Xiao-Qing Liu

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4327432/publications.pdf

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25 papers

6,055 citations

361413 20 h-index 25 g-index

25 all docs

25 docs citations

25 times ranked

8707 citing authors

#	Article	IF	CITATIONS
1	Functional impact of global rare copy number variation in autism spectrum disorders. Nature, 2010, 466, 368-372.	27.8	1,803
2	Mapping autism risk loci using genetic linkage and chromosomal rearrangements. Nature Genetics, 2007, 39, 319-328.	21.4	1,272
3	A genome-wide linkage and association scan reveals novel loci for autism. Nature, 2009, 461, 802-808.	27.8	570
4	A genome-wide scan for common alleles affecting risk for autism. Human Molecular Genetics, 2010, 19, 4072-4082.	2.9	538
5	Dimensions of Religiosity and Their Relationship to Lifetime Psychiatric and Substance Use Disorders. American Journal of Psychiatry, 2003, 160, 496-503.	7.2	425
6	Individual common variants exert weak effects on the risk for autism spectrum disorders. Human Molecular Genetics, 2012, 21, 4781-4792.	2.9	334
7	The Novel Neuronal Ceroid Lipofuscinosis Gene MFSD8 Encodes a Putative Lysosomal Transporter. American Journal of Human Genetics, 2007, 81, 136-146.	6.2	180
8	Sex differences in repetitive stereotyped behaviors in autism: Implications for genetic liability. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 5-12.	1.7	154
9	Genome-Wide Linkage Scan of a Large Family with IgA Nephropathy Localizes a Novel Susceptibility Locus to Chromosome 2q36. Journal of the American Society of Nephrology: JASN, 2007, 18, 2408-2415.	6.1	112
10	Multiple Superoxide Dismutase 1/Splicing Factor Serine Alanine 15 Variants Are Associated With the Development and Progression of Diabetic Nephropathy. Diabetes, 2008, 57, 218-228.	0.6	89
11	Multiple Variants in Vascular Endothelial Growth Factor (VEGFA) Are Risk Factors for Time to Severe Retinopathy in Type 1 Diabetes. Diabetes, 2007, 56, 2161-2168.	0.6	88
12	Genome-wide Linkage Analyses of Quantitative and Categorical Autism Subphenotypes. Biological Psychiatry, 2008, 64, 561-570.	1.3	80
13	ABCG2 null alleles define the Jr(aâ^²) blood group phenotype. Nature Genetics, 2012, 44, 131-132.	21.4	77
14	Stepwise formation of a SMAD activity gradient during dorsal-ventral patterning of the Drosophila embryo. Development (Cambridge), 2003, 130, 5705-5716.	2.5	66
15	Genetic Variation of DKK3 May Modify Renal Disease Severity in ADPKD. Journal of the American Society of Nephrology: JASN, 2010, 21, 1510-1520.	6.1	59
16	Assessing the validity of the association between the SUMO4 M55V variant and risk of type 1 diabetes. Nature Genetics, 2005, 37, 111-112.	21.4	47
17	IL5RA and TNFRSF6B Gene Variants Are Associated With Sporadic IgA Nephropathy. Journal of the American Society of Nephrology: JASN, 2008, 19, 1025-1033.	6.1	30
18	Postpartum Septic Sacroiliitis Misdiagnosed as Sciatic Neuropathy. American Journal of the Medical Sciences, 2010, 339, 292-295.	1.1	26

#	Article	IF	CITATION
19	Identification of Genetic Loci Underlying the Phenotypic Constructs of Autism Spectrum Disorders. Journal of the American Academy of Child and Adolescent Psychiatry, 2011, 50, 687-696.e13.	0.5	26
20	The role of Self-Defined Race/Ethnicity in Population Structure Control. Annals of Human Genetics, 2006, 70, 496-505.	0.8	24
21	The Effects of Anemia and Blood Transfusion on Patients With Stage III-IV Ovarian Cancer. International Journal of Gynecological Cancer, 2013, 23, 1569-1576.	2.5	19
22	Genetic analysis of common factors underlying cardiovascular disease-related traits. BMC Genetics, 2003, 4, S56.	2.7	11
23	A genome scan for parent-of-origin linkage effects in alcoholism. BMC Genetics, 2005, 6, S160.	2.7	11
24	Measurement equivalence of the autism symptom phenotype in children and youth. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2013, 54, 1346-1355.	5.2	10
25	Identity-by-descent mapping for diastolic blood pressure in unrelated Mexican Americans. BMC Proceedings, 2016, 10, 263-267.	1.6	4