# Felicity K Boardman

List of Publications by Year
in descending order



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|---|---|---|---|
| 38 papers | 2,590 citations | 516710 16 h-index | 315739 38 g-index |
| 38 all docs | 38 docs citations | 38 times ranked | 3241 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | The Mixed Methods Appraisal Tool (MMAT) version 2018 for information professionals and researchers. Education for Information, 2018, 34, 285-291. | 0.5 | 1,312 |
| 2 | Improving the content validity of the mixed methods appraisal tool: aÂmodified e-Delphi study. Journal of Clinical Epidemiology, 2019, 111, 49-59.e1. | 5.0 | 441 |
| 3 | Social networks â€“ The future for health care delivery. Social Science and Medicine, 2012, 75, 2233-2241. | 3.8 | 157 |
| 4 | Evaluating recovery following hip fracture: a qualitative interview study of what is important to patients. BMJ Open, 2015, 5, e005406-e005406. | 1.9 | 69 |
| 5 | The Warwick Patient Experiences Framework: patient-based evidence in clinical guidelines. International Journal for Quality in Health Care, 2014, 26, 151-157. | 1.8 | 66 |
| 6 | The expressivist objection to prenatal testing: The experiences of families living with genetic disease. Social Science and Medicine, 2014, 107, 18-25. | 3.8 | 46 |
| 7 | Resilience as a response to the stigma of depression: A mixed methods analysis. Journal of Affective Disorders, 2011, 135, 267-276. | 4.1 | 39 |
| 8 | Becoming pregnant: exploring the perspectives of women living with diabetes. British Journal of General Practice, 2008, 58, 184-190. | 1.4 | 37 |
| 9 | Population screening for spinal muscular atrophy: A mixed methods study of the views of affected families. American Journal of Medical Genetics, Part A, 2017, 173, 421-434. | 1.2 | 34 |
| 10 | Accessing the field: Disability and the research process. Social Science and Medicine, 2011, 72, 23-30. | 3.8 | 33 |
| 11 | Knowledge is power? The role of experiential knowledge in genetically â€˜riskyâ€™ reproductive decisions. Sociology of Health and Illness, 2014, 36, 137-150. | 2.1 | 32 |
| 12 | How do genetically disabled adults view selective reproduction? Impairment, identity, and genetic screening. Molecular Genetics & Genomic Medicine, 2018, 6, 941-956. | 1.2 | 29 |
| 13 | What is a â€˜seriousâ€™ genetic condition? The perceptions of people living with genetic conditions. European Journal of Human Genetics, 2022, 30, 160-169. | 2.8 | 22 |
| 14 | Preventing lives affected by hemophilia: A mixed methods study of the views of adults with hemophilia and their families toward genetic screening. Molecular Genetics & Genomic Medicine, 2019, 7, e618. | 1.2 | 21 |
| 15 | Experience as knowledge: Disability, distillation and (reprogenetic) decision-making. Social Science and Medicine, 2017, 191, 186-193. | 3.8 | 20 |
| 16 | Newborn genetic screening for spinal muscular atrophy in the <scp>UK</scp>: The views of the general population. Molecular Genetics & Genomic Medicine, 2018, 6, 99-108. | 1.2 | 20 |
| 17 | Responsibility, identity, and genomic sequencing: A comparison of published recommendations and patient perspectives on accepting or declining incidental findings. Molecular Genetics & Genomic Medicine, 2018, 6, 1079-1096. | 1.2 | 18 |
| 18 | Social and cultural influences on genetic screening programme acceptability: A mixedâ€methods study of the views of adults, carriers, and family members living with thalassemia in the UK. Journal of Genetic Counseling, 2020, 29, 1026-1040. | 1.6 | 18 |

