

Myfanwy Morgan

List of Publications by Year in descending order

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Version: 2024-02-01

174
papers

9,254
citations

61984

43
h-index

42399

92
g-index

177
all docs

177
docs citations

177
times ranked

10089
citing authors

#	ARTICLE	IF	CITATIONS
1	Should the family have a role in deceased organ donation decision-making? A systematic review of public knowledge and attitudes towards organ procurement policies in Europe. <i>Transplantation Reviews</i> , 2022, 36, 100673.	2.9	9
2	Identifying the psychosocial predictors of ultraviolet exposure to the face in patients with xeroderma pigmentosum: a study of the behavioural factors affecting clinical outcomes in this genetic disease. <i>Journal of Medical Genetics</i> , 2022, 59, 1095-1103.	3.2	2
3	The Personal Sociability and Connections Scale (PeSCS): Development and initial assessment at a primary care facility. <i>International Journal of Social Psychiatry</i> , 2021, , 002076402199351.	3.1	4
4	Food-related quality of life is impaired in inflammatory bowel disease and associated with reduced intake of key nutrients. <i>American Journal of Clinical Nutrition</i> , 2021, 113, 832-844.	4.7	26
5	Community-based short-term integrated palliative and supportive care reduces symptom distress for older people with chronic noncancer conditions compared with usual care: A randomised controlled single-blind mixed method trial. <i>International Journal of Nursing Studies</i> , 2021, 120, 103978.	5.6	18
6	Response to Zhou (2021) "Comment on Evans et al (2021) "Community-based short-term integrated palliative and supportive care reduces symptom distress for older people with chronic noncancer conditions compared with usual care" International Journal of Nursing Studies, 2021, 125, 104119.	5.6	0
7	Ultraviolet exposure to the face in patients with xeroderma pigmentosum and healthy controls: applying a novel methodology to define photoprotection behaviour. <i>British Journal of Dermatology</i> , 2021, , .	1.5	3
8	Approaches to Photoprotection and Normalization in Highly Adherent Families of Children With Xeroderma Pigmentosum in the United Kingdom. <i>Qualitative Health Research</i> , 2020, 30, 1275-1286.	2.1	5
9	Perceptions and psychosocial impact of food, nutrition, eating and drinking in people with inflammatory bowel disease: a qualitative investigation of food-related quality of life. <i>Journal of Human Nutrition and Dietetics</i> , 2020, 33, 115-127.	2.5	58
10	Forms, interactions, and responses to social support: A qualitative study of support and adherence to photoprotection amongst patients with Xeroderma Pigmentosum. <i>British Journal of Health Psychology</i> , 2020, 25, 89-106.	3.5	8
11	Why? What? How? Using an Intervention Mapping approach to develop a personalised intervention to improve adherence to photoprotection in patients with Xeroderma Pigmentosum. <i>Health Psychology and Behavioral Medicine</i> , 2020, 8, 475-500.	1.8	6
12	Improving photoprotection in adults with xeroderma pigmentosum: personalisation and tailoring in the "XPAND" intervention. <i>Health Psychology and Behavioral Medicine</i> , 2020, 8, 543-572.	1.8	5
13	Patient views on use of emergency and alternative care services for adult epilepsy: A qualitative study. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2020, 80, 56-62.	2.0	5
14	Primary care service use by end-of-life cancer patients: a nationwide population-based cohort study in the United Kingdom. <i>BMC Family Practice</i> , 2020, 21, 76.	2.9	11
15	Seizure First Aid Training For people with Epilepsy (SAFE) frequently attending emergency departments and their significant others: results of a UK multi-centre randomised controlled pilot trial. <i>BMJ Open</i> , 2020, 10, e035516.	1.9	1
16	Access to and interventions to improve maternity care services for immigrant women: a narrative synthesis systematic review. <i>Health Services and Delivery Research</i> , 2020, 8, 1-122.	1.4	5
17	Seizure first aid training for people with epilepsy attending emergency departments and their significant others: the SAFE intervention and feasibility RCT. <i>Health Services and Delivery Research</i> , 2020, 8, 1-190.	1.4	2
18	The influence of perceived medical risks and psychosocial concerns on photoprotection behaviours among adults with xeroderma pigmentosum: a qualitative interview study in the UK. <i>BMJ Open</i> , 2019, 9, e024445.	1.9	11

