

Vikki A Entwistle

List of Publications by Year in descending order

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Version: 2024-02-01

179
papers

13,547
citations

47006

47
h-index

27406

106
g-index

183
all docs

183
docs citations

183
times ranked

15303
citing authors

#	ARTICLE	IF	CITATIONS
1	What is an adequate sample size? Operationalising data saturation for theory-based interview studies. <i>Psychology and Health</i> , 2010, 25, 1229-1245.	2.2	1,833
2	Developing a quality criteria framework for patient decision aids: online international Delphi consensus process. <i>BMJ: British Medical Journal</i> , 2006, 333, 417-0.	2.3	1,373
3	Decision aids for patients facing health treatment or screening decisions: systematic review. <i>BMJ: British Medical Journal</i> , 1999, 319, 731-734.	2.3	718
4	What influences recruitment to randomised controlled trials? A review of trials funded by two UK funding agencies. <i>Trials</i> , 2006, 7, 9.	1.6	712
5	Sharing decisions with patients: is the information good enough?. <i>BMJ: British Medical Journal</i> , 1999, 318, 318-322.	2.3	519
6	Decision aids for people facing health treatment or screening decisions. , 2009, , CD001431.		481
7	Decision aids for people facing health treatment or screening decisions. , 2003, , CD001431.		388
8	Supporting Patient Autonomy: The Importance of Clinician-patient Relationships. <i>Journal of General Internal Medicine</i> , 2010, 25, 741-745.	2.6	339
9	Personalised care planning for adults with chronic or long-term health conditions. <i>The Cochrane Library</i> , 2015, , CD010523.	2.8	329
10	Interventions for providers to promote a patient-centred approach in clinical consultations. , 2001, , CD003267.		324
11	Lay perspectives: advantages for health research. <i>BMJ: British Medical Journal</i> , 1998, 316, 463-466.	2.3	287
12	Treating Patients as Persons: A Capabilities Approach to Support Delivery of Person-Centered Care. <i>American Journal of Bioethics</i> , 2013, 13, 29-39.	0.9	276
13	Do Patient Decision Aids Meet Effectiveness Criteria of the International Patient Decision Aid Standards Collaboration? A Systematic Review and Meta-analysis. <i>Medical Decision Making</i> , 2007, 27, 554-574.	2.4	237
14	Patient involvement in treatment decision-making: The case for a broader conceptual framework. <i>Patient Education and Counseling</i> , 2006, 63, 268-278.	2.2	224
15	Reasons for participating in randomised controlled trials: conditional altruism and considerations for self. <i>Trials</i> , 2010, 11, 31.	1.6	219
16	Three questions that patients can ask to improve the quality of information physicians give about treatment options: A cross-over trial. <i>Patient Education and Counseling</i> , 2011, 84, 379-385.	2.2	189
17	Evidence-Informed Patient Choice: Practical Issues of Involving Patients in Decisions About Health Care Technologies. <i>International Journal of Technology Assessment in Health Care</i> , 1998, 14, 212-225.	0.5	163
18	Rationalising the "irrational": a think aloud study of discrete choice experiment responses. <i>Health Economics (United Kingdom)</i> , 2009, 18, 321-336.	1.7	163

