Myfanwy Morgan

List of Publications by Year in descending order

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174 papers 9,254 citations

43 h-index 92 g-index

177 all docs

177 docs citations

177 times ranked

10089 citing authors

#	Article	IF	CITATIONS
1	Using meta ethnography to synthesise qualitative research: a worked example. Journal of Health Services Research and Policy, 2002, 7, 209-215.	1.7	833
2	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
3	Resisting medicines: a synthesis of qualitative studies of medicine taking. Social Science and Medicine, 2005, 61, 133-155.	3.8	737
4	What does 'access to health care' mean?. Journal of Health Services Research and Policy, 2002, 7, 186-188.	1.7	687
5	The impact of chronic urticaria on the quality of life. British Journal of Dermatology, 1997, 136, 197-201.	1.5	419
6	Evaluating meta-ethnography: systematic analysis and synthesis of qualitative research. Health Technology Assessment, 2011, 15, 1-164.	2.8	417
7	The SF-36 Health Survey Questionnaire: Is it Suitable for use with Older Adults?. Age and Ageing, 1995, 24, 120-125.	1.6	303
8	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
9	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. BMC Medicine, 2013, 11, 111.	5.5	256
10	Sickness absence and 'working through' illness: a comparison of two professional groups. Journal of Public Health, 1997, 19, 295-300.	1.8	222
11	What is 'continuity of care'?. Journal of Health Services Research and Policy, 2006, 11, 248-250.	1.7	207
12	The impact of chronic urticaria on the quality of life. British Journal of Dermatology, 1997, 136, 197-201.	1.5	124
13	Why do Mothers Consult when their Children Cough?. Family Practice, 1993, 10, 193-196.	1.9	101
14	Dermatology quality of life scales –a measure of the impact of skin diseases. British Journal of Dermatology, 1997, 136, 202-206.	1.5	97
15	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
16	Hospital inpatients' experiences of access to food: a qualitative interview and observational study. Health Expectations, 2008, 11, 294-303.	2.6	91
17	Effect of social deprivation on blood pressure monitoring and control in England: a survey of data from the quality and outcomes framework. BMJ: British Medical Journal, 2008, 337, a2030-a2030.	2.3	89
18	Managing hypertension: beliefs and responses to medication among cultural groups Sociology of Health and Illness, 1988, 10, 561-578.	2.1	87

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19	Cultural meanings of pain: a qualitative study of Black Caribbean and White British patients with advanced cancer. Palliative Medicine, 2008, 22, 350-359.	3.1	74
20	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. Age and Ageing, 2016, 45, 863-873.	1.6	70
21	Dermatology quality of life scales -a measure of the impact of skin diseases. British Journal of Dermatology, 1997, 136, 202-206.	1.5	69
22	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. Ethnicity and Health, 2013, 18, 367-390.	2.5	69
23	Patients' perceptions and experiences of 'continuity of care' in diabetes. Health Expectations, 2006, 9, 118-129.	2.6	68
24	"l know he controls cancer†The meanings of religion among Black Caribbean and White British patients with advanced cancer. Social Science and Medicine, 2008, 67, 780-789.	3.8	68
25	Immigrant women's experiences of maternity-care services in Canada: a systematic review using a narrative synthesis. Systematic Reviews, 2015, 4, 13.	5.3	68
26	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
27	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
28	Illness doesn't belong to us. Journal of the Royal Society of Medicine, 1997, 90, 491-495.	2.0	60
29	Experience of continuity of care of patients with multiple long-term conditions in England. Journal of Health Services Research and Policy, 2009, 14, 82-87.	1.7	60
30	Psychosocial support and change in the health status of physically disabled people. Social Science and Medicine, 1986, 22, 1347-1354.	3.8	59
31	Perceptions and psychosocial impact of food, nutrition, eating and drinking in people with inflammatory bowel disease: a qualitative investigation of foodâ€related quality of life. Journal of Human Nutrition and Dietetics, 2020, 33, 115-127.	2.5	58
32	Pathways to HIV testing and care by black African and white patients in London. Sexually Transmitted Infections, 2002, 78, 37-39.	1.9	57
33	Ratio of plasma alpha amino-n-butyric acid to leucine as an empirical marker of alcoholism: diagnostic value. Science, 1977, 197, 1183-1185.	12.6	55
34	ACORN group, social class, and child health Journal of Epidemiology and Community Health, 1983, 37, 196-203.	3.7	54
35	Patient pressure for referral for headache: a qualitative study of GPs' referral behaviour. British Journal of General Practice, 2007, 57, 29-35.	1.4	54
36	Continuity of care and intermediate outcomes of type 2 diabetes mellitus. Family Practice, 2007, 24, 245-251.	1.9	53

