

# 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	Talking it better: conversations and normative complexity in healthcare improvement. Medical Humanities, 2022, 48, 85-93.	1.2	11
2	Understanding and investigating relationality in the capability approach. Journal for the Theory of Social Behaviour, 2022, 52, 86-104.	1.2	5
3	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
4	Implementing shared decision-making in UK: Progress 2017â€“2022. Zeitschrift Fur Evidenz, Fortbildung Und Qualitat Im Gesundheitswesen, 2022, 171, 139-143.	0.9	10
5	Unifying and Universalising Personalised Care? An Analysis of a National Curriculum With Implications for Policy and Education Relating to Person-centred Care. Patient Education and Counseling, 2022, , .	2.2	4
6	How should the â€˜privilegeâ€™ 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
7	Pushing poverty off limits: quality improvement and the architecture of healthcare values. BMC Medical Ethics, 2021, 22, 91.	2.4	1
8	An international core outcome set for evaluating interventions to improve informed consent to clinical trials: The ELICIT Study. Journal of Clinical Epidemiology, 2021, 137, 14-22.	5.0	5
9	Made to Measure: The Ethics of Routine Measurement for Healthcare Improvement. Health Care Analysis, 2021, 29, 39-58.	2.2	5
10	Health professionalsâ€™ experiences and perspectives on food insecurity and long-term conditions: A qualitative investigation. Health and Social Care in the Community, 2020, 28, 404-413.	1.6	19
11	Revisiting the equity debate in COVID-19: ICU is no panacea. Journal of Medical Ethics, 2020, 46, 641-645.	1.8	22
12	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. BMC Family Practice, 2020, 21, 114.	2.9	5
13	Some Unresolved Ethical Challenges in Healthcare Decision-Making: Navigating Family Involvement. Asian Bioethics Review, 2020, 12, 27-36.	1.3	9
14	What does â€˜qualityâ€™ add? Towards an ethics of healthcare improvement. Journal of Medical Ethics, 2020, 46, 118-122.	1.8	11
15	Risk, Overdiagnosis and Ethical Justifications. Health Care Analysis, 2019, 27, 231-248.	2.2	9
16	Defining What is Good: Pluralism and Healthcare Quality. Kennedy Institute of Ethics Journal, 2019, 29, 367-388.	0.5	5
17	â€œThe more you know, the more you realise it is really challenging to doâ€ Tensions and uncertainties in person-centred support for people with long-term conditions. Patient Education and Counseling, 2018, 101, 1460-1467.	2.2	38
18	Exploring the impact and use of patientsâ€™ feedback about their care experiences in general practice settingsâ€”a realist synthesis. Family Practice, 2018, 35, 13-21.	1.9	23

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19	Primary goals, information-giving and men's understanding: a qualitative study of Australian and UK doctors' varied communication about PSA screening. <i>BMJ Open</i> , 2018, 8, e018009.	1.9	10
20	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
21	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
22	Addressing Deficits and Injustices: The Potential Epistemic Contributions of Patients to Research. <i>Health Care Analysis</i> , 2017, 25, 386-403.	2.2	18
23	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
24	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
25	Ethical issues raised by thyroid cancer overdiagnosis: A matter for public health?. <i>Bioethics</i> , 2017, 31, 590-598.	1.4	20
26	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
27	“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
28	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
29	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. <i>Health Expectations</i> , 2017, 20, 243-259.	2.6	76
30	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
31	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
32	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
33	Reframing the Debate Around State Responses to Infertility: Considering the Harms of Subfertility and Involuntary Childlessness. <i>Public Health Ethics</i> , 2016, 9, 290-300.	1.0	8
34	Can consumers learn to ask three questions to improve shared decision making? A feasibility study of the ASK (AskShareKnow) Patient-Clinician Communication Model <sup>®</sup> intervention in a primary health-care setting. <i>Health Expectations</i> , 2016, 19, 1160-1168.	2.6	75
35	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
36	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

