

# Deepti Anand

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8004749/publications.pdf>

Version: 2024-02-01

35  
papers

1,179  
citations

361413

20  
h-index

414414

32  
g-index

37  
all docs

37  
docs citations

37  
times ranked

1460  
citing authors

| #  | ARTICLE  | IF   | CITATIONS |
|----|--|------|-----------|
| 1  | Variant analyses of candidate genes in orofacial clefts in multiethnic populations. <i>Oral Diseases</i> , 2022, 28, 1921-1935.  | 3.0  | 3         |
| 2  | Whole-genome sequencing reveals de-novo mutations associated with nonsyndromic cleft lip/palate. <i>Scientific Reports</i> , 2022, 12, .   | 3.3  | 11        |
| 3  | Genome-Wide Analysis of Differentially Expressed miRNAs and Their Associated Regulatory Networks in Lenses Deficient for the Congenital Cataract-Linked Tudor Domain Containing Protein TDRD7. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 615761. | 3.7  | 12        |
| 4  | A large multiethnic GWAS meta-analysis of cataract identifies new risk loci and sex-specific effects. <i>Nature Communications</i> , 2021, 12, 3595.   | 12.8 | 39        |
| 5  | MS/MS in silico subtraction-based proteomic profiling as an approach to facilitate disease gene discovery: application to lens development and cataract. <i>Human Genetics</i> , 2020, 139, 151-184.   | 3.8  | 12        |
| 6  | Six2 regulates Pax9 expression, palatogenesis and craniofacial bone formation. <i>Developmental Biology</i> , 2020, 458, 246-256.  | 2.0  | 13        |
| 7  | The Tudor-domain protein TDRD7, mutated in congenital cataract, controls the heat shock protein HSPB1 (HSP27) and lens fiber cell morphology. <i>Human Molecular Genetics</i> , 2020, 29, 2076-2097.   | 2.9  | 27        |
| 8  | High-throughput transcriptome analysis reveals that the loss of Pten activates a novel NKX6-1/RASGRP1 regulatory module to rescue microphthalmia caused by Fgfr2-deficient lenses. <i>Human Genetics</i> , 2019, 138, 1391-1407.                                     | 3.8  | 14        |
| 9  | Missense Pathogenic variants in KIF4A Affect Dental Morphogenesis Resulting in X-linked Taurodontism, Microdontia and Dens-Invaginatus. <i>Frontiers in Genetics</i> , 2019, 10, 800.  | 2.3  | 7         |
| 10 | Molecular characterization of the human lens epithelium-derived cell line SRA01/04. <i>Experimental Eye Research</i> , 2019, 188, 107787.  | 2.6  | 14        |
| 11 | Mutations in GDF11 and the extracellular antagonist, Follistatin, as a likely cause of Mendelian forms of orofacial clefting in humans. <i>Human Mutation</i> , 2019, 40, 1813-1825.   | 2.5  | 26        |
| 12 | A systematic genetic analysis and visualization of phenotypic heterogeneity among orofacial cleft GWAS signals. <i>Genetic Epidemiology</i> , 2019, 43, 704-716.   | 1.3  | 36        |
| 13 | Pathogenic variants in <i>PLOD3</i> result in a Stickler syndrome-like connective tissue disorder with vascular complications. <i>Journal of Medical Genetics</i> , 2019, 56, 629-638.   | 3.2  | 23        |
| 14 | Genomic analyses in African populations identify novel risk loci for cleft palate. <i>Human Molecular Genetics</i> , 2019, 28, 1038-1051.  | 2.9  | 61        |
| 15 | Identification of OAF and PVRL1 as candidate genes for an ocular anomaly characterized by Peters anomaly type 2 and ectopia lentis. <i>Experimental Eye Research</i> , 2018, 168, 161-170.   | 2.6  | 5         |
| 16 | Mutation update of transcription factor genes <i>FOXE3</i> , <i>HSF4</i> , <i>MAF</i> , and <i>PITX3</i> causing cataracts and other developmental ocular defects. <i>Human Mutation</i> , 2018, 39, 471-494.  | 2.5  | 60        |
| 17 | iSyTE 2.0: a database for expression-based gene discovery in the eye. <i>Nucleic Acids Research</i> , 2018, 46, D875-D885.   | 14.5 | 71        |
| 18 | RNA sequencing-based transcriptomic profiles of embryonic lens development for cataract gene discovery. <i>Human Genetics</i> , 2018, 137, 941-954.  | 3.8  | 29        |

