

Fiona M Ulph

List of Publications by Year in descending order

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Version: 2024-02-01

57
papers

1,045
citations

394421

19
h-index

501196

28
g-index

60
all docs

60
docs citations

60
times ranked

1115
citing authors

#	ARTICLE	IF	CITATIONS
1	Communication of carrier status information following universal newborn screening for sickle cell disorders and cystic fibrosis: qualitative study of experience and practice. <i>Health Technology Assessment</i> , 2009, 13, 1-82, iii.	2.8	58
2	Effect of hopelessness on the links between psychiatric symptoms and suicidality in a vulnerable population at risk of suicide. <i>Psychiatry Research</i> , 2015, 230, 464-471.	3.3	57
3	Cognitive-behavioural suicide prevention for male prisoners: a pilot randomized controlled trial. <i>Psychological Medicine</i> , 2015, 45, 3441-3451.	4.5	47
4	Personality functioning: the influence of stature. <i>Archives of Disease in Childhood</i> , 2004, 89, 17-21.	1.9	46
5	Parents' responses to receiving sickle cell or cystic fibrosis carrier results for their child following newborn screening. <i>European Journal of Human Genetics</i> , 2015, 23, 459-465.	2.8	46
6	The moderating effects of coping and self-esteem on the relationship between defeat, entrapment and suicidality in a sample of prisoners at high risk of suicide. <i>European Psychiatry</i> , 2015, 30, 988-994.	0.2	38
7	What are the benefits and harms of risk stratified screening as part of the NHS breast screening Programme? Study protocol for a multi-site non-randomised comparison of BC-predict versus usual screening (NCT04359420). <i>BMC Cancer</i> , 2020, 20, 570.	2.6	37
8	Developing an intervention to facilitate family communication about inherited genetic conditions, and training genetic counsellors in its delivery. <i>European Journal of Human Genetics</i> , 2016, 24, 794-802.	2.8	35
9	Syria: refugee parents' experiences and need for parenting support in camps and humanitarian settings. <i>Vulnerable Children and Youth Studies</i> , 2018, 13, 19-29.	1.1	34
10	Imparting carrier status results detected by universal newborn screening for sickle cell and cystic fibrosis in England: a qualitative study of current practice and policy challenges. <i>BMC Health Services Research</i> , 2007, 7, 203.	2.2	31
11	Psychological Impact on Parents of an Inconclusive Diagnosis Following Newborn Bloodspot Screening for Cystic Fibrosis: A Qualitative Study. <i>International Journal of Neonatal Screening</i> , 2019, 5, 23.	3.2	31
12	Rapid Point-of-Care Genotyping to Avoid Aminoglycoside-Induced Ototoxicity in Neonatal Intensive Care. <i>JAMA Pediatrics</i> , 2022, 176, 486.	6.2	30
13	Provision of information about newborn screening antenatally: a sequential exploratory mixed-methods project. <i>Health Technology Assessment</i> , 2017, 21, 1-240.	2.8	26
14	Risks and benefits of nanotechnology: How young adults perceive possible advances in nanomedicine compared with conventional treatments. <i>Health, Risk and Society</i> , 2007, 9, 159-171.	1.7	24
15	The introduction of risk stratified screening into the NHS breast screening Programme: views from British-Pakistani women. <i>BMC Cancer</i> , 2020, 20, 452.	2.6	23
16	Engagement barriers and service inequities in the NHS Breast Screening Programme: Views from British-Pakistani women. <i>Journal of Medical Screening</i> , 2020, 27, 130-137.	2.3	21
17	How should risk be communicated to children: a cross-sectional study comparing different formats of probability information. <i>BMC Medical Informatics and Decision Making</i> , 2009, 9, 26.	3.0	20
18	Informing Children of Their Newborn Screening Carrier Result for Sickle Cell or Cystic Fibrosis: Qualitative Study of Parents' Intentions, Views and Support Needs. <i>Journal of Genetic Counseling</i> , 2014, 23, 409-420.	1.6	20

