

Phyllis Butow

List of Publications by Year in descending order

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Version: 2024-02-01

627
papers

35,052
citations

4345

89
h-index

8433

152
g-index

637
all docs

637
docs citations

637
times ranked

28104
citing authors

#	ARTICLE	IF	CITATIONS
1	Developing a quality criteria framework for patient decision aids: online international Delphi consensus process. <i>BMJ: British Medical Journal</i> , 2006, 333, 417-0.	2.4	1,373
2	What are the unmet supportive care needs of people with cancer? A systematic review. <i>Supportive Care in Cancer</i> , 2009, 17, 1117-1128.	1.0	729
3	Communicating With Realism and Hope: Incurable Cancer Patients' Views on the Disclosure of Prognosis. <i>Journal of Clinical Oncology</i> , 2005, 23, 1278-1288.	0.8	463
4	Sharing decisions in cancer care. <i>Social Science and Medicine</i> , 2001, 52, 1865-1878.	1.8	445
5	The dynamics of change: Cancer patients' preferences for information, involvement and support. <i>Annals of Oncology</i> , 1997, 8, 857-863.	0.6	441
6	A Systematic Review of Prognostic/End-of-Life Communication with Adults in the Advanced Stages of a Life-Limiting Illness: Patient/Caregiver Preferences for the Content, Style, and Timing of Information. <i>Journal of Pain and Symptom Management</i> , 2007, 34, 81-93.	0.6	437
7	Cancer Patient Preferences for Communication of Prognosis in the Metastatic Setting. <i>Journal of Clinical Oncology</i> , 2004, 22, 1721-1730.	0.8	428
8	Communicating prognosis in cancer care: a systematic review of the literature. <i>Annals of Oncology</i> , 2005, 16, 1005-1053.	0.6	394
9	Clinical trials in children. <i>Lancet, The</i> , 2004, 364, 803-811.	6.3	381
10	Truth-telling in discussing prognosis in advanced life-limiting illnesses: a systematic review. <i>Palliative Medicine</i> , 2007, 21, 507-517.	1.3	357
11	Psychological responses of patients receiving a diagnosis of cancer. <i>Annals of Oncology</i> , 2003, 14, 48-56.	0.6	347
12	From normal response to clinical problem: definition and clinical features of fear of cancer recurrence. <i>Supportive Care in Cancer</i> , 2016, 24, 3265-3268.	1.0	334
13	Randomized Controlled Trial of a Prompt List to Help Advanced Cancer Patients and Their Caregivers to Ask Questions About Prognosis and End-of-Life Care. <i>Journal of Clinical Oncology</i> , 2007, 25, 715-723.	0.8	318
14	When the diagnosis is cancer: Patient communication experiences and preferences. <i>Cancer</i> , 1996, 77, 2630-2637.	2.0	305
15	Physicianâ€“patientâ€“companion communication and decision-making: A systematic review of triadic medical consultations. <i>Patient Education and Counseling</i> , 2013, 91, 3-13.	1.0	304
16	When the Treatment Goal Is Not Cure: Are Cancer Patients Equipped to Make Informed Decisions?. <i>Journal of Clinical Oncology</i> , 2002, 20, 503-513.	0.8	296
17	A systematic review on communicating with patients about evidence. <i>Journal of Evaluation in Clinical Practice</i> , 2006, 12, 13-23.	0.9	282
18	Oncologists' reactions to cancer patients' verbal cues. <i>Psycho-Oncology</i> , 2002, 11, 47-58.	1.0	274