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|---|---|---|---|
| 19 | The effect of strategies of personal resilience on depression recovery in an Australian cohort: A mixed methods study. Health (United Kingdom), 2015, 19, 86-106. | 1.5 | 17 |
| 20 | Newborn screening for spinal muscular atrophy: The views of affected families and adults. American Journal of Medical Genetics, Part A, 2017, 173, 1546-1561. | 1.2 | 17 |
| 21 | The role of experiential knowledge within attitudes towards genetic carrier screening: A comparison of people with and without experience of spinal muscular atrophy. Health Expectations, 2018, 21, 201-211. | 2.6 | 17 |
| 22 | Impairment Experiences, Identity and Attitudes Towards Genetic Screening: the Views of People with Spinal Muscular Atrophy. Journal of Genetic Counseling, 2018, 27, 69-84. | 1.6 | 16 |
| 23 | Human genome editing and the identity politics of genetic disability. Journal of Community Genetics, 2020, 11, 125-127. | 1.2 | 15 |
| 24 | Which types of conditions should be included in reproductive genetic carrier screening? Views of parents of children with a genetic condition. European Journal of Medical Genetics, 2020, 63, 104075. | 1.3 | 14 |
| 25 | Attitudes toward population screening among people living with fragile X syndrome in the UK: â€˜I wouldnâ€™t wish him away, Iâ€™d just wish his fragile X syndrome awayâ€™. Journal of Genetic Counseling, 2021, 30, 85-97. | 1.6 | 11 |
| 26 | Experiential knowledge of disability, impairment and illness: The reproductive decisions of families genetically at risk. Health (United Kingdom), 2014, 18, 476-492. | 1.5 | 10 |
| 27 | â€œI didnâ€™t take it too seriously because Iâ€™d just never heard of itâ€¦ Experiential knowledge and genetic screening for thalassaemia in the UK. Journal of Genetic Counseling, 2019, 28, 141-154. | 1.6 | 9 |
| 28 | Absorbing it all: A meta-ethnography of parentsâ€™ unfolding experiences of newborn screening. Social Science and Medicine, 2021, 287, 114367. | 3.8 | 8 |
| 29 | Newborn screening for haemophilia: The views of families and adults living with haemophilia in the UK. Haemophilia, 2019, 25, 276-282. | 2.1 | 7 |
| 30 | â€˜We're kind of like genetic nomads': Parents' experiences of biographical disruption and uncertainty following in/conclusive results from newborn cystic fibrosis screening. Social Science and Medicine, 2022, 301, 114972. | 3.8 | 7 |
| 31 | EarLy Surveillance for Autoimmune diabetes: protocol for a qualitative study of general population and stakeholder perspectives on screening for type 1 diabetes in the UK (ELSA 1). BMJ Open Diabetes Research and Care, 2022, 10, e002750. | 2.8 | 6 |
| 32 | VP26 A Critical Appraisal Tool For Systematic Mixed Studies Reviews. International Journal of Technology Assessment in Health Care, 2018, 34, 166-166. | 0.5 | 5 |
| 33 | Children's perspectives and experiences of health, diet, physical activity and weight in an urban, multiâ€ethnic UK population: A qualitative study. Child: Care, Health and Development, 2021, 47, 597-607. | 1.7 | 4 |
| 34 | Expanding the notion of â€œbenefitâ€: comparing public, parent, and professional attitudes towards whole genome sequencing in newborns. New Genetics and Society, 2022, 41, 96-115. | 1.2 | 4 |
| 35 | Whose life is worth preserving? Disabled people and the expressivist objection to neonatology. Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 391-393. | 1.5 | 3 |
| 36 | Exploring trust in (bio)medical and experiential knowledge of birth: The perspectives of pregnant women, new mothers and maternity care providers. Midwifery, 2022, 107, 103272. | 2.3 | 3 |

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| 37 | Letter to the editor. Gene editing and disabled people: a response to lÃ±igo de Miguel Beriain. Journal of Community Genetics, 2020, 11, 245-247. | 1.2 | 2 |
| 38 | Enabling women to access preferred methods of contraception: a rapid review and behavioural analysis. BMC Public Health, 2021, 21, 2176. | 2.9 | 1 |