Eliza Courtney

List of Publications by Year
in descending order

21 papers	415 citations	1162889 8 h-index	752573 20 g-index
21 all docs	21 docs citations	21 times ranked	751 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	Understanding patients' views and willingness toward the use of telehealth in a cancer genetics service in Asia. Journal of Genetic Counseling, 2021, 30, 1658-1670.	0.9	5
3	Evaluating empowerment in genetic counseling using patientâ€reported outcomes. Clinical Genetics, 2020, 97, 246-256.	1.0	19
4	An in-depth exploration of the post-test informational needs of BRCA1 and BRCA2 pathogenic variant carriers in Asia. Hereditary Cancer in Clinical Practice, 2020, 18, 22.	0.6	8
5	Early-onset breast cancer in a woman with a germline mobile element insertion resulting in BRCA2 disruption: a case report. Human Genome Variation, 2020, 7, 24.	0.4	3
6	Predictive Testing for Tumor Predisposition Syndromes in Pediatric Relatives: An Asian Experience. Frontiers in Pediatrics, 2020, 8, 568528.	0.9	1
7	Biallelic NF1 inactivation in high grade serous ovarian cancers from patients with neurofibromatosis type 1. Familial Cancer, 2020, 19, 353-358.	0.9	3
8	Use of telephone intake for family history taking at a cancer genetics service in Asia. Journal of Genetic Counseling, 2020, 29, 1192-1199.	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	Impact of free cancer predisposition cascade genetic testing on uptake in Singapore. Npj Genomic Medicine, 2019, 4, 22.	1.7	26
12	The Global State of the Genetic Counseling Profession. European Journal of Human Genetics, 2019, 27, 183-197.	1.4	215
13	Risk management adherence following genetic testing for hereditary cancer syndromes: a Singaporean experience. Familial Cancer, 2018, 17, 621-626.	0.9	6
14	Factors influencing the decision to share cancer genetic results among family members: An inâ€depth interview study of women in an Asian setting. Psycho-Oncology, 2018, 27, 998-1004.	1.0	22
15	Evaluation of the relative effectiveness of the 2017 updated Manchester scoring system for predicting BRCA1/2 mutations in a Southeast Asian country. Journal of Medical Genetics, 2018, 55, 344-350.	1.5	5
16	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
17	A delayed diagnosis of Pallister-Hall syndrome in an adult male following the incidental detection of a hypothalamic hamartoma. Human Genome Variation, 2018, 5, 31.	0.4	3
18	The influence of Malay cultural beliefs on breast cancer screening and genetic testing: A focus group study. Psycho-Oncology, 2018, 27, 2855-2861.	1.0	30

#	Article	IF	Citations
19	Impact of Appointment Waiting Time on Attendance Rates at a Clinical Cancer Genetics Service. Journal of Genetic Counseling, 2018, 27, 1473-1481.	0.9	15
20	Predictors of next-generation sequencing panel selection using a shared decision-making approach. Npj Genomic Medicine, 2018, 3, 11.	1.7	9
21	Clinical management of pheochromocytoma and paraganglioma in Singapore: missed opportunities for genetic testing. Molecular Genetics & Genomic Medicine, 2017, 5, 602-607.	0.6	7