

AnikÃ³ Bozsik

List of Publications by Year in descending order

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Version: 2024-02-01

11
papers

532
citations

1163117

8
h-index

1281871

11
g-index

11
all docs

11
docs citations

11
times ranked

1795
citing authors

#	ARTICLE	IF	CITATIONS
1	Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. <i>Nature Genetics</i> , 2017, 49, 1767-1778.	21.4	289
2	Polygenic risk scores and breast and epithelial ovarian cancer risks for carriers of BRCA1 and BRCA2 pathogenic variants. <i>Genetics in Medicine</i> , 2020, 22, 1653-1666.	2.4	82
3	Common alleles at 6q25.1 and 1p11.2 are associated with breast cancer risk for BRCA1 and BRCA2 mutation carriers. <i>Human Molecular Genetics</i> , 2011, 20, 3304-3321.	2.9	68
4	The use of morphomutants to investigate septum formation and cell separation in <i>Schizosaccharomyces pombe</i> . <i>Archives of Microbiology</i> , 2000, 174, 386-392.	2.2	22
5	Breast and Prostate Cancer Risks for Male BRCA1 and BRCA2 Pathogenic Variant Carriers Using Polygenic Risk Scores. <i>Journal of the National Cancer Institute</i> , 2022, 114, 109-122.	6.3	19
6	Modulation of cancer pathways by inhibitors of guanylate metabolism. <i>Advances in Enzyme Regulation</i> , 2006, 46, 176-190.	2.6	16
7	Marker construction and cloning of acut1-like sequence with ARS activity in the fission yeast <i>Schizosaccharomyces japonicus</i> . <i>Yeast</i> , 2002, 19, 485-498.	1.7	13
8	Complex Characterization of Germline Large Genomic Rearrangements of the BRCA1 and BRCA2 Genes in High-Risk Breast Cancer Patients: Novel Variants from a Large National Center. <i>International Journal of Molecular Sciences</i> , 2020, 21, 4650.	4.1	10
9	Germline Structural Variations in Cancer Predisposition Genes. <i>Frontiers in Genetics</i> , 2021, 12, 634217.	2.3	7
10	Application of Multilayer Evidence for Annotation of C-Terminal BRCA2 Variants. <i>Cancers</i> , 2021, 13, 881.	3.7	3
11	Surprising genetic and pathological findings in a patient with giant bilateral perirenal tumours: PEComas and mutations of PTCH1 in Gorlin-Goltz syndrome. <i>Journal of Medical Genetics</i> , 2022, 59, 916-919.	3.2	3