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19	Assessing Treatment Fidelity within an Epilepsy Randomized Controlled Trial: Seizure First Aid Training for People with Epilepsy Who Visit Emergency Departments. <i>Behavioural Neurology</i> , 2019, 2019, 1-11.	2.1	5
20	Developing patient-centred, feasible alternative care for adult emergency department users with epilepsy: protocol for the mixed-methods observational "Collaborate"™ project. <i>BMJ Open</i> , 2019, 9, e031696.	1.9	8
21	Experience of and access to maternity care in the UK by immigrant women: a narrative synthesis systematic review. <i>BMJ Open</i> , 2019, 9, e029478.	1.9	40
22	Public knowledge and attitudes towards consent policies for organ donation in Europe. A systematic review. <i>Transplantation Reviews</i> , 2019, 33, 1-8.	2.9	25
23	Experiences of stigma over the lifetime of people with xeroderma pigmentosum: A qualitative interview study in the United Kingdom. <i>Journal of Health Psychology</i> , 2019, 24, 2031-2041.	2.3	19
24	Evaluation of a personalised adherence intervention to improve photoprotection in adults with Xeroderma Pigmentosum (XP): protocol for the trial of XPAND. <i>BMJ Open</i> , 2019, 9, e028577.	1.9	5
25	Organ donation attitudes and general self-efficacy: exploratory views from a rural primary care setting. <i>Rural and Remote Health</i> , 2019, 19, 5241.	0.5	7
26	Bereaved donor families' experiences of organ and tissue donation, and perceived influences on their decision making. <i>Journal of Critical Care</i> , 2018, 45, 82-89.	2.2	37
27	The effectiveness of a group self-management education course for adults with poorly controlled epilepsy, SMILE (UK): A randomized controlled trial. <i>Epilepsia</i> , 2018, 59, 1048-1061.	5.1	20
28	A Population-Based Conceptual Framework for Evaluating the Role of Healthcare Services in Place of Death. <i>Healthcare (Switzerland)</i> , 2018, 6, 107.	2.0	13
29	Self-Management education for adults with poorly controlled epilepsy [SMILE (UK)]: a randomised controlled trial. <i>Health Technology Assessment</i> , 2018, 22, 1-142.	2.8	23
30	People with epilepsy obtain added value from education in groups: results of a qualitative study. <i>European Journal of Neurology</i> , 2017, 24, 609-616.	3.3	15
31	Adjustment modes in the trajectory of progressive multiple sclerosis: a qualitative study and conceptual model. <i>Psychology and Health</i> , 2017, 32, 343-360.	2.2	13
32	Developing and assessing the acceptability of an epilepsy first aid training intervention for patients who visit UK emergency departments: A multi-method study of patients and professionals. <i>Epilepsy and Behavior</i> , 2017, 68, 177-185.	1.7	6
33	An investigation of the predictors of photoprotection and UVR dose to the face in patients with XP: a protocol using observational mixed methods. <i>BMJ Open</i> , 2017, 7, e018364.	1.9	15
34	PO051...Seizure first aid training for people with epilepsy attending emergency departments, and informal carers. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2017, 88, A25.2-A25.	1.9	1
35	Interventions that improve maternity care for immigrant women in the UK: protocol for a narrative synthesis systematic review. <i>BMJ Open</i> , 2017, 7, e016988.	1.9	5
36	Quantifying benefit-risk preferences for new medicines in rare disease patients and caregivers. <i>Orphanet Journal of Rare Diseases</i> , 2016, 11, 70.	2.7	39