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19	The development and evaluation of a measure to assess cancer survivors' unmet supportive care needs: the CaSUN (Cancer Survivors' Unmet Needs measure). <i>Psycho-Oncology</i> , 2007, 16, 796-804.	1.0	268
20	Cancer Patient Disclosure and Patient-Doctor Communication of Complementary and Alternative Medicine Use: A Systematic Review. <i>Oncologist</i> , 2012, 17, 1475-1481.	1.9	265
21	A one-item question with a Likert or Visual Analog Scale adequately measured current anxiety. <i>Journal of Clinical Epidemiology</i> , 2007, 60, 356-360.	2.4	262
22	Clinical practice guidelines for communicating prognosis and end-of-life issues with adults in the advanced stages of a life-limiting illness, and their caregivers. <i>Medical Journal of Australia</i> , 2007, 186, S77-S105.	0.8	254
23	Effect of a Patient-Centered Communication Intervention on Oncologist-Patient Communication, Quality of Life, and Health Care Utilization in Advanced Cancer. <i>JAMA Oncology</i> , 2017, 3, 92-100.	3.4	249
24	Breast cancer survivors'™ supportive care needs 2-10 years after diagnosis. <i>Supportive Care in Cancer</i> , 2007, 15, 515-523.	1.0	242
25	Fostering coping and nurturing hope when discussing the future with terminally ill cancer patients and their caregivers. <i>Cancer</i> , 2005, 103, 1965-1975.	2.0	239
26	Effective health communication - a key factor in fighting the COVID-19 pandemic. <i>Patient Education and Counseling</i> , 2020, 103, 873-876.	1.0	239
27	How to recognize and manage psychological distress in cancer patients. <i>European Journal of Cancer Care</i> , 2005, 14, 7-15.	0.7	229
28	Hospital-based interventions: a systematic review of staff-reported barriers and facilitators to implementation processes. <i>Implementation Science</i> , 2018, 13, 36.	2.5	228
29	Randomized Clinical Trials in Oncology: Understanding and Attitudes Predict Willingness to Participate. <i>Journal of Clinical Oncology</i> , 2001, 19, 3554-3561.	0.8	221
30	Long-term survival from gynecologic cancer: Psychosocial outcomes, supportive care needs and positive outcomes. <i>Gynecologic Oncology</i> , 2007, 104, 381-389.	0.6	220
31	Communicating prognosis to patients with metastatic disease: what do they really want to know?. <i>Supportive Care in Cancer</i> , 2002, 10, 161-168.	1.0	219
32	Choosing between the EORTC QLQ-C30 and FACT-G for measuring health-related quality of life in cancer clinical research: issues, evidence and recommendations. <i>Annals of Oncology</i> , 2011, 22, 2179-2190.	0.6	219
33	Review of Adherence-Related Issues in Adolescents and Young Adults With Cancer. <i>Journal of Clinical Oncology</i> , 2010, 28, 4800-4809.	0.8	216
34	Psychosocial well-being and supportive care needs of cancer patients living in urban and rural/regional areas: a systematic review. <i>Supportive Care in Cancer</i> , 2012, 20, 1-22.	1.0	207
35	Patient participation in the cancer consultation: Evaluation of a question prompt sheet. <i>Annals of Oncology</i> , 1994, 5, 199-204.	0.6	196
36	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.	1.0	189

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37	Do patients want doctors to talk about spirituality? A systematic literature review. <i>Patient Education and Counseling</i> , 2015, 98, 1320-1328.	1.0	185
38	Survivorship care plans in cancer: a systematic review of care plan outcomes. <i>British Journal of Cancer</i> , 2014, 111, 1899-1908.	2.9	183
39	Effect of medical Qigong on cognitive function, quality of life, and a biomarker of inflammation in cancer patients: a randomized controlled trial. <i>Supportive Care in Cancer</i> , 2012, 20, 1235-1242.	1.0	179
40	Impact of Medical Qigong on quality of life, fatigue, mood and inflammation in cancer patients: a randomized controlled trial. <i>Annals of Oncology</i> , 2010, 21, 608-614.	0.6	176
41	A Research Agenda for Communication Between Health Care Professionals and Patients Living With Serious Illness. <i>JAMA Internal Medicine</i> , 2017, 177, 1361.	2.6	175
42	Fear of cancer recurrence in young women with a history of early-stage breast cancer: a cross-sectional study of prevalence and association with health behaviours. <i>Supportive Care in Cancer</i> , 2012, 20, 2651-2659.	1.0	168
43	Psychological outcomes and risk perception after genetic testing and counselling in breast cancer: a systematic review. <i>Medical Journal of Australia</i> , 2003, 178, 77-81.	0.8	166
44	Advance care planning for cancer patients: a systematic review of perceptions and experiences of patients, families, and healthcare providers. <i>Psycho-Oncology</i> , 2016, 25, 362-386.	1.0	166
45	Fear of cancer recurrence: a theoretical review and novel cognitive processing formulation. <i>Journal of Cancer Survivorship</i> , 2016, 10, 663-673.	1.5	166
46	Misunderstanding in cancer patients: Why shoot the messenger?. <i>Annals of Oncology</i> , 1999, 10, 39-46.	0.6	164
47	The characteristics and effectiveness of Question Prompt List interventions in oncology: a systematic review of the literature. <i>Psycho-Oncology</i> , 2015, 24, 245-252.	1.0	163
48	Psychological morbidity and quality of life of ethnic minority patients with cancer: a systematic review and meta-analysis. <i>Lancet Oncology</i> , The, 2011, 12, 1240-1248.	5.1	162
49	Computer-based interaction analysis of the cancer consultation. <i>British Journal of Cancer</i> , 1995, 71, 1115-1121.	2.9	157
50	Parents' attitudes to children's participation in randomized controlled trials. <i>Journal of Pediatrics</i> , 2003, 142, 554-559.	0.9	155
51	Clinical pathway for the screening, assessment and management of anxiety and depression in adult cancer patients: Australian guidelines. <i>Psycho-Oncology</i> , 2015, 24, 987-1001.	1.0	148
52	Randomized Trial of ConquerFear: A Novel, Theoretically Based Psychosocial Intervention for Fear of Cancer Recurrence. <i>Journal of Clinical Oncology</i> , 2017, 35, 4066-4077.	0.8	148
53	Communication styles in the cancer consultation: preferences for a patient-centred approach. , 2000, 9, 147-156.		147
54	Can a "œprompt list"empower cancer patients to ask relevant questions?. <i>Cancer</i> , 2008, 113, 225-237.	2.0	147