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37	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
38	Experiences of food access in hospital. A new questionnaire measure. Clinical Nutrition, 2009, 28, 625-630.	5.0	51
39	Measuring Continuity of Care in Diabetes Mellitus: An Experience-Based Measure. Annals of Family Medicine, 2006, 4, 548-555.	1.9	48
40	Promoting self-care in epilepsy: the views of patients on the advice they had received from specialists, family doctors and an epilepsy nurse. Patient Education and Counseling, 1999, 37, 43-47.	2.2	47
41	How do patients referred to neurologists for headache differ from those managed in primary care?. British Journal of General Practice, 2007, 57, 388-95.	1.4	47
42	Relational and management continuity survey in patients with multiple long-term conditions. Journal of Health Services Research and Policy, 2011, 16, 67-74.	1.7	45
43	â€īl'll be in a safe place': a qualitative study of the decisions taken by people with advanced cancer to seek emergency department care. BMJ Open, 2016, 6, e012134.	1.9	45
44	Social networks and psychosocial support among disabled people. Social Science and Medicine, 1984, 19, 489-497.	3.8	44
45	Organ Donation Knowledge and Attitudes among Health Science Students in Greece: Emerging Interprofessional Needs. International Journal of Medical Sciences, 2014, 11, 634-640.	2.5	44
46	Can hospital use be a measure of need for health care?. Journal of Epidemiology and Community Health, 1987, 41, 269-274.	3.7	43
47	Quality of midwifery led care: assessing the effects of different models of continuity for women's satisfaction. Quality and Safety in Health Care, 1998, 7, 77-82.	2.5	43
48	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. BMC Medicine, 2013, 11, 213.	5.5	41
49	Food-related Quality of Life in Inflammatory Bowel Disease: Development and Validation of a Questionnaire. Journal of Crohn's and Colitis, 2016, 10, 194-201.	1.3	40
50	Experience of and access to maternity care in the UK by immigrant women: a narrative synthesis systematic review. BMJ Open, 2019, 9, e029478.	1.9	40
51	Ethnicity and registration as a kidney donor: The significance of identity and belonging. Social Science and Medicine, 2008, 66, 147-158.	3.8	39
52	What are effective approaches to increasing rates of organ donor registration among ethnic minority populations: a systematic review. BMJ Open, 2013, 3, e003453.	1.9	39
53	Quantifying benefit-risk preferences for new medicines in rare disease patients and caregivers. Orphanet Journal of Rare Diseases, $2016, 11, 70$.	2.7	39
54	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

