegeneration</i> , 2010, 5, 1.	4.4	51
28	Haploinsufficiency of the autism-associated Shank3 gene leads to deficits in synaptic function, social interaction, and social communication. <i>Molecular Autism</i> , 2010, 1, 15.	2.6	521
29	Relationship of psychopathological symptoms and cognitive function to subjective quality of life in patients with chronic schizophrenia. <i>Psychiatry and Clinical Neurosciences</i> , 2010, 64, 62-69.	1.0	44
30	Strong synaptic transmission impact by copy number variations in schizophrenia. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 10584-10589.	3.3	212
31	Slc25a12 Disruption Alters Myelination and Neurofilaments: A Model for a Hypomyelination Syndrome and Childhood Neurodevelopmental Disorders. <i>Biological Psychiatry</i> , 2010, 67, 887-894.	0.7	47
32	Gene-wide association study between the methylenetetrahydrofolate reductase gene (MTHFR) and schizophrenia in the Japanese population, with an updated meta-analysis on currently available data. <i>Schizophrenia Research</i> , 2010, 124, 216-222.	1.1	28
33	Linking white and grey matter in schizophrenia: Oligodendrocyte and neuron pathology in the prefrontal cortex. <i>Frontiers in Neuroanatomy</i> , 2009, 3, 9.	0.9	67
34	Common genetic variants on 5p14.1 associate with autism spectrum disorders. <i>Nature</i> , 2009, 459, 528-533.	13.7	912
35	No association between the protein tyrosine phosphatase, receptor-type, Z Polypeptide 1 ( <i>PTPRZ1</i> ) gene and schizophrenia in the Japanese population. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2008, 147B, 1013-1018.	1.1	4
36	No association between tagging SNPs of SNARE complex genes (STX1A, VAMP2 and SNAP25) and schizophrenia in a Japanese population. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2008, 147B, 1327-1331.	1.1	18

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37	Genetic analysis of the gene coding for DARPP-32 (PPP1R1B) in Japanese patients with schizophrenia or bipolar disorder. <i>Schizophrenia Research</i> , 2008, 100, 334-341.	1.1	15
38	Failure to replicate the association between NRG1 and schizophrenia using Japanese large sample. <i>Schizophrenia Research</i> , 2008, 101, 1-8.	1.1	30
39	Relationship between three serotonin receptor subtypes (HTR3A, HTR2A and HTR4) and treatment-resistant schizophrenia in the Japanese population. <i>Neuroscience Letters</i> , 2008, 435, 95-98.	1.0	35
40	Association between the brain-derived neurotrophic factor Val66Met polymorphism and brain morphology in a Japanese sample of schizophrenia and healthy comparisons. <i>Neuroscience Letters</i> , 2008, 435, 34-39.	1.0	42
41	A genetic association study of the FXYD domain containing ion transport regulator 6 (FXYD6) gene, encoding phosphohippolin, in susceptibility to schizophrenia in a Japanese population. <i>Neuroscience Letters</i> , 2008, 438, 70-75.	1.0	11
42	The association of genotypic combination of the DRD3 and BDNF polymorphisms on the adhesion interthalamic and medial temporal lobe structures. <i>Progress in Neuro-Psychopharmacology and Biological Psychiatry</i> , 2008, 32, 1236-1242.	2.5	28
43	Pathway-based association analysis of genome-wide screening data suggest that genes associated with the $\Gamma^3$ -aminobutyric acid receptor signaling pathway are involved in neuroleptic-induced, treatment-resistant tardive dyskinesia. <i>Pharmacogenetics and Genomics</i> , 2008, 18, 317-323.	0.7	95
44	An association study of tachykinin receptor 3 gene with schizophrenia in the Japanese population. <i>NeuroReport</i> , 2008, 19, 471-473.	0.6	7
45	Association of SOX10 with schizophrenia in the Japanese population. <i>Psychiatric Genetics</i> , 2007, 17, 227-231.	0.6	27
46	Association study between the transferrin gene and schizophrenia in the Japanese population. <i>NeuroReport</i> , 2007, 18, 517-520.	0.6	6
47	Gene-gene interaction analysis of personality traits in a Japanese population using an electrochemical DNA array chip analysis. <i>Neuroscience Letters</i> , 2007, 414, 209-212.	1.0	27
48	Gap junction coding genes and schizophrenia: a genetic association study. <i>Journal of Human Genetics</i> , 2007, 52, 498-501.	1.1	28
49	Association between chromogranin A gene polymorphism and schizophrenia in the Japanese population. <i>Schizophrenia Research</i> , 2006, 83, 179-183.	1.1	16
50	The $2\text{-},3\text{-}$ cyclic nucleotide $3\text{-}$ phosphodiesterase and oligodendrocyte lineage transcription factor 2 genes do not appear to be associated with schizophrenia in the Japanese population. <i>Schizophrenia Research</i> , 2006, 88, 245-250.	1.1	16
51	Association between gene polymorphisms of SLC22A3 and methamphetamine use disorder. <i>International Clinical Psychopharmacology</i> , 2006, 21, A32.	0.9	0
52	Association Between Gene Polymorphisms of SLC22A3 and Methamphetamine Use Disorder. <i>Alcoholism: Clinical and Experimental Research</i> , 2006, 30, 1644-1649.	1.4	49
53	Association study between kynurenine 3-monooxygenase gene and schizophrenia in the Japanese population. <i>Genes, Brain and Behavior</i> , 2006, 5, 364-368.	1.1	42
54	Association Study between Vesicle-Associated Membrane Protein 2 Gene Polymorphisms and Fluvoxamine Response in Japanese Major Depressive Patients. <i>Neuropsychobiology</i> , 2006, 54, 226-230.	0.9	15

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55	No association was found between a functional SNP in ZDHHC8 and schizophrenia in a Japanese caseâ€“control population. <i>Neuroscience Letters</i> , 2005, 374, 21-24.	1.0	38
56	No association between monoamine oxidase A promoter polymorphism and personality traits in Japanese females. <i>Neuroscience Letters</i> , 2005, 389, 121-123.	1.0	13
57	The inter-rater reliability of the Japanese version of the Montgomeryâ€“Asberg depression rating scale(MADRS) using a structured interview guide for MADRS(SIGMA). <i>Human Psychopharmacology</i> , 2004, 19, 187-192.	0.7	39