

Fiona Wood,, Ba, Reader

List of Publications by Year in descending order

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59
papers

1,392
citations

430874

18
h-index

377865

34
g-index

64
all docs

64
docs citations

64
times ranked

2027
citing authors

#	ARTICLE	IF	CITATIONS
1	â€Distributed health literacyâ€™: longitudinal qualitative analysis of the roles of health literacy mediators and social networks of people living with a longâ€™term health condition. Health Expectations, 2015, 18, 1180-1193.	2.6	256
2	Socially responsible antibiotic choices in primary care: a qualitative study of GPs' decisions to prescribe broad-spectrum and fluoroquinolone antibiotics. Family Practice, 2007, 24, 427-434.	1.9	96
3	Influences of cancer symptom knowledge, beliefs and barriers on cancer symptom presentation in relation to socioeconomic deprivation: a systematic review. BMC Cancer, 2015, 15, 1000.	2.6	87
4	Supporting shared decision making using an Option Grid for osteoarthritis of the knee in an interface musculoskeletal clinic: A stepped wedge trial. Patient Education and Counseling, 2016, 99, 571-577.	2.2	69
5	A multi-country qualitative study of clinicians' and patients' views on point of care tests for lower respiratory tract infection. Family Practice, 2011, 28, 661-669.	1.9	58
6	Protection by exclusion? The (lack of) inclusion of adults who lack capacity to consent to research in clinical trials in the UK. Trials, 2019, 20, 474.	1.6	54
7	Primary care clinicians' perceptions of antibiotic resistance: a multi-country qualitative interview study. Journal of Antimicrobial Chemotherapy, 2013, 68, 237-243.	3.0	51
8	Use of Mobile Devices to Help Cancer Patients Meet Their Information Needs in Non-Inpatient Settings: Systematic Review. JMIR MHealth and UHealth, 2018, 6, e10026.	3.7	47
9	A descriptive model of shared decision making derived from routine implementation in clinical practice (â€™Implement-SDMâ€™): Qualitative study. Patient Education and Counseling, 2019, 102, 1774-1785.	2.2	44
10	Coproduction and health: Public and cliniciansâ€™ perceptions of the barriers and facilitators. Health Expectations, 2019, 22, 93-101.	2.6	42
11	A Question of Balance: A Qualitative Study of Mothers' Interpretations of Dietary Recommendations. Annals of Family Medicine, 2010, 8, 51-57.	1.9	33
12	Doctorsâ€™ perspectives of informed consent for nonâ€™emergency surgical procedures: a qualitative interview study. Health Expectations, 2016, 19, 751-761.	2.6	33
13	What adolescents living with long-term conditions say about being involved in decision-making about their healthcare: A systematic review and narrative synthesis of preferences and experiences. Patient Education and Counseling, 2018, 101, 1725-1735.	2.2	28
14	â€™Itâ€™s a tough decisionâ€™: a qualitative study of proxy decision-making for research involving adults who lack capacity to consent in UK. Age and Ageing, 2019, 48, 903-909.	1.6	28
15	Probiotics for Antibiotic-Associated Diarrhoea (PAAD): a prospective observational study of antibiotic-associated diarrhoea (including Clostridium difficile-associated diarrhoea) in care homes. Health Technology Assessment, 2014, 18, 1-84.	2.8	27
16	Early impact of COVID-19 social distancing measures on reported sexual behaviour of HIV pre-exposure prophylaxis users in Wales. Sexually Transmitted Infections, 2021, 97, 85-87.	1.9	23
17	A pilot randomised controlled trial of community-led Antipsychotic Drug REduction for Adults with Learning Disabilities. Health Technology Assessment, 2017, 21, 1-92.	2.8	23
18	Consent, including advanced consent, of older adults to research in care homes: a qualitative study of stakeholdersâ€™ views in South Wales. Trials, 2013, 14, 247.	1.6	22