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55	Do personal stories make patient decision aids more effective? A critical review of theory and evidence. <i>BMC Medical Informatics and Decision Making</i> , 2013, 13, S9.	1.5	147
56	When and How to Initiate Discussion About Prognosis and End-of-Life Issues with Terminally Ill Patients. <i>Journal of Pain and Symptom Management</i> , 2005, 30, 132-144.	0.6	146
57	Psychological Responses and Coping Strategies Among Patients With Malignant Melanoma. <i>Archives of Dermatology</i> , 2009, 145, 1415-27.	1.7	146
58	Parental adjustment to the completion of their child's cancer treatment. <i>Pediatric Blood and Cancer</i> , 2011, 56, 524-531.	0.8	145
59	Making hard choices easier: a prospective, multicentre study to assess the efficacy of a fertility-related decision aid in young women with early-stage breast cancer. <i>British Journal of Cancer</i> , 2012, 106, 1053-1061.	2.9	144
60	Does Age Really Matter? Recall of Information Presented to Newly Referred Patients With Cancer. <i>Journal of Clinical Oncology</i> , 2008, 26, 5450-5457.	0.8	143
61	Symptom Clusters in Patients With Advanced Cancer: A Systematic Review of Observational Studies. <i>Journal of Pain and Symptom Management</i> , 2014, 48, 411-450.	0.6	142
62	Cancer Consultation Preparation Package: Changing Patients but Not Physicians Is Not Enough. <i>Journal of Clinical Oncology</i> , 2004, 22, 4401-4409.	0.8	140
63	Doctors discussing religion and spirituality: A systematic literature review. <i>Palliative Medicine</i> , 2016, 30, 327-337.	1.3	140
64	Analysis of cancer risk and BRCA1 and BRCA2 mutation prevalence in the kConFab familial breast cancer resource. <i>Breast Cancer Research</i> , 2006, 8, R12.	2.2	135
65	Epidemiological evidence for a relationship between life events, coping style, and personality factors in the development of breast cancer. <i>Journal of Psychosomatic Research</i> , 2000, 49, 169-181.	1.2	133
66	Increasing oncologists' skills in eliciting and responding to emotional cues: evaluation of a communication skills training program. <i>Psycho-Oncology</i> , 2008, 17, 209-218.	1.0	131
67	A review and recommendations for optimal outcome measures of anxiety, depression and general distress in studies evaluating psychosocial interventions for English-speaking adults with heterogeneous cancer diagnoses. <i>Supportive Care in Cancer</i> , 2010, 18, 1241-1262.	1.0	131
68	Asking questions can help: development and preliminary evaluation of a question prompt list for palliative care patients. <i>British Journal of Cancer</i> , 2003, 89, 2069-2077.	2.9	130
69	The needs of terminally ill cancer patients versus those of caregivers for information regarding prognosis and end-of-life issues. <i>Cancer</i> , 2005, 103, 1957-1964.	2.0	130
70	Discrepant Perceptions About End-of-Life Communication: A Systematic Review. <i>Journal of Pain and Symptom Management</i> , 2007, 34, 190-200.	0.6	130
71	Meeting the Decision-Making Preferences of Patients With Breast Cancer in Oncology Consultations: Impact on Decision-Related Outcomes. <i>Journal of Clinical Oncology</i> , 2012, 30, 857-862.	0.8	128
72	Psychosocial Predictors of Survival in Metastatic Melanoma. <i>Journal of Clinical Oncology</i> , 1999, 17, 2256-2256.	0.8	126