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19	Health in the 'hidden population' of people with low literacy. A systematic review of the literature. BMC Public Health, 2010, 10, 459.	2.9	122
20	Which Experiences of Health Care Delivery Matter to Service Users and Why? A Critical Interpretive Synthesis and Conceptual Map. Journal of Health Services Research and Policy, 2012, 17, 70-78.	1.7	112
21	How information about other people's personal experiences can help with healthcare decision-making: A qualitative study. Patient Education and Counseling, 2011, 85, e291-e298.	2.2	110
22	“Unilateral” and “bilateral” practitioner approaches in decision-making about treatment. Social Science and Medicine, 2005, 61, 2611-2627.	3.8	109
23	A Systematic Review of the Effectiveness of Health Service Interventions Aimed at Reducing Inequalities in Health. Journal of Health Services Research and Policy, 1996, 1, 93-103.	1.7	107
24	How the stigma of low literacy can impair patient-professional spoken interactions and affect health: insights from a qualitative investigation. BMC Health Services Research, 2013, 13, 319.	2.2	100
25	Shared decision making: trade-offs between narrower and broader conceptions. Health Expectations, 2011, 14, 210-219.	2.6	96
26	Trust in the health-care provider-patient relationship: a systematic mapping review of the evidence base. International Journal for Quality in Health Care, 2013, 25, 682-688.	1.8	95
27	Evaluating Interventions to Promote Patient Involvement in Decision-Making: By What Criteria Should Effectiveness be Judged?. Journal of Health Services Research and Policy, 1998, 3, 100-107.	1.7	92
28	Advising Patients About Patient Safety: Current Initiatives Risk Shifting Responsibility. Joint Commission Journal on Quality and Patient Safety, 2005, 31, 483-494.	0.7	91
29	Patient choice modules for summaries of clinical effectiveness: a proposal. BMJ: British Medical Journal, 2001, 322, 664-667.	2.3	88
30	“Avoiding harm to others” considerations in relation to parental measles, mumps and rubella (MMR) vaccination discussions “ An analysis of an online chat forum. Social Science and Medicine, 2008, 67, 1382-1390.	3.8	85
31	Involvement in treatment decision-making: Its meaning to people with diabetes and implications for conceptualisation. Social Science and Medicine, 2008, 66, 362-375.	3.8	81
32	Promoting research participation: Why not advertise altruism?. Social Science and Medicine, 2008, 66, 1451-1456.	3.8	81
33	Speaking up about safety concerns: multi-setting qualitative study of patients' views and experiences. BMJ Quality and Safety, 2010, 19, e33-e33.	3.7	79
34	We need to talk about purpose: a critical interpretive synthesis of health and social care professionals’ approaches to self-management support for people with long-term conditions. Health Expectations, 2017, 20, 243-259.	2.6	76
35	Can consumers learn to ask three questions to improve shared decision making? A feasibility study of the <sc>ASK</sc> (AskShareKnow) Patient-Clinician Communication Model [®] intervention in a primary health-care setting. Health Expectations, 2016, 19, 1160-1168.	2.6	75
36	Scoping Review and Approach to Appraisal of Interventions Intended to Involve Patients in Patient Safety. Journal of Health Services Research and Policy, 2010, 15, 17-25.	1.7	74

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37	Addressing complex healthcare problems in diverse settings: Insights from activity theory. <i>Social Science and Medicine</i> , 2012, 74, 305-312.	3.8	73
38	Communicating about screening. <i>BMJ: British Medical Journal</i> , 2008, 337, a1591-a1591.	2.3	73
39	Effectiveness of interventions designed to promote patient involvement to enhance safety: a systematic review. <i>BMJ Quality and Safety</i> , 2010, 19, e10-e10.	3.7	72
40	Barriers and facilitators to smoking cessation in a cancer context: A qualitative study of patient, family and professional views. <i>BMC Cancer</i> , 2017, 17, 348.	2.6	66
41	Decisions about treatment: interpretations of two measures of control by women having a hysterectomy. <i>Social Science and Medicine</i> , 2001, 53, 721-732.	3.8	62
42	A review of health literacy: Definitions, interpretations, and implications for policy initiatives. <i>Journal of Public Health Policy</i> , 2016, 37, 334-352.	2.0	62
43	Toward estimating the impact of changes in immigrants' insurance eligibility on hospital expenditures for uncompensated care. <i>BMC Health Services Research</i> , 2003, 3, 1.	2.2	60
44	Assessing patients'™ participation and quality of decision-making: insights from a study of routine practice in diverse settings. <i>Patient Education and Counseling</i> , 2004, 55, 105-113.	2.2	59
45	The requirement for prior consent to participate on survey response rates: a population-based survey in Grampian. <i>BMC Health Services Research</i> , 2003, 3, 21.	2.2	58
46	The TRIO Framework: Conceptual insights into family caregiver involvement and influence throughout cancer treatment decision-making. <i>Patient Education and Counseling</i> , 2017, 100, 2035-2046.	2.2	57
47	Why Health and Social Care Support for People with Long-Term Conditions Should be Oriented Towards Enabling Them to Live Well. <i>Health Care Analysis</i> , 2018, 26, 48-65.	2.2	57
48	Developing Information Materials to Present the Findings of Technology Assessments to Consumers: <i>The Experience of the NHS Centre for Reviews and Dissemination</i>. <i>International Journal of Technology Assessment in Health Care</i> , 1998, 14, 47-70.	0.5	56
49	Enhancing Citizen Engagement in Cancer Screening Through Deliberative Democracy. <i>Journal of the National Cancer Institute</i> , 2013, 105, 380-386.	6.3	56
50	Trust in the context of patient safety problems. <i>Journal of Health Organization and Management</i> , 2006, 20, 397-416.	1.3	51
51	Empirical relationships between health literacy and treatment decision making: A scoping review of the literature. <i>Patient Education and Counseling</i> , 2015, 98, 296-309.	2.2	51
52	Consumer involvement in decisions about what health-related research is funded. <i>Health Policy</i> , 2004, 70, 281-290.	3.0	50
53	The significance for decision-making of information that is not exchanged by patients and health professionals during consultations. <i>Social Science and Medicine</i> , 2006, 63, 2065-2078.	3.8	47
54	Managing demand: A patient led NHS: managing demand at the interface between lay and primary care. <i>BMJ: British Medical Journal</i> , 1998, 316, 1816-1819.	2.3	46

