# T Conrad Gilliam

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/11738073/publications.pdf

Version: 2024-02-01

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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 | Strong Association of De Novo Copy Number Mutations with Autism. Science, 2007, 316, 445-449. | 12.6 | 2,497 |
| 2 | Large-Scale Copy Number Polymorphism in the Human Genome. Science, 2004, 305, 525-528. | 12.6 | 2,293 |
| 3 | Mapping autism risk loci using genetic linkage and chromosomal rearrangements. Nature Genetics, 2007, 39, 319-328. | 21.4 | 1,272 |
| 4 | Recurrent 16p11.2 microdeletions in autism. Human Molecular Genetics, 2007, 17, 628-638. | 2.9 | 614 |
| 5 | Mutations in LGI1 cause autosomal-dominant partial epilepsy with auditory features. Nature Genetics, 2002, 30, 335-341. | 21.4 | 555 |
| 6 | 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 |
| 7 | Differential SMN2 expression associated with SMA severity. Nature Genetics, 1998, 20, 230-231. | 21.4 | 304 |
| 8 | A possible vulnerability locus for bipolar affective disorder on chromosome 21q22.3. Nature Genetics, 1994, 8, 291-296. | 21.4 | 297 |
| 9 | A Genomewide Screen for Autism Susceptibility Loci. American Journal of Human Genetics, 2001, 69, 327-340. | 6.2 | 287 |
| 10 | 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 |
| 11 | Novel Submicroscopic Chromosomal Abnormalities Detected in Autism Spectrum Disorder. Biological Psychiatry, 2008, 63, 1111-1117. | 1.3 | 268 |
| 12 | 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 |
| 13 | A Genomewide Screen of 345 Families for Autism-Susceptibility Loci. American Journal of Human Genetics, 2003, 73, 886-897. | 6.2 | 247 |
| 14 | Catechol-O-Methyltransferase (COMT) Genotypes and Working Memory: Associations with Differing Cognitive Operations. Biological Psychiatry, 2005, 58, 901-907. | 1.3 | 237 |
| 15 | 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 |
| 16 | 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 |
| 17 | Evidence for Sex-Specific Risk Alleles in Autism Spectrum Disorder. American Journal of Human Genetics, 2004, 75, 1117-1123. | 6.2 | 165 |
| 18 | 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 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | TULP1 mutation in two extended Dominican kindreds with autosomal recessive Retinitis pigmentosa. Nature Genetics, 1998, 18, 177-179. | 21.4 | 151 |
| 20 | 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 |
| 21 | 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 |
| 22 | Genomewide Linkage Analysis of Celiac Disease in Finnish Families. American Journal of Human Genetics, 2002, 70, 51-59. | 6.2 | 90 |
| 23 | DRD2 C957T polymorphism interacts with the COMT Val158Met polymorphism in human working memory ability. Schizophrenia Research, 2007, 90, 104-107. | 2.0 | 71 |
| 24 | Population-based study of SR-BI genetic variation and lipid profile. Atherosclerosis, 2004, 175, 159-168. | 0.8 | 69 |
| 25 | Genetic-linkage mapping of complex hereditary disorders to a whole-genome molecular-interaction network. Genome Research, 2008, 18, 1150-1162. | 5.5 | 63 |
| 26 | 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 |
| 27 | 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 |
| 28 | 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 |
| 29 | 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 |
| 30 | 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 |
| 31 | 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 |
| 32 | Extensive DNA deletion associated with severe disease alleles on spinal muscular atrophy homologues. Annals of Neurology, 1997, 42, 41-49. | 5.3 | 17 |
| 33 | Arm tremor secondary to Wilson's disease. Movement Disorders, 1998, 13, 351-353. | 3.9 | 11 |
| 34 | Genetic disorders of copper metabolism. Current Opinion in Pediatrics, 1994, 6, 698-701. | 2.0 | 9 |
| 35 | Is the Spinal Muscular Atrophy gene found?. Nature Medicine, 1995, 1, 124-127. | 30.7 | 7 |