Ming T Tsuang

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71 papers 11,749 26 p-index 9-index 76 ext. papers ext. citations 9.7 avg, IF L-index

#	Paper	IF	Citations
71	Analysis of protein-coding genetic variation in 60,706 humans. <i>Nature</i> , 2016 , 536, 285-91	50.4	6940
70	Hyperactivity and hyperconnectivity of the default network in schizophrenia and in first-degree relatives of persons with schizophrenia. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009 , 106, 1279-84	11.5	1067
69	Genome scan meta-analysis of schizophrenia and bipolar disorder, part II: Schizophrenia. <i>American Journal of Human Genetics</i> , 2003 , 73, 34-48	11	985
68	Comparative gene expression analysis of blood and brain provides concurrent validation of SELENBP1 up-regulation in schizophrenia. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005 , 102, 15533-8	11.5	271
67	Genome scan of European-American schizophrenia pedigrees: Results of the NIMH genetics initiative and millennium consortium 1998 , 81, 290-295		212
66	NIMH genetics initiative millennium schizophrenia consortium: Linkage analysis of African-American pedigrees 1998 , 81, 282-289		208
65	Analysis of 94 candidate genes and 12 endophenotypes for schizophrenia from the Consortium on the Genetics of Schizophrenia. <i>American Journal of Psychiatry</i> , 2011 , 168, 930-46	11.9	201
64	Association of Thalamic Dysconnectivity and Conversion to Psychosis in Youth and Young Adults at Elevated Clinical Risk. <i>JAMA Psychiatry</i> , 2015 , 72, 882-91	14.5	199
63	Assessing the validity of blood-based gene expression profiles for the classification of schizophrenia and bipolar disorder: a preliminary report. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2005 , 133B, 1-5	3.5	180
62	Linkage disequilibrium for schizophrenia at the chromosome 15q13-14 locus of the 1 -nicotinic acetylcholine receptor subunit gene (CHRNA7). <i>American Journal of Medical Genetics Part A</i> , 2001 , 105, 20-22		164
61	Altered brain activation in dorsolateral prefrontal cortex in adolescents and young adults at genetic risk for schizophrenia: an fMRI study of working memory. <i>Schizophrenia Research</i> , 2006 , 85, 58-7	.3.6	116
60	Evidence for the multigenic inheritance of schizophrenia. <i>American Journal of Medical Genetics Part A</i> , 2001 , 105, 794-800		88
59	Cerebello-thalamo-cortical hyperconnectivity as a state-independent functional neural signature for psychosis prediction and characterization. <i>Nature Communications</i> , 2018 , 9, 3836	17.4	83
58	Examination of genetic linkage of chromosome 15 to schizophrenia in a large Veterans Affairs Cooperative Study sample*. <i>American Journal of Medical Genetics Part A</i> , 2001 , 105, 662-668		69
57	Genotypes, phenotypes, and the brain. A search for connections in schizophrenia. <i>British Journal of Psychiatry</i> , 1993 , 163, 299-307	5.4	67
56	Genome scan of Han Chinese schizophrenia families from Taiwan: confirmation of linkage to 10q22.3. <i>American Journal of Psychiatry</i> , 2006 , 163, 1760-6	11.9	65
55	How do genes influence marijuana use? The role of subjective effects. <i>Addiction</i> , 1997 , 92, 409-417	4.6	57

54	Genetic association analysis of behavioral inhibition using candidate loci from mouse models. American Journal of Medical Genetics Part A, 2001 , 105, 226-35		56	
53	Genetic assessment of additional endophenotypes from the Consortium on the Genetics of Schizophrenia Family Study. <i>Schizophrenia Research</i> , 2016 , 170, 30-40	3.6	46	
52	Convergent Functional Genomics of bipolar disorder: from animal model pharmacogenomics to human genetics and biomarkers. <i>Neuroscience and Biobehavioral Reviews</i> , 2007 , 31, 897-903	9	46	
51	Regional prefrontal cortex gray matter volumes in youth at familial risk for schizophrenia from the Harvard Adolescent High Risk Study. <i>Schizophrenia Research</i> , 2010 , 123, 15-21	3.6	43	
50	RASD2, MYH9, and CACNG2 genes at chromosome 22q12 associated with the subgroup of schizophrenia with non-deficit in sustained attention and executive function. <i>Biological Psychiatry</i> , 2008 , 64, 789-96	7.9	43	
49	Genetic liability, prenatal health, stress and family environment: risk factors in the Harvard Adolescent Family High Risk for schizophrenia study. <i>Schizophrenia Research</i> , 2014 , 157, 142-8	3.6	37	
48	Epidemiology of Mood and Anxiety Disorders in Children and Adolescents657-704		36	
47	Genome-wide linkage scan of schizophrenia: a cross-isolate study. <i>Genomics</i> , 2007 , 89, 167-77	4.3	35	
46	Mapping genomic loci implicates genes and synaptic biology in schizophrenia Nature, 2022,	50.4	35	
45	Defining alternative phenotypes for genetic studies: What can we learn from studies of schizophrenia?. <i>American Journal of Medical Genetics Part A</i> , 2001 , 105, 8-10		26	
44	Five NOTCH4 polymorphisms show weak evidence for association with schizophrenia: evidence from meta-analyses. <i>Schizophrenia Research</i> , 2005 , 73, 281-90	3.6	23	
43	The National Comorbidity Survey343-362		23	
42	Symptom Scales and Diagnostic Schedules in Adult Psychiatry273-332		22	
41	Medial temporal lobe default mode functioning and hippocampal structure as vulnerability indicators for schizophrenia: a MRI study of non-psychotic adolescent first-degree relatives. <i>Schizophrenia Research</i> , 2014 , 159, 426-34	3.6	21	
40	Comparison of schizotypal relatives of schizophrenic versus affective probands. <i>American Journal of Medical Genetics Part A</i> , 1994 , 54, 279-85		21	
39	ANXA7, PPP3CB, DNAJC9, and ZMYND17 genes at chromosome 10q22 associated with the subgroup of schizophrenia with deficits in attention and executive function. <i>Biological Psychiatry</i> , 2011 , 70, 51-8	7.9	16	
38	Department of Veterans Affairs Cooperative Studies Program genetic linkage study of schizophrenia: ascertainment methods and sample description. <i>American Journal of Medical Genetics Part A</i> , 2000 , 96, 342-7		16	
37	Epidemiology of suicide. <i>International Review of Psychiatry</i> , 1992 , 4, 117-129	3.6	16	