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55	Explanations given by people with epilepsy for using emergency medical services: A qualitative study. Epilepsy and Behavior, 2012, 25, 529-533.	1.7	38
56	Implementing â€~translational' biomedical research: Convergence and divergence among clinical and basic scientists. Social Science and Medicine, 2011, 73, 945-952.	3.8	37
57	A nurse-led self-management intervention for people who attend emergency departments with epilepsy: the patients' view. Journal of Neurology, 2013, 260, 1022-1030.	3.6	37
58	Bereaved donor families' experiences of organ and tissue donation, and perceived influences on their decision making. Journal of Critical Care, 2018, 45, 82-89.	2.2	37
59	Endodermal cyst of the oculomotor nerve. Neuroradiology, 2001, 43, 1063-1066.	2.2	36
60	Dermatology quality of life scales-a measure of the impact of skin diseases. British Journal of Dermatology, 1997, 136, 202-6.	1.5	36
61	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
62	Parental attitudes towards the management of asthma in ethnic minorities. Archives of Disease in Childhood, 2007, 92, 1082-1087.	1.9	34
63	Coâ€construction of chronic illness narratives by older stroke survivors and their spouses. Sociology of Health and Illness, 2013, 35, 993-1007.	2.1	34
64	Factors Associated with Transition from Community Settings to Hospital as Place of Death for Adults Aged 75 and Older: A Populationâ€Based Mortality Followâ€Back Survey. Journal of the American Geriatrics Society, 2016, 64, 2210-2217.	2.6	34
65	Service use and costs for people with headache: a UK primary care study. Journal of Headache and Pain, 2011, 12, 617-623.	6.0	32
66	The value of uncertainty in critical illness? An ethnographic study of patterns and conflicts in care and decision-making trajectories. BMC Anesthesiology, 2015, 16, 11.	1.8	32
67	Self-Management education for adults with poorly controlled epILEpsy (SMILE (UK)): a randomised controlled trial protocol. BMC Neurology, 2014, 14, 69.	1.8	30
68	Vulnerability in palliative care research: findings from a qualitative study of black Caribbean and white British patients with advanced cancer. Journal of Medical Ethics, 2009, 35, 440-444.	1.8	29
69	Resisting psychotropic medicines: a synthesis of qualitative studies of medicine-taking. Advances in Psychiatric Treatment, 2010, 16, 207-218.	0.5	29
70	â€ît 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: FigureÂ1. BMJ Open, 2015, 5, e008242.	1.9	28
71	Working to establish â€`normality' post-transplant: A qualitative study of kidney transplant patients. Chronic Illness, 2014, 10, 247-258.	1.5	26
72	Food-related quality of life is impaired in inflammatory bowel disease and associated with reduced intake of key nutrients. American Journal of Clinical Nutrition, 2021, 113, 832-844.	4.7	26

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73	Length of stay for common surgical procedures: Variation among districts. British Journal of Surgery, 2005, 74, 884-889.	0.3	25
74	Anomalous patients: the experiences of doctors with an illness. Sociology of Health and Illness, 1997, 19, 644-667.	2.1	25
75	Public knowledge and attitudes towards consent policies for organ donation in Europe. A systematic review. Transplantation Reviews, 2019, 33, 1-8.	2.9	25
76	Measuring social inequality: occupational classifications and their alternatives. Community Medicine, 1983, 5, 116-24.	0.4	24
77	Self-Management education for adults with poorly controlled epILEpsy [SMILE (UK)]: a randomised controlled trial. Health Technology Assessment, 2018, 22, 1-142.	2.8	23
78	Marital status, health, illness andservice use. Social Science & Medicine Part A, Medical Sociology, 1980, 14, 633-643.	0.1	22
79	Doctors' Interviewing Technique and Its Response to Different Booking Time. Family Practice, 1992, 9, 57-60.	1.9	22
80	Self-management for people with poorly controlled epilepsy: Participants' views of the UK Self-Management in epILEpsy (SMILE) program. Epilepsy and Behavior, 2015, 52, 159-164.	1.7	22
81	Ethnic Differences in Behavioural Risk Factors for Stroke: Implications for Health Promotion. Ethnicity and Health, 2001, 6, 95-103.	2.5	21
82	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
83	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
84	The effectiveness of a group selfâ€management education course for adults with poorly controlled epilepsy, SMILE (UK): A randomized controlled trial. Epilepsia, 2018, 59, 1048-1061.	5.1	20
85	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). Programme Grants for Applied Research, 2016, 4, 1-196.	1.0	20
86	Patients' perceptions of changes in their blood pressure. Journal of Human Hypertension, 1997, 11, 221-225.	2.2	19
87	†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. Psycho-Oncology, 2012, 21, 400-408.	2.3	19
88	Experiences of stigma over the lifetime of people with xeroderma pigmentosum: A qualitative interview study in the United Kingdom. Journal of Health Psychology, 2019, 24, 2031-2041.	2.3	19
89	Attitudes to kidney donation among primary care patients in rural Crete, Greece. BMC Public Health, 2009, 9, 54.	2.9	18
90	Patients' experiences of a behavioural intervention for migraine headache: a qualitative study. Journal of Headache and Pain, 2016, 17, 16.	6.0	18