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55	Recruitment to clinical trials: a meta-ethnographic synthesis of studies of reasons for participation. <i>Journal of Health Services Research and Policy</i> , 2013, 18, 233-241.	1.7	44
56	Shared decision-making: enhancing the clinical relevance. <i>Journal of the Royal Society of Medicine</i> , 2012, 105, 416-421.	2.0	43
57	Offering informed choice about breast screening. <i>Journal of Medical Screening</i> , 2014, 21, 194-200.	2.3	43
58	Ethical Justifications for Access to Unapproved Medical Interventions: An Argument for (Limited) Patient Obligations. <i>American Journal of Bioethics</i> , 2014, 14, 3-15.	0.9	42
59	Supporting consumer involvement in decision making: what constitutes quality in consumer health information?. <i>International Journal for Quality in Health Care</i> , 1996, 8, 425.	1.8	42
60	Consumer involvement in research projects: the activities of research funders. <i>Health Policy</i> , 2004, 69, 229-238.	3.0	41
61	Differing perspectives on patient involvement in patient safety. <i>Quality and Safety in Health Care</i> , 2007, 16, 82-83.	2.5	41
62	Marketing and clinical trials: a case study. <i>Trials</i> , 2007, 8, 37.	1.6	41
63	Women's perceptions of decision-making about hysterectomy. <i>BJOG: an International Journal of Obstetrics and Gynaecology</i> , 2004, 111, 133-142.	2.3	39
64	Which surgical decisions should patients participate in and how? Reflections on women's recollections of discussions about variants of hysterectomy. <i>Social Science and Medicine</i> , 2006, 62, 499-509.	3.8	39
65	Developing a placebo-controlled trial in surgery: Issues of design, acceptability and feasibility. <i>Trials</i> , 2011, 12, 50.	1.6	38
66	Supporting positive experiences and sustained participation in clinical trials: looking beyond information provision. <i>Journal of Medical Ethics</i> , 2012, 38, 751-756.	1.8	38
67	Use of a Birth Plan within Woman-held Maternity Records: A Qualitative Study with Women and Staff in Northeast Scotland. <i>Birth</i> , 2014, 41, 283-289.	2.2	38
68	"The more you know, the more you realise it is really challenging to do" Tensions and uncertainties in person-centred support for people with long-term conditions. <i>Patient Education and Counseling</i> , 2018, 101, 1460-1467.	2.2	38
69	Health and medical coverage in the UK national press. <i>Public Understanding of Science</i> , 1992, 1, 367-382.	2.8	37
70	How personal experiences feature in women's accounts of use of information for decisions about antenatal diagnostic testing for foetal abnormality. <i>Social Science and Medicine</i> , 2011, 72, 755-762.	3.8	36
71	The value of different aspects of person-centred care: a series of discrete choice experiments in people with long-term conditions. <i>BMJ Open</i> , 2017, 7, e015689.	1.9	36
72	Media coverage of the Child B case. <i>BMJ: British Medical Journal</i> , 1996, 312, 1587-1591.	2.3	36

