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List of Publications by Year in descending order

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Version: 2024-02-01

17
papers

298
citations

1477746

6
h-index

940134

16
g-index

17
all docs

17
docs citations

17
times ranked

844
citing authors

#	ARTICLE	IF	CITATIONS
1	SweGen: a whole-genome data resource of genetic variability in a cross-section of the Swedish population. <i>European Journal of Human Genetics</i> , 2017, 25, 1253-1260.	1.4	148
2	Whole-genome sequencing of synchronous thyroid carcinomas identifies aberrant DNA repair in thyroid cancer dedifferentiation. <i>Journal of Pathology</i> , 2020, 250, 183-194.	2.1	40
3	Exome sequencing in one family with gastric- and rectal cancer. <i>BMC Genetics</i> , 2016, 17, 41.	2.7	31
4	Exome sequencing in 51 early onset non-familial CRC cases. <i>Molecular Genetics & Genomic Medicine</i> , 2019, 7, e605.	0.6	17
5	Frequent GU wobble pairings reduce translation efficiency in <i>Plasmodium falciparum</i> . <i>Scientific Reports</i> , 2017, 7, 723.	1.6	14
6	<i>PHIP</i> - a novel candidate breast cancer susceptibility locus on 6q14.1. <i>Oncotarget</i> , 2017, 8, 102769-102782.	0.8	9
7	pyCancerSig: subclassifying human cancer with comprehensive single nucleotide, structural and microsatellite mutational signature deconstruction from whole genome sequencing. <i>BMC Bioinformatics</i> , 2020, 21, 128.	1.2	7
8	Cancer risk susceptibility loci in a Swedish population. <i>Oncotarget</i> , 2017, 8, 110300-110310.	0.8	7
9	Genetic analyses supporting colorectal, gastric, and prostate cancer syndromes. <i>Genes Chromosomes and Cancer</i> , 2019, 58, 775-782.	1.5	6
10	Linkage analysis revealed risk loci on 6p21 and 18p11.2-q11.2 in familial colon and rectal cancer, respectively. <i>European Journal of Human Genetics</i> , 2019, 27, 1286-1295.	1.4	4
11	Two novel colorectal cancer risk loci in the region on chromosome 9q22.32. <i>Oncotarget</i> , 2018, 9, 11170-11179.	0.8	4
12	Massive parallel sequencing in a family with rectal cancer. <i>Hereditary Cancer in Clinical Practice</i> , 2021, 19, 23.	0.6	3
13	A search for modifying genetic factors in CHEK2:c.1100delC breast cancer patients. <i>Scientific Reports</i> , 2021, 11, 14763.	1.6	3
14	Sequencing for germline mutations in Swedish breast cancer families reveals novel breast cancer risk genes. <i>Scientific Reports</i> , 2021, 11, 14737.	1.6	2
15	Colorectal cancer risk susceptibility loci in a Swedish population. <i>Molecular Carcinogenesis</i> , 2021, , .	1.3	2
16	A Swedish Genome-Wide Haplotype Association Analysis Identifies a Novel Breast Cancer Susceptibility Locus in 8p21.2 and Characterizes Three Loci on Chromosomes 10, 11 and 16. <i>Cancers</i> , 2022, 14, 1206.	1.7	1
17	Identification of known and novel familial cancer genes in Swedish colorectal cancer families. <i>International Journal of Cancer</i> , 2021, 149, 627-634.	2.3	0