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19	Improving the quality of prison research: A qualitative study of ex-offender service user involvement in prison suicide prevention research. <i>Health Expectations</i> , 2018, 21, 100-109.	2.6	20
20	An Exploration of Parents' Perceptions and Beliefs About Changes Following Participation in a Family Skill Training Program: a Qualitative Study in a Developing Country. <i>Prevention Science</i> , 2015, 16, 674-684.	2.6	19
21	Young children's experiences of living with a parent with bipolar disorder: Understanding the child's perspective. <i>Psychology and Psychotherapy: Theory, Research and Practice</i> , 2017, 90, 212-228.	2.5	17
22	Diagnosing and Preventing Hearing Loss in the Genomic Age. <i>Trends in Hearing</i> , 2019, 23, 233121651987898.	1.3	16
23	Psychological Characteristics and Predictors of Suicide Probability in High-Risk Prisoners. <i>Criminal Justice and Behavior</i> , 2017, 44, 321-335.	1.8	15
24	Receiving results of uncertain clinical relevance from population genetic screening: systematic review & meta-synthesis of qualitative research. <i>European Journal of Human Genetics</i> , 2022, 30, 520-531.	2.8	15
25	Familial influences on antenatal and newborn haemoglobinopathy screening. <i>Ethnicity and Health</i> , 2011, 16, 361-375.	2.5	14
26	Exploration of Mechanisms behind Changes after Participation in a Parenting Intervention: A Qualitative Study in a Low-Resource Setting. <i>American Journal of Community Psychology</i> , 2016, 57, 181-189.	2.5	14
27	Eliciting Preferences for Information Provision in Newborn Bloodspot Screening Programs. <i>Value in Health</i> , 2017, 20, 651-661.	0.3	14
28	The prioritization of symptom beliefs over illness beliefs: The development and validation of the Pain Perception Questionnaire for Young People. <i>British Journal of Health Psychology</i> , 2018, 23, 68-87.	3.5	14
29	A qualitative study exploring genetic counsellors' experiences of counselling children. <i>European Journal of Human Genetics</i> , 2010, 18, 1090-1094.	2.8	13
30	Preferences for aspects of antenatal and newborn screening: a systematic review. <i>BMC Pregnancy and Childbirth</i> , 2019, 19, 131.	2.4	12
31	Qualitative exploration of health professionals' experiences of communicating positive newborn bloodspot screening results for nine conditions in England. <i>BMJ Open</i> , 2020, 10, e037081.	1.9	12
32	The Role of Information Provision in Economic Evaluations of Newborn Bloodspot Screening: A Systematic Review. <i>Applied Health Economics and Health Policy</i> , 2015, 13, 615-626.	2.1	11
33	Training Genetic Counsellors to Deliver an Innovative Therapeutic Intervention: their Views and Experience of Facilitating Multi-Family Discussion Groups. <i>Journal of Genetic Counseling</i> , 2017, 26, 199-214.	1.6	11
34	Ethical issues in research into conflict and displacement. <i>Lancet, The</i> , 2013, 382, 764-765.	18.7	10
35	The Strengthening Families Program 10-14 in Panama: Parents' perceptions of cultural fit. <i>Professional Psychology: Research and Practice</i> , 2016, 47, 56-65.	1.0	10
36	A qualitative study to explore how professionals in the United Kingdom make decisions to test children for a sickle cell carrier status. <i>European Journal of Human Genetics</i> , 2016, 24, 164-170.	2.8	9