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73	Life after cancer: couples' and partners' psychological adjustment and supportive care needs. <i>Supportive Care in Cancer</i> , 2007, 15, 405-415.	1.0	121
74	Handling missing items in the Hospital Anxiety and Depression Scale (HADS): a simulation study. <i>BMC Research Notes</i> , 2016, 9, 479.	0.6	121
75	Psychological impact of genetic testing in women from high-risk breast cancer families. <i>European Journal of Cancer</i> , 2002, 38, 2025-2031.	1.3	120
76	Developing ethical strategies to assist oncologists in seeking informed consent to cancer clinical trials. <i>Social Science and Medicine</i> , 2004, 58, 379-390.	1.8	119
77	A systematic review of evidence for end-of-life communication interventions: Who do they target, how are they structured and do they work?. <i>Patient Education and Counseling</i> , 2016, 99, 3-16.	1.0	119
78	A review of the barriers to using Patient-Reported Outcomes (PROs) and Patient-Reported Outcome Measures (PROMs) in routine cancer care. <i>Journal of Medical Radiation Sciences</i> , 2021, 68, 186-195.	0.8	117
79	Promoting fruit and vegetable consumption. Testing an intervention based on the theory of planned behaviour. <i>Appetite</i> , 2012, 58, 997-1004.	1.8	113
80	Intention to Undergo Prophylactic Bilateral Mastectomy in Women at Increased Risk of Developing Hereditary Breast Cancer. <i>Journal of Clinical Oncology</i> , 2000, 18, 2250-2257.	0.8	112
81	Family involvement in cancer treatment decision-making: A qualitative study of patient, family, and clinician attitudes and experiences. <i>Patient Education and Counseling</i> , 2016, 99, 1146-1155.	1.0	111
82	What is important in cancer care coordination? A qualitative investigation. <i>European Journal of Cancer Care</i> , 2011, 20, 220-227.	0.7	109
83	Information booklets about cancer. <i>Patient Education and Counseling</i> , 1998, 33, 129-141.	1.0	108
84	Lay constructions of decision-making in cancer. <i>Psycho-Oncology</i> , 2002, 11, 295-306.	1.0	105
85	Evaluation of a novel individualised communication-skills training intervention to improve doctors' confidence and skills in end-of-life communication. <i>Palliative Medicine</i> , 2013, 27, 236-243.	1.3	104
86	A systematic review of decision aids for patients making a decision about treatment for early breast cancer. <i>Breast</i> , 2016, 26, 31-45.	0.9	104
87	The use of unproven methods of treatment by cancer patients. <i>Supportive Care in Cancer</i> , 1998, 6, 337-347.	1.0	101
88	Attitudes and information needs of Chinese migrant cancer patients and their relatives. <i>Australian and New Zealand Journal of Medicine</i> , 1999, 29, 207-213.	0.5	100
89	Hearing the bad news of a cancer diagnosis: The Australian melanoma patients' perspective. <i>Annals of Oncology</i> , 2001, 12, 365-371.	0.6	100
90	Family Perspectives on Deceased Organ Donation: Thematic Synthesis of Qualitative Studies. <i>American Journal of Transplantation</i> , 2014, 14, 923-935.	2.6	99

