Valer Gotea

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	A novel role for nucleolin in splice site selection. RNA Biology, 2022, 19, 333-352.	1.5	3
2	Characterization and clustering of kinase isoform expression in metastatic melanoma. PLoS Computational Biology, 2022, 18, e1010065.	1.5	4
3	Assessing predictions of the impact of variants on splicing in CACI5. Human Mutation, 2019, 40, 1215-1224.	1.1	18
4	Transposable Elements: Classification, Identification, and Their Use As a Tool For Comparative Genomics. Methods in Molecular Biology, 2019, 1910, 177-207.	0.4	74
5	CAGI experiments: Modeling sequence variant impact on gene splicing using predictions from computational tools. Human Mutation, 2019, 40, 1252-1260.	1.1	2
6	Significant associations between driver gene mutations and DNA methylation alterations across many cancer types. PLoS Computational Biology, 2017, 13, e1005840.	1.5	39
7	CpG island methylator phenotype in adenocarcinomas from the digestive tract: Methods, conclusions, and controversies. World Journal of Gastrointestinal Oncology, 2017, 9, 105.	0.8	9
8	A Case of IL-7R Deficiency Caused by a Novel Synonymous Mutation and Implications for Mutation Screening in SCID Diagnosis. Frontiers in Immunology, 2016, 7, 443.	2.2	15
9	Pan-cancer stratification of solid human epithelial tumors and cancer cell lines reveals commonalities and tissue-specific features of the CpG island methylator phenotype. Epigenetics and Chromatin, 2015, 8, 14.	1.8	42
10	The functional relevance of somatic synonymous mutations in melanoma and other cancers. Pigment Cell and Melanoma Research, 2015, 28, 673-684.	1.5	47
11	Ascertaining regions affected by GC-biased gene conversion through weak-to-strong mutational hotspots. Genomics, 2014, 103, 349-356.	1.3	5
12	Whole-genome sequencing identifies a recurrent functional synonymous mutation in melanoma. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 13481-13486.	3.3	147
13	Recurrent patterns of DNA methylation in the <i>ZNF154,CASP8</i> , and <i>VHL</i> promoters across a wide spectrum of human solid epithelial tumors and cancer cell lines. Epigenetics, 2013, 8, 1355-1372.	1.3	52
14	Bidirectional Promoters as Important Drivers for the Emergence of Species-Specific Transcripts. PLoS ONE, 2013, 8, e57323.	1.1	25
15	Transposable Elements and Their Identification. Methods in Molecular Biology, 2012, 855, 337-359.	0.4	26
16	Functional analysis of synonymous substitutions predicted to affect splicing of the CFTR gene. Journal of Cystic Fibrosis, 2012, 11, 511-517.	0.3	17
17	Genome-wide detection of a TFIID localization element from an initial human disease mutation. Nucleic Acids Research, 2011, 39, 2175-2187.	6.5	26
18	Homotypic clusters of transcription factor binding sites are a key component of human promoters and enhancers. Genome Research, 2010, 20, 565-577.	2.4	203

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19	Tissue-Specific and Ubiquitous Expression Patterns from Alternative Promoters of Human Genes. PLoS ONE, 2010, 5, e12274.	1.1	30
20	DiRE: identifying distant regulatory elements of co-expressed genes. Nucleic Acids Research, 2008, 36, W133-W139.	6.5	118
21	Evolution of genes and genomes on the Drosophila phylogeny. Nature, 2007, 450, 203-218.	13.7	1,886
22	Transposable elements as a significant source of transcription regulating signals. Gene, 2006, 365, 104-110.	1.0	116
23	Do transposable elements really contribute to proteomes?. Trends in Genetics, 2006, 22, 260-267.	2.9	81
24	Spliceosomal small nuclear RNA genes in 11 insect genomes. Rna, 2006, 13, 5-14.	1.6	33
25	Mastering seeds for genomic size nucleotide BLAST searches. Nucleic Acids Research, 2003, 31, 6935-6941.	6.5	25