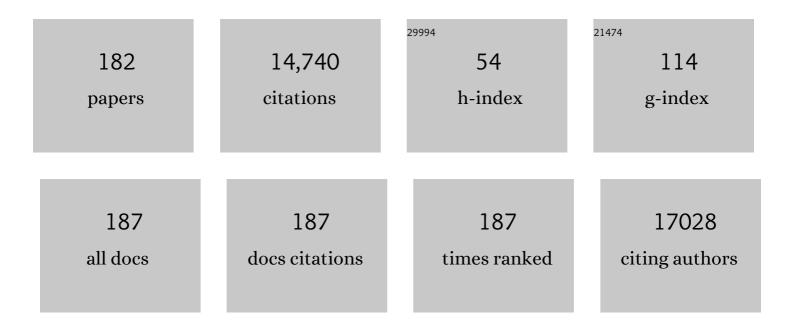
Sue Ziebland

List of Publications by Year in descending order

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SHE ZIERLAND

#	Article	lF	CITATIONS
1	What happens when patients say "no―to offers of referral for weight loss? - Results and recommendations from a conversation analysis of primary care interactions. Patient Education and Counseling, 2022, 105, 524-533.	1.0	9
2	Cultivating Doctors' Gut Feeling: Experience, Temporality and Politics of Gut Feelings in Family Medicine. Culture, Medicine and Psychiatry, 2022, 46, 564-581.	0.7	9
3	Unintended consequences of online consultations: a qualitative study in UK primary care. British Journal of General Practice, 2022, 72, e128-e137.	0.7	38
4	Building the case for the use of gut feelings in cancer referrals: perspectives of patients referred to a non-specific symptoms pathway. British Journal of General Practice, 2022, 72, e43-e50.	0.7	2
5	Challenges of safeguarding via remote consulting during the COVID-19 pandemic: a qualitative interview study. British Journal of General Practice, 2022, 72, e199-e208.	0.7	20
6	Polyphonic perspectives on health and care: Reflections from two decades of the DIPEx project. Journal of Health Services Research and Policy, 2021, 26, 133-140.	0.8	22
7	A systematic review and thematic synthesis of qualitative studies exploring GPs' and nurses' perspectives on discussing weight with patients with overweight and obesity in primary care. Obesity Reviews, 2021, 22, e13151.	3.1	32
8	Combining patient talk about internet use during primary care consultations with retrospective accounts. A qualitative analysis of interactional and interview data. Social Science and Medicine, 2021, 272, 113703.	1.8	7
9	Conversations about FGM in primary care: a realist review on how, why and under what circumstances FGM is discussed in general practice consultations. BMJ Open, 2021, 11, e039809.	0.8	12
10	Shifting research culture to address the mismatch between where trials recruit and where populations with the most disease live: a qualitative study. BMC Medical Research Methodology, 2021, 21, 80.	1.4	8
11	Qualitative study: patients' enduring concerns about discussing internet use in general practice consultations. BMJ Open, 2021, 11, e047508.	0.8	5
12	People with weightâ€related longâ€term conditions want support from GPs : A qualitative interview study. Clinical Obesity, 2021, 11, e12471.	1.1	5
13	Gender in the consolidated criteria for reporting qualitative research (COREQ) checklist. International Journal for Quality in Health Care, 2021, 33, .	0.9	9
14	Power, paradox and pessimism: On the unintended consequences of digital health technologies in primary care. Social Science and Medicine, 2021, 289, 114419.	1.8	26
15	GPs' use of gut feelings when assessing cancer risk: a qualitative study in UK primary care. British Journal of General Practice, 2021, 71, e356-e363.	0.7	12
16	Discussing weight loss opportunistically and effectively in family practice: a qualitative study of clinical interactions using conversation analysis in UK family practice. Family Practice, 2021, 38, 321-328.	0.8	12
17	Are some feasibility studies more feasible than others? A review of the outcomes of feasibility studies on the ISRCTN registry. Pilot and Feasibility Studies, 2021, 7, 195.	0.5	7
18	Caring as sharing. Negotiating the moral boundaries of receiving care. Critical Public Health, 2020, 30, 567-576.	1.4	7

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19	Understanding the role of GPs' gut feelings in diagnosing cancer in primary care: a systematic review and meta-analysis of existing evidence. British Journal of General Practice, 2020, 70, e612-e621.	0.7	48
20	Face-to-Face Compared With Online Collected Accounts of Health and Illness Experiences: A Scoping Review. Qualitative Health Research, 2020, 30, 2092-2102.	1.0	77
