

# Phyllis Butow

## List of Publications by Citations

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598  
papers

26,995  
citations

85  
h-index

136  
g-index

637  
ext. papers

31,201  
ext. citations

3.6  
avg, IF

7.11  
L-index

#	Paper	IF	Citations
598	Developing a quality criteria framework for patient decision aids: online international Delphi consensus process. <i>BMJ, The</i> , <b>2006</b> , 333, 417	5.9	1095
597	What are the unmet supportive care needs of people with cancer? A systematic review. <i>Supportive Care in Cancer</i> , <b>2009</b> , 17, 1117-28	3.9	587
596	Sharing decisions in cancer care. <i>Social Science and Medicine</i> , <b>2001</b> , 52, 1865-78	5.1	393
595	The dynamics of change: cancer patients' preferences for information, involvement and support. <i>Annals of Oncology</i> , <b>1997</b> , 8, 857-63	10.3	392
594	Communicating with realism and hope: incurable cancer patients' views on the disclosure of prognosis. <i>Journal of Clinical Oncology</i> , <b>2005</b> , 23, 1278-88	2.2	388
593	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> , <b>2007</b> , 34, 81-93	4.8	361
592	Cancer patient preferences for communication of prognosis in the metastatic setting. <i>Journal of Clinical Oncology</i> , <b>2004</b> , 22, 1721-30	2.2	351
591	Communicating prognosis in cancer care: a systematic review of the literature. <i>Annals of Oncology</i> , <b>2005</b> , 16, 1005-53	10.3	326
590	Clinical trials in children. <i>Lancet, The</i> , <b>2004</b> , 364, 803-11	40	308
589	Truth-telling in discussing prognosis in advanced life-limiting illnesses: a systematic review. <i>Palliative Medicine</i> , <b>2007</b> , 21, 507-17	5.5	284
588	Psychological responses of patients receiving a diagnosis of cancer. <i>Annals of Oncology</i> , <b>2003</b> , 14, 48-56	10.3	273
587	When the diagnosis is cancer: patient communication experiences and preferences. <i>Cancer</i> , <b>1996</b> , 77, 2630-7	6.4	267
586	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> , <b>2007</b> , 25, 715-23	2.2	257
585	When the treatment goal is not cure: are cancer patients equipped to make informed decisions?. <i>Journal of Clinical Oncology</i> , <b>2002</b> , 20, 503-13	2.2	256
584	Oncologists' reactions to cancer patients' verbal cues. <i>Psycho-Oncology</i> , <b>2002</b> , 11, 47-58	3.9	235
583	A systematic review on communicating with patients about evidence. <i>Journal of Evaluation in Clinical Practice</i> , <b>2006</b> , 12, 13-23	2.5	232
582	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> , <b>2007</b> , 16, 796-804	3.9	218

581	Physician-patient-companion communication and decision-making: a systematic review of triadic medical consultations. <i>Patient Education and Counseling</i> , <b>2013</b> , 91, 3-13	3.1	215
580	A one-item question with a Likert or Visual Analog Scale adequately measured current anxiety. <i>Journal of Clinical Epidemiology</i> , <b>2007</b> , 60, 356-60	5.7	215
579	Breast cancer survivors' supportive care needs 2-10 years after diagnosis. <i>Supportive Care in Cancer</i> , <b>2007</b> , 15, 515-23	3.9	211
578	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> , <b>2007</b> , 186, S77-S105	4	209
577	From normal response to clinical problem: definition and clinical features of fear of cancer recurrence. <i>Supportive Care in Cancer</i> , <b>2016</b> , 24, 3265-8	3.9	208
576	How to recognize and manage psychological distress in cancer patients. <i>European Journal of Cancer Care</i> , <b>2005</b> , 14, 7-15	2.4	197
575	Fostering coping and nurturing hope when discussing the future with terminally ill cancer patients and their caregivers. <i>Cancer</i> , <b>2005</b> , 103, 1965-75	6.4	195
574	Randomized clinical trials in oncology: understanding and attitudes predict willingness to participate. <i>Journal of Clinical Oncology</i> , <b>2001</b> , 19, 3554-61	2.2	195
573	Long-term survival from gynecologic cancer: psychosocial outcomes, supportive care needs and positive outcomes. <i>Gynecologic Oncology</i> , <b>2007</b> , 104, 381-9	4.9	191
572	Cancer patient disclosure and patient-doctor communication of complementary and alternative medicine use: a systematic review. <i>Oncologist</i> , <b>2012</b> , 17, 1475-81	5.7	189
571	Communicating prognosis to patients with metastatic disease: what do they really want to know?. <i>Supportive Care in Cancer</i> , <b>2002</b> , 10, 161-8	3.9	189
570	Review of adherence-related issues in adolescents and young adults with cancer. <i>Journal of Clinical Oncology</i> , <b>2010</b> , 28, 4800-9	2.2	174
569	Patient participation in the cancer consultation: evaluation of a question prompt sheet. <i>Annals of Oncology</i> , <b>1994</b> , 5, 199-204	10.3	171
568	Effect of a Patient-Centered Communication Intervention on Oncologist-Patient Communication, Quality of Life, and Health Care Utilization in Advanced Cancer: The VOICE Randomized Clinical Trial. <i>JAMA Oncology</i> , <b>2017</b> , 3, 92-100	13.4	170
567	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> , <b>2011</b> , 22, 2179-90	10.3	162
566	Survivorship care plans in cancer: a systematic review of care plan outcomes. <i>British Journal of Cancer</i> , <b>2014</b> , 111, 1899-908	8.7	155
565	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> , <b>2012</b> , 20, 1-22	3.9	154
564	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> , <b>2012</b> , 20, 1235-42	3.9	152

