Jennifer Hammond

List of Publications by Year
in descending order

13	148	1307594 7	1199594 12
papers	citations	h-index	g-index

13	13	13	192
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Living with osteogenesis imperfecta: A qualitative study exploring experiences and psychosocial impact from the perspective of patients, parents and professionals. Disability and Health Journal, 2022, 15, 101168.	2.8	6
2	Assessing womenâ€™s preferences towards tests that may reveal uncertain results from prenatal genomic testing: Development of attributes for a discrete choice experiment, using a mixed-methods design. PLoS ONE, 2022, 17, e0261898.	2.5	4
3	â€œItâ€™s okay to not know â€¦â€•a qualitative exploration of faculty approaches to working with uncertainty. BMC Medical Education, 2022, 22, 135.	2.4	6
4	Factors that impact on women's decisionâ€•making around prenatal genomic tests: An international discrete choice survey. Prenatal Diagnosis, 2022, 42, 934-946.	2.3	5
5	Parental experiences of uncertainty following an abnormal fetal anomaly scan: Insights using Hanâ€™s taxonomy of uncertainty. Journal of Genetic Counseling, 2021, 30, 198-210.	1.6	20
6	Dealing with uncertain results from chromosomal microarray and exome sequencing in the prenatal setting: An international crossâ€•sectional study with healthcare professionals. Prenatal Diagnosis, 2021, 41, 720-732.	2.3	13
7	Animation or leaflet: Does it make a difference when educating young people about genome sequencing?. Patient Education and Counseling, 2021, 104, 2522-2530.	2.2	2
8	How to deal with uncertainty in prenatal genomics: A systematic review of guidelines and policies. Clinical Genetics, 2021, 100, 647-658.	2.0	15
9	Young people's understanding, attitudes and involvement in decision-making about genome sequencing for rare diseases: A qualitative study with participants in the UK 100, 000 Genomes Project. European Journal of Medical Genetics, 2020, 63, 104043.	1.3	13
10	Couples experiences of receiving uncertain results following prenatal microarray or exome sequencing: A mixedâ€•methods systematic review. Prenatal Diagnosis, 2020, 40, 1028-1039.	2.3	25
11	Delivering genome sequencing for rapid genetic diagnosis in critically ill children: parent and professional views, experiences and challenges. European Journal of Human Genetics, 2020, 28, 1529-1540.	2.8	29
12	Development and mixed-methods evaluation of an online animation for young people about genome sequencing. European Journal of Human Genetics, 2020, 28, 896-906.	2.8	10
13	56â€…Parent and health professional experiences and views of genome sequencing for rapid diagnosis in critically ill children. , 2019, , .		0