21	Standardising definitions for the pre-eclampsia core outcome set: A consensus development study. Pregnancy Hypertension, 2020, 21, 208-217.	0.6	9
22	If social determinants of health are so important, shouldn't we ask patients about them?. BMJ, The, 2020, 371, m4150.	3.0	42
23	Assessing health research grant applications: A retrospective comparative review of a one-stage versus a two-stage application assessment process. PLoS ONE, 2020, 15, e0230118.	1.1	4
24	The convivial and the pastoral in patient–doctor relationships: a multiâ€country study of patient stories of care, choice and medical authority in cancer diagnostic processes. Sociology of Health and Illness, 2020, 42, 844-861.	1.1	6
25	How do frontline staff use patient experience data for service improvement? Findings from an ethnographic case study evaluation. Journal of Health Services Research and Policy, 2020, 25, 151-161.	0.8	23
26	Understanding how front-line staff use patient experience data for service improvement: an exploratory case study evaluation. Health Services and Delivery Research, 2020, 8, 1-170.	1.4	7
27	Supporting patients with female genital mutilation in primary care: a qualitative study exploring the perspectives of GPs' working in England. British Journal of General Practice, 2020, 70, e749-e756.	0.7	10
28	A systematic review of reasons for and against asking patients about their socioeconomic contexts. International Journal for Equity in Health, 2019, 18, 112.	1.5	30
29	General Practitioner's use of online resources during medical visits: managing the boundary between inside and outside the clinic. Sociology of Health and Illness, 2019, 41, 65-81.	1.1	14
30	Communication with children and adolescents about the diagnosis of their own life-threatening condition. Lancet, The, 2019, 393, 1150-1163.	6.3	100
31	Communication with children and adolescents about the diagnosis of a life-threatening condition in their parent. Lancet, The, 2019, 393, 1164-1176.	6.3	99
32	How do GPs and patients share the responsibility for cancer safety netting follow-up actions? A qualitative interview study of GPs and patients in Oxfordshire, UK. BMJ Open, 2019, 9, e029316.	0.8	9
33	Tackling poorly selected, collected, andÂreported outcomes in obstetrics andÂgynecology research. American Journal of Obstetrics and Gynecology, 2019, 220, 71.e1-71.e4.	0.7	41
34	How wide is the Goldilocks Zone in your health system?. Journal of Health Services Research and Policy, 2019, 24, 52-56.	0.8	14
35	Depression at work, authenticity in question: Experiencing, concealing and revealing. Health (United) Tj ETQq1	1 0.784314	4 rgBT /Overlo
36	Using online patient feedback to improve NHS services: the INQUIRE multimethod study. Health Services and Delivery Research, 2019, 7, 1-150.	1.4	22

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37	Quality improvements of safety-netting guidelines for cancer in UK primary care: insights from a qualitative interview study of GPs. British Journal of General Practice, 2019, 69, e819-e826.	0.7	11
38	Alternatives to the face-to-face consultation in general practice: focused ethnographic case study. British Journal of General Practice, 2018, 68, e293-e300.	0.7	74
39	Methodological and Practical Issues in Cross-National Qualitative Research: Lessons From the Literature and a Comparative Study of the Experiences of People Receiving a Diagnosis of Cancer. Qualitative Health Research, 2018, 28, 789-799.	1.0	37
40	Pain and the Internet: Transforming the Experience?. , 2018, , 129-155.		3
41	Qualitative cross-country comparison of whether, when and how people diagnosed with lung cancer talk about cigarette smoking in narrative interviews. BMJ Open, 2018, 8, e023934.	0.8	5