563	Psychological outcomes and risk perception after genetic testing and counselling in breast cancer: a systematic review. <i>Medical Journal of Australia</i> , <b>2003</b> , 178, 77-81	4	149
562	Impact of medical Qigong on quality of life, fatigue, mood and inflammation in cancer patients: a randomized controlled trial. <i>Annals of Oncology</i> , <b>2010</b> , 21, 608-614	10.3	143
561	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> , <b>2012</b> , 20, 2651-9	3.9	138
560	Misunderstanding in cancer patients: why shoot the messenger?. <i>Annals of Oncology</i> , <b>1999</b> , 10, 39-46	10.3	138
559	Computer-based interaction analysis of the cancer consultation. <i>British Journal of Cancer</i> , <b>1995</b> , 71, 1115-1121	8.1	138
558	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> , <b>2011</b> , 84, 379-85	3.1	136
557	Parents' attitudes to children's participation in randomized controlled trials. <i>Journal of Pediatrics</i> , <b>2003</b> , 142, 554-9	3.6	134
556	Psychological morbidity and quality of life of ethnic minority patients with cancer: a systematic review and meta-analysis. <i>Lancet Oncology</i> , <b>2011</b> , 12, 1240-8	21.7	130
555	Do patients want doctors to talk about spirituality? A systematic literature review. <i>Patient Education and Counseling</i> , <b>2015</b> , 98, 1320-8	3.1	129
554	Analysis of cancer risk and BRCA1 and BRCA2 mutation prevalence in the kConFab familial breast cancer resource. <i>Breast Cancer Research</i> , <b>2006</b> , 8, R12	8.3	125
553	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> , <b>2005</b> , 30, 132-44	4.8	124
552	Hospital-based interventions: a systematic review of staff-reported barriers and facilitators to implementation processes. <i>Implementation Science</i> , <b>2018</b> , 13, 36	8.4	122
551	Cancer consultation preparation package: changing patients but not physicians is not enough. <i>Journal of Clinical Oncology</i> , <b>2004</b> , 22, 4401-9	2.2	121
550	Communication styles in the cancer consultation: preferences for a patient-centred approach. <i>Psycho-Oncology</i> , <b>2000</b> , 9, 147-56	3.9	121
549	A Research Agenda for Communication Between Health Care Professionals and Patients Living With Serious Illness. <i>JAMA Internal Medicine</i> , <b>2017</b> , 177, 1361-1366	11.5	120
548	Psychological responses and coping strategies among patients with malignant melanoma: a systematic review of the literature. <i>Archives of Dermatology</i> , <b>2009</b> , 145, 1415-27		120
547	Can a "prompt list" empower cancer patients to ask relevant questions?. <i>Cancer</i> , <b>2008</b> , 113, 225-37	6.4	120
546	Fear of cancer recurrence: a theoretical review and novel cognitive processing formulation. <i>Journal of Cancer Survivorship</i> , <b>2016</b> , 10, 663-73	5.1	118