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73	Valuing patients' experiences of healthcare processes: Towards broader applications of existing methods. <i>Social Science and Medicine</i> , 2014, 106, 194-203.	3.8	35
74	Provision of cancer information as a "support for navigating the knowledge landscape": Findings from a critical interpretive literature synthesis. <i>European Journal of Oncology Nursing</i> , 2013, 17, 360-369.	2.1	33
75	Regulatory withdrawal of medicines marketed with uncertain benefits: the bevacizumab case study. <i>Journal of Pharmaceutical Policy and Practice</i> , 2015, 8, 25.	2.4	33
76	Trust relations in health care: an agenda for future research. <i>Journal of Health Organization and Management</i> , 2006, 20, 477-484.	1.3	32
77	Virtue, progress and practice. <i>Journal of Evaluation in Clinical Practice</i> , 2011, 17, 839-846.	1.8	31
78	Communicating good care: A qualitative study of what people with urological cancer value in interactions with health care providers. <i>European Journal of Oncology Nursing</i> , 2014, 18, 35-40.	2.1	31
79	Disseminating information about healthcare effectiveness: a survey of consumer health information services. <i>Quality and Safety in Health Care</i> , 1998, 7, 124-129.	2.5	29
80	Interventions for improving communication with children and adolescents about their cancer. , 2003, , CD002969.		29
81	Recordings or summaries of consultations for people with cancer. , 2003, , CD001539.		29
82	Consumer involvement in setting the health services research agenda: Persistent questions of value. <i>Journal of Health Services Research and Policy</i> , 2008, 13, 76-81.	1.7	28
83	Vaginal birth after caesarean section: why is uptake so low? Insights from a meta-ethnographic synthesis of women's accounts of their birth choices. <i>BMJ Open</i> , 2016, 6, e008881.	1.9	28
84	Ethical, Legal, and Social Issues in Health Technology Assessment for Prenatal/Preconceptional and Newborn Screening: A Workshop Report. <i>Public Health Genomics</i> , 2009, 12, 4-10.	1.0	25
85	General Practitioners'™ Experiences of, and Responses to, Uncertainty in Prostate Cancer Screening: Insights from a Qualitative Study. <i>PLoS ONE</i> , 2016, 11, e0153299.	2.5	25
86	Guidance for considering ethical, legal, and social issues in health technology assessment: Application to genetic screening. <i>International Journal of Technology Assessment in Health Care</i> , 2008, 24, 412-422.	0.5	24
87	Enabling mutual helping? Examining variable needs for facilitated peer support. <i>Patient Education and Counseling</i> , 2011, 85, e120-e125.	2.2	24
88	Giving tape recordings or written summaries of consultations to people with cancer: a systematic review. <i>Health Expectations</i> , 2001, 4, 162-169.	2.6	23
89	Nursing shortages and patient safety problems in hospital care: is clinical monitoring by families part of the solution?. <i>Health Expectations</i> , 2004, 7, 1-5.	2.6	23
90	Relational conceptions of paternalism: a way to rebut nanny-state accusations and evaluate public health interventions. <i>Public Health</i> , 2015, 129, 1021-1029.	2.9	23