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37	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
38	Evaluation of interventions for informed consent for randomised controlled trials (ELICIT): developing a core outcome set. <i>Trials</i> , 2015, 16, .	1.6	0
39	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
40	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
41	Personalised care planning for adults with chronic or long-term health conditions. <i>The Cochrane Library</i> , 2015, , CD010523.	2.8	329
42	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
43	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
44	Special Access Programs Warrant Further Critical Attention: Authors'™ Response to Open Peer Commentaries on "Ethical Justifications for Access to Unapproved Medical Interventions: An Argument for (Limited) Patient Obligations". <i>American Journal of Bioethics</i> , 2014, 14, W1-W2.	0.9	0
45	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
46	Offering informed choice about breast screening. <i>Journal of Medical Screening</i> , 2014, 21, 194-200.	2.3	43
47	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
48	Involving Patients in Their Care. <i>Current Breast Cancer Reports</i> , 2014, 6, 211-218.	1.0	7
49	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
50	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
51	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
52	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
53	How the stigma of low literacy can impair patient-professional spoken interactions and affect health: insights from a qualitative investigation. <i>BMC Health Services Research</i> , 2013, 13, 319.	2.2	100
54	The science of health communication: Impressions from the International Conference on Communication in Healthcare in St Andrews, Scotland, UK. <i>Patient Education and Counseling</i> , 2013, 92, 283-285.	2.2	0

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55	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
56	Recruitment to trials: insights from a meta-ethnography of qualitative studies. <i>Trials</i> , 2013, 14, .	1.6	0
57	Trust in the health-care provider-patient relationship: a systematic mapping review of the evidence base. <i>International Journal for Quality in Health Care</i> , 2013, 25, 682-688.	1.8	95
58	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?. <i>Primary Health Care Research and Development</i> , 2013, 14, 90-102.	1.2	6
59	Enhancing Citizen Engagement in Cancer Screening Through Deliberative Democracy. <i>Journal of the National Cancer Institute</i> , 2013, 105, 380-386.	6.3	56
60	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
61	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
62	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
63	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
64	Asking women to complete health and maternal histories for maternity records: A qualitative study. <i>British Journal of Midwifery</i> , 2013, 21, 793-799.	0.4	2
65	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
66	Shared decision-making: enhancing the clinical relevance. <i>Journal of the Royal Society of Medicine</i> , 2012, 105, 416-421.	2.0	43
67	Which Experiences of Health Care Delivery Matter to Service Users and Why? A Critical Interpretive Synthesis and Conceptual Map. <i>Journal of Health Services Research and Policy</i> , 2012, 17, 70-78.	1.7	112
68	Reason and value: making reasoning fit for practice. <i>Journal of Evaluation in Clinical Practice</i> , 2012, 18, 929-937.	1.8	15
69	Devolution and Patient Choice: Policy Rhetoric versus Experience in Practice. <i>Social Policy and Administration</i> , 2012, 46, 199-218.	3.0	16
70	Addressing complex healthcare problems in diverse settings: Insights from activity theory. <i>Social Science and Medicine</i> , 2012, 74, 305-312.	3.8	73
71	Philosophy, health services and research: the importance of keeping conversations open. <i>Health Expectations</i> , 2011, 14, 178-181.	2.6	2
72	Shared decision making: trade-offs between narrower and broader conceptions. <i>Health Expectations</i> , 2011, 14, 210-219.	2.6	96

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73	Pay-for-virtue: an option to improve pay-for-performance?. <i>Journal of Evaluation in Clinical Practice</i> , 2011, 17, 894-898.	1.8	3
74	Virtue, progress and practice. <i>Journal of Evaluation in Clinical Practice</i> , 2011, 17, 839-846.	1.8	31
75	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
76	Enabling mutual helping? Examining variable needs for facilitated peer support. <i>Patient Education and Counseling</i> , 2011, 85, e120-e125.	2.2	24
77	How information about other people's personal experiences can help with healthcare decision-making: A qualitative study. <i>Patient Education and Counseling</i> , 2011, 85, e291-e298.	2.2	110
78	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
79	Patient involvement: from guidelines to good practice. <i>The Prescriber</i> , 2011, 22, 30-36.	0.3	0
80	Developing a placebo-controlled trial in surgery: Issues of design, acceptability and feasibility. <i>Trials</i> , 2011, 12, 50.	1.6	38
81	Shared Health Governance: The Potential Danger of Oppressive "Healthism". <i>American Journal of Bioethics</i> , 2011, 11, 57-59.	0.9	12
82	Enabling consultations: the facilitative significance of relational aspects of interpersonal communication. <i>Health Expectations</i> , 2010, 13, 1-3.	2.6	3
83	Involving service users in qualitative analysis: approaches and assessment. <i>Health Expectations</i> , 2010, 13, 111-112.	2.6	4
84	Supporting Patient Autonomy: The Importance of Clinician-patient Relationships. <i>Journal of General Internal Medicine</i> , 2010, 25, 741-745.	2.6	339
85	Health in the 'hidden population' of people with low literacy. A systematic review of the literature. <i>BMC Public Health</i> , 2010, 10, 459.	2.9	122
86	Reasons for participating in randomised controlled trials: conditional altruism and considerations for self. <i>Trials</i> , 2010, 11, 31.	1.6	219
87	Brief reflections " from Vikki. <i>Health Expectations</i> , 2010, 13, 331-332.	2.6	0
88	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
89	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
90	Scoping Review and Approach to Appraisal of Interventions Intended to Involve Patients in Patient Safety. <i>Journal of Health Services Research and Policy</i> , 2010, 15, 17-25.	1.7	74