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91	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. International Journal of Nursing Studies, 2021, 120, 103978.	5.6	18
92	Immigrant women's experiences of postpartum depression in Canada: a protocol for systematic review using a narrative synthesis. Systematic Reviews, 2013, 2, 65.	5. 3	17
93	â€~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. BMJ Open, 2015, 5, e009040.	1.9	17
94	Measles immunisation: feasibility of a 90% target uptake Archives of Disease in Childhood, 1987, 62, 1209-1214.	1.9	16
95	Meanings of happiness among two ethnic groups living with advanced cancer in south London: a qualitative study. Psycho-Oncology, 2013, 22, 1096-1103.	2.3	16
96	Shifting towards an Opt-Out System in Greece: A General Practice Based Pilot Study. International Journal of Medical Sciences, 2013, 10, 1547-1551.	2.5	16
97	Self-management education for adults with poorly controlled epilepsy (SMILE (UK)): statistical, economic and qualitative analysis plan for a randomised controlled trial. Trials, 2015, 16, 269.	1.6	16
98	People with epilepsy obtain added value from education in groups: results of a qualitative study. European Journal of Neurology, 2017, 24, 609-616.	3.3	15
99	An investigation of the predictors of photoprotection and UVR dose to the face in patients with XP: a protocol using observational mixed methods. BMJ Open, 2017, 7, e018364.	1.9	15
100	â€~Nudging' registration as an organ donor: Implications of changes in choice contexts for socio-cultural groups. Current Sociology, 2015, 63, 714-728.	1.4	14
101	How can a nurse intervention help people with newly diagnosed epilepsy? A qualitative study (of) Tj ETQq $1\ 1\ 0.7$	'84314 rgE 2.0	3T <u>/</u> gverlock
102	Adjustment modes in the trajectory of progressive multiple sclerosis: a qualitative study and conceptual model. Psychology and Health, 2017, 32, 343-360.	2.2	13
103	A Population-Based Conceptual Framework for Evaluating the Role of Healthcare Services in Place of Death. Healthcare (Switzerland), 2018, 6, 107.	2.0	13
104	Understanding kidney transplant patients' treatment choices: The interaction of emotion with medical and social influences on risk preferences. Social Science and Medicine, 2016, 155, 43-50.	3.8	12
105	Sociological Approaches to Health and Medicine. Canadian Journal of Sociology, 1986, 11, 473.	0.2	11
106	Human resources policies and continuity of care. Journal of Health Organization and Management, 2003, 17, 102-121.	1.3	11
107	Lay and Professional Constructions of Time: Implications for Illness Behaviour and Management of a Chronic Condition. Sociology, 2009, 43, 555-572.	2.5	11
108	Ethnicity and attitudes to deceased kidney donation: a survey in Barbados and comparison with Black Caribbean people in the United Kingdom. BMC Public Health, 2010, 10, 266.	2.9	11

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109	Kidney organ donation knowledge and attitudes among health care professionals: Findings from a Greek general hospital. Applied Nursing Research, 2012, 25, 283-290.	2.2	11
110	Immigrant women's experiences of maternity-care services in Canada: a protocol for systematic review using a narrative synthesis. Systematic Reviews, 2012, 1, 27.	5. 3	11
111	The influence of perceived medical risks and psychosocial concerns on photoprotection behaviours among adults with xeroderma pigmentosum: a qualitative interview study in the UK. BMJ Open, 2019, 9, e024445.	1.9	11
112	Primary care service use by end-of-life cancer patients: a nationwide population-based cohort study in the United Kingdom. BMC Family Practice, 2020, 21, 76.	2.9	11
113	Measuring social inequality: Occupational classifications and their alternatives. , 1983, 5, 116.		11
114	Marital status and hospital use Journal of Epidemiology and Community Health, 1977, 31, 192-198.	3.7	10
115	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
116	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. Health Services and Delivery Research, 2013, 1, 1-128.	1.4	10
117	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. Transplantation Reviews, 2022, 36, 100673.	2.9	9
118	Audit from preschool developmental surveillance of vision, hearing, and language referrals Archives of Disease in Childhood, 1991, 66, 921-926.	1.9	8
119	Developing patient-centred, feasible alternative care for adult emergency department users with epilepsy: protocol for the mixed-methods observational  Collaborate' project. BMJ Open, 2019, 9, e031696.	1.9	8
120	Forms, interactions, and responses to social support: A qualitative study of support and adherence to photoprotection amongst patients with Xeroderma Pigmentosum. British Journal of Health Psychology, 2020, 25, 89-106.	3. 5	8
121	Marital status, health, illness and service use. Social Science & Medicine Medical Psychology & Medical Sociology, 1980, 14, 633-643.	0.1	8
122	Organ donation attitudes and general self-efficacy: exploratory views from a rural primary care setting. Rural and Remote Health, 2019, 19, 5241.	0.5	7
123	Parents' attitudes to measles immunization. The Journal of the Royal College of General Practitioners, 1987, 37, 25-7.	0.3	7
124	Referral patterns after school medical examinations Archives of Disease in Childhood, 1989, 64, 829-833.	1.9	6
125	GPs' employment of locum doctors and satisfaction with their service. Family Practice, 2000, 17, 53-55.	1.9	6
126	Kidney organ donation: developing family practice initiatives to reverse inertia. BMC Health Services Research, 2010, 10, 127.	2.2	6