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37	“I’ll be in a safe place”: a qualitative study of the decisions taken by people with advanced cancer to seek emergency department care. <i>BMJ Supportive and Palliative Care</i> , 2016, 6, 394.2-394.	1.6	3
38	“Nudging” Deceased Donation Through an Opt-Out System: A Libertarian Approach or Manipulation?. <i>American Journal of Bioethics</i> , 2016, 16, 25-28.	0.9	5
39	“I’ll be in a safe place”: a qualitative study of the decisions taken by people with advanced cancer to seek emergency department care. <i>BMJ Open</i> , 2016, 6, e012134.	1.9	45
40	Understanding kidney transplant patients' treatment choices: The interaction of emotion with medical and social influences on risk preferences. <i>Social Science and Medicine</i> , 2016, 155, 43-50.	3.8	12
41	Factors Associated with Transition from Community Settings to Hospital as Place of Death for Adults Aged 75 and Older: A Population-Based Mortality Follow-Back Survey. <i>Journal of the American Geriatrics Society</i> , 2016, 64, 2210-2217.	2.6	34
42	Developing a model of short-term integrated palliative and supportive care for frail older people in community settings: perspectives of older people, carers and other key stakeholders. <i>Age and Ageing</i> , 2016, 45, 863-873.	1.6	70
43	Patients’ experiences of a behavioural intervention for migraine headache: a qualitative study. <i>Journal of Headache and Pain</i> , 2016, 17, 16.	6.0	18
44	Food-related Quality of Life in Inflammatory Bowel Disease: Development and Validation of a Questionnaire. <i>Journal of Crohn's and Colitis</i> , 2016, 10, 194-201.	1.3	40
45	Increasing the acceptability and rates of organ donation among minority ethnic groups: a programme of observational and evaluative research on Donation, Transplantation and Ethnicity (DonaTE). <i>Programme Grants for Applied Research</i> , 2016, 4, 1-196.	1.0	20
46	The value of uncertainty in critical illness? An ethnographic study of patterns and conflicts in care and decision-making trajectories. <i>BMC Anesthesiology</i> , 2015, 16, 11.	1.8	32
47	Perceptions and experiences of epilepsy among patients from black ethnic groups in South London. <i>Primary Health Care Research and Development</i> , 2015, 16, 450-460.	1.2	5
48	“Nudging” registration as an organ donor: Implications of changes in choice contexts for socio-cultural groups. <i>Current Sociology</i> , 2015, 63, 714-728.	1.4	14
49	“Seizure First Aid Training” for people with epilepsy who attend emergency departments, and their family and friends: study protocol for intervention development and a pilot randomised controlled trial. <i>BMJ Open</i> , 2015, 5, e009040.	1.9	17
50	Immigrant women’s experiences of maternity-care services in Canada: a systematic review using a narrative synthesis. <i>Systematic Reviews</i> , 2015, 4, 13.	5.3	68
51	Self-management education for adults with poorly controlled epilepsy (SMILE (UK)): statistical, economic and qualitative analysis plan for a randomised controlled trial. <i>Trials</i> , 2015, 16, 269.	1.6	16
52	Organ Donation and New Policies. <i>JAMA Internal Medicine</i> , 2015, 175, 1999.	5.1	1
53	“It doesn't do the care for you”: a qualitative study of health care professionals' perceptions of the benefits and harms of integrated care pathways for end of life care: Figure 1. <i>BMJ Open</i> , 2015, 5, e008242.	1.9	28
54	Self-management for people with poorly controlled epilepsy: Participants' views of the UK Self-Management in epilePSy (SMILE) program. <i>Epilepsy and Behavior</i> , 2015, 52, 159-164.	1.7	22

