

Darek A Kedra

List of Publications by Year in descending order

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Version: 2024-02-01

25
papers

2,570
citations

361045
20
h-index

580395
25
g-index

25
all docs

25
docs citations

25
times ranked

3265
citing authors

#	ARTICLE	IF	CITATIONS
1	The DNA sequence of human chromosome 22. <i>Nature</i> , 1999, 402, 489-495.	13.7	1,086
2	Deregulation of the platelet-derived growth factor β -chain gene via fusion with collagen gene COL1A1 in dermatofibrosarcoma protuberans and giant-cell fibroblastoma. <i>Nature Genetics</i> , 1997, 15, 95-98.	9.4	510
3	Genome and transcriptome analysis of the Mesoamerican common bean and the role of gene duplications in establishing tissue and temporal specialization of genes. <i>Genome Biology</i> , 2016, 17, 32.	3.8	166
4	Third Report on Chicken Genes and Chromosomes 2015. <i>Cytogenetic and Genome Research</i> , 2015, 145, 78-179.	0.6	97
5	Haplotype selection as an adaptive mechanism in the protozoan pathogen <i>Leishmania donovani</i> . <i>Nature Ecology and Evolution</i> , 2017, 1, 1961-1969.	3.4	95
6	Role of L-arginine, a substrate for nitric oxide-synthase, in gastroprotection and ulcer healing. <i>Journal of Gastroenterology</i> , 1997, 32, 442-452.	2.3	63
7	Characterization of a second human clathrin heavy chain polypeptide gene (CLH-22) from chromosome 22q11. <i>Human Molecular Genetics</i> , 1996, 5, 625-631.	1.4	54
8	Characterization of the human synaptogyrin gene family. <i>Human Genetics</i> , 1998, 103, 131-141.	1.8	54
9	Protein identification and quantification by two-dimensional infrared spectroscopy: Implications for an all-optical proteomic platform. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 15352-15357.	3.3	50
10	Characterization of the human NIPSNAP1 gene from 22q12: a member of a novel gene family. <i>Gene</i> , 1998, 212, 13-20.	1.0	48
11	Mutations in short stature homeobox containing gene (SHOX) in dyschondrosteosis but not in hypochondroplasia. <i>Human Genetics</i> , 2000, 107, 145-149.	1.8	37
12	The LZTFL1 Gene Is a Part of a Transcriptional Map Covering 250 kb within the Common Eliminated Region 1 (C3CER1) in 3p21.3. <i>Genomics</i> , 2001, 73, 10-19.	1.3	33
13	Duplications on Human Chromosome 22 Reveal a Novel Ret Finger Protein-Like Gene Family with Sense and Endogenous Antisense Transcripts. <i>Genome Research</i> , 1999, 9, 803-814.	2.4	32
14	The germinal center kinase gene and a novel CDC25-like gene are located in the vicinity of the PYGM gene on 11q13. <i>Human Genetics</i> , 1997, 100, 611-619.	1.8	28
15	A 1-Mb PAC Contig Spanning the Common Eliminated Region 1 (CER1) in Microcell Hybrid-Derived SCID Tumors. <i>Genomics</i> , 1999, 62, 147-155.	1.3	28
16	Structure of the Promoter and Genomic Organization of the Human β -Adaptin Gene (BAM22) from Chromosome 22q12. <i>Genomics</i> , 1996, 36, 112-117.	1.3	27
17	The Mouse Ortholog of the Human SMARCB1 Gene Encodes Two Splice Forms. <i>Biochemical and Biophysical Research Communications</i> , 1999, 257, 886-890.	1.0	27
18	A novel gene containing LIM domains (LIMD1) is located within the common eliminated region 1 (C3CER1) in 3p21.3. <i>Human Genetics</i> , 1999, 105, 552-559.	1.8	26

#	ARTICLE	IF	CITATIONS
19	TOM1Genes Map to Human Chromosome 22q13.1 and Mouse Chromosome 8C1 and Encode Proteins Similar to the Endosomal Proteins HGS and STAM. Genomics, 1999, 57, 380-388.	1.3	26
20	Fine mapping of the constitutional translocation t(11;22)(q23;q11). Human Genetics, 2000, 106, 506-516.	1.8	20
21	ShiftDetector: detection of shift mutations. Bioinformatics, 2002, 18, 1137-1138.	1.8	18
22	Genomic Structure, 5' Flanking Sequences, and Precise Localization in 1P31.1 of the Human Prostaglandin F Receptor Gene. Biochemical and Biophysical Research Communications, 1999, 254, 413-416.	1.0	17
23	Characterization of Five Novel Human Genes in the 11q13-q22 Region. Biochemical and Biophysical Research Communications, 2000, 273, 90-94.	1.0	13
24	Sequence and Expression of the Mouse Homologue to Human Phospholipase C β 3 Neighboring Gene. Biochemical and Biophysical Research Communications, 1996, 223, 335-340.	1.0	8
25	Variable degree of mosaicism for tetrasomy 18p in phenotypically discordant monozygotic twins"Diagnostic implications. Molecular Genetics & Genomic Medicine, 2021, 9, e1526.	0.6	7