

Carla Lintas

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7760061/publications.pdf>

Version: 2024-02-01

36
papers

3,179
citations

393982

19
h-index

360668

35
g-index

36
all docs

36
docs citations

36
times ranked

5967
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|---|------|-----------|
| 1 | Melanoma Cell Resistance to Vemurafenib Modifies Inter-Cellular Communication Signals. <i>Biomedicines</i> , 2021, 9, 79. | 1.4 | 10 |
| 2 | Molecular biomarkers to track clinical improvement following an integrative treatment model in autistic toddlers. <i>Acta Neuropsychiatrica</i> , 2021, 33, 267-272. | 1.0 | 2 |
| 3 | Reevaluation of Serum Arylesterase Activity in Neurodevelopmental Disorders. <i>Antioxidants</i> , 2021, 10, 164. | 2.2 | 5 |
| 4 | Genotype-Phenotype Correlations in Relation to Newly Emerging Monogenic Forms of Autism Spectrum Disorder and Associated Neurodevelopmental Disorders: The Importance of Phenotype Reevaluation after Pangenomic Results. <i>Journal of Clinical Medicine</i> , 2021, 10, 5060. | 1.0 | 4 |
| 5 | Phenotypic spectrum of <i>NRXN1</i> mono- and bi-allelic deficiency: A systematic review. <i>Clinical Genetics</i> , 2020, 97, 125-137. | 1.0 | 38 |
| 6 | Huntingtin gene CAG repeat size affects autism risk: Family-based and case-control association study. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2020, 183, 341-351. | 1.1 | 5 |
| 7 | FAR1 deletion is associated with lack of response to autism treatment by early start denver model in a multiplex family. <i>Molecular Genetics & Genomic Medicine</i> , 2020, 8, e1373. | 0.6 | 10 |
| 8 | Genetic and epigenetic <i>MTHFR</i> gene variants in the mothers of attention-deficit/hyperactivity disorder affected children as possible risk factors for neurodevelopmental disorders. <i>Epigenomics</i> , 2020, 12, 813-823. | 1.0 | 2 |
| 9 | Appropriateness of array-CGH in the ADHD clinics: A comparative study. <i>Genes, Brain and Behavior</i> , 2020, 19, e12651. | 1.1 | 4 |
| 10 | Large-Scale Exome Sequencing Study Implicates Both Developmental and Functional Changes in the Neurobiology of Autism. <i>Cell</i> , 2020, 180, 568-584.e23. | 13.5 | 1,422 |
| 11 | Linking genetics to epigenetics: The role of folate and folate-related pathways in neurodevelopmental disorders. <i>Clinical Genetics</i> , 2019, 95, 241-252. | 1.0 | 32 |
| 12 | Theophylline induces differentiation and modulates cytoskeleton dynamics and cytokines secretion in human melanoma-initiating cells. <i>Life Sciences</i> , 2019, 230, 121-131. | 2.0 | 14 |
| 13 | Evidence that ITGB3 promoter variants increase serotonin blood levels by regulating platelet serotonin transporter trafficking. <i>Human Molecular Genetics</i> , 2019, 28, 1153-1161. | 1.4 | 10 |
| 14 | Unraveling molecular pathways shared by Kabuki and Kabuki-like syndromes. <i>Clinical Genetics</i> , 2018, 94, 283-295. | 1.0 | 32 |
| 15 | An Interstitial 17q11.2 de novo Deletion Involving the CDK5R1 Gene in a High-Functioning Autistic Patient. <i>Molecular Syndromology</i> , 2018, 9, 247-252. | 0.3 | 2 |
| 16 | Copy number variation in 19 Italian multiplex families with autism spectrum disorder: Importance of synaptic and neurite elongation genes. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2017, 174, 547-556. | 1.1 | 7 |
| 17 | Recurrent 15q11.2 BP1-BP2 microdeletions and microduplications in the etiology of neurodevelopmental disorders. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2016, 171, 1088-1098. | 1.1 | 41 |
| 18 | Differential methylation at the RELN gene promoter in temporal cortex from autistic and typically developing post-puberal subjects. <i>Journal of Neurodevelopmental Disorders</i> , 2016, 8, 18. | 1.5 | 35 |