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91	Speaking up about safety concerns: multi-setting qualitative study of patients' views and experiences. <i>BMJ Quality and Safety</i> , 2010, 19, e33-e33.	3.7	79
92	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
93	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
94	Public involvement in health service governance and development: questions of potential for influence. <i>Health Expectations</i> , 2009, 12, 1-3.	2.6	12
95	When service users' support obscures problems with care: the need for rigorous research into patients' experiences. <i>Health Expectations</i> , 2009, 12, 117-119.	2.6	1
96	Editorial. <i>Health Expectations</i> , 2009, 12, 345-346.	2.6	1
97	Decision aids for people facing health treatment or screening decisions. , 2009, , CD001431.		481
98	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
99	Offering choices. <i>Health Expectations</i> , 2008, 11, 1-2.	2.6	1
100	Health literacy: the need to consider images as well as words. <i>Health Expectations</i> , 2008, 11, 99-101.	2.6	18
101	Supporting participation in clinical research: decision aids for trial recruitment?. <i>Health Expectations</i> , 2008, 11, 205-207.	2.6	10
102	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
103	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
104	Involvement in treatment decision-making: Its meaning to people with diabetes and implications for conceptualisation. <i>Social Science and Medicine</i> , 2008, 66, 362-375.	3.8	81
105	Promoting research participation: Why not advertise altruism?. <i>Social Science and Medicine</i> , 2008, 66, 1451-1456.	3.8	81
106	Placing evidence in context: A response to Fry's commentary. <i>Social Science and Medicine</i> , 2008, 66, 1461-1462.	3.8	4
107	"Avoiding harm to others" considerations in relation to parental measles, mumps and rubella (MMR) vaccination discussions " An analysis of an online chat forum. <i>Social Science and Medicine</i> , 2008, 67, 1382-1390.	3.8	85
108	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

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109	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
110	Communicating about screening. <i>BMJ: British Medical Journal</i> , 2008, 337, a1591-a1591.	2.3	73
111	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
112	Differing perspectives on patient involvement in patient safety. <i>Quality and Safety in Health Care</i> , 2007, 16, 82-83.	2.5	41
113	Marketing and clinical trials: a case study. <i>Trials</i> , 2007, 8, 37.	1.6	41
114	Considering 'balance' in information. <i>Health Expectations</i> , 2007, 10, 307-308.	2.6	8
115	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
116	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
117	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
118	Considerations of 'fit' and patient involvement in decision making. <i>Health Expectations</i> , 2006, 9, 95-97.	2.6	5
119	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
120	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
121	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
122	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?. <i>BMC Health Services Research</i> , 2006, 6, 71.	2.2	8
123	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
124	Trust in the context of patient safety problems. <i>Journal of Health Organization and Management</i> , 2006, 20, 397-416.	1.3	51
125	Advising Patients About Patient Safety: Current Initiatives Risk Shifting Responsibility. <i>Joint Commission Journal on Quality and Patient Safety</i> , 2005, 31, 483-494.	0.7	91
126	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



