Simon L Girard

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/4353749/publications.pdf

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42 papers	2,780 citations	304743 22 h-index	276875 41 g-index
48 all docs	48 docs citations	48 times ranked	6429 citing authors

#	Article	IF	Citations
1	Association of ultraâ€rare coding variants with genetic generalized epilepsy: A caseâ€“control whole exome sequencing study. Epilepsia, 2022, 63, 723-735.	5.1	8
2	Association of Essential Tremor With Novel Risk Loci. JAMA Neurology, 2022, 79, 185.	9.0	17
3	The role of common genetic variation in presumed monogenic epilepsies. EBioMedicine, 2022, 81, 104098.	6.1	12
4	Assessment of burden and segregation profiles of <scp>CNVs</scp> in patients with epilepsy. Annals of Clinical and Translational Neurology, 2022, 9, 1050-1058.	3.7	2
5	Assessing the role of rare genetic variants in drugâ€resistant, nonâ€lesional focal epilepsy. Annals of Clinical and Translational Neurology, 2021, 8, 1376-1387.	3.7	16
6	Polygenic risk score for atopic dermatitis in the Canadian population. Journal of Allergy and Clinical Immunology, 2021, 147, 406-409.	2.9	12
7	Testing association of rare genetic variants with resistance to three common antiseizure medications. Epilepsia, 2020, 61, 657-666.	5.1	22
8	Polygenic risk scores of several subtypes of epilepsies in a founder population. Neurology: Genetics, 2020, 6, e416.	1.9	17
9	Genome-wide estimates of heritability and genetic correlations in essential tremor. Parkinsonism and Related Disorders, 2019, 64, 262-267.	2.2	10
10	Whole genome sequencing and variant discovery in the ASPIRE autism spectrum disorder cohort. Clinical Genetics, 2019, 96, 199-206.	2.0	18
11	Impact of Paternal Age at Conception on Human Health. Clinical Chemistry, 2019, 65, 146-152.	3.2	25
12	Investigating the association and causal relationship between restless legs syndrome and essential tremor. Parkinsonism and Related Disorders, 2019, 61, 238-240.	2.2	9
13	Exome sequencing of sporadic childhoodâ€onset schizophrenia suggests the contribution of Xâ€linked genes in males. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2019, 180, 335-340.	1.7	8
14	Human copy number variants are enriched in regions of low mappability. Nucleic Acids Research, 2018, 46, 7236-7249.	14.5	36
15	Global characterization of copy number variants in epilepsy patients from whole genome sequencing. PLoS Genetics, 2018, 14, e1007285.	3.5	50
16	Rare coding variants in genes encoding GABAA receptors in genetic generalised epilepsies: an exome-based case-control study. Lancet Neurology, The, 2018, 17, 699-708.	10.2	67
17	High Rate of Recurrent De Novo Mutations in Developmental and Epileptic Encephalopathies. American Journal of Human Genetics, 2017, 101, 664-685.	6.2	337
18	No rare deleterious variants from STK32B , PPARGC1A , and CTNNA3 are associated with essential tremor. Neurology: Genetics, 2017, 3, e195.	1.9	5

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19	Paternal Age Explains a Major Portion of De Novo Germline Mutation Rate Variability in Healthy Individuals. PLoS ONE, 2016, 11, e0164212.	2.5	41
20	The role of the melanoma gene MC1R in Parkinson disease and REM sleep behavior disorder. Neurobiology of Aging, 2016, 43, 180.e7-180.e13.	3.1	12
21	Genome-wide association study in essential tremor identifies three new loci. Brain, 2016, 139, 3163-3169.	7.6	78
22	De novo variants in sporadic cases of childhood onset schizophrenia. European Journal of Human Genetics, 2016, 24, 944-948.	2.8	77
23	Comparison of Sequencing Based CNV Discovery Methods Using Monozygotic Twin Quartets. PLoS ONE, 2015, 10, e0122287.	2.5	22
24	Mutation Burden of Rare Variants in Schizophrenia Candidate Genes. PLoS ONE, 2015, 10, e0128988.	2.5	17
25	Functional variants of POC5 identified in patients with idiopathic scoliosis. Journal of Clinical Investigation, 2015, 125, 1124-1128.	8.2	87
26	Cross-Disorder Genome-Wide Analyses Suggest a Complex Genetic Relationship Between Tourette's Syndrome and OCD. American Journal of Psychiatry, 2015, 172, 82-93.	7.2	117
27	Parkinson's Disease Genetic Loci in Rapid Eye Movement Sleep Behavior Disorder. Journal of Molecular Neuroscience, 2015, 56, 617-622.	2.3	42
28	Deleterious mutations in the essential mRNA metabolism factor, hGle1, in amyotrophic lateral sclerosis. Human Molecular Genetics, 2015, 24, 1363-1373.	2.9	122
29	Genome-wide association study in FTD: divide to conquer. Lancet Neurology, The, 2014, 13, 643-644.	10.2	4
30	Genome-wide association study of Tourette's syndrome. Molecular Psychiatry, 2013, 18, 721-728.	7.9	161
31	Investigation of rare variants in LRP1, KPNA1, ALS2CL and ZNF480 genes in schizophrenia patients reflects genetic heterogeneity of the disease. Behavioral and Brain Functions, 2013, 9, 9.	3.3	15
32	Whole-Exome Sequencing Reveals a Rapid Change in the Frequency of Rare Functional Variants in a Founding Population of Humans. PLoS Genetics, 2013, 9, e1003815.	3.5	70
33	Partitioning the Heritability of Tourette Syndrome and Obsessive Compulsive Disorder Reveals Differences in Genetic Architecture. PLoS Genetics, 2013, 9, e1003864.	3.5	241
34	Exome Sequencing Identifies FUS Mutations as a Cause of Essential Tremor. American Journal of Human Genetics, 2012, 91, 313-319.	6.2	176
35	Schizophrenia Genetics: Putting All the Pieces Together. Current Neurology and Neuroscience Reports, 2012, 12, 261-266.	4.2	37
36	Increased exonic de novo mutation rate in individuals with schizophrenia. Nature Genetics, 2011, 43, 860-863.	21.4	392

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37	Where are the missing pieces of the schizophrenia genetics puzzle?. Current Opinion in Genetics and Development, 2011, 21, 310-316.	3.3	22
38	How de novo mutation studies will change our view of the genetics of neurological and psychiatric disorders. Future Neurology, 2011, 6, 719-721.	0.5	0
39	Restless legs syndromeâ€associated MEIS1 risk variant influences iron homeostasis. Annals of Neurology, 2011, 70, 170-175.	5.3	81
40	Mutations in DCC Cause Congenital Mirror Movements. Science, 2010, 328, 592-592.	12.6	161
41	Autosomalâ€dominant locus for restless legs syndrome in Frenchâ€Canadians on chromosome 16p12.1. Movement Disorders, 2009, 24, 40-50.	3.9	44
42	MEIS1 intronic risk haplotype associated with restless legs syndrome affects its mRNA and protein expression levels. Human Molecular Genetics, 2009, 18, 1065-1074.	2.9	85