42	Protocol paper for the â€~Harnessing resources from the internet to maximise outcomes from GP consultations (HaRI)' study: a mixed qualitative methods study. BMJ Open, 2018, 8, e024188.	0.8	8
43	GP-delivered brief weight loss interventions: a cohort study of patient responses and subsequent actions, using conversation analysis in UK primary care. British Journal of General Practice, 2018, 68, e646-e653.	0.7	19
44	Receptionists' role in new approaches to consultations in primary care: a focused ethnographic study. British Journal of General Practice, 2018, 68, e478-e486.	0.7	13
45	GPs' understanding and practice of safety netting for potential cancer presentations: a qualitative study in primary care. British Journal of General Practice, 2018, 68, e505-e511.	0.7	35
46	The potential of alternatives to face-to-face consultation in general practice, and the impact on different patient groups: a mixed-methods case study. Health Services and Delivery Research, 2018, 6, 1-200.	1.4	59
47	Core outcome sets in women's and newborn health: a systematic review. BJOG: an International Journal of Obstetrics and Gynaecology, 2017, 124, 1481-1489.	1.1	131
48	Do we all agree what "good health care―looks like? Views from those who are "seldom heard―in health research, policy and service improvement. Health Expectations, 2017, 20, 878-885.	1.1	17
49	A systematic review of primary outcomes and outcome measure reporting in randomized trials evaluating treatments for preâ€eclampsia. International Journal of Gynecology and Obstetrics, 2017, 139, 262-267.	1.0	26
50	Advancing gender equality through the Athena SWAN Charter for Women in Science: an exploratory study of women's and men's perceptions. Health Research Policy and Systems, 2017, 15, 12.	1.1	99
51	Patients' initial steps to cancer diagnosis in Denmark, England and Sweden: what can a qualitative, cross-country comparison of narrative interviews tell us about potentially modifiable factors?. BMJ Open, 2017, 7, e018210.	0.8	23
52	Narratives of experience of mental health and illness on healthtalk.org. BJPsych Bulletin, 2016, 40, 273-276.	0.7	2
53	A joint effort over a period of time: factors affecting use of urate-lowering therapy for long-term treatment of gout. BMC Musculoskeletal Disorders, 2016, 17, 249.	0.8	12
54	Randomised feasibility study of a novel experience-based internet intervention to support self-management in chronic asthma. BMJ Open, 2016, 6, e013401.	0.8	21

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55	What do we need to consider when planning, implementing and researching the use of alternatives to face-to-face consultations in primary healthcare?. Digital Health, 2016, 2, 205520761667555.	0.9	34
56	Prescribing antibiotics to â€~at-risk' children with influenza-like illness in primary care: qualitative study: Table 1. BMJ Open, 2016, 6, e011497.	0.8	23
57	What patients do and their impact on implementation. Journal of Health Organization and Management, 2016, 30, 258-278.	0.6	50
58	â€~l knew before I was told': Breaches, cues and clues in the diagnostic assemblage. Social Science and Medicine, 2016, 154, 85-92.	1.8	28
59	A protocol for developing, disseminating, and implementing a core outcome set for pre-eclampsia. Pregnancy Hypertension, 2016, 6, 274-278.	0.6	48
60	Using alternatives to face-to-face consultations: a survey of prevalence and attitudes in general practice. British Journal of General Practice, 2016, 66, e460-e466.	0.7	100
61	A novel experience-based internet intervention for smoking cessation: feasibility randomised controlled trial. BMC Public Health, 2016, 16, 1156.	1.2	8
62	An open letter to <i>The BMJ</i> editors on qualitative research. BMJ, The, 2016, 352, i563.	3.0	234
63	Proliferation of private online healthcare companies. BMJ, The, 2016, 352, i1076.	3.0	6
64	Drawing on Accounts of Long-Term Urinary Catheter Use. Qualitative Health Research, 2016, 26, 154-163.	1.0	9