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55	Organ Donation Knowledge and Attitudes among Health Science Students in Greece: Emerging Interprofessional Needs. <i>International Journal of Medical Sciences</i> , 2014, 11, 634-640.	2.5	44
56	SELF-MANAGEMENT EDUCATION FOR EPILEPSY: AN RCT PROTOCOL. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2014, 85, e4.190-e4.	1.9	2
57	Working to establish "normality" post-transplant: A qualitative study of kidney transplant patients. <i>Chronic Illness</i> , 2014, 10, 247-258.	1.5	26
58	AN INTERVENTION TRIAL FOR PEOPLE WITH EPILEPSY ATTENDING A&E. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2014, 85, e4.189-e4.	1.9	0
59	OPTIMISING PALLIATIVE CARE FOR FRAIL ELDERLY IN COMMUNITY SETTINGS; PREFERENCES FOR CARE AND PALLIATIVE OUTCOMES IN THE LAST MONTHS OF LIFE. <i>BMJ Supportive and Palliative Care</i> , 2014, 4, 111.4-112.	1.6	2
60	PATIENTS' IDEAS AND CHOICES ABOUT THE MANAGEMENT OF HEADACHE. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2014, 85, e4.10-e4.	1.9	1
61	Self-Management education for adults with poorly controlled epilepsy (SMILE (UK)): a randomised controlled trial protocol. <i>BMC Neurology</i> , 2014, 14, 69.	1.8	30
62	Headache Diagnosis in Primary Care. <i>Journal of Neurology and Neurosurgery</i> , 2014, 01, .	0.3	1
63	Headache: The Patient's View. <i>Journal of Neurology and Neurosurgery</i> , 2014, 01, .	0.3	1
64	Meanings of happiness among two ethnic groups living with advanced cancer in south London: a qualitative study. <i>Psycho-Oncology</i> , 2013, 22, 1096-1103.	2.3	16
65	Evaluating complex interventions in End of Life Care: the MORECare Statement on good practice generated by a synthesis of transparent expert consultations and systematic reviews. <i>BMC Medicine</i> , 2013, 11, 111.	5.5	256
66	A nurse-led self-management intervention for people who attend emergency departments with epilepsy: the patients' view. <i>Journal of Neurology</i> , 2013, 260, 1022-1030.	3.6	37
67	Development and evaluation of the feasibility and effects on staff, patients, and families of a new tool, the Psychosocial Assessment and Communication Evaluation (PACE), to improve communication and palliative care in intensive care and during clinical uncertainty. <i>BMC Medicine</i> , 2013, 11, 213.	5.5	41
68	The ideas of people referred to neurologists about managing their headaches: A qualitative study. <i>Journal of Headache and Pain</i> , 2013, 14, .	6.0	0
69	Immigrant women's experiences of postpartum depression in Canada: a protocol for systematic review using a narrative synthesis. <i>Systematic Reviews</i> , 2013, 2, 65.	5.3	17
70	Attitudes to deceased organ donation and registration as a donor among minority ethnic groups in North America and the UK: a synthesis of quantitative and qualitative research. <i>Ethnicity and Health</i> , 2013, 18, 367-390.	2.5	69
71	"Having a different conversation around death": diverse hospital chaplains' views on end-of-life care. <i>Ethnicity and Health</i> , 2013, 18, 530-543.	2.5	6
72	What are effective approaches to increasing rates of organ donor registration among ethnic minority populations: a systematic review. <i>BMJ Open</i> , 2013, 3, e003453.	1.9	39