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91	Shared decision making in the UK: Moving towards wider uptake. <i>Zeitschrift Fur Evidenz, Fortbildung Und Qualitat Im Gesundheitswesen</i> , 2017, 123-124, 99-103.	0.9	23
92	Exploring the impact and use of patients' feedback about their care experiences in general practice settings—a realist synthesis. <i>Family Practice</i> , 2018, 35, 13-21.	1.9	23
93	Researching experiences of cancer: the importance of methodology. <i>European Journal of Cancer Care</i> , 2002, 11, 232-237.	1.5	22
94	Financial considerations in the conduct of multi-centre randomised controlled trials: evidence from a qualitative study. <i>Trials</i> , 2006, 7, 34.	1.6	22
95	Revisiting the equity debate in COVID-19: ICU is no panacea. <i>Journal of Medical Ethics</i> , 2020, 46, 641-645.	1.8	22
96	Trust and shared decision-making: an emerging research agenda. <i>Health Expectations</i> , 2004, 7, 271-273.	2.6	21
97	Evaluation of interventions for informed consent for randomised controlled trials (ELICIT): protocol for a systematic review of the literature and identification of a core outcome set using a Delphi survey. <i>Trials</i> , 2015, 16, 484.	1.6	21
98	Should women aged 70–74 be invited to participate in screening mammography? A report on two Australian community juries. <i>BMJ Open</i> , 2018, 8, e021174.	1.9	21
99	Supporting and resourcing treatment decision-making: some policy considerations. <i>Health Expectations</i> , 2000, 3, 77-85.	2.6	20
100	Using the theory of planned behaviour to develop targets for interventions to enhance patient communication during pharmacy consultations for non-prescription medicines. <i>International Journal of Pharmacy Practice</i> , 2014, 22, 386-396.	0.6	20
101	Doctors' perspectives on PSA testing illuminate established differences in prostate cancer screening rates between Australia and the UK: a qualitative study. <i>BMJ Open</i> , 2016, 6, e011932.	1.9	20
102	Ethical issues raised by thyroid cancer overdiagnosis: A matter for public health?. <i>Bioethics</i> , 2017, 31, 590-598.	1.4	20
103	The case of Norplant as an example of media coverage over the life of a new health technology. <i>Lancet, The</i> , 2000, 355, 1633-1636.	13.7	19
104	The potential contribution of decision aids to screening programmes. <i>Health Expectations</i> , 2001, 4, 109-115.	2.6	19
105	Health professionals' experiences and perspectives on food insecurity and long-term conditions: A qualitative investigation. <i>Health and Social Care in the Community</i> , 2020, 28, 404-413.	1.6	19
106	Health literacy: the need to consider images as well as words. <i>Health Expectations</i> , 2008, 11, 99-101.	2.6	18
107	Addressing Deficits and Injustices: The Potential Epistemic Contributions of Patients to Research. <i>Health Care Analysis</i> , 2017, 25, 386-403.	2.2	18
108	Devolution and Patient Choice: Policy Rhetoric versus Experience in Practice. <i>Social Policy and Administration</i> , 2012, 46, 199-218.	3.0	16

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109	Reason and value: making reasoning fit for practice. <i>Journal of Evaluation in Clinical Practice</i> , 2012, 18, 929-937.	1.8	15
110	An exploration of the implementation of open disclosure of adverse events in the UK: a scoping review and qualitative exploration. <i>Health Services and Delivery Research</i> , 2014, 2, 1-196.	1.4	15
111	Patients's experiences with partial dentures: a qualitative study. <i>Gerodontology</i> , 2005, 22, 187-192.	2.0	14
112	The information and support needs of patients discharged after a short hospital stay for treatment of low-risk Community Acquired Pneumonia: implications for treatment without admission. <i>BMC Pulmonary Medicine</i> , 2008, 8, 11.	2.0	14
113	Hurtful comments are harmful comments: respectful communication is not just an optional extra in healthcare. <i>Health Expectations</i> , 2008, 11, 319-320.	2.6	14
114	A meta-ethnography of organisational culture in primary care medical practice. <i>Journal of Health Organization and Management</i> , 2014, 28, 21-40.	1.3	14
115	Informing, communicating and sharing decisions with people who have cancer. <i>Quality in Health Care: QHC</i> , 2001, 10, 193-196.	1.2	14
116	Public involvement in health service governance and development: questions of potential for influence. <i>Health Expectations</i> , 2009, 12, 1-3.	2.6	12
117	Shared Health Governance: The Potential Danger of Oppressive "Healthism". <i>American Journal of Bioethics</i> , 2011, 11, 57-59.	0.9	12
118	A Capabilities Approach to Person-Centered Care: Response to Open Peer Commentaries on "Treating Patients as Persons: A Capabilities Approach to Support Delivery of Person-Centered Care". <i>American Journal of Bioethics</i> , 2013, 13, W1-W4.	0.9	12
119	"Was that a success or not a success?" a qualitative study of health professionals' perspectives on support for people with long-term conditions. <i>BMC Family Practice</i> , 2017, 18, 39.	2.9	12
120	Judging journalism: how should the quality of news reporting about clinical interventions be assessed and improved?. <i>Quality and Safety in Health Care</i> , 1999, 8, 172-176.	2.5	11
121	Ethical tensions associated with the promotion of public health policy in health visiting: a qualitative investigation of health visitors' views. <i>Primary Health Care Research and Development</i> , 2013, 14, 200-211.	1.2	11
122	Being open about unanticipated problems in health care: the challenges of uncertainties. <i>Journal of Health Services Research and Policy</i> , 2015, 20, 54-60.	1.7	11
123	What does "quality" add? Towards an ethics of healthcare improvement. <i>Journal of Medical Ethics</i> , 2020, 46, 118-122.	1.8	11
124	Talking it better: conversations and normative complexity in healthcare improvement. <i>Medical Humanities</i> , 2022, 48, 85-93.	1.2	11
125	Public opinion on systems for feeding back views to the National Health Service. <i>Quality and Safety in Health Care</i> , 2003, 12, 435-442.	2.5	10
126	Interventions for improving communication with children and adolescents about a family member's cancer. <i>The Cochrane Library</i> , 2003, , CD004511.	2.8	10

