# Uppala Radhakrishna

List of Publications by Year
in descending order



| 16 | 1,932 | 567281 15 | 996975 15 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 16 | 16 | 16 | 1969 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Duplications ofBHLHA9are associated with ectrodactyly and tibia hemimelia inherited in non-Mendelian fashion. Journal of Medical Genetics, 2012, 49, 119-125. | 3.2 | 81 |
| 2 | Genomewide Linkage Scan for Splitâ€“Hand/Foot Malformation with Long-Bone Deficiency in a Large Arab Family Identifies Two Novel Susceptibility Loci on Chromosomes 1q42.2-q43 and 6q14.1. American Journal of Human Genetics, 2007, 80, 105-111. | 6.2 | 30 |
| 3 | Autosomal Dominant Nonsyndromic Cleft Lip and Palate: Significant Evidence of Linkage at 18q21.1. American Journal of Human Genetics, 2007, 81, 180-188. | 6.2 | 17 |
| 4 | Molecular and Clinical Characteristics in 46 Families Affected with Peutzâ€“Jeghers Syndrome. Digestive Diseases and Sciences, 2007, 52, 1924-1933. | 2.3 | 56 |
| 5 | Genomewide Scan for Nonsyndromic Cleft Lip and Palate in Multigenerational Indian Families Reveals Significant Evidence of Linkage at 13q33.1-34. American Journal of Human Genetics, 2006, 79, 580-585. | 6.2 | 29 |
| 6 | Split-hand/split-foot malformation 3 (SHFM3) at 10q24, development of rapid diagnostic methods and gene expression from the region. American Journal of Medical Genetics, Part A, 2006, 140A, 1384-1395. | 1.2 | 33 |
| 7 | A genomic rearrangement resulting in a tandem duplication is associated with split hand-split foot malformation 3 (SHFM3) at 10q24. Human Molecular Genetics, 2003, 12, 1959-1971. | 2.9 | 88 |
| 8 | Insertion of Î²-satellite repeats identifies a transmembrane protease causing both congenital and childhood onset autosomal recessive deafness. Nature Genetics, 2001, 27, 59-63. | 21.4 | 208 |
| 9 | Novel mutations of TMPRSS3 in four DFNB8/B10 families segregating congenital autosomal recessive deafness. Journal of Medical Genetics, 2001, 38, 396-400. | 3.2 | 55 |
| 10 | Mutations in GJB6 cause hidrotic ectodermal dysplasia. Nature Genetics, 2000, 26, 142-144. | 21.4 | 270 |
| 11 | Clouston hidrotic ectodermal dysplasia (HED): genetic homogeneity, presence of a founder effect in the French Canadian population and fine genetic mapping. European Journal of Human Genetics, 2000, 8, 372-380. | 2.8 | 43 |
| 12 | Lack of linkage or association between schizophrenia and the polymorphic trinucleotide repeat within the KCNN3 gene on chromosome 1q21. , 1999, 88, 348-351. | | 33 |
| 13 | The Phenotypic Spectrum of GLI3 Morphopathies Includes Autosomal Dominant Preaxial Polydactyly Type-IV and Postaxial Polydactyly Type-A/B; No Phenotype Prediction from the Position of GLI3 Mutations. American Journal of Human Genetics, 1999, 65, 645-655. | 6.2 | 175 |
| 14 | Schizophrenia susceptibility loci on chromosomes 13q32 and 8p21. Nature Genetics, 1998, 20, 70-73. | 21.4 | 506 |
| 15 | Peutz-Jeghers Syndrome: Confirmation of Linkage to Chromosome 19p13.3 and Identification of a Potential Second Locus, on 19q13.4. American Journal of Human Genetics, 1997, 61, 1327-1334. | 6.2 | 145 |
| 16 | Mutation in GLI3 in postaxial polydactyly type A. Nature Genetics, 1997, 17, 269-271. | 21.4 | 163 |