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127	†Having a different conversation around death': diverse hospital chaplains' views on end-of-life care. Ethnicity and Health, 2013, 18, 530-543.	2.5	6
128	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. Epilepsy and Behavior, 2017, 68, 177-185.	1.7	6
129	Why? What? How? Using an Intervention Mapping approach to develop a personalised intervention to improve adherence to photoprotection in patients with Xeroderma Pigmentosum. Health Psychology and Behavioral Medicine, 2020, 8, 475-500.	1.8	6
130	Perceptions and experiences of epilepsy among patients from black ethnic groups in South London. Primary Health Care Research and Development, 2015, 16, 450-460.	1.2	5
131	"Nudging―Deceased Donation Through an Opt-Out System: A Libertarian Approach or Manipulation?. American Journal of Bioethics, 2016, 16, 25-28.	0.9	5
132	Interventions that improve maternity care for immigrant women in the UK: protocol for a narrative synthesis systematic review. BMJ Open, 2017, 7, e016988.	1.9	5
133	Assessing Treatment Fidelity within an Epilepsy Randomized Controlled Trial: Seizure First Aid Training for People with Epilepsy Who Visit Emergency Departments. Behavioural Neurology, 2019, 2019, 1-11.	2.1	5
134	Approaches to Photoprotection and Normalization in Highly Adherent Families of Children With Xeroderma Pigmentosum in the United Kingdom. Qualitative Health Research, 2020, 30, 1275-1286.	2.1	5
135	Improving photoprotection in adults with xeroderma pigmentosum: personalisation and tailoring in the †XPAND' intervention. Health Psychology and Behavioral Medicine, 2020, 8, 543-572.	1.8	5
136	Patient views on use of emergency and alternative care services for adult epilepsy: A qualitative study. Seizure: the Journal of the British Epilepsy Association, 2020, 80, 56-62.	2.0	5
137	Evaluation of a personalised adherence intervention to improve photoprotection in adults with Xeroderma Pigmentosum (XP): protocol for the trial of XPAND. BMJ Open, 2019, 9, e028577.	1.9	5
138	Access to and interventions to improve maternity care services for immigrant women: a narrative synthesis systematic review. Health Services and Delivery Research, 2020, 8, 1-122.	1.4	5
139	Measuring social inequality: Occupational classifications and their alternatives. Journal of Public Health, 1983, 5, 116-124.	1.8	4
140	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
141	Anomalous patients: the experiences of doctors with an illness Sociology of Health and Illness, 1997, 19, 644-667.	2.1	4
142	Measuring process and outcomes: professional compliance, professional opinions and patients' well-being. Family Practice, 2000, 17, 21S-25.	1.9	4
143	From gluttony to obesity: moral discourses on apoplexy and stroke. Sociology of Health and Illness, 2010, 32, 730-744.	2.1	4
144	Handwritten 'post-it' notes, questionnaire formats and response to a postal questionnaire survey. International Journal of Epidemiology, 2011, 40, 254-255.	1.9	4

