

Susan L Christian

List of Publications by Year in descending order

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Version: 2024-02-01

21
papers

3,449
citations

430874

18
h-index

713466

21
g-index

22
all docs

22
docs citations

22
times ranked

5156
citing authors

#	ARTICLE	IF	CITATIONS
1	The Influence of Microdeletions and Microduplications of 16p11.2 on Global Transcription Profiles. <i>Journal of Child Neurology</i> , 2015, 30, 1947-1953.	1.4	13
2	De Novo Mutations in the Beta-Tubulin Gene TUBB2A Cause Simplified Gyral Patterning and Infantile-Onset Epilepsy. <i>American Journal of Human Genetics</i> , 2014, 94, 634-641.	6.2	99
3	Copy number variants and infantile spasms: evidence for abnormalities in ventral forebrain development and pathways of synaptic function. <i>European Journal of Human Genetics</i> , 2011, 19, 1238-1245.	2.8	74
4	Microcephaly, sensorineural deafness and Currarino triad with duplication-deletion of distal 7q. <i>European Journal of Pediatrics</i> , 2010, 169, 475-481.	2.7	22
5	Duplication 16p11.2 in a child with infantile seizure disorder. <i>American Journal of Medical Genetics, Part A</i> , 2010, 152A, 1567-1574.	1.2	37
6	A de novo 1p34.2 microdeletion identifies the synaptic vesicle gene RIMS3 as a novel candidate for autism. <i>Journal of Medical Genetics</i> , 2010, 47, 81-90.	3.2	52
7	Association and Mutation Analyses of 16p11.2 Autism Candidate Genes. <i>PLoS ONE</i> , 2009, 4, e4582.	2.5	80
8	Genetics of autism spectrum disorders. <i>Current Neurology and Neuroscience Reports</i> , 2009, 9, 188-197.	4.2	125
9	Microduplications of 16p11.2 are associated with schizophrenia. <i>Nature Genetics</i> , 2009, 41, 1223-1227.	21.4	646
10	Copy number and sequence variants implicate <i>APBA2</i> as an autism candidate gene. <i>Autism Research</i> , 2009, 2, 359-364.	3.8	32
11	Targeted loss of Arx results in a developmental epilepsy mouse model and recapitulates the human phenotype in heterozygous females. <i>Brain</i> , 2009, 132, 1563-1576.	7.6	178
12	Mutations of CASK cause an X-linked brain malformation phenotype with microcephaly and hypoplasia of the brainstem and cerebellum. <i>Nature Genetics</i> , 2008, 40, 1065-1067.	21.4	252
13	Novel Submicroscopic Chromosomal Abnormalities Detected in Autism Spectrum Disorder. <i>Biological Psychiatry</i> , 2008, 63, 1111-1117.	1.3	268
14	Recurrent 16p11.2 microdeletions in autism. <i>Human Molecular Genetics</i> , 2007, 17, 628-638.	2.9	614
15	AUTISM AS A PARADIGMATIC COMPLEX GENETIC DISORDER. <i>Annual Review of Genomics and Human Genetics</i> , 2004, 5, 379-405.	6.2	237
16	Polymorphisms at the G72/G30 Gene Locus, on 13q33, Are Associated with Bipolar Disorder in Two Independent Pedigree Series*. <i>American Journal of Human Genetics</i> , 2003, 72, 1131-1140.	6.2	253
17	An Evaluation of the Assembly of an Approximately 15-Mb Region on Human Chromosome 13q32-q33 Linked to Bipolar Disorder and Schizophrenia. <i>Genomics</i> , 2002, 79, 635-658.	2.9	18
18	Fine mapping supports previous linkage evidence for a bipolar disorder susceptibility locus on 13q32. <i>American Journal of Medical Genetics Part A</i> , 2001, 105, 375-380.	2.4	33

#	ARTICLE	IF	CITATIONS
19	Somatic segregation errors predominantly contribute to the gain or loss of a paternal chromosome leading to uniparental disomy for chromosome 15. <i>Clinical Genetics</i> , 2000, 57, 349-358.	2.0	58
20	Methylation-specific PCR simplifies imprinting analysis. <i>Nature Genetics</i> , 1997, 16, 16-17.	21.4	265
21	Validation studies of SNRPN methylation as a diagnostic test for Prader-Willi syndrome. , 1996, 66, 77-80.		87