36	Rare coding variants in ten genes confer substantial risk for schizophrenia Nature, 2022,	50.4	16
35	The multidimensionality of schizotypy in nonpsychotic relatives of patients with schizophrenia and its applications in ordered subsets linkage analysis of schizophrenia. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2010 , 153B, 1-9	3.5	14
34	Epidemiology of Bipolar Disorder427-444		14
33	Epidemiology of Depressive and Anxiety Disorders389-426		14
32	Suggestive linkage of chromosome 10p to schizophrenia is not due to transmission ratio distortion. <i>American Journal of Medical Genetics Part A</i> , 1999 , 88, 607-8		14
31	Progressive reconfiguration of resting-state brain networks as psychosis develops: Preliminary results from the North American Prodrome Longitudinal Study (NAPLS) consortium. <i>Schizophrenia Research</i> , 2020 , 226, 30-37	3.6	14
30	Medial prefrontal cortical activation during working memory differentiates schizophrenia and bipolar psychotic patients: a pilot FMRI study. <i>Schizophrenia Research</i> , 2011 , 129, 208-10	3.6	13
29	Family-based association study of SELENBP1 in schizophrenia. Schizophrenia Research, 2009, 113, 268-7	723.6	13
28	New phenotype definition of attention deficit hyperactivity disorder in relatives for genetic analyses. <i>American Journal of Medical Genetics Part A</i> , 1996 , 67, 369-77		11
27	Evaluating the impact of cannabis use on thalamic connectivity in youth at clinical high risk of psychosis. <i>BMC Psychiatry</i> , 2015 , 15, 276	4.2	10
26	The Epidemiology of Child and Adolescent Mental Disorders629-655		10
25	Epidemiology of Schizophrenia 2011 , 263-287		7
24	Neurocognitive and clinical dysfunction in adult Chinese, nonpsychotic relatives of patients with schizophrenia: Findings from the Changsha study and evidence for schizotaxia. <i>Asian Journal of Psychiatry</i> , 2012 , 5, 83-92	6.7	5
23	Epidemiology of Drug Dependence479-561		5
22	Validity: Definitions and Applications to Psychiatric Research149-163		5
21	The Epidemiology of First-Onset Mania445-458		5
20	Analysis of Categorized Data: Use of the Odds Ratio as a Measure of Association35-63		4
19	Birth and Development of Psychiatric Interviews257-271		4

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18	Genome scan of European-American schizophrenia pedigrees: Results of the NIMH genetics initiative and millennium consortium 1998 , 81, 290		4
17	A Frameshift Variant in the CHST9 Gene Identified by Family-Based Whole Genome Sequencing Is Associated with Schizophrenia in Chinese Population. <i>Scientific Reports</i> , 2019 , 9, 12717	4.9	2
16	DSM-IV and Psychiatric Epidemiology333-342		2
15	Mental Health Services Research165-179		2
14	Studying the Natural History of Psychopathology213-238		2
13	Introduction to Epidemiologic Research Methods1-33		2
12	The Developmental Epidemiology of Psychiatric Disorders239-255		2
11	Methods in Psychiatric Genetics65-130		2
10	The Pharmacoepidemiology of Psychiatric Medications181-194		2
9	Conceptualization of the liability for schizophrenia: clinical implications. <i>Dialogues in Clinical Neuroscience</i> , 1999 , 1, 153-64	5.7	2
8	Schizophrenia: family studies and treatment of spectrum disorders. <i>Dialogues in Clinical Neuroscience</i> , 2000 , 2, 381-91	5.7	2
7	Personality Disorders: Epidemiological Findings, Methods, and Concepts563-599		1
6	Epidemiology and Geriatric Psychiatry601-628		1
5	The Epidemiology of Alcohol Use, Abuse, and Dependence459-477		1
4	Epidemiology of Psychosis with Special Reference to Schizophrenia363-387		1
3	Mental Health Etiology: Biological and Genetic Determinants 2017 , 63-68		
2	Peering into the Future of Psychiatric Epidemiology195-211		
1	Schizophrenia: vulnerability versus disease. <i>Dialogues in Clinical Neuroscience</i> , 2000 , 2, 257-66	5.7	