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	The Global State of the Genetic Counseling Profession. European Journal of Human Genetics, 2019, 27, 183-197.	1.4	215
2	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
3	Impact of free cancer predisposition cascade genetic testing on uptake in Singapore. Npj Genomic Medicine, 2019, 4, 22.	1.7	26
4	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
5	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
6	Evaluating empowerment in genetic counseling using patientâ€reported outcomes. Clinical Genetics, 2020, 97, 246-256.	1.0	19
7	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
8	Predictors of next-generation sequencing panel selection using a shared decision-making approach. Npj Genomic Medicine, 2018, 3, 11.	1.7	9
9	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
10	Clinical management of pheochromocytoma and paraganglioma in Singapore: missed opportunities for genetic testing. Molecular Genetics & Genomic Medicine, 2017, 5, 602-607.	0.6	7
11	Risk management adherence following genetic testing for hereditary cancer syndromes: a Singaporean experience. Familial Cancer, 2018, 17, 621-626.	0.9	6
12	Functional analysis of clinical BARD1 germline variants. Journal of Physical Education and Sports Management, 2019, 5, a004093.	0.5	6
13	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
14	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
15	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
16	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
17	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
18	Biallelic NF1 inactivation in high grade serous ovarian cancers from patients with neurofibromatosis type 1. Familial Cancer, 2020, 19, 353-358.	0.9	3

#	Article	IF	Citations
19	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
20	Missense PALB2 germline variant disrupts nuclear localization of PALB2 in a patient with breast cancer. Familial Cancer, 2020, 19, 123-131.	0.9	3
21	Predictive Testing for Tumor Predisposition Syndromes in Pediatric Relatives: An Asian Experience. Frontiers in Pediatrics, 2020, 8, 568528.	0.9	1