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73	Co-€construction of chronic illness narratives by older stroke survivors and their spouses. <i>Sociology of Health and Illness</i> , 2013, 35, 993-1007.	2.1	34
74	PWE-092-€...Psychosocial Impact of Food and Nutrition in People with IBD: A Qualitative Study. <i>Gut</i> , 2013, 62, A168.1-A168.	12.1	4
75	Shifting towards an Opt-Out System in Greece: A General Practice Based Pilot Study. <i>International Journal of Medical Sciences</i> , 2013, 10, 1547-1551.	2.5	16
76	Can an epilepsy nurse specialist-led self-management intervention reduce attendance at emergency departments and promote well-being for people with severe epilepsy? A non-randomised trial with a nested qualitative phase. <i>Health Services and Delivery Research</i> , 2013, 1, 1-128.	1.4	10
77	053-€...The characteristics of emergency attendees for epilepsy in London hospitals. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2012, 83, e1.219-e1.	1.9	1
78	Explanations given by people with epilepsy for using emergency medical services: A qualitative study. <i>Epilepsy and Behavior</i> , 2012, 25, 529-533.	1.7	38
79	Kidney organ donation knowledge and attitudes among health care professionals: Findings from a Greek general hospital. <i>Applied Nursing Research</i> , 2012, 25, 283-290.	2.2	11
80	Immigrant women-€™s experiences of maternity-care services in Canada: a protocol for systematic review using a narrative synthesis. <i>Systematic Reviews</i> , 2012, 1, 27.	5.3	11
81	-€™The greatest thing in the world is the family-€™: the meaning of social support among Black Caribbean and White British patients living with advanced cancer. <i>Psycho-Oncology</i> , 2012, 21, 400-408.	2.3	19
82	Implementing -€™translational-€™ biomedical research: Convergence and divergence among clinical and basic scientists. <i>Social Science and Medicine</i> , 2011, 73, 945-952.	3.8	37
83	Service use and costs for people with headache: a UK primary care study. <i>Journal of Headache and Pain</i> , 2011, 12, 617-623.	6.0	32
84	Handwritten 'post-it' notes, questionnaire formats and response to a postal questionnaire survey. <i>International Journal of Epidemiology</i> , 2011, 40, 254-255.	1.9	4
85	Evaluating meta-ethnography: systematic analysis and synthesis of qualitative research. <i>Health Technology Assessment</i> , 2011, 15, 1-164.	2.8	417
86	Relational and management continuity survey in patients with multiple long-term conditions. <i>Journal of Health Services Research and Policy</i> , 2011, 16, 67-74.	1.7	45
87	How far can Foucault take us? An analysis of the changing discourses and limitations of the medical treatment of apoplexy and stroke. <i>Health (United Kingdom)</i> , 2011, 15, 369-384.	1.5	3
88	Compromised food access in hospital among older patients and those with multiple morbidity: the results from a survey of four hospitals. <i>Proceedings of the Nutrition Society</i> , 2010, 69, .	1.0	0
89	Ethnicity and attitudes to deceased kidney donation: a survey in Barbados and comparison with Black Caribbean people in the United Kingdom. <i>BMC Public Health</i> , 2010, 10, 266.	2.9	11
90	Kidney organ donation: developing family practice initiatives to reverse inertia. <i>BMC Health Services Research</i> , 2010, 10, 127.	2.2	6

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91	From gluttony to obesity: moral discourses on apoplexy and stroke. <i>Sociology of Health and Illness</i> , 2010, 32, 730-744.	2.1	4
92	Resisting psychotropic medicines: a synthesis of qualitative studies of medicine-taking. <i>Advances in Psychiatric Treatment</i> , 2010, 16, 207-218.	0.5	29
93	Experience of continuity of care of patients with multiple long-term conditions in England. <i>Journal of Health Services Research and Policy</i> , 2009, 14, 82-87.	1.7	60
94	Lay and Professional Constructions of Time: Implications for Illness Behaviour and Management of a Chronic Condition. <i>Sociology</i> , 2009, 43, 555-572.	2.5	11
95	Vulnerability in palliative care research: findings from a qualitative study of black Caribbean and white British patients with advanced cancer. <i>Journal of Medical Ethics</i> , 2009, 35, 440-444.	1.8	29
96	Attitudes to kidney donation among primary care patients in rural Crete, Greece. <i>BMC Public Health</i> , 2009, 9, 54.	2.9	18
97	Experiences of food access in hospital. A new questionnaire measure. <i>Clinical Nutrition</i> , 2009, 28, 625-630.	5.0	51
98	11.5 Ethnic minorities and indigenous peoples. , 2009, , .		0
99	7.1 Sociology and psychology in public health. , 2009, , .		0
100	Hospital inpatients's experiences of access to food: a qualitative interview and observational study. <i>Health Expectations</i> , 2008, 11, 294-303.	2.6	91
101	Ethnicity and registration as a kidney donor: The significance of identity and belonging. <i>Social Science and Medicine</i> , 2008, 66, 147-158.	3.8	39
102	“I know he controls cancer”: The meanings of religion among Black Caribbean and White British patients with advanced cancer. <i>Social Science and Medicine</i> , 2008, 67, 780-789.	3.8	68
103	Cultural meanings of pain: a qualitative study of Black Caribbean and White British patients with advanced cancer. <i>Palliative Medicine</i> , 2008, 22, 350-359.	3.1	74
104	Effect of social deprivation on blood pressure monitoring and control in England: a survey of data from the quality and outcomes framework. <i>BMJ: British Medical Journal</i> , 2008, 337, a2030-a2030.	2.3	89
105	Parental attitudes towards the management of asthma in ethnic minorities. <i>Archives of Disease in Childhood</i> , 2007, 92, 1082-1087.	1.9	34
106	Continuity of care and intermediate outcomes of type 2 diabetes mellitus. <i>Family Practice</i> , 2007, 24, 245-251.	1.9	53
107	Patient pressure for referral for headache: a qualitative study of GPs' referral behaviour. <i>British Journal of General Practice</i> , 2007, 57, 29-35.	1.4	54
108	How do patients referred to neurologists for headache differ from those managed in primary care?. <i>British Journal of General Practice</i> , 2007, 57, 388-95.	1.4	47