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127	Supporting participation in clinical research: decision aids for trial recruitment?. Health Expectations, 2008, 11, 205-207.	2.6	10
128	Primary goals, information-giving and men's understanding: a qualitative study of Australian and UK doctors' varied communication about PSA screening. BMJ Open, 2018, 8, e018009.	1.9	10
129	Implementing shared decision-making in UK: Progress 2017-2022. Zeitschrift Fur Evidenz, Fortbildung Und Qualitat Im Gesundheitswesen, 2022, 171, 139-143.	0.9	10
130	Risk, Overdiagnosis and Ethical Justifications. Health Care Analysis, 2019, 27, 231-248.	2.2	9
131	Some Unresolved Ethical Challenges in Healthcare Decision-Making: Navigating Family Involvement. Asian Bioethics Review, 2020, 12, 27-36.	1.3	9
132	What, how, when and who of trial results summaries for trial participants: stakeholder-informed guidance from the RECAP project. BMJ Open, 2022, 12, e057019.	1.9	9
133	Informed choice initiative: an example of reaching users with evidence-based information. Journal of Clinical Effectiveness, 1996, 1, 143-145.	0.2	8
134	Protocol for stage 1 of the GaP study (Genetic testing acceptability for Paget's disease of bone): an interview study about genetic testing and preventive treatment: would relatives of people with Paget's disease want testing and treatment if they were available?. BMC Health Services Research, 2006, 6, 71.	2.2	8
135	Considering "balance" in information. Health Expectations, 2007, 10, 307-308.	2.6	8
136	Reframing the Debate Around State Responses to Infertility: Considering the Harms of Subfertility and Involuntary Childlessness. Public Health Ethics, 2016, 9, 290-300.	1.0	8
137	Vagueness and variety in person-centred care. Wellcome Open Research, 0, 7, 170.	1.8	8
138	Information to facilitate patient involvement in decision-making " some issues. Journal of Clinical Effectiveness, 1997, 2, 69-72.	0.2	7
139	Research funding organisations and consumer involvement. Journal of Health Services Research and Policy, 2003, 8, 129-131.	1.7	7
140	Involving Patients in Their Care. Current Breast Cancer Reports, 2014, 6, 211-218.	1.0	7
141	Editorial. Health Expectations, 1999, 2, 75-77.	2.6	6
142	Health visitor professional education and post-qualification clinical supervision: how well does it equip practitioners for dealing with ethical tensions associated with promoting the public health agenda to individual clients?. Primary Health Care Research and Development, 2013, 14, 90-102.	1.2	6
143	How should the "privilege" in therapeutic privilege be conceived when considering the decision-making process for patients with borderline capacity?. Journal of Medical Ethics, 2021, 47, 47-50.	1.8	6
144	Letter to the Editor. Patient Education and Counseling, 2001, 42, 295-296.	2.2	5

