# Claire S Leblond

List of Publications by Year
in descending order



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|---|---|---|---|
| 14 papers | 1,300 citations | 840776 11 h-index | 996975 15 g-index |
| 16 all docs | 16 docs citations | 16 times ranked | 2333 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Genetic correlates of phenotypic heterogeneity in autism. Nature Genetics, 2022, 54, 1293-1304. | 21.4 | 51 |
| 2 | Decreased phenol sulfotransferase activities associated with hyperserotonemia in autism spectrum disorders. Translational Psychiatry, 2021, 11, 23. | 4.8 | 11 |
| 3 | Operative list of genes associated with autism and neurodevelopmental disorders based on database review. Molecular and Cellular Neurosciences, 2021, 113, 103623. | 2.2 | 51 |
| 4 | Oligogenicity, C9orf72 expansion, and variant severity in ALS. Neurogenetics, 2020, 21, 227-242. | 1.4 | 13 |
| 5 | Social and non-social autism symptoms and trait domains are genetically dissociable. Communications Biology, 2019, 2, 328. | 4.4 | 57 |
| 6 | Both rare and common genetic variants contribute to autism in the Faroe Islands. Npj Genomic Medicine, 2019, 4, 1. | 3.8 | 72 |
| 7 | Somatic expansion of the C9orf72 hexanucleotide repeat does not occur in ALS spinal cord tissues. Neurology: Genetics, 2019, 5, e317. | 1.9 | 8 |
| 8 | Altered spinogenesis in iPSC-derived cortical neurons from patients with autism carrying de novo SHANK3 mutations. Scientific Reports, 2019, 9, 94. | 3.3 | 51 |
| 9 | Homozygous 2p11.2 deletion supports the implication of ELMOD3 in hearing loss and reveals the potential association of CAPG with ASD/ID etiology. Journal of Applied Genetics, 2019, 60, 49-56. | 1.9 | 11 |
| 10 | Heritability of the melatonin synthesis variability in autism spectrum disorders. Scientific Reports, 2017, 7, 17746. | 3.3 | 28 |
| 11 | A framework to identify contributing genes in patients with Phelan-McDermid syndrome. Npj Genomic Medicine, 2017, 2, 32. | 3.8 | 58 |
| 12 | 11q24.2â€25 microâ€rearrangements in autism spectrum disorders: Relation to brain structures. American Journal of Medical Genetics, Part A, 2015, 167, 3019-3030. | 1.2 | 25 |
| 13 | Meta-analysis of SHANK Mutations in Autism Spectrum Disorders: A Gradient of Severity in Cognitive Impairments. PLoS Genetics, 2014, 10, e1004580. | 3.5 | 501 |
| 14 | Genetic and Functional Analyses of SHANK2 Mutations Suggest a Multiple Hit Model of Autism Spectrum Disorders. PLoS Genetics, 2012, 8, e1002521. | 3.5 | 358 |