65	Examining the role of patients' experiences as a resource for choice and decision-making in health care: a creative, interdisciplinary mixed-method study in digital health. Programme Grants for Applied Research, 2016, 4, 1-214.	0.4	17
66	Barriers to shared decisions in the most serious of cancers: a qualitative study of patients with pancreatic cancer treated in the <scp>UK</scp> . Health Expectations, 2015, 18, 3302-3312.	1.1	39
67	Mapping patients' experiences from initial symptoms to gout diagnosis: a qualitative exploration: TableÂ1. BMJ Open, 2015, 5, e008323.	0.8	30
68	"Why me? I don't fit the mould … I am a freak of nature― a qualitative study of women's experience gout. BMC Women's Health, 2015, 15, 122.	e of 0.8	23
69	Taboo and the different death? Perceptions of those bereaved by suicide or other traumatic death. Sociology of Health and Illness, 2015, 37, 610-625.	1.1	77
70	On interviewing people with pets: reflections from qualitative research on people with longâ€ŧerm conditions. Sociology of Health and Illness, 2015, 37, 67-80.	1.1	38
71	The role of the Internet for people with chronic pain: examples from the DIPEx International Project. British Journal of Pain, 2015, 9, 62-64.	0.7	25
72	Measuring the effects of online health information: Scale validation for the e-Health Impact Questionnaire. Patient Education and Counseling, 2015, 98, 1418-1424.	1.0	55

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73	Mike Bury: Biographical Disruption and Long-Term and Other Health Conditions. , 2015, , 582-598.		15
74	An Overview of Self-Administered Health Literacy Instruments. PLoS ONE, 2014, 9, e109110.	1.1	43
75	"lt can't be very important because it comes and goesâ€â€"patients' accounts of intermittent symptoms preceding a pancreatic cancer diagnosis: a qualitative study: TableÂ1. BMJ Open, 2014, 4, e004215.	0.8	61
76	Collecting data on patient experience is not enough: they must be used to improve care. BMJ, The, 2014, 348, g2225-g2225.	3.0	298
77	Using secondary analysis of qualitative data of patient experiences of health care to inform health services research and policy. Journal of Health Services Research and Policy, 2014, 19, 177-182.	0.8	70
78	Using a national archive of patient experience narratives to promote local patient-centered quality improvement: an ethnographic process evaluation of †̃accelerated' experience-based co-design. Journal of Health Services Research and Policy, 2014, 19, 200-207.	0.8	84
79	User-Generated Online Health Content: A Survey of Internet Users in the United Kingdom. Journal of Medical Internet Research, 2014, 16, e118.	2.1	37
80	Testing accelerated experience-based co-design: a qualitative study of using a national archive of patient experience narrative interviews to promote rapid patient-centred service improvement. Health Services and Delivery Research, 2014, 2, 1-122.	1.4	71
81	Informing the development of NICE (National Institute for Health and Care Excellence) quality standards through secondary analysis of qualitative narrative interviews on patients' experiences. Health Services and Delivery Research, 2014, 2, 1-206.	1.4	7
82	Personal identity and the role of â€~carer' among relatives and friends of people with multiple sclerosis. Social Science and Medicine, 2013, 96, 78-85.	1.8	81
83	Measuring the effects of online health information for patients: Item generation for an e-health impact questionnaire. Patient Education and Counseling, 2013, 93, 433-438.	1.0	51
84	What parents say about disclosing the end of their pregnancy due to fetal abnormality. Midwifery, 2013, 29, 24-32.	1.0	19
85	Narrative interviewing. , 2013, , 38-48.		13
86	How people bereaved by suicide perceive newspaper reporting: qualitative study. British Journal of Psychiatry, 2013, 203, 228-232.	1.7	31
