

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	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
2	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
3	Novel phenotypes and loci identified through clinical genomics approaches to pediatric cataract. <i>Human Genetics</i> , 2017, 136, 205-225.	3.8	73
4	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
5	Genomic analyses in African populations identify novel risk loci for cleft palate. <i>Human Molecular Genetics</i> , 2019, 28, 1038-1051.	2.9	61
6	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
7	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
8	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
9	Compound mouse mutants of bZIP transcription factors <i>Mafg</i> and <i>Mafk</i> reveal a regulatory network of non-crystallin genes associated with cataract. <i>Human Genetics</i> , 2015, 134, 717-735.	3.8	47
10	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
11	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
12	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
13	N-myc regulates growth and fiber cell differentiation in lens development. <i>Developmental Biology</i> , 2017, 429, 105-117.	2.0	37
14	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
15	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
16	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
17	RNA sequencing-based transcriptomic profiles of embryonic lens development for cataract gene discovery. <i>Human Genetics</i> , 2018, 137, 941-954.	3.8	29
18	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

#	ARTICLE	IF	CITATIONS
19	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
20	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
21	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
22	Î²1-Integrin Deletion From the Lens Activates Cellular Stress Responses Leading to Apoptosis and Fibrosis. , 2017, 58, 3896.		19
23	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
24	Molecular characterization of the human lens epithelium-derived cell line SRA01/04. <i>Experimental Eye Research</i> , 2019, 188, 107787.	2.6	14
25	Six2 regulates Pax9 expression, palatogenesis and craniofacial bone formation. <i>Developmental Biology</i> , 2020, 458, 246-256.	2.0	13
26	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
27	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
28	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
29	Carbon nanotube-textured sand for controlling bioavailability of contaminated sediments. <i>Nano Research</i> , 2010, 3, 412-422.	10.4	11
30	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
31	Whole-genome sequencing reveals de-novo mutations associated with nonsyndromic cleft lip/palate. <i>Scientific Reports</i> , 2022, 12, .	3.3	11
32	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
33	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
34	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
35	Variant analyses of candidate genes in orofacial clefts in multi-ethnic populations. <i>Oral Diseases</i> , 2022, 28, 1921-1935.	3.0	3