T Conrad Gilliam

List of Publications by Year
in descending order

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35 papers	11,389 citations	172457 29 h-index	361022 35 g-index
35 all docs	35 docs citations	35 times ranked	12254 citing authors

#	Article	IF	Citations
1	Novel Submicroscopic Chromosomal Abnormalities Detected in Autism Spectrum Disorder. Biological Psychiatry, 2008, 63, 1111-1117.	1.3	268
2	Genetic-linkage mapping of complex hereditary disorders to a whole-genome molecular-interaction network. Genome Research, 2008, 18, 1150-1162.	5.5	63
3	Recurrent 16p11.2 microdeletions in autism. Human Molecular Genetics, 2007, 17, 628-638.	2.9	614
4	Genomewide Scan for Linkage Reveals Evidence of Several Susceptibility Loci for Alopecia Areata. American Journal of Human Genetics, 2007, 80, 316-328.	6.2	132
5	Strong Association of De Novo Copy Number Mutations with Autism. Science, 2007, 316, 445-449.	12.6	2,497
6	DRD2 C957T polymorphism interacts with the COMT Val158Met polymorphism in human working memory ability. Schizophrenia Research, 2007, 90, 104-107.	2.0	71
7	Mapping autism risk loci using genetic linkage and chromosomal rearrangements. Nature Genetics, 2007, 39, 319-328.	21.4	1,272
8	Catechol-O-Methyltransferase (COMT) Genotypes and Working Memory: Associations with Differing Cognitive Operations. Biological Psychiatry, 2005, 58, 901-907.	1.3	237
9	Large-Scale Copy Number Polymorphism in the Human Genome. Science, 2004, 305, 525-528.	12.6	2,293
10	Evidence for Sex-Specific Risk Alleles in Autism Spectrum Disorder. American Journal of Human Genetics, 2004, 75, 1117-1123.	6.2	165
11	Population-based study of SR-BI genetic variation and lipid profile. Atherosclerosis, 2004, 175, 159-168.	0.8	69
12	Association between lipoprotein lipase (LPL) gene and blood lipids: A common variant for a common trait?. Genetic Epidemiology, 2003, 24, 309-321.	1.3	25
13	A Genomewide Screen of 345 Families for Autism-Susceptibility Loci. American Journal of Human Genetics, 2003, 73, 886-897.	6.2	247
14	Association of extreme blood lipid profile phenotypic variation with 11 reverse cholesterol transport genes and 10 non-genetic cardiovascular disease risk factors. Human Molecular Genetics, 2003, 12, 2733-2743.	2.9	34
15	Evidence for a Language Quantitative Trait Locus on Chromosome 7q in Multiplex Autism Families. American Journal of Human Genetics, 2002, 70, 60-71.	6.2	253
16	Genomewide Linkage Analysis of Celiac Disease in Finnish Families. American Journal of Human Genetics, 2002, 70, 51-59.	6.2	90
17	Four New Families with Autosomal Dominant Partial Epilepsy with Auditory Features: Clinical Description and Linkage to Chromosome 10q24. Epilepsia, 2002, 43, 60-67.	5.1	59
18	Mutations in LGI1 cause autosomal-dominant partial epilepsy with auditory features. Nature Genetics, 2002, 30, 335-341.	21.4	555

#	Article	IF	Citations
19	A follow-up linkage study supports evidence for a bipolar affective disorder locus on chromosome 21q22. American Journal of Medical Genetics Part A, 2001, 105, 189-194.	2.4	43
20	A Genomewide Screen for Autism Susceptibility Loci. American Journal of Human Genetics, 2001, 69, 327-340.	6.2	287
21	Null Mutation of the Murine ATP7B (Wilson Disease) Gene Results in Intracellular Copper Accumulation and Late-Onset Hepatic Nodular Transformation. Human Molecular Genetics, 1999, 8, 1665-1671.	2.9	186
22	The neuronal ceroid lipofuscinoses in human EPMR and mnd mutant mice are associated with mutations in CLN8. Nature Genetics, 1999, 23, 233-236.	21.4	277
23	A Comprehensive Linkage Analysis of Chromosome 21q22 Supports Prior Evidence for a Putative Bipolar Affective Disorder Locus. American Journal of Human Genetics, 1999, 64, 210-217.	6.2	104
24	Differential SMN2 expression associated with SMA severity. Nature Genetics, 1998, 20, 230-231.	21.4	304
25	TULP1 mutation in two extended Dominican kindreds with autosomal recessive Retinitis pigmentosa. Nature Genetics, 1998, 18, 177-179.	21.4	151
26	Arm tremor secondary to Wilson's disease. Movement Disorders, 1998, 13, 351-353.	3.9	11
27	No Evidence for Significant Linkage between Bipolar Affective Disorder and Chromosome 18 Pericentromeric Markers in a Large Series of Multiplex Extended Pedigrees. American Journal of Human Genetics, 1998, 62, 916-924.	6.2	36
28	High-Resolution Mapping and Transcript Identification at the Progressive Epilepsy with Mental Retardation Locus on Chromosomeâ€‰8p. Genome Research, 1997, 7, 887-896.	5.5	26
29	Identification and Analysis of Mutations in the Wilson Disease Gene (ATP7B): Population Frequencies, Genotype-Phenotype Correlation, and Functional Analyses. American Journal of Human Genetics, 1997, 61, 317-328.	6.2	346
30	Extensive DNA deletion associated with severe disease alleles on spinal muscular atrophy homologues. Annals of Neurology, 1997, 42, 41-49.	5.3	17
31	Is the Spinal Muscular Atrophy gene found?. Nature Medicine, 1995, 1, 124-127.	30.7	7
32	A possible vulnerability locus for bipolar affective disorder on chromosome 21q22.3. Nature Genetics, 1994, 8, 291-296.	21.4	297
33	Genetic disorders of copper metabolism. Current Opinion in Pediatrics, 1994, 6, 698-701.	2.0	9
34	Diminished support for linkage between manic depressive illness and Xâ€“chromosome markers in three Israeli pedigrees. Nature Genetics, 1993, 3, 49-55.	21.4	158
35	Linkage of Tunisian autosomal recessive Duchenneâ€“like muscular dystrophy to the pericentromeric region of chromosome 13q. Nature Genetics, 1992, 2, 315-317.	21.4	186