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List of Publications by Year in descending order

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Version: 2024-02-01

59
papers

1,392
citations

430442

18
h-index

377514

34
g-index

64
all docs

64
docs citations

64
times ranked

2027
citing authors

#	ARTICLE	IF	CITATIONS
1	Changing sexual behaviours amongst MSM during the COVID-19 restrictions in Wales: a mixed methods study. BMC Public Health, 2022, 22, 396.	1.2	5
2	Doing qualitative health services research remotely. Communication and Medicine, 2022, 17, 194-198.	0.1	2
3	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 COMaEB framework. Health Expectations, 2022, , .	1.1	2
4	Experiences of men who have sex with men when initiating, implementing and persisting with HIV pre-exposure prophylaxis. Health Expectations, 2022, 25, 1332-1341.	1.1	5
5	Trials using deferred consent in the emergency setting: a systematic review and narrative synthesis of stakeholders' attitudes. Trials, 2022, 23, 411.	0.7	9
6	Perceived threat of COVID-19, attitudes towards vaccination, and vaccine hesitancy: A prospective longitudinal study in the UK. British Journal of Health Psychology, 2022, 27, 1354-1381.	1.9	19
7	Monitoring and Managing Lifestyle Behaviors Using Wearable Activity Trackers: Mixed Methods Study of Views From the Huntington Disease Community. JMIR Formative Research, 2022, 6, e36870.	0.7	4
8	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.	0.8	23
9	"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. Dementia, 2021, 20, 308-325.	1.0	8
10	Development of a decision support intervention for family members of adults who lack capacity to consent to trials. BMC Medical Informatics and Decision Making, 2021, 21, 30.	1.5	13
11	Conducting focus groups in neurodegenerative disease populations: ethical and methodological considerations. BMJ Open, 2021, 11, e041869.	0.8	1
12	Web-based physical activity intervention for people with progressive multiple sclerosis: application of consensus-based intervention development guidance. BMJ Open, 2021, 11, e045378.	0.8	7
13	Cancer Clinicians' Views Regarding an App That Helps Patients With Cancer Meet Their Information Needs: Qualitative Interview Study. JMIR Cancer, 2021, 7, e23671.	0.9	1
14	Lifestyle, exercise and activity package for people living with progressive multiple sclerosis (LEAP-MS): protocol for a single-arm feasibility study. Pilot and Feasibility Studies, 2021, 7, 111.	0.5	3
15	F33...Perceptions, motivators and barriers to the acceptance of wearable activity trackers in people with huntington's disease. , 2021, , .		0
16	Cohort profile: The UK COVID-19 Public Experiences (COPE) prospective longitudinal mixed-methods study of health and well-being during the SARSCoV2 coronavirus pandemic. PLoS ONE, 2021, 16, e0258484.	1.1	5
17	A web-based Life-style, Exercise and Activity intervention for People with Progressive Multiple Sclerosis: Results of a Single-Arm Feasibility Study. Multiple Sclerosis and Related Disorders, 2021, 57, 103388.	0.9	0
18	Constructing authentic decisions: proxy decision making for research involving adults who lack capacity to consent. Journal of Medical Ethics, 2021, 47, e42-e42.	1.0	11

#	ARTICLE	IF	CITATIONS
19	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.	1.4	9
20	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.	1.1	10
21	Protection by exclusion? The (lack of) inclusion of adults who lack capacity to consent to research in clinical trials in the UK. <i>Trials</i> , 2019, 20, 474.	0.7	54
22	‘I’d Like to Have More of a Say Because It’s My Body’: Adolescents’ Perceptions Around Barriers and Facilitators to Shared Decision-Making. <i>Journal of Adolescent Health</i> , 2019, 65, 633-642.	1.2	12
23	A descriptive model of shared decision making derived from routine implementation in clinical practice (‘Implement-SDM’): Qualitative study. <i>Patient Education and Counseling</i> , 2019, 102, 1774-1785.	1.0	44
24	‘It’s a tough decision’: a qualitative study of proxy decision-making for research involving adults who lack capacity to consent in UK. <i>Age and Ageing</i> , 2019, 48, 903-909.	0.7	28
25	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.	1.1	3
26	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.	0.7	19
27	‘What would you recommend doctor?’ Discourse analysis of a moment of dissonance when sharing decisions in clinical consultations. <i>Health Expectations</i> , 2019, 22, 547-554.	1.1	12
28	Coproduction and health: Public and clinicians’ perceptions of the barriers and facilitators. <i>Health Expectations</i> , 2019, 22, 93-101.	1.1	42
29	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.	1.8	13
30	Understanding sciatica: illness and treatment beliefs in a lumbar radicular pain population. A qualitative interview study. <i>BJGP Open</i> , 2019, 3, bjgpopen19X101654.	0.9	13
31	Shifting mindsets: a realist synthesis of evidence from self-management support training. <i>Medical Education</i> , 2018, 52, 274-287.	1.1	16
32	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.	0.9	19
33	Healthcare professionals’ 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.	1.0	21
34	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.	0.8	18
35	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. <i>Patient Education and Counseling</i> , 2018, 101, 1725-1735.	1.0	28
36	Feasibility and acceptability of a cancer symptom awareness intervention for adults living in socioeconomically deprived communities. <i>BMC Public Health</i> , 2018, 18, 695.	1.2	11