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 19 | Autism genetics: Methodological issues and experimental design. <i>Science China Life Sciences</i> , 2015, 58, 946-957. | 2.3 | 1 |
| 20 | Xp22.33p22.12 Duplication in a Patient with Intellectual Disability and Dysmorphic Facial Features. <i>Molecular Syndromology</i> , 2015, 6, 236-241. | 0.3 | 10 |
| 21 | Age-Dependent Decrease and Alternative Splicing of Methionine Synthase mRNA in Human Cerebral Cortex and an Accelerated Decrease in Autism. <i>PLoS ONE</i> , 2013, 8, e56927. | 1.1 | 54 |
| 22 | Genome-wide expression studies in Autism spectrum disorder, Rett syndrome, and Down syndrome. <i>Neurobiology of Disease</i> , 2012, 45, 57-68. | 2.1 | 81 |
| 23 | Lack of Infection with XMRV or Other MLV-Related Viruses in Blood, Post-Mortem Brains and Paternal Gametes of Autistic Individuals. <i>PLoS ONE</i> , 2011, 6, e16609. | 1.1 | 16 |
| 24 | Neocortical RELN promoter methylation increases significantly after puberty. <i>NeuroReport</i> , 2010, 21, 114-118. | 0.6 | 40 |
| 25 | Association of autism with polyomavirus infection in postmortem brains. <i>Journal of NeuroVirology</i> , 2010, 16, 141-149. | 1.0 | 42 |
| 26 | Decreased serum arylesterase activity in autism spectrum disorders. <i>Psychiatry Research</i> , 2010, 180, 105-113. | 1.7 | 33 |
| 27 | Genomic and epigenetic evidence for oxytocin receptor deficiency in autism. <i>BMC Medicine</i> , 2009, 7, 62. | 2.3 | 497 |
| 28 | Involvement of the PRKCB1 gene in autistic disorder: significant genetic association and reduced neocortical gene expression. <i>Molecular Psychiatry</i> , 2009, 14, 705-718. | 4.1 | 75 |
| 29 | An 8-year-old boy with autoimmune hepatitis and Candida onychosis as the first symptoms of autoimmune polyglandular syndrome (APS1): identification of a new homozygous mutation in the autoimmune regulator gene (aire). <i>European Journal of Pediatrics</i> , 2008, 167, 949-953. | 1.3 | 20 |
| 30 | Immune transcriptome alterations in the temporal cortex of subjects with autism. <i>Neurobiology of Disease</i> , 2008, 30, 303-311. | 2.1 | 344 |
| 31 | Autistic phenotypes and genetic testing: state-of-the-art for the clinical geneticist. <i>Journal of Medical Genetics</i> , 2008, 46, 1-8. | 1.5 | 146 |
| 32 | Reelin Gene Polymorphisms in Autistic Disorder. , 2008, , 385-399. | | 1 |
| 33 | Do mutations of RAG genes have a role in human autoimmunity? The Notarangelo's hypothesis revisited. <i>Diabetes/Metabolism Research and Reviews</i> , 2006, 22, 108-110. | 1.7 | 1 |
| 34 | Methylation profile in tumor and sputum samples of lung cancer patients detected by spiral computed tomography: A nested case-control study. <i>International Journal of Cancer</i> , 2006, 118, 1248-1253. | 2.3 | 49 |
| 35 | Mutations that affect the ability of the vnd/NK-2 homeoprotein to regulate gene expression: Transgenic alterations and tertiary structure. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003, 100, 3119-3124. | 3.3 | 26 |
| 36 | SPATIAL VARIATION IN THE FAUNA ASSOCIATED WITH MYTILUS EDULIS ON A WAVE-EXPOSED ROCKY SHORE. <i>Journal of Molluscan Studies</i> , 1994, 60, 165-174. | 0.4 | 68 |