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19	Healthcare professionalsâ€™ understanding of the legislation governing research involving adults lacking mental capacity in England and Wales: a national survey. <i>Journal of Medical Ethics</i> , 2018, 44, 632-637.	1.8	21
20	Achieving online consent to participation in large-scale gene-environment studies: a tangible destination. <i>Journal of Medical Ethics</i> , 2011, 37, 487-492.	1.8	19
21	On a learning curve for shared decision making: Interviews with clinicians using the knee osteoarthritis Option Grid. <i>Journal of Evaluation in Clinical Practice</i> , 2018, 24, 56-64.	1.8	19
22	Research involving adults lacking capacity to consent: a content analysis of participant information sheets for consultees and legal representatives in England and Wales. <i>Trials</i> , 2019, 20, 233.	1.6	19
23	Perceived threat of COVID-19, attitudes towards vaccination, and vaccine hesitancy: A prospective longitudinal study in the UK. <i>British Journal of Health Psychology</i> , 2022, 27, 1354-1381.	3.5	19
24	Ethical understandings of proxy decision making for research involving adults lacking capacity: A systematic review (framework synthesis) of empirical research. <i>AJOB Empirical Bioethics</i> , 2018, 9, 267-286.	1.6	18
25	Patientsâ€™ views on the use of an Option Grid for knee osteoarthritis in physiotherapy clinical encounters: An interview study. <i>Health Expectations</i> , 2017, 20, 1302-1310.	2.6	16
26	Shifting mindsets: a realist synthesis of evidence from self-management support training. <i>Medical Education</i> , 2018, 52, 274-287.	2.1	16
27	Option Grids to facilitate shared decision making for patients with Osteoarthritis of the knee: protocol for a single site, efficacy trial. <i>BMC Health Services Research</i> , 2014, 14, 160.	2.2	15
28	Patients' opinions of the use of psychiatric case-finding questionnaires in general practice. <i>Health Expectations</i> , 2002, 5, 282-288.	2.6	14
29	What constitutes consent when parents and daughters have different views about having the HPV vaccine: qualitative interviews with stakeholders. <i>Journal of Medical Ethics</i> , 2011, 37, 466-471.	1.8	14
30	Development of a decision support intervention for family members of adults who lack capacity to consent to trials. <i>BMC Medical Informatics and Decision Making</i> , 2021, 21, 30.	3.0	13
31	The Preferences of Patients With Cancer Regarding Apps to Help Meet Their Illness-Related Information Needs: Qualitative Interview Study. <i>JMIR MHealth and UHealth</i> , 2019, 7, e14187.	3.7	13
32	Understanding sciatica: illness and treatment beliefs in a lumbar radicular pain population. A qualitative interview study. <i>BJGP Open</i> , 2019, 3, bjgpopen19X101654.	1.8	13
33	â€œI'd Like to Have More of a Say Because It's My Bodyâ€• Adolescents' Perceptions Around Barriers and Facilitators to Shared Decision-Making. <i>Journal of Adolescent Health</i> , 2019, 65, 633-642.	2.5	12
34	â€œWhat would you recommend doctor?â€• Discourse analysis of a moment of dissonance when sharing decisions in clinical consultations. <i>Health Expectations</i> , 2019, 22, 547-554.	2.6	12
35	Feasibility and acceptability of a cancer symptom awareness intervention for adults living in socioeconomically deprived communities. <i>BMC Public Health</i> , 2018, 18, 695.	2.9	11
36	Constructing authentic decisions: proxy decision making for research involving adults who lack capacity to consent. <i>Journal of Medical Ethics</i> , 2021, 47, e42-e42.	1.8	11