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145	PWE-092â€Psychosocial Impact of Food and Nutrition in People with IBD: A Qualitative Study. Gut, 2013, 62, A168.1-A168.	12.1	4
146	The Personal Sociability and Connections Scale (PeSCS): Development and initial assessment at a primary care facility. International Journal of Social Psychiatry, 2021, , 002076402199351.	3.1	4
147	Surgeons' views of day surgery: is there a consensus among providers?. Journal of Public Health Medicine, 1992, 14, 192-8.	0.4	4
148	Immigrant Families in an Industrial City: a Study of Households in Holyoke, 1880. Journal of Family History, 1979, 4, 59-68.	0.5	3
149	How far can Foucault take us? An analysis of the changing discourses and limitations of the medical treatment of apoplexy and stroke. Health (United Kingdom), 2011, 15, 369-384.	1.5	3
150	"l'll be in a safe place― a qualitative study of the decisions taken by people with advanced cancer to seek emergency department care. BMJ Supportive and Palliative Care, 2016, 6, 394.2-394.	1.6	3
151	Anomalous Patients: The Experiences of Doctors with an Illness. Sociology of Health and Illness, 1997, 19, 644-667.	2.1	3
152	Ultraviolet exposure to the face in patients with xeroderma pigmentosum and healthy controls: applying a novel methodology to define photoprotection behaviour. British Journal of Dermatology, 2021, , .	1.5	3
153	Continuity of care. Family Medicine, 2005, 37, 687-8; author reply 688.	0.5	3
154	SELF-MANAGEMENT EDUCATION FOR EPILEPSY: AN RCT PROTOCOL. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, e4.190-e4.	1.9	2
155	OPTIMISING PALLIATIVE CARE FOR FRAIL ELDERS IN COMMUNITY SETTINGS; PREFERENCES FOR CARE AND PALLIATIVE OUTCOMES IN THE LAST MONTHS OF LIFE. BMJ Supportive and Palliative Care, 2014, 4, 111.4-112.	1.6	2
156	Seizure first aid training for people with epilepsy attending emergency departments and their significant others: the SAFE intervention and feasibility RCT. Health Services and Delivery Research, 2020, 8, 1-190.	1.4	2
157	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. Journal of Medical Genetics, 2022, 59, 1095-1103.	3.2	2
158	Midwifery group practices: what do hospital based midwives think?. British Journal of Midwifery, 1998, 6, 430-433.	0.4	1
159	053â€The characteristics of emergency attendees for epilepsy in London hospitals. Journal of Neurology, Neurosurgery and Psychiatry, 2012, 83, e1.219-e1.	1.9	1
160	PATIENTS' IDEAS AND CHOICES ABOUT THE MANAGEMENT OF HEADACHE. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, e4.10-e4.	1.9	1
161	Organ Donation and New Policies. JAMA Internal Medicine, 2015, 175, 1999.	5.1	1
162	PO051â€Seizure first aid training for people with epilepsy attending emergency departments, and informal carers. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, A25.2-A25.	1.9	1

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163	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. BMJ Open, 2020, 10, e035516.	1.9	1
164	Headache Diagnosis in Primary Care. Journal of Neurology and Neurosurgery, 2014, 01, .	0.3	1
165	Headache: The Patient's View. Journal of Neurology and Neurosurgery, 2014, 01, .	0.3	1
166	Sociological Approaches to Health and Medicine Contemporary Sociology, 1987, 16, 409.	0.0	0
167	Social Support and Health (Book) Sociology of Health and Illness, 1987, 9, 98-99.	2.1	0
168	Handbook of Life Stress, Cognition and Health (Book) Sociology of Health and Illness, 1989, 11, 195-196.	2.1	0
169	Compromised food access in hospital among older patients and those with multiple morbidity: the results from a survey of four hospitals. Proceedings of the Nutrition Society, 2010, 69, .	1.0	0
170	The ideas of people referred to neurologists about managing their headaches: A qualitative study. Journal of Headache and Pain, 2013, 14, .	6.0	0
171	AN INTERVENTION TRIAL FOR PEOPLE WITH EPILEPSY ATTENDING A&E. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, e4.189-e4.	1.9	O
172	11.5 Ethnic minorities and indigenous peoples. , 2009, , .		0
173	7.1 Sociology and psychology in public health. , 2009, , .		0
174	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