

Mojca Strazisar

List of Publications by Year in descending order

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Version: 2024-02-01

27
papers

867
citations

567281

15
h-index

552781

26
g-index

33
all docs

33
docs citations

33
times ranked

2158
citing authors

#	ARTICLE	IF	CITATIONS
1	Scarless excision of an insertion sequence restores capsule production and virulence in <i>Acinetobacter baumannii</i> . ISME Journal, 2022, 16, 1473-1477.	9.8	18
2	Long-Read Sequencing to Unravel Complex Structural Variants of CEP78 Leading to Cone-Rod Dystrophy and Hearing Loss. Frontiers in Cell and Developmental Biology, 2021, 9, 664317.	3.7	11
3	Methplotlib: analysis of modified nucleotides from nanopore sequencing. Bioinformatics, 2020, 36, 3236-3238.	4.1	23
4	Critical length in long-read resequencing. NAR Genomics and Bioinformatics, 2020, 2, lqz027.	3.2	4
5	Structural variants identified by Oxford Nanopore PromethION sequencing of the human genome. Genome Research, 2019, 29, 1178-1187.	5.5	143
6	Loss of DPP6 in neurodegenerative dementia: a genetic player in the dysfunction of neuronal excitability. Acta Neuropathologica, 2019, 137, 901-918.	7.7	37
7	NanoSatellite: accurate characterization of expanded tandem repeat length and sequence through whole genome long-read sequencing on PromethION. Genome Biology, 2019, 20, 239.	8.8	47
8	miRVaS: a tool to predict the impact of genetic variants on miRNAs. Nucleic Acids Research, 2016, 44, e23-e23.	14.5	19
9	Genetic variants in microRNA genes: impact on microRNA expression, function, and disease. Frontiers in Genetics, 2015, 6, 186.	2.3	106
10	Schizophrenia-Associated MIR204 Regulates Noncoding RNAs and Affects Neurotransmitter and Ion Channel Gene Sets. PLoS ONE, 2015, 10, e0144428.	2.5	12
11	MIR137 variants identified in psychiatric patients affect synaptogenesis and neuronal transmission gene sets. Molecular Psychiatry, 2015, 20, 472-481.	7.9	73
12	Identification of rare copy number variants in high burden schizophrenia families. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2013, 162, 273-282.	1.7	23
13	Less Cognitive and Neurological Deficits in Schizophrenia Patients Carrying Risk Variant in <i>ZNF804A</i> . Neuropsychobiology, 2012, 66, 158-166.	1.9	15
14	Co-occurrence of Marfan syndrome and schizophrenia: What can be learned?. European Journal of Medical Genetics, 2012, 55, 252-255.	1.3	11
15	Rare copy number variants in neuropsychiatric disorders: Specific phenotype or not?. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 812-822.	1.7	34
16	BBC3 is down-regulated with increased tumor size independently of p53 expression in head and neck cancer. Cancer Biomarkers, 2012, 11, 197-208.	1.7	9
17	Identification of a <i>CACNA2D4</i> deletion in late onset bipolar disorder patients and implications for the involvement of voltage-dependent calcium channels in psychiatric disorders. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 465-475.	1.7	27
18	LATS2 (LATS, large tumor suppressor, homolog 2 (Drosophila)). Atlas of Genetics and Cytogenetics in Oncology and Haematology, 2011, , .	0.1	0

#	ARTICLE	IF	CITATIONS
19	Absence of Pathogenic Mutations in VSX1 and SOD1 Genes in Patients With Keratoconus. <i>Cornea</i> , 2010, 29, 172-176.	1.7	66
20	Darier disease in Slovenia: spectrum of ATP2A2 mutations and relation to patients' phenotypes. <i>European Journal of Dermatology</i> , 2010, 20, 271-275.	0.6	8
21	Oligonucleotide DNA Microarray Profiling of Lung Adenocarcinoma Revealed Significant Downregulation and Deletions of Vasoactive Intestinal Peptide Receptor 1. <i>Cancer Investigation</i> , 2010, 28, 487-494.	1.3	9
22	The expression of COX-2, hTERT, MDM2, LATS2 and S100A2 in different types of non-small cell lung cancer (NSCLC). <i>Cellular and Molecular Biology Letters</i> , 2009, 14, 442-56.	7.0	21
23	Frequent polymorphic variations but rare tumour specific mutations of the S100A2 on 1q21 in non-small cell lung cancer. <i>Lung Cancer</i> , 2009, 63, 354-359.	2.0	9
24	LATS2 tumour specific mutations and down-regulation of the gene in non-small cell carcinoma. <i>Lung Cancer</i> , 2009, 64, 257-262.	2.0	49
25	Somatic Alterations of the Serine/Threonine Kinase LKB1 Gene in Squamous Cell (SCC) and Large Cell (LCC) Lung Carcinoma. <i>Cancer Investigation</i> , 2009, 27, 407-416.	1.3	21
26	Polymorphisms in COL4A3 and COL4A4 genes associated with keratoconus. <i>Molecular Vision</i> , 2009, 15, 2848-60.	1.1	62
27	K-RAS and P53 Mutations in Association with COX-2 and Htert Expression and Clinico-Pathological Status of Nsclc Patients. <i>Disease Markers</i> , 2008, 25, 97-106.	1.3	2