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109	What is 'continuity of care'?. Journal of Health Services Research and Policy, 2006, 11, 248-250.	1.7	207
110	Patients' perceptions and experiences of 'continuity of care' in diabetes. Health Expectations, 2006, 9, 118-129.	2.6	68
111	Attitudes to kidney donation and registering as a donor among ethnic groups in the UK. Journal of Public Health, 2006, 28, 226-234.	1.8	67
112	Measuring Continuity of Care in Diabetes Mellitus: An Experience-Based Measure. Annals of Family Medicine, 2006, 4, 548-555.	1.9	48
113	Resisting medicines: a synthesis of qualitative studies of medicine taking. Social Science and Medicine, 2005, 61, 133-155.	3.8	737
114	Length of stay for common surgical procedures: Variation among districts. British Journal of Surgery, 2005, 74, 884-889.	0.3	25
115	Barriers to Uptake and Adherence with Malaria Prophylaxis by the African Community in London, England: Focus Group Study. Ethnicity and Health, 2005, 10, 355-372.	2.5	64
116	Continuity of care. Family Medicine, 2005, 37, 687-8; author reply 688.	0.5	3
117	Women's views and experiences of outpatient hysteroscopy: Implications for a patient-centered service. Australian Journal of Cancer Nursing, 2004, 6, 315-320.	1.6	21
118	Implementing the 2-week wait rule for cancer referral in the UK: general practitioners' views and practices.. European Journal of Cancer Care, 2004, 13, 82-87.	1.5	34
119	Evaluating meta-ethnography: a synthesis of qualitative research on lay experiences of diabetes and diabetes care. Social Science and Medicine, 2003, 56, 671-684.	3.8	779
120	How can a nurse intervention help people with newly diagnosed epilepsy?. Seizure: the Journal of the British Epilepsy Association, 2003, 12, 69-73.	2.0	38
121	Human resources policies and continuity of care. Journal of Health Organization and Management, 2003, 17, 102-121.	1.3	11
122	Using meta ethnography to synthesise qualitative research: a worked example. Journal of Health Services Research and Policy, 2002, 7, 209-215.	1.7	833
123	Pathways to HIV testing and care by black African and white patients in London. Sexually Transmitted Infections, 2002, 78, 37-39.	1.9	57
124	Hearing children's voices: methodological issues in conducting focus groups with children aged 7-11 years. Qualitative Research, 2002, 2, 5-20.	3.5	276
125	What does 'access to health care' mean?. Journal of Health Services Research and Policy, 2002, 7, 186-188.	1.7	687
126	How can a nurse intervention help people with newly diagnosed epilepsy? A qualitative study (of) Tj ETQq0 0 0 rgBTJ/Overlock_10 Tf 50 6	2.0	13