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91	Psychological intervention targeting distress for cancer patients: a meta-analytic study investigating uptake and adherence. <i>Psycho-Oncology</i> , 2016, 25, 882-890.	1.0	98
92	A matter of trust - patient's views on decision-making in colorectal cancer. <i>Health Expectations</i> , 2004, 7, 104-114.	1.1	96
93	Conquer fear: protocol of a randomised controlled trial of a psychological intervention to reduce fear of cancer recurrence. <i>BMC Cancer</i> , 2013, 13, 201.	1.1	92
94	Caring for women with ovarian cancer in the last year of life: A longitudinal study of caregiver quality of life, distress and unmet needs. <i>Gynecologic Oncology</i> , 2014, 132, 690-697.	0.6	92
95	The impact of communication on adherence in pain management. <i>Pain</i> , 2013, 154, S101-S107.	2.0	91
96	Psychoeducational Intervention to Reduce Fear of Cancer Recurrence in People at High Risk of Developing Another Primary Melanoma: Results of a Randomized Controlled Trial. <i>Journal of Clinical Oncology</i> , 2016, 34, 4405-4414.	0.8	91
97	Seeking informed consent to cancer clinical trials. <i>Social Science and Medicine</i> , 2004, 58, 2445-2457.	1.8	89
98	Advance Care Planning for Adults With CKD: A Systematic Integrative Review. <i>American Journal of Kidney Diseases</i> , 2014, 63, 761-770.	2.1	88
99	Comparison of group and individual cognitive-behavioral therapy for patients with bulimia nervosa. <i>International Journal of Eating Disorders</i> , 2003, 33, 241-254.	2.1	85
100	A Comparison of Nutritional Management with Stress Management in the Treatment of Bulimia Nervosa. <i>British Journal of Psychiatry</i> , 1991, 159, 250-261.	1.7	84
101	Fear of cancer recurrence and death anxiety. <i>Psycho-Oncology</i> , 2018, 27, 2559-2565.	1.0	84
102	Return to work after a cancer diagnosis: a meta-review of reviews and a meta-synthesis of recent qualitative studies. <i>Journal of Cancer Survivorship</i> , 2020, 14, 114-134.	1.5	84
103	Discussing Adjuvant Cancer Therapy. <i>Journal of Clinical Oncology</i> , 2001, 19, 1768-1778.	0.8	83
104	Genetic testing for melanoma risk: a prospective cohort study of uptake and outcomes among Australian families. <i>Genetics in Medicine</i> , 2009, 11, 265-278.	1.1	83
105	From inside the bubble: migrants' perceptions of communication with the cancer team. <i>Supportive Care in Cancer</i> , 2011, 19, 281-290.	1.0	83
106	Prevalence and predictors of anxiety and depression in women with invasive ovarian cancer and their caregivers. <i>Medical Journal of Australia</i> , 2010, 193, S52-7.	0.8	82
107	Discussing end-of-life issues with terminally ill cancer patients and their carers: a qualitative study. <i>Supportive Care in Cancer</i> , 2005, 13, 589-599.	1.0	81
108	Physician-Identified Factors Affecting Patient Participation in Reaching Treatment Decisions. <i>Journal of Clinical Oncology</i> , 2008, 26, 1724-1731.	0.8	81

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109	Family information needs at childhood cancer treatment completion. <i>Pediatric Blood and Cancer</i> , 2012, 58, 621-626.	0.8	81
110	Immigrants' perceptions of the quality of their cancer care: an Australian comparative study, identifying potentially modifiable factors. <i>Annals of Oncology</i> , 2014, 25, 1643-1649.	0.6	81
111	Encouraging early discussion of life expectancy and end-of-life care: A randomised controlled trial of a nurse-led communication support program for patients and caregivers. <i>International Journal of Nursing Studies</i> , 2017, 67, 31-40.	2.5	81
112	Communicating risk information: The influence of graphical display format on quantitative information perception—Accuracy, comprehension and preferences. <i>Patient Education and Counseling</i> , 2007, 69, 121-128.	1.0	80
113	Psychosocial predictors of survival: Metastatic breast cancer. <i>Annals of Oncology</i> , 2000, 11, 469-474.	0.6	79
114	Discussing life expectancy with terminally ill cancer patients and their carers: a qualitative study. <i>Supportive Care in Cancer</i> , 2005, 13, 733-742.	1.0	79
115	Anxiety in the context of cancer: A systematic review and development of an integrated model. <i>Clinical Psychology Review</i> , 2017, 56, 40-54.	6.0	79
116	A bridge between cultures: interpreters' perspectives of consultations with migrant oncology patients. <i>Supportive Care in Cancer</i> , 2012, 20, 235-244.	1.0	78
117	Long-term outcomes of genetic counseling in women at increased risk of developing hereditary breast cancer. <i>Patient Education and Counseling</i> , 2001, 44, 215-225.	1.0	76
118	The role of psychosocial factors in the development of breast carcinoma: Part II. <i>Cancer</i> , 2001, 91, 686-697.	2.0	76
119	Therapeutic Life Review in Palliative Care: A Systematic Review of Quantitative Evaluations. <i>Journal of Pain and Symptom Management</i> , 2015, 49, 747-761.	0.6	75
120	Adapting the nominal group technique for priority setting of evidence-practice gaps in implementation science. <i>BMC Medical Research Methodology</i> , 2016, 16, 110.	1.4	75
121	Can consumers learn to ask three questions to improve shared decision making? A feasibility study of the <sc>ASK</sc> (AskShareKnow) Patient—Clinician Communication Model [®] intervention in a primary health care setting. <i>Health Expectations</i> , 2016, 19, 1160-1168.	1.1	75
122	Exploring the screening capacity of the Fear of Cancer Recurrence Inventory—Short Form for clinical levels of fear of cancer recurrence. <i>Psycho-Oncology</i> , 2018, 27, 492-499.	1.0	75
123	On the receiving end IV: Validation of quality of life indicators. <i>Annals of Oncology</i> , 1991, 2, 597-603.	0.6	74
124	Pediatricians' attitudes toward randomized controlled trials involving children. <i>Journal of Pediatrics</i> , 2002, 141, 798-803.	0.9	74
125	Survivorship care after breast cancer treatment — Experiences and preferences of Australian women. <i>Breast</i> , 2011, 20, 271-277.	0.9	74
126	Promoting End-of-Life Discussions in Advanced Cancer: Effects of Patient Coaching and Question Prompt Lists. <i>Journal of Clinical Oncology</i> , 2017, 35, 842-851.	0.8	74