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37	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.	1.8	47
38	Interventions to improve the self-management support health professionals provide for people with progressive neurological conditions: protocol for a realist synthesis. BMJ Open, 2017, 7, e014575.	0.8	6
39	Patients's views on the use of an Option Grid for knee osteoarthritis in physiotherapy clinical encounters: An interview study. Health Expectations, 2017, 20, 1302-1310.	1.1	16
40	Working with interpreters: The challenges of introducing Option Grid patient decision aids. Patient Education and Counseling, 2017, 100, 456-464.	1.0	8
41	A pilot randomised controlled trial of community-led Antipsychotic Drug Reduction for Adults with Learning Disabilities. Health Technology Assessment, 2017, 21, 1-92.	1.3	23
42	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.	1.0	69
43	Doctors' perspectives of informed consent for non-emergency surgical procedures: a qualitative interview study. Health Expectations, 2016, 19, 751-761.	1.1	33
44	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.	1.1	87
45	147. Patient Perspectives on the Option Grid for Osteoarthritis of the Knee. Rheumatology, 2015, , .	0.9	0
46	"Distributed health literacy": longitudinal qualitative analysis of the roles of health literacy mediators and social networks of people living with a long-term health condition. Health Expectations, 2015, 18, 1180-1193.	1.1	256
47	Adapting the coping in deliberation (CODE) framework: A multi-method approach in the context of familial ovarian cancer risk management. Patient Education and Counseling, 2014, 97, 200-210.	1.0	5
48	Option Grids to facilitate shared decision making for patients with Osteoarthritis of the knee: protocol for a single site, efficacy trial. BMC Health Services Research, 2014, 14, 160.	0.9	15
49	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.	1.3	27
50	Consent, including advanced consent, of older adults to research in care homes: a qualitative study of stakeholders' views in South Wales. Trials, 2013, 14, 247.	0.7	22
51	Primary care clinicians' perceptions of antibiotic resistance: a multi-country qualitative interview study. Journal of Antimicrobial Chemotherapy, 2013, 68, 237-243.	1.3	51
52	Achieving online consent to participation in large-scale gene-environment studies: a tangible destination. Journal of Medical Ethics, 2011, 37, 487-492.	1.0	19
53	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.	0.8	58
54	What constitutes consent when parents and daughters have different views about having the HPV vaccine: qualitative interviews with stakeholders. Journal of Medical Ethics, 2011, 37, 466-471.	1.0	14

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55	A Question of Balance: A Qualitative Study of Mothers' Interpretations of Dietary Recommendations. <i>Annals of Family Medicine</i> , 2010, 8, 51-57.	0.9	33
56	Socially responsible antibiotic choices in primary care: a qualitative study of GPs' decisions to prescribe broad-spectrum and fluoroquinolone antibiotics. <i>Family Practice</i> , 2007, 24, 427-434.	0.8	96
57	Borna disease virus: The generation and review of a scientific study. <i>Social Science and Medicine</i> , 2006, 63, 1072-1083.	1.8	1
58	Can peer review help the marking experience?. <i>Medical Education</i> , 2005, 39, 1156-1157.	1.1	2
59	Patients' opinions of the use of psychiatric case-finding questionnaires in general practice. <i>Health Expectations</i> , 2002, 5, 282-288.	1.1	14