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127	Endodermal cyst of the oculomotor nerve. <i>Neuroradiology</i> , 2001, 43, 1063-1066.	2.2	36
128	Ethnic Differences in Behavioural Risk Factors for Stroke: Implications for Health Promotion. <i>Ethnicity and Health</i> , 2001, 6, 95-103.	2.5	21
129	GPs' employment of locum doctors and satisfaction with their service. <i>Family Practice</i> , 2000, 17, 53-55.	1.9	6
130	Measuring process and outcomes: professional compliance, professional opinions and patients' well-being. <i>Family Practice</i> , 2000, 17, 21S-25.	1.9	4
131	Promoting self-care in epilepsy: the views of patients on the advice they had received from specialists, family doctors and an epilepsy nurse. <i>Patient Education and Counseling</i> , 1999, 37, 43-47.	2.2	47
132	Quality of midwifery led care: assessing the effects of different models of continuity for women's satisfaction. <i>Quality and Safety in Health Care</i> , 1998, 7, 77-82.	2.5	43
133	Midwifery group practices: what do hospital based midwives think?. <i>British Journal of Midwifery</i> , 1998, 6, 430-433.	0.4	1
134	Anomalous patients: the experiences of doctors with an illness.. <i>Sociology of Health and Illness</i> , 1997, 19, 644-667.	2.1	4
135	Sickness absence and 'working through' illness: a comparison of two professional groups. <i>Journal of Public Health</i> , 1997, 19, 295-300.	1.8	222
136	Illness doesn't belong to us. <i>Journal of the Royal Society of Medicine</i> , 1997, 90, 491-495.	2.0	60
137	Patients'™ perceptions of changes in their blood pressure. <i>Journal of Human Hypertension</i> , 1997, 11, 221-225.	2.2	19
138	The impact of chronic urticaria on the quality of life. <i>British Journal of Dermatology</i> , 1997, 136, 197-201.	1.5	124
139	Dermatology quality of life scales -a measure of the impact of skin diseases. <i>British Journal of Dermatology</i> , 1997, 136, 202-206.	1.5	69
140	Anomalous patients: the experiences of doctors with an illness. <i>Sociology of Health and Illness</i> , 1997, 19, 644-667.	2.1	25
141	Anomalous Patients: The Experiences of Doctors with an Illness. <i>Sociology of Health and Illness</i> , 1997, 19, 644-667.	2.1	3
142	The impact of chronic urticaria on the quality of life. <i>British Journal of Dermatology</i> , 1997, 136, 197-201.	1.5	419
143	Dermatology quality of life scales "a measure of the impact of skin diseases. <i>British Journal of Dermatology</i> , 1997, 136, 202-206.	1.5	97
144	Dermatology quality of life scales--a measure of the impact of skin diseases. <i>British Journal of Dermatology</i> , 1997, 136, 202-6.	1.5	36

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145	The SF-36 Health Survey Questionnaire: Is it Suitable for use with Older Adults?. Age and Ageing, 1995, 24, 120-125.	1.6	303
146	The Significance of Ethnicity for Health Promotion: Patients' use of Anti-Hypertensive Drugs in Inner London. International Journal of Epidemiology, 1995, 24, S79-S84.	1.9	51
147	Why do Mothers Consult when their Children Cough?. Family Practice, 1993, 10, 193-196.	1.9	101
148	Doctors' Interviewing Technique and Its Response to Different Booking Time. Family Practice, 1992, 9, 57-60.	1.9	22
149	Constraints on innovatory practice: The case of day surgery in the NHS. International Journal of Health Planning and Management, 1992, 7, 133-148.	1.7	10
150	Surgeons' views of day surgery: is there a consensus among providers?. Journal of Public Health Medicine, 1992, 14, 192-8.	0.4	4
151	Audit from preschool developmental surveillance of vision, hearing, and language referrals.. Archives of Disease in Childhood, 1991, 66, 921-926.	1.9	8
152	Variations in lengths of stay and rates of day case surgery: implications for the efficiency of surgical management.. Journal of Epidemiology and Community Health, 1990, 44, 90-105.	3.7	93
153	Changes in diet and coronary heart disease mortality among social classes in Great Britain.. Journal of Epidemiology and Community Health, 1989, 43, 162-167.	3.7	20
154	Referral patterns after school medical examinations.. Archives of Disease in Childhood, 1989, 64, 829-833.	1.9	6
155	Handbook of Life Stress, Cognition and Health (Book).. Sociology of Health and Illness, 1989, 11, 195-196.	2.1	0
156	Managing hypertension: beliefs and responses to medication among cultural groups.. Sociology of Health and Illness, 1988, 10, 561-578.	2.1	87
157	The role of information flow between health professionals and the Child Health Computer System in the uptake of measles immunisation. Journal of Public Health, 1988, 10, 40-47.	1.8	4
158	Can hospital use be a measure of need for health care?. Journal of Epidemiology and Community Health, 1987, 41, 269-274.	3.7	43
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