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145	Considerations of 'fit' and patient involvement in decision making. <i>Health Expectations</i> , 2006, 9, 95-97.	2.6	5
146	Defining What is Good: Pluralism and Healthcare Quality. <i>Kennedy Institute of Ethics Journal</i> , 2019, 29, 367-388.	0.5	5
147	The Patient Activation through Community Empowerment/Engagement for Diabetes Management (PACE-D) protocol: a non-randomised controlled trial of personalised care and support planning for persons living with diabetes. <i>BMC Family Practice</i> , 2020, 21, 114.	2.9	5
148	Understanding and investigating relationality in the capability approach. <i>Journal for the Theory of Social Behaviour</i> , 2022, 52, 86-104.	1.2	5
149	An international core outcome set for evaluating interventions to improve informed consent to clinical trials: The ELICIT Study. <i>Journal of Clinical Epidemiology</i> , 2021, 137, 14-22.	5.0	5
150	Made to Measure: The Ethics of Routine Measurement for Healthcare Improvement. <i>Health Care Analysis</i> , 2021, 29, 39-58.	2.2	5
151	Editorial. <i>Health Expectations</i> , 1999, 2, 1-2.	2.6	4
152	Placing evidence in context: A response to Fry's commentary. <i>Social Science and Medicine</i> , 2008, 66, 1461-1462.	3.8	4
153	Involving service users in qualitative analysis: approaches and assessment. <i>Health Expectations</i> , 2010, 13, 111-112.	2.6	4
154	Unifying and Universalising Personalised Care? An Analysis of a National Curriculum With Implications for Policy and Education Relating to Person-centred Care. <i>Patient Education and Counseling</i> , 2022, , .	2.2	4
155	Editorial. <i>Health Expectations</i> , 2000, 3, 87-89.	2.6	3
156	Who are your public? A survey comparing the views of a population-based sample with those of a community-based public forum in Scotland. <i>Health and Social Care in the Community</i> , 2005, 13, 164-169.	1.6	3
157	Protocol for stage 2 of the GaP study (genetic testing acceptability for Paget's disease of bone): A questionnaire study to investigate whether relatives of people with Paget's disease would accept genetic testing and preventive treatment if they were available. <i>BMC Health Services Research</i> , 2008, 8, 116.	2.2	3
158	Enabling consultations: the facilitative significance of relational aspects of interpersonal communication. <i>Health Expectations</i> , 2010, 13, 1-3.	2.6	3
159	Payâ€forâ€virtue: an option to improve payâ€forâ€performance?. <i>Journal of Evaluation in Clinical Practice</i> , 2011, 17, 894-898.	1.8	3
160	Sharing outcomes information with consumers: a new course for health librarians. <i>Health Libraries Review</i> , 1994, 11, 279-282.	0.3	2
161	Participation in screening programmes. <i>Health Expectations</i> , 2001, 4, 79-80.	2.6	2
162	Putting participation in the picture. <i>Health Expectations</i> , 2002, 5, 93-94.	2.6	2

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163	Review: decision aids increase patients' knowledge and realistic expectations and reduce decisional conflict. Evidence-Based Medicine, 2004, 9, 21-21.	0.6	2
164	A centralised public information resource for randomised trials: a scoping study to explore desirability and feasibility. BMC Health Services Research, 2005, 5, 39.	2.2	2
165	Philosophy, health services and research: the importance of keeping conversations open. Health Expectations, 2011, 14, 178-181.	2.6	2
166	Asking women to complete health and maternal histories for maternity records: A qualitative study. British Journal of Midwifery, 2013, 21, 793-799.	0.4	2
167	Offering choices. Health Expectations, 2008, 11, 1-2.	2.6	1
168	When service usersâ€™ support obscures problems with care: the need for rigorous research into patientsâ€™ experiences. Health Expectations, 2009, 12, 117-119.	2.6	1
169	Editorial. Health Expectations, 2009, 12, 345-346.	2.6	1
170	Pushing poverty off limits: quality improvement and the architecture of healthcare values. BMC Medical Ethics, 2021, 22, 91.	2.4	1
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