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37	Training in health coaching skills for health professionals who work with people with progressive neurological conditions: A realist evaluation. <i>Health Expectations</i> , 2020, 23, 919-933.	2.6	10
38	Recruitment and retention of participants from socioeconomically deprived communities: lessons from the Awareness and Beliefs About Cancer (ABACus3) Randomised Controlled Trial. <i>BMC Medical Research Methodology</i> , 2020, 20, 272.	3.1	9
39	Trials using deferred consent in the emergency setting: a systematic review and narrative synthesis of stakeholders' attitudes. <i>Trials</i> , 2022, 23, 411.	1.6	9
40	Working with interpreters: The challenges of introducing Option Grid patient decision aids. <i>Patient Education and Counseling</i> , 2017, 100, 456-464.	2.2	8
41	'There's more to life than money and health': Family caregivers' views on the role of Power of Attorney in proxy decisions about research participation for people living with dementia. <i>Dementia</i> , 2021, 20, 308-325.	2.0	8
42	Web-based physical activity intervention for people with progressive multiple sclerosis: application of consensus-based intervention development guidance. <i>BMJ Open</i> , 2021, 11, e045378.	1.9	7
43	Interventions to improve the self-management support health professionals provide for people with progressive neurological conditions: protocol for a realist synthesis. <i>BMJ Open</i> , 2017, 7, e014575.	1.9	6
44	Adapting the coping in deliberation (CODE) framework: A multi-method approach in the context of familial ovarian cancer risk management. <i>Patient Education and Counseling</i> , 2014, 97, 200-210.	2.2	5
45	Cohort profile: The UK COVID-19 Public Experiences (COPE) prospective longitudinal mixed-methods study of health and well-being during the SARS-CoV2 coronavirus pandemic. <i>PLoS ONE</i> , 2021, 16, e0258484.	2.5	5
46	Changing sexual behaviours amongst MSM during the COVID-19 restrictions in Wales: a mixed methods study. <i>BMC Public Health</i> , 2022, 22, 396.	2.9	5
47	Experiences of men who have sex with men when initiating, implementing and persisting with HIV pre-exposure prophylaxis. <i>Health Expectations</i> , 2022, 25, 1332-1341.	2.6	5
48	Monitoring and Managing Lifestyle Behaviors Using Wearable Activity Trackers: Mixed Methods Study of Views From the Huntington Disease Community. <i>JMIR Formative Research</i> , 2022, 6, e36870.	1.4	4
49	A critical discourse analysis of how public participants and their evidence are presented in health impact assessment reports in Wales. <i>Health Expectations</i> , 2019, 22, 585-593.	2.6	3
50	Lifestyle, exercise and activity package for people living with progressive multiple sclerosis (LEAP-MS): protocol for a single-arm feasibility study. <i>Pilot and Feasibility Studies</i> , 2021, 7, 111.	1.2	3
51	Can peer review help the marking experience?. <i>Medical Education</i> , 2005, 39, 1156-1157.	2.1	2
52	Doing qualitative health services research remotely. <i>Communication and Medicine</i> , 2022, 17, 194-198.	0.2	2
53	Barriers and facilitators to the use of personal information documents in health and social care settings for people living with dementia: A thematic synthesis and mapping to the COMEB framework. <i>Health Expectations</i> , 2022, , .	2.6	2
54	Borna disease virus: The generation and review of a scientific study. <i>Social Science and Medicine</i> , 2006, 63, 1072-1083.	3.8	1

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55	Conducting focus groups in neurodegenerative disease populations: ethical and methodological considerations. <i>BMJ Open</i> , 2021, 11, e041869.	1.9	1
56	Cancer Clinicians's Views Regarding an App That Helps Patients With Cancer Meet Their Information Needs: Qualitative Interview Study. <i>JMIR Cancer</i> , 2021, 7, e23671.	2.4	1
57	147. Patient Perspectives on the Option Grid for Osteoarthritis of the Knee. <i>Rheumatology</i> , 2015, , .	1.9	0
58	F33...Perceptions, motivators and barriers to the acceptance of wearable activity trackers in people with huntington's disease. , 2021, , .		0
59	A web-based Life-style, Exercise and Activity intervention for People with Progressive Multiple Sclerosis: Results of a Single-Arm Feasibility Study. <i>Multiple Sclerosis and Related Disorders</i> , 2021, 57, 103388.	2.0	0