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127	Management of common clinical problems experienced by survivors of cancer. <i>Lancet, The</i> , 2022, 399, 1537-1550.	6.3	74
128	Prevalence and Predictors of Sleep Difficulty in a National Cohort of Women With Primary Breast Cancer Three to Four Months Postsurgery. <i>Journal of Pain and Symptom Management</i> , 2011, 42, 710-720.	0.6	73
129	Developing a roadmap for the translation of e-mental health services for depression. <i>Australian and New Zealand Journal of Psychiatry</i> , 2015, 49, 776-784.	1.3	73
130	Responding to the active and passive patient: flexibility is the key. <i>Health Expectations</i> , 2002, 5, 236-245.	1.1	72
131	Current approaches to managing fear of cancer recurrence; a descriptive survey of psychosocial and clinical health professionals. <i>Psycho-Oncology</i> , 2014, 23, 390-396.	1.0	72
132	When the safety net of treatment has been removed: Patients' unmet needs at the completion of treatment for haematological malignancies. <i>Patient Education and Counseling</i> , 2009, 77, 103-108.	1.0	71
133	Online group-based cognitive-behavioural therapy for adolescents and young adults after cancer treatment: A multicenter randomised controlled trial of Recapture Life-AYA. <i>BMC Cancer</i> , 2012, 12, 339.	1.1	71
134	Implementing patient question-prompt lists into routine cancer care. <i>Patient Education and Counseling</i> , 2012, 86, 252-258.	1.0	70
135	A question prompt list for patients with advanced cancer in the final year of life: Development and cross-cultural evaluation. <i>Palliative Medicine</i> , 2013, 27, 779-788.	1.3	69
136	Improving decision making about clinical trial participation – a randomised controlled trial of a decision aid for women considering participation in the IBIS-II breast cancer prevention trial. <i>British Journal of Cancer</i> , 2014, 111, 1-7.	2.9	69
137	Attitudes toward Prophylactic Oophorectomy and Screening Utilization in Women at Increased Risk of Developing Hereditary Breast/Ovarian Cancer. <i>Gynecologic Oncology</i> , 1999, 75, 122-129.	0.6	68
138	Health professional and consumer views on involving breast cancer patients in the multidisciplinary discussion of their disease and treatment plan. <i>Cancer</i> , 2007, 110, 1937-1944.	2.0	68
139	Medical Qigong for Cancer Patients: Pilot Study of Impact on Quality of Life, Side Effects of Treatment and Inflammation. <i>The American Journal of Chinese Medicine</i> , 2008, 36, 459-472.	1.5	68
140	Patient perspectives regarding communication about prognosis and end-of-life issues: How can it be optimised?. <i>Patient Education and Counseling</i> , 2013, 90, 307-314.	1.0	68
141	Psychological morbidity and stress but not social factors influence level of fear of cancer recurrence in young women with early breast cancer: results of a cross-sectional study. <i>Psycho-Oncology</i> , 2013, 22, 2797-2806.	1.0	68
142	Attentional bias and metacognitions in cancer survivors with high fear of cancer recurrence. <i>Psycho-Oncology</i> , 2015, 24, 416-423.	1.0	68
143	Symptom Clusters in Advanced Cancer Patients: An Empirical Comparison of Statistical Methods and the Impact on Quality of Life. <i>Journal of Pain and Symptom Management</i> , 2016, 51, 88-98.	0.6	68
144	Attitudes to randomized clinical trials amongst outpatients attending a medical oncology clinic. <i>Health Expectations</i> , 1999, 2, 33-43.	1.1	67