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37	Pharmacogenetics to Avoid Loss of Hearing (PALOH) trial: a protocol for a prospective observational implementation trial. <i>BMJ Open</i> , 2021, 11, e044457.	1.9	9
38	Process evaluation of co-designed interventions to improve communication of positive newborn bloodspot screening results. <i>BMJ Open</i> , 2021, 11, e050773.	1.9	9
39	Communication with children about sickle cell disease: A qualitative study of parent experience. <i>British Journal of Health Psychology</i> , 2018, 23, 685-700.	3.5	8
40	Rethinking Strategies for Positive Newborn Screening Result (NBS+) Delivery (ReSPoND): a process evaluation of co-designing interventions to minimise impact on parental emotional well-being and stress. <i>Pilot and Feasibility Studies</i> , 2019, 5, 108.	1.2	8
41	Processing of positive newborn screening results: a qualitative exploration of current practice in England. <i>BMJ Open</i> , 2020, 10, e044755.	1.9	8
42	The feasibility of implementing risk stratification into a national breast cancer screening programme: a focus group study investigating the perspectives of healthcare personnel responsible for delivery. <i>BMC Women's Health</i> , 2022, 22, 142.	2.0	8
43	Young Adults' Pre-existing Knowledge of Cystic Fibrosis and Sickle Cell Diseases: Implications for Newborn Screening. <i>Journal of Genetic Counseling</i> , 2014, 23, 121-130.	1.6	7
44	Consent for newborn screening: screening professionals' and parents' views. <i>Public Health</i> , 2020, 178, 151-158.	2.9	6
45	Understanding Midwives' Preferences for Providing Information About Newborn Bloodspot Screening. <i>MDM Policy and Practice</i> , 2018, 3, 238146831774617.	0.9	5
46	Disparities in Current and Future Childhood and Newborn Carrier Identification. <i>Journal of Genetic Counseling</i> , 2014, 23, 701-707.	1.6	4
47	Facilitators and Barriers to Participating in a Randomized Controlled Trial of a Psychological Therapy for Substance Use. <i>Journal of Nervous and Mental Disease</i> , 2019, 207, 487-496.	1.0	4
48	Preventing interpersonal violence in Panama: is a parenting intervention developed in Australia culturally appropriate?. <i>International Journal of Public Health</i> , 2016, 61, 915-922.	2.3	3
49	Constructing a Bioethical Framework to Evaluate and Optimise Newborn Bloodspot Screening for Cystic Fibrosis. <i>International Journal of Neonatal Screening</i> , 2020, 6, 40.	3.2	2
50	Role of religion/spirituality in the context of genetic counseling: health professionals' experiences in an Islamic country. <i>Journal of Biochemical and Clinical Genetics</i> , 0, , 60-70.	0.1	2
51	<i>I just want to be normal</i>: A qualitative investigation of adolescents' coping goals when dealing with pain related to arthritis and the underlying parent-adolescent personal models. <i>Paediatric and Neonatal Pain</i> , 2022, 4, 96-109.	1.7	2
52	<i>We want to live a little longer and our family want[s] us around</i>: A summative content analysis of adherence to COVID-19-related guidelines using the Theoretical Domains Framework. <i>British Journal of Health Psychology</i> , 2022, 27, 1119-1152.	3.5	2
53	Consent for newborn screening and storage of blood samples. <i>British Journal of Midwifery</i> , 2017, 25, 730-732.	0.4	1
54	Exploring the role of Islam on the lived experience of patients with Long QT Syndrome in Saudi Arabia. <i>Journal of Genetic Counseling</i> , 2022, 31, 922-936.	1.6	1

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55	Communicating Carrier Status Information to Children: Their Ability to Understand Concepts Related to Genetic Testing. Australian Journal of Cancer Nursing, 2007, 9, 250-250.	1.6	0
56	WS20-2 Parents' psychological adaptation to an unclear diagnosis after positive newborn bloodspot screening for cystic fibrosis: a qualitative study. Journal of Cystic Fibrosis, 2019, 18, S37.	0.7	0
57	The parenting process in Syrian refugees: A grounded theory study.. Peace and Conflict, 2021, 27, 67-80.	0.4	0