87	Emotions and chronic illness. Chronic Illness, 2012, 8, 159-162.	0.6	8
88	Talk of frustration in the narratives of people with chronic pain. Chronic Illness, 2012, 8, 176-191.	0.6	86
89	Why Listening to Health Care Users Really Matters. Journal of Health Services Research and Policy, 2012, 17, 68-69.	0.8	8
90	A qualitative exploration of the role of primary care in supporting colorectal cancer patients. Supportive Care in Cancer, 2012, 20, 3071-3078.	1.0	19

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91	An Alarming Prognosis: How People Affected by Pancreatic Cancer Use (and Avoid) Internet Information. Policy and Internet, 2012, 4, 1.	2.0	18
92	Women's experience of coronary heart disease: why is it different?. British Journal of Cardiac Nursing, 2012, 7, 165-170.	0.0	5
93	Reassured or fobbed off? Perspectives on infertility consultations in primary care: a qualitative study. British Journal of General Practice, 2012, 62, e438-e445.	0.7	15
94	Incurable, invisible and inconclusive: watchful waiting for chronic lymphocytic leukaemia and implications for doctor-patient communication. European Journal of Cancer Care, 2012, 21, 67-77.	0.7	45
95	Imagined futures: how experiential knowledge of disability affects parents' decision making about fetal abnormality. Health Expectations, 2012, 15, 139-156.	1.1	21
96	Understanding depression through a â€~coming out' framework. Sociology of Health and Illness, 2012, 34, 730-745.	1.1	21
97	Health and Illness in a Connected World: How Might Sharing Experiences on the Internet Affect People's Health?. Milbank Quarterly, 2012, 90, 219-249.	2.1	377
98	A Proper, Fitting Explanation?. Crisis, 2012, 33, 230-238.	0.9	15
99	Patients' needs following colorectal cancer diagnosis: where does primary care fit in?. British Journal of General Practice, 2011, 61, e692-e699.	0.7	35
100	Exceptionally good? Positive experiences of NHS care and treatment surprises lymphoma patients: a qualitative interview study. Health Expectations, 2011, 14, 21-28.	1.1	5
101	Nurses should be encouraged and helped to monitor patients' medicines. Journal of Nursing Management, 2011, 19, 393-394.	1.4	1
102	How personal experiences feature in women's accounts of use of information for decisions about antenatal diagnostic testing for foetal abnormality. Social Science and Medicine, 2011, 72, 755-762.	1.8	36
103	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.	1.0	110
104	Women's Distress about Unexpected Dcis Uncertainties and Information Provision (Response to Pryke) Tj ETQq(0.0 rgBT 1.1	/Oyerlock 10
105	How the Internet is changing the experience of bereavement by suicide: A qualitative study in the UK. Health (United Kingdom), 2011, 15, 173-187.	0.9	42
106	The role of spirituality and religion for those bereaved due to a traumatic death. Mortality, 2011, 16, 1-19.	0.3	27
107	Infertility; isolation and the Internet: A qualitative interview study. Patient Education and Counseling, 2010, 81, 436-441.	1.0	91
108	Factors contributing to the time taken to consult with symptoms of lung cancer: a cross-sectional study. Thorax, 2009, 64, 523-531.	2.7	91

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109	Patients' views of a multimedia resource featuring experiences of rheumatoid arthritis: pilot evaluation of www.healthtalkonline.org. Health Informatics Journal, 2009, 15, 147-159.	1.1	34
110	Disclosing a Cancer Diagnosis to Friends and Family: A Gendered Analysis of Young Men's and Women's Experiences. Qualitative Health Research, 2009, 19, 744-754.	1.0	77
111	†Getting through' not †going under': A qualitative study of gender and spousal support after diagnosi with colorectal cancer. Social Science and Medicine, 2009, 68, 1169-1175.	⁵ 1.8	68
112	â€~My brain couldn't move from planning a birth to planning a funeral': A qualitative study of parents' experiences of decisions after ending a pregnancy for fetal abnormality. International Journal of Nursing Studies, 2009, 46, 1111-1121.	2.5	44
