Elizabeth Ormondroyd

List of Publications by Year
in descending order

28 papers	1,631 citations	516710 16 h-index	526287 27 g-index
28 all docs	28 docs citations	28 times ranked	3851 citing authors

#	Article	IF	Citations
1	Genomic health data generation in the UK: a 360 view. European Journal of Human Genetics, 2022, 30, 782-789.	2.8	8
2	Provocation Testing and Therapeutic Response in a Newly Described Channelopathy: RyR2 Calcium Release Deficiency Syndrome. Circulation Genomic and Precision Medicine, 2022, 15, CIRCGEN121003589.	3.6	15
3	Genomic sequencing in oncology: Considerations for integration in routine cancer care. European Journal of Cancer Care, 2022, 31, e13584.	1.5	4
4	Common genetic variants and modifiable risk factors underpin hypertrophic cardiomyopathy susceptibility and expressivity. Nature Genetics, 2021, 53, 135-142.	21.4	165
5	Bi-allelic MCM10 variants associated with immune dysfunction and cardiomyopathy cause telomere shortening. Nature Communications, 2021, 12, 1626.	12.8	22
6	Maximal Wall Thickness Measurement in Hypertrophic Cardiomyopathy. JACC: Cardiovascular Imaging, 2021, 14, 2123-2134.	5.3	18
7	Taking it to the bank: the ethical management of individual findings arising in secondary research. Journal of Medical Ethics, 2021, 47, 689-696.	1.8	4
8	Incremental value of left atrial booster and reservoir strain in predicting atrial fibrillation in patients with hypertrophic cardiomyopathy: a cardiovascular magnetic resonance study. Journal of Cardiovascular Magnetic Resonance, 2021, 23, 109.	3.3	14
9	Secondary findings in inherited heart conditions: a genotype-first feasibility study to assess phenotype, behavioural and psychosocial outcomes. European Journal of Human Genetics, 2020, 28, 1486-1496.	2.8	13
10	Whole-genome sequencing of patients with rare diseases in a national health system. Nature, 2020, 583, 96-102.	27.8	338
11	Reevaluation of the South Asian MYBPC3 ^{Î”25bp} Intronic Deletion in Hypertrophic Cardiomyopathy. Circulation Genomic and Precision Medicine, 2020, 13, e002783.	3.6	31
12	ImplementationÂof a genomic medicine multi-disciplinary team approach for rare diseaseÂin the clinical setting: a prospective exome sequencingÂcase series. Genome Medicine, 2019, 11, 46.	8.2	25
13	Do health professionals value genomic testing? A discrete choice experiment in inherited cardiovascular disease. European Journal of Human Genetics, 2019, 27, 1639-1648.	2.8	11
14	6â€…RV function deteriorates earlier than LV function and predicts adverse cardiovascular outcomes. , 2019, , .		0
15	Analysis of 51 proposed hypertrophic cardiomyopathy genes from genome sequencing data in sarcomere negative cases has negligible diagnostic yield. Genetics in Medicine, 2019, 21, 1576-1584.	2.4	44
16	Views of rare disease participants in aÂUK whole-genome sequencing study towards secondary findings: a qualitative study. European Journal of Human Genetics, 2018, 26, 652-659.	2.8	30
17	â€œNot pathogenic until proven otherwiseâ€: perspectives of UK clinical genomics professionals toward secondary findings in context of a Genomic Medicine Multidisciplinary Team and the 100,000 Genomes Project. Genetics in Medicine, 2018, 20, 320-328.	2.4	56
18	From Genotype to Phenotype. Circulation Genomic and Precision Medicine, 2018, 11, e002316.	3.6	8

#	Article	IF	Citations
19	Distinct ECG Phenotypes Identified in Hypertrophic Cardiomyopathy Using Machine Learning Associate With Arrhythmic Risk Markers. Frontiers in Physiology, 2018, 9, 213.	2.8	57
20	Insights from early experience of a Rare Disease Genomic Medicine Multidisciplinary Team: a qualitative study. European Journal of Human Genetics, 2017, 25, 680-686.	2.8	24
21	Stakeholder views on secondary findings in whole-genome and whole-exome sequencing: a systematic review of quantitative and qualitative studies. Genetics in Medicine, 2017, 19, 283-293.	2.4	119
22	Exploring the potential duty of care in clinical genomics under UK law. Medical Law International, 2017, 17, 158-182.	1.1	17
23	Factors influencing success of clinical genome sequencing across a broad spectrum of disorders. Nature Genetics, 2015, 47, 717-726.	21.4	310
24	Pre-symptomatic genetic testing for inherited cardiac conditions: a qualitative exploration of psychosocial and ethical implications. European Journal of Human Genetics, 2014, 22, 88-93.	2.8	65
25	Attitudes to reproductive genetic testing in women who had a positive BRCA test before having children: a qualitative analysis. European Journal of Human Genetics, 2012, 20, 4-10.	2.8	68
26	Genome Wide Identification of Recessive Cancer Genes by Combinatorial Mutation Analysis. PLoS ONE, 2008, 3, e3380.	2.5	12
27	Molecular Cloning, cDNA Sequence, and Chromosomal Localization of the Human Phosphatidylinositol 3-Kinase p110Î± (PIK3CA) Gene. Genomics, 1994, 24, 472-477.	2.9	107
28	Mapping of cosmid clones in Huntington's disease region of chromosome 4. Somatic Cell and Molecular Genetics, 1991, 17, 83-91.	0.7	46