| #  | ARTICLE  | IF   | CITATIONS |
|----|--|------|-----------|
| 19 | Mutations in the Epithelial Cadherin-p120-Catenin Complex Cause Mendelian Non-Syndromic Cleft Lip with or without Cleft Palate. <i>American Journal of Human Genetics</i> , 2018, 102, 1143-1157.                                    | 6.2  | 94        |
| 20 | The RNA-binding protein Celf1 post-transcriptionally regulates p27Kip1 and Dnase2b to control fiber cell nuclear degradation in lens development. <i>PLoS Genetics</i> , 2018, 14, e1007278.   | 3.5  | 43        |
| 21 | Systems biology of lens development: A paradigm for disease gene discovery in the eye. <i>Experimental Eye Research</i> , 2017, 156, 22-33.  | 2.6  | 39        |
| 22 | Exome sequencing provides additional evidence for the involvement of <i>ARHGAP29</i> in Mendelian orofacial clefting and extends the phenotypic spectrum to isolated cleft palate. <i>Birth Defects Research</i> , 2017, 109, 27-37. | 1.5  | 49        |
| 23 | N-myc regulates growth and fiber cell differentiation in lens development. <i>Developmental Biology</i> , 2017, 429, 105-117.  | 2.0  | 37        |
| 24 | Novel phenotypes and loci identified through clinical genomics approaches to pediatric cataract. <i>Human Genetics</i> , 2017, 136, 205-225.   | 3.8  | 73        |
| 25 | Î²1-Integrin Deletion From the Lens Activates Cellular Stress Responses Leading to Apoptosis and Fibrosis. , 2017, 58, 3896.   |      | 19        |
| 26 | Prox1 and fibroblast growth factor receptors form a novel regulatory loop controlling lens fiber differentiation and gene expression. <i>Development (Cambridge)</i> , 2015, 143, 318-28.  | 2.5  | 59        |
| 27 | Molecular characterization of mouse lens epithelial cell lines and their suitability to study RNA granules and cataract associated genes. <i>Experimental Eye Research</i> , 2015, 131, 42-55.                                       | 2.6  | 29        |
| 28 | Compound mouse mutants of bZIP transcription factors Mafg and Mafk reveal a regulatory network of non-crystallin genes associated with cataract. <i>Human Genetics</i> , 2015, 134, 717-735.   | 3.8  | 47        |
| 29 | An integrative approach to analyze microarray datasets for prioritization of genes relevant to lens biology and disease. <i>Genomics Data</i> , 2015, 5, 223-227.  | 1.3  | 27        |
| 30 | Novel InDel variation in GS3 locus and development of InDel based marker for marker assisted breeding of short grain aromatic rices. <i>Journal of Plant Biochemistry and Biotechnology</i> , 2015, 24, 120-127.                     | 1.7  | 8         |
| 31 | Validation of gene based marker-QTL association for grain dimension traits in rice. <i>Journal of Plant Biochemistry and Biotechnology</i> , 2013, 22, 467-473.  | 1.7  | 12        |
| 32 | Analysis of molecular diversity and fingerprinting of commercially grown Indian rice hybrids. <i>Journal of Plant Biochemistry and Biotechnology</i> , 2012, 21, 173-179.  | 1.7  | 11        |
| 33 | Adsorption and Desorption of Chlorinated Compounds from Pristine and Thermally Treated Multiwalled Carbon Nanotubes. <i>Journal of Physical Chemistry C</i> , 2011, 115, 4552-4557.  | 3.1  | 35        |
| 34 | Carbon nanotube-textured sand for controlling bioavailability of contaminated sediments. <i>Nano Research</i> , 2010, 3, 412-422.  | 10.4 | 11        |
| 35 | Marker-assisted improvement of bacterial blight resistance in parental lines of Pusa RH10, a superfine grain aromatic rice hybrid. <i>Molecular Breeding</i> , 2010, 26, 293-305.  | 2.1  | 122       |