113	Breast cancer in young families: a qualitative interview study of fathers and their role and communication with their children following the diagnosis of maternal breast cancer. Psycho-Oncology, 2009, 18, 96-103.	1.0	58
114	Biographical disruption, abruption and repair in the context of Motor Neurone Disease. Sociology of Health and Illness, 2009, 31, 1043-1058.	1.1	98
115	Have men been overlooked? A comparison of young men and women's experiences of chemotherapyâ€induced alopecia. Psycho-Oncology, 2008, 17, 577-583.	1.0	113
116	How patients' experiences contribute to decision making: illustrations from DIPEx (personal) Tj ETQq0 0 0 rgB	Г /Qverloc 1.4	k 10 Tf 50 4
117	What affects the uptake of screening for bowel cancer using a faecal occult blood test (FOBt): A qualitative study. Social Science and Medicine, 2008, 66, 2425-2435.	1.8	122
118	Why Men in the United Kingdom Still Want the Prostate Specific Antigen Test. Qualitative Health Research, 2008, 18, 56-64.	1.0	15
119	The Effect of Joint Interviewing on the Performance of Gender. Field Methods, 2008, 20, 107-128.	0.5	55
120	Fertility Issues: The Perceptions and Experiences of Young Men Recently Diagnosed and Treated for Cancer. Journal of Adolescent Health, 2007, 40, 69-75.	1.2	97
121	Exploring men's and women's experiences of depression and engagement with health professionals: more similarities than differences? A qualitative interview study. BMC Family Practice, 2007, 8, 43.	2.9	59
122	Does it matter if clinicians recruiting for a trial don't understand what the trial is really about? Qualitative study of surgeons' experiences of participation in a pragmatic multi-centre RCT. Trials, 2007, 8, 4.	0.7	66
123	Positive prevention: Contemporary issues facing HIV positive people negotiating sex in the UK. Social Science and Medicine, 2007, 65, 755-770.	1.8	16
124	Making sense of qualitative data analysis: an introduction with illustrations from DIPEx (personal) Tj ETQq0 0 0 rgE	3T /Overloo	ck 10 Tf 50 456

125	The choice is yours? How women with ovarian cancer make sense of treatment choices. Patient Education and Counseling, 2006, 62, 361-367.	1.0	38
126	The specialist palliative care nurse: A qualitative study of the patients' perspective. International Journal of Nursing Studies, 2006, 43, 1011-1022.	2.5	32

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127	Men's accounts of depression: Reconstructing or resisting hegemonic masculinity?. Social Science and Medicine, 2006, 62, 2246-2257.	1.8	327
128	Gender, cancer experience and internet use: A comparative keyword analysis of interviews and online cancer support groups. Social Science and Medicine, 2006, 62, 2577-2590.	1.8	252
129	Breast cancer in the family—children's perceptions of their mother's cancer and its initial treatment: qualitative study. BMJ: British Medical Journal, 2006, 332, 998-1003.	2.4	96
130	Why do General Practitioners prescribe antibiotics for acute infective conjunctivitis in children? Qualitative interviews with GPs and a questionnaire survey of parents and teachers. Family Practice, 2006, 23, 226-232.	0.8	36
131	Minimizing delays in ovarian cancer diagnosis: an expansion of Andersen's model of 'total patient delay'. Family Practice, 2006, 24, 48-55.	0.8	74
132	'Can I come off the tablets now?' A qualitative analysis of heart failure patients' understanding of their medication. Family Practice, 2006, 23, 624-630.	0.8	39
133	What people close to death say about euthanasia and assisted suicide: a qualitative study. Journal of Medical Ethics, 2006, 32, 706-710.	1.0	69
134	"The Old Me Could Never Have Done That― How People Give Meaning to Recovery Following Depression. Qualitative Health Research, 2006, 16, 1038-1053.	1.0	73