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127	Patients' experiences with partial dentures: a qualitative study. <i>Gerodontology</i> , 2005, 22, 187-192.	2.0	14
128	'Unilateral' and 'bilateral' practitioner approaches in decision-making about treatment. <i>Social Science and Medicine</i> , 2005, 61, 2611-2627.	3.8	109
129	A centralised public information resource for randomised trials: a scoping study to explore desirability and feasibility. <i>BMC Health Services Research</i> , 2005, 5, 39.	2.2	2
130	Methods of hysterectomy: should women have a say?. <i>BMJ: British Medical Journal</i> , 2005, 331, 351.3-352.	2.3	0
131	Review: decision aids increase patients' knowledge and realistic expectations and reduce decisional conflict. <i>Evidence-Based Medicine</i> , 2004, 9, 21-21.	0.6	2
132	Trust and shared decision-making: an emerging research agenda. <i>Health Expectations</i> , 2004, 7, 271-273.	2.6	21
133	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
134	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
135	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
136	Consumer involvement in research projects: the activities of research funders. <i>Health Policy</i> , 2004, 69, 229-238.	3.0	41
137	Consumer involvement in decisions about what health-related research is funded. <i>Health Policy</i> , 2004, 70, 281-290.	3.0	50
138	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
139	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
140	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
141	Research funding organisations and consumer involvement. <i>Journal of Health Services Research and Policy</i> , 2003, 8, 129-131.	1.7	7
142	Interventions for improving communication with children and adolescents about a family member's cancer. <i>The Cochrane Library</i> , 2003, , CD004511.	2.8	10
143	Interventions for improving communication with children and adolescents about their cancer. , 2003, , CD002969.		29
144	Decision aids for people facing health treatment or screening decisions. , 2003, , CD001431.		388

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145	Recordings or summaries of consultations for people with cancer. , 2003, , CD001539.		29
146	Researching experiences of cancer: the importance of methodology. European Journal of Cancer Care, 2002, 11, 232-237.	1.5	22
147	Putting participation in the picture. Health Expectations, 2002, 5, 93-94.	2.6	2
148	Interventions for providers to promote a patient-centred approach in clinical consultations. , 2001, , CD003267.		324
149	Patient choice modules for summaries of clinical effectiveness: a proposal. BMJ: British Medical Journal, 2001, 322, 664-667.	2.3	88
150	Letter to the Editor. Patient Education and Counseling, 2001, 42, 295-296.	2.2	5
151	Giving tape recordings or written summaries of consultations to people with cancer: a systematic review. Health Expectations, 2001, 4, 162-169.	2.6	23
152	The potential contribution of decision aids to screening programmes. Health Expectations, 2001, 4, 109-115.	2.6	19
153	Participation in screening programmes. Health Expectations, 2001, 4, 79-80.	2.6	2
154	Decisions about treatment: interpretations of two measures of control by women having a hysterectomy. Social Science and Medicine, 2001, 53, 721-732.	3.8	62
155	Informing, communicating and sharing decisions with people who have cancer. Quality in Health Care: QHC, 2001, 10, 193-196.	1.2	14
156	Discussing wisdom teeth with patients: a critique of information materials. Health Bulletin, 2001, 59, 171-7.	0.1	0
157	Supporting and resourcing treatment decision-making: some policy considerations. Health Expectations, 2000, 3, 77-85.	2.6	20
158	Editorial. Health Expectations, 2000, 3, 87-89.	2.6	3
159	The case of Norplant as an example of media coverage over the life of a new health technology. Lancet, The, 2000, 355, 1633-1636.	13.7	19
160	Judging journalism: how should the quality of news reporting about clinical interventions be assessed and improved?. Quality and Safety in Health Care, 1999, 8, 172-176.	2.5	11
161	Decision aids for patients facing health treatment or screening decisions: systematic review. BMJ: British Medical Journal, 1999, 319, 731-734.	2.3	718
162	Editorial. Health Expectations, 1999, 2, 1-2.	2.6	4

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163	Editorial. Health Expectations, 1999, 2, 75-77.	2.6	6
164	Sharing decisions with patients: is the information good enough?. BMJ: British Medical Journal, 1999, 318, 318-322.	2.3	519
165	Disseminating information about healthcare effectiveness: a survey of consumer health information services. Quality and Safety in Health Care, 1998, 7, 124-129.	2.5	29
166	Evidence-Informed Patient Choice: Practical Issues of Involving Patients in Decisions About Health Care Technologies. International Journal of Technology Assessment in Health Care, 1998, 14, 212-225.	0.5	163
167	Developing Information Materials to Present the Findings of Technology Assessments to Consumers: The Experience of the NHS Centre for Reviews and Dissemination. International Journal of Technology Assessment in Health Care, 1998, 14, 47-70.	0.5	56
168	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
169	Lay perspectives: advantages for health research. BMJ: British Medical Journal, 1998, 316, 463-466.	2.3	287
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