Ashley Crook

List of Publications by Year
in descending order

22 papers	469 citations	1039406 9 h-index	713013 21 g-index
23 all docs	23 docs citations	23 times ranked	942 citing authors

#	Article	IF	Citations
1	Genetic counseling and testing practices for late-onset neurodegenerative disease: a systematic review. Journal of Neurology, 2022, 269, 676-692.	1.8	11
2	The psychological impact and experience of breast cancer screening in young women with an increased risk of breast cancer due to neurofibromatosis type 1. Familial Cancer, 2022, 21, 241-253.	0.9	6
3	Genetic counselling and testing for neurodegenerative disorders using a proposed standard of practice for ALS/MND: diagnostic testing comes first. European Journal of Human Genetics, 2022, 30, 394-395.	1.4	4
4	Surveillance Improves Outcomes for Carriers of SDHB Pathogenic Variants: A Multicenter Study. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e1907-e1916.	1.8	11
5	Incorporating patient perspectives in the development of a core outcome set for reproductive genetic carrier screening: a sequential systematic review. European Journal of Human Genetics, 2022, , .	1.4	3
6	Toward genetic counseling practice standards for diagnostic testing in amyotrophic lateral sclerosis and frontotemporal dementia. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2022, 23, 562-574.	1.1	1
7	Predictive genetic testing for Motor neuron disease: time for a guideline?. European Journal of Human Genetics, 2022, 30, 635-636.	1.4	2
8	Genetic counseling and diagnostic genetic testing for familial amyotrophic lateral sclerosis and/or frontotemporal dementia: A qualitative study of client experiences. Journal of Genetic Counseling, 2022, 31, 1206-1218.	0.9	5
9	Implementing gene curation for hereditary cancer susceptibility in Australia: achieving consensus on genes with clinical utility. Journal of Medical Genetics, 2021, 58, 853-858.	1.5	3
10	Helping young children understand inherited cancer predisposition syndromes using bibliotherapy. Journal of Genetic Counseling, 2021, 30, 1119-1132.	0.9	3
11	Health system interventions to integrate genetic testing in routine oncology services: A systematic review. PLoS ONE, 2021, 16, e0250379.	1.1	13
12	Patient and Relative Experiences and Decision-making About Genetic Testing and Counseling for Familial ALS and FTD. Alzheimer Disease and Associated Disorders, 2021, Publish Ahead of Print, 374-385.	0.6	9
13	Systematic review of outcomes in studies of reproductive genetic carrier screening: Towards development of a core outcome set. Genetics in Medicine, 2021, , .	1.1	9
14	The C9orf72 hexanucleotide repeat expansion presents a challenge for testing laboratories and genetic counseling. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 310-316.	1.1	16
15	Bayesian approach to determining penetrance of pathogenic SDH variants. Journal of Medical Genetics, 2018, 55, 729-734.	1.5	44
16	Predictive genetic testing for amyotrophic lateral sclerosis and frontotemporal dementia: genetic counselling considerations. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2017, 18, 475-485.	1.1	26
17	Patient-centered decision making in amyotrophic lateral sclerosis: where are we?. Neurodegenerative Disease Management, 2017, 7, 377-386.	1.2	6
18	Lessons learnt from implementation of a Lynch syndrome screening program for patients with gynaecological malignancy. Pathology, 2017, 49, 457-464.	0.3	34

#	Article	IF	Citations
19	Fumarate Hydrataseâ€“deficient Uterine Leiomyomas Occur in Both the Syndromic and Sporadic Settings. American Journal of Surgical Pathology, 2016, 40, 599-607.	2.1	102
20	Connecting patients, researchers and clinical genetics services: the experiences of participants in the Australian Ovarian Cancer Study (AOCS). European Journal of Human Genetics, 2015, 23, 152-158.	1.4	14
21	The responses of research participants and their next of kin to receiving feedback of genetic test results following participation in the Australian Ovarian Cancer Study. Genetics in Medicine, 2013, 15, 458-465.	1.1	21
22	BRAFV600E Immunohistochemistry Facilitates Universal Screening of Colorectal Cancers for Lynch Syndrome. American Journal of Surgical Pathology, 2013, 37, 1592-1602.	2.1	125