135	What happened when Scottish women were given advance supplies of emergency contraception? A survey and qualitative study of women's views and experiences. Social Science and Medicine, 2005, 60, 1767-1779.	1.8	35
136	"Not that sort of practice― the views and behaviour of primary care practitioners in a study of advance provision of emergency contraception. Family Practice, 2005, 22, 280-286.	0.8	12
137	In the absence of evidence, who chooses? A qualitative study of patients' needs after treatment for colorectal cancer. Journal of Health Services Research and Policy, 2004, 9, 159-164.	0.8	40
138	Expressions of Loss of Adulthood in the Narratives of People with Colorectal Cancer. Qualitative Health Research, 2004, 14, 187-203.	1.0	70
139	How the internet affects patients' experience of cancer: a qualitative study. BMJ: British Medical Journal, 2004, 328, 564.	2.4	387
140	Reported barriers to eating more fruit and vegetables before and after participation in a randomized controlled trial: a qualitative study. Health Education Research, 2004, 19, 165-174.	1.0	76
141	The Role of Humor for Men with Testicular Cancer. Qualitative Health Research, 2004, 14, 1123-1139.	1.0	71
142	The importance of being expert: the quest for cancer information on the Internet. Social Science and Medicine, 2004, 59, 1783-1793.	1.8	180
143	What do patients with prostate or breast cancer want from an Internet site? A qualitative study of information needs. Patient Education and Counseling, 2004, 53, 57-64.	1.0	232
144	Advanced provision of emergency contraception does not reduce abortion rates. Contraception, 2004, 69, 361-366.	0.8	147

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145	Emergency contraception: why can't you give it away? Qualitative findings from an evaluation of advance provision of emergency contraception. Contraception, 2004, 70, 25-29.	0.8	37
146	Qualitative study of men's perceptions of why treatment delays occur in the UK for those with testicular cancer. British Journal of General Practice, 2004, 54, 25-32.	0.7	58
147	Lung cancer patients' perceptions of access to financial benefits: a qualitative study. British Journal of General Practice, 2004, 54, 589-94.	0.7	30
148	Ethics and dementia: mapping the literature by bibliometric analysis. International Journal of Geriatric Psychiatry, 2003, 18, 41-54.	1.3	62
149	DIPEx: Fresh Insights for Medical Practice. Journal of the Royal Society of Medicine, 2003, 96, 209-210.	1.1	6
150	DIPEx: fresh insights for medical practice. Journal of the Royal Society of Medicine, 2003, 96, 209-210.	1.1	9
151	Development and validation of the Day in the Life Questionnaire (DILQ) as a measure of fruit and vegetable questionnaire for 7-9 year olds. Health Education Research, 2002, 17, 211-220.	1.0	88
152	Why men with prostate cancer want wider access to prostate specific antigen testing: qualitative study. BMJ: British Medical Journal, 2002, 325, 737-737.	2.4	46
153	Dietary effect on blood pressure. Lancet, The, 2002, 360, 1786.	6.3	0
154	Effects of fruit and vegetable consumption on plasma antioxidant concentrations and blood pressure: a randomised controlled trial. Lancet, The, 2002, 359, 1969-1974.	6.3	438
155	Carers, ethics and dementia: a survey and review of the literature. International Journal of Geriatric Psychiatry, 2002, 17, 35-40.	1.3	26
156	Is â€~watchful waiting' a real choice for men with prostate cancer? A qualitative study. BJU International, 2002, 90, 257-264.	1.3	91
157	â€~People sometimes react funny if they're not told enough': women's views about the risks of diagnostic laparoscopy. Health Expectations, 2002, 5, 302-309.	1.1	25
158	Body image and weight change in middle age: a qualitative study. International Journal of Obesity, 2002, 26, 1083-1091.	1.6	32
159	Prostate cancer: embodied experience and perceptions of masculinity. Sociology of Health and Illness, 2002, 24, 820-841.	1.1	220
