Sock Hoai Chan

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/6703364/publications.pdf

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20 papers	365 citations	1039406 9 h-index	839053 18 g-index
22 all docs	22 docs citations	22 times ranked	752 citing authors

#	Article	IF	Citations
1	Investigation into the origins of an ancient BRCA1 founder mutation identified among Chinese families in Singapore. International Journal of Cancer, 2021, 148, 637-645.	2.3	5
2	Family history assessment significantly enhances delivery of precision medicine in the genomics era. Genome Medicine, 2021, 13, 3.	3.6	19
3	Impact of Variant Reclassification in Cancer Predisposition Genes on Clinical Care. JCO Precision Oncology, 2021, 5, 577-584.	1.5	18
4	CDKN2A germline alterations and the relevance of genotype-phenotype associations in cancer predisposition. Hereditary Cancer in Clinical Practice, 2021, 19, 21.	0.6	36
5	Spectrum of Germline Mutations Within Fanconi Anemiaâ€“Associated Genes Across Populations of Varying Ancestry. JNCI Cancer Spectrum, 2021, 5, .	1.4	3
6	Multiple neoplasia in a patient with Gitelman syndrome harboring germline monoallelic MUTYH mutation. Npj Genomic Medicine, 2020, 5, 39.	1.7	3
7	Functional characterisation guides classification of novel BAP1 germline variants. Npj Genomic Medicine, 2020, 5, 50.	1.7	3
8	Biallelic NF1 inactivation in high grade serous ovarian cancers from patients with neurofibromatosis type 1. Familial Cancer, 2020, 19, 353-358.	0.9	3
9	Missense PALB2 germline variant disrupts nuclear localization of PALB2 in a patient with breast cancer. Familial Cancer, 2020, 19, 123-131.	0.9	3
10	Functional analysis of clinical BARD1 germline variants. Journal of Physical Education and Sports Management, 2019, 5, a004093.	0.5	6
11	Advances in Sarcoma Genomics and Therapeutic Management. , 2019, , 609-621.		0
12	Clinical relevance of screening checklists for detecting cancer predisposition syndromes in Asian childhood tumours. Npj Genomic Medicine, 2018, 3, 30.	1.7	13
13	Germline Pathogenic Variants in Homologous Recombination and DNA Repair Genes in an Asian Cohort of Young-Onset Colorectal Cancer. JNCI Cancer Spectrum, 2018, 2, pky054.	1.4	21
14	Whole exome sequencing identifies recessive germline mutations in FAM160A1 in familial NK/T cell lymphoma. Blood Cancer Journal, 2018, 8, 111.	2.8	5
15	Germline Mutations in Cancer Predisposition Genes are Frequent in Sporadic Sarcomas. Scientific Reports, 2017, 7, 10660.	1.6	52
16	Impact of subsidies on cancer genetic testing uptake in Singapore. Journal of Medical Genetics, 2017, 54, 254-259.	1.5	24
17	Germline mutation contribution to chromosomal instability. Endocrine-Related Cancer, 2017, 24, T33-T46.	1.6	13
18	Germline hemizygous deletion of CDKN2Aâ€“CDKN2B locus in a patient presenting with Liâ€“Fraumeni syndrome. Npj Genomic Medicine, 2016, 1, 16015.	1.7	9

#	Article	IF	Citations
19	Tissue Microbiome Profiling Identifies an Enrichment of Specific Enteric Bacteria in Opisthorchis viverrini Associated Cholangiocarcinoma. EBioMedicine, 2016, 8, 195-202.	2.7	94
20	Mutation spectrum of POLE and POLD1 mutations in South East Asian women presenting with grade 3 endometrioid endometrial carcinomas. Gynecologic Oncology, 2016, 141, 113-120.	0.6	34