# Aparna Prasad

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/4985468/publications.pdf

Version: 2024-02-01

| | | 567281 | 839539 |
|---|---|---|---|
| 18 | 3,798 | 15 | 18 |
| papers | citations | h-index | g-index |

| | | | |
|---|---|---|---|
| 18 | 18 | 18 | 7625 |
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | The Temple Grandin Genome: Comprehensive Analysis in a Scientist with High-Functioning Autism. Journal of Personalized Medicine, 2021, 11, 21. | 2.5 | 5 |
| 2 | Critical exon indexing improves clinical interpretation of copy number variants in neurodevelopmental disorders. Neurology: Genetics, 2019, 5, e378. | 1.9 | 4 |
| 3 | Clinical utility of exome sequencing in individuals with large homozygous regions detected by chromosomal microarray analysis. BMC Medical Genetics, 2018, 19, 46. | 2.1 | 18 |
| 4 | Identification of Gene Mutations and Fusion Genes in Patients with SÃ©zary Syndrome. Journal of Investigative Dermatology, 2016, 136, 1490-1499. | 0.7 | 77 |
| 5 | Rare exonic deletions implicate the synaptic organizer Gephyrin (GPHN) in risk for autism, schizophrenia and seizures. Human Molecular Genetics, 2013, 22, 2055-2066. | 2.9 | 139 |
| 6 | A Discovery Resource of Rare Copy Number Variations in Individuals with Autism Spectrum Disorder. G3: Genes, Genomes, Genetics, 2012, 2, 1665-1685. | 1.8 | 175 |
| 7 | Rare Deletions at the Neurexin 3 Locus in Autism Spectrum Disorder. American Journal of Human Genetics, 2012, 90, 133-141. | 6.2 | 182 |
| 8 | SHANK1 Deletions in Males with Autism Spectrum Disorder. American Journal of Human Genetics, 2012, 90, 879-887. | 6.2 | 292 |
| 9 | Comprehensive assessment of array-based platforms and calling algorithms for detection of copy number variants. Nature Biotechnology, 2011, 29, 512-520. | 17.5 | 384 |
| 10 | Variant detection and the Autism sequencing project. BMC Bioinformatics, 2011, 12, . | 2.6 | 1 |
| 11 | A 2cM genome-wide scan of European Holstein cattle affected by classical BSE. BMC Genetics, 2010, 11, 20. | 2.7 | 22 |
| 12 | Functional impact of global rare copy number variation in autism spectrum disorders. Nature, 2010, 466, 368-372. | 27.8 | 1,803 |
| 13 | A genome-wide scan for common alleles affecting risk for autism. Human Molecular Genetics, 2010, 19, 4072-4082. | 2.9 | 538 |
| 14 | A first generation whole genome RH map of the river buffalo with comparison to domestic cattle. BMC Genomics, 2008, 9, 631. | 2.8 | 78 |
| 15 | High resolution radiation hybrid maps of bovine chromosomes 19 and 29: comparison with the bovine genome sequence assembly. BMC Genomics, 2007, 8, 310. | 2.8 | 16 |
| 16 | A second generation radiation hybrid map to aid the assembly of the bovine genome sequence. BMC Genomics, 2006, 7, 283. | 2.8 | 26 |
| 17 | Fate of SRY, PABY, DYS1, DYZ3 and DYZ1 loci in Indian patients harbouring sex chromosomal anomalies. Molecular Human Reproduction, 2005, 11, 117-127. | 2.8 | 19 |
| 18 | Organizational Variation of DYZ1 Repeat Sequences on the Human Y Chromosome and Its Diagnostic Potentials. DNA and Cell Biology, 2004, 23, 561-571. | 1.9 | 19 |