160	Smoking cessation in pregnancy: What's a man to do?. Health Education Journal, 2001, 60, 232-240.	0.6	7
161	Update on DIPEx: a Database of Individual Patients' Experience. BMC News and Views, 2001, 1, .	0.0	1
162	The use of patients' stories by self-help groups: a survey of voluntary organizations in the UK on the register of the College of Health. Health Expectations, 2000, 3, 176-181.	1.1	11

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163	Qualitative research in health care: Analysing qualitative data. BMJ: British Medical Journal, 2000, 320, 114-116.	2.4	4,793
164	An introduction to qualitative methods for health professionals. Master classes in primary care research No. 1 Family Practice, 2000, 17, 281-281.	0.8	0
165	Database of patients' experiences (DIPEx): a multi-media approach to sharing experiences and information. Lancet, The, 2000, 355, 1540-1543.	6.3	124
166	Emergency contraception: an anomalous position in the family planning repertoire?. Social Science and Medicine, 1999, 49, 1409-1417.	1.8	36
167	Lack of willpower or lack of wherewithal? "Internal―and "external―barriers to changing diet and exercise in a three year follow-up of participants in a health check. Social Science and Medicine, 1998, 46, 461-465.	1.8	58
168	Patients' views of routine hospital follow-up: a qualitative study of women with breast cancer in remission. , 1998, 7, 436-439.		52
169	A 'five-a-day' fruit and vegetable pack for primary school children. Part I : development and pre-testing. Health Education Journal, 1998, 57, 97-104.	0.6	3
170	Concerns and cautions about prescribing and deregulating emergency contraception: a qualitative study of GPs using telephone interviews. Family Practice, 1998, 15, 449-456.	0.8	33
171	How important is the smoking status of the woman's partner as a predictor of smoking cessation in pregnancy? A literature review. Health Education Journal, 1998, 57, 70-80.	0.6	8
172	Not a â€~proper' solution? The Gap between Professional Guidelines and Users' Views about the Safety of Using Emergency Contraception. Journal of Health Services Research and Policy, 1998, 3, 12-19.	0.8	10
173	Patients' views of routine hospital follow-up: a qualitative study of women with breast cancer in remission. , 1998, 7, 436.		2
174	Smoking cessation interventions for dental patients–attitudes and reported practices of dentists in the Oxford region. British Dental Journal, 1997, 183, 359-364.	0.3	38
175	Software for Analysing Textual Data. Journal of Health Services Research and Policy, 1996, 1, 247-249.	0.8	0
176	Desire for the body normal: body image and discrepancies between self reported and measured height and weight in a British population Journal of Epidemiology and Community Health, 1996, 50, 105-106.	2.0	84
177	Qualitative methods in health and medicine (Book) Sociology of Health and Illness, 1995, 17, 430-431.	1.1	0
178	The short form 36 health status questionnaire: clues from the Oxford region's normative data about its usefulness in measuring health gain in population surveys Journal of Epidemiology and Community Health, 1995, 49, 102-105.	2.0	36
179	Tacit models of disability underlying health status instruments. Social Science and Medicine, 1993, 37, 69-75.	1.8	29
180	Comparison of two approaches to measuring change in health status in rheumatoid arthritis: the Health Assessment Questionnaire (HAQ) and modified HAQ Annals of the Rheumatic Diseases, 1992, 51, 1202-1205.	0.5	74

#	Article	IF	CITATIONS
181	The DIPEx Project: Collecting Personal Experiences of Illness and Health Care. , 0, , 115-131.		22
182	What are the mechanisms that enable the reciprocal involvement of seldom heard groups in health and social care research? A rapid realist review protocol. HRB Open Research, 0, 1, 7.	0.3	13