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145	Women's preferences for discussion of prognosis in early breast cancer. <i>Health Expectations</i> , 2001, 4, 48-57.	1.1	67
146	Towards better communication in cancer care: a framework for developing evidence-based interventions. <i>Patient Education and Counseling</i> , 2004, 55, 32-39.	1.0	67
147	A Qualitative Analysis of Reasons for Leaving, or Not Attending, a Cancer Support Group. <i>Social Work in Health Care</i> , 2008, 47, 14-29.	0.8	67
148	Grappling with cultural differences; Communication between oncologists and immigrant cancer patients with and without interpreters. <i>Patient Education and Counseling</i> , 2011, 84, 398-405.	1.0	67
149	The motherhood choice: Themes arising in the decision-making process for women with multiple sclerosis. <i>Multiple Sclerosis Journal</i> , 2008, 14, 701-704.	1.4	66
150	Returning to School After Adolescent Cancer: A Qualitative Examination of Australian Survivors' and Their Families' Perspectives. <i>Journal of Adolescent and Young Adult Oncology</i> , 2011, 1, 87-94.	0.7	66
151	Interpretation in Consultations With Immigrant Patients With Cancer: How Accurate Is It?. <i>Journal of Clinical Oncology</i> , 2011, 29, 2801-2807.	0.8	66
152	Current state and future prospects of research on fear of cancer recurrence. <i>Psycho-Oncology</i> , 2017, 26, 424-427.	1.0	66
153	Consultation audio tapes: an underused cancer patient information aid and clinical research tool. <i>Lancet Oncology</i> , The, 2002, 3, 431-437.	5.1	65
154	Analyzing the Process and Content of Genetic Counseling in Familial Breast Cancer Consultations. <i>Journal of Genetic Counseling</i> , 2004, 13, 403-424.	0.9	64
155	Improving doctors' letters. <i>Medical Journal of Australia</i> , 2002, 177, 516-520.	0.8	63
156	Changes in quality of life in patients with advanced cancer. <i>Journal of Psychosomatic Research</i> , 2005, 58, 497-504.	1.2	63
157	A Randomized Controlled Trial of a Standardized Educational Intervention for Patients with Cancer Pain. <i>Journal of Pain and Symptom Management</i> , 2010, 40, 49-59.	0.6	62
158	Sharing vs. caring—The relative impact of sharing decisions versus managing emotions on patient outcomes. <i>Patient Education and Counseling</i> , 2011, 82, 233-239.	1.0	62
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623	Predictors of the benefits women consider necessary to make adjuvant chemotherapy (ACT) worthwhile for early breast cancer (EBC). <i>Journal of Clinical Oncology</i> , 2004, 22, 787-787.	0.8	0
624	Reproductive Choices of Women With Multiple Sclerosis. <i>International Journal of MS Care</i> , 2007, 9, 9-12.	0.4	0
625	A pilot randomised controlled trial examining the feasibility, acceptability and impact of giving information on personalised genomic risk of melanoma to the public, for motivating preventive behaviours.. <i>Journal of Clinical Oncology</i> , 2016, 34, 1556-1556.	0.8	0
626	Body Image Scale: Evaluation of the Psychometric Properties in Three Indian Head and Neck Cancer Language Groups. <i>Frontiers in Psychology</i> , 2022, 13, .	1.1	0
627	Further validation of the Perceptions of Uncertainties in Genome Sequencing scale among patients with cancer undergoing tumor sequencing. <i>Clinical Genetics</i> , 2022, 102, 110-116.	1.0	0