545	Does age really matter? Recall of information presented to newly referred patients with cancer. <i>Journal of Clinical Oncology</i> , <b>2008</b> , 26, 5450-7	2.2	116
544	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> , <b>2000</b> , 49, 169-81	4.1	115
543	The needs of terminally ill cancer patients versus those of caregivers for information regarding prognosis and end-of-life issues. <i>Cancer</i> , <b>2005</b> , 103, 1957-64	6.4	113
542	Psychosocial predictors of survival in metastatic melanoma. <i>Journal of Clinical Oncology</i> , <b>1999</b> , 17, 2256-63	6.3	113
541	Do personal stories make patient decision aids more effective? A critical review of theory and evidence. <i>BMC Medical Informatics and Decision Making</i> , <b>2013</b> , 13 Suppl 2, S9	3.6	112
540	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> , <b>2012</b> , 106, 1053-61	8.7	112
539	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> , <b>2010</b> , 18, 1241-62	3.9	112
538	Parental adjustment to the completion of their child's cancer treatment. <i>Pediatric Blood and Cancer</i> , <b>2011</b> , 56, 524-31	3	111
537	Increasing oncologists' skills in eliciting and responding to emotional cues: evaluation of a communication skills training program. <i>Psycho-Oncology</i> , <b>2008</b> , 17, 209-18	3.9	111
536	Discrepant perceptions about end-of-life communication: a systematic review. <i>Journal of Pain and Symptom Management</i> , <b>2007</b> , 34, 190-200	4.8	109
535	Psychological impact of genetic testing in women from high-risk breast cancer families. <i>European Journal of Cancer</i> , <b>2002</b> , 38, 2025-31	7.5	109
534	Advance care planning for cancer patients: a systematic review of perceptions and experiences of patients, families, and healthcare providers. <i>Psycho-Oncology</i> , <b>2016</b> , 25, 362-86	3.9	108
533	Clinical pathway for the screening, assessment and management of anxiety and depression in adult cancer patients: Australian guidelines. <i>Psycho-Oncology</i> , <b>2015</b> , 24, 987-1001	3.9	106
532	Developing ethical strategies to assist oncologists in seeking informed consent to cancer clinical trials. <i>Social Science and Medicine</i> , <b>2004</b> , 58, 379-90	5.1	104
531	Intention to undergo prophylactic bilateral mastectomy in women at increased risk of developing hereditary breast cancer. <i>Journal of Clinical Oncology</i> , <b>2000</b> , 18, 2250-7	2.2	104
530	Meeting the decision-making preferences of patients with breast cancer in oncology consultations: impact on decision-related outcomes. <i>Journal of Clinical Oncology</i> , <b>2012</b> , 30, 857-62	2.2	103
529	Life after cancer: couples' and partners' psychological adjustment and supportive care needs. <i>Supportive Care in Cancer</i> , <b>2007</b> , 15, 405-15	3.9	103
528	Doctors discussing religion and spirituality: A systematic literature review. <i>Palliative Medicine</i> , <b>2016</b> , 30, 327-37	5.5	101

527	The characteristics and effectiveness of Question Prompt List interventions in oncology: a systematic review of the literature. <i>Psycho-Oncology</i> , <b>2015</b> , 24, 245-52	3.9	101
526	Asking questions can help: development and preliminary evaluation of a question prompt list for palliative care patients. <i>British Journal of Cancer</i> , <b>2003</b> , 89, 2069-77	8.7	98
525	Symptom clusters in patients with advanced cancer: a systematic review of observational studies. <i>Journal of Pain and Symptom Management</i> , <b>2014</b> , 48, 411-50	4.8	97
524	Information booklets about cancer: factors influencing patient satisfaction and utilization. <i>Patient Education and Counseling</i> , <b>1998</b> , 33, 129-41	3.1	94
523	Promoting fruit and vegetable consumption. Testing an intervention based on the theory of planned behaviour. <i>Appetite</i> , <b>2012</b> , 58, 997-1004	4.5	93
522	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> , <b>2016</b> , 99, 3-16	3.1	92
521	Lay constructions of decision-making in cancer. <i>Psycho-Oncology</i> , <b>2002</b> , 11, 295-306	3.9	92
520	Hearing the bad news of a cancer diagnosis: the Australian melanoma patient's perspective. <i>Annals of Oncology</i> , <b>2001</b> , 12, 365-71	10.3	92
519	Randomized Trial of ConquerFear: A Novel, Theoretically Based Psychosocial Intervention for Fear of Cancer Recurrence. <i>Journal of Clinical Oncology</i> , <b>2017</b> , 35, 4066-4077	2.2	91
518	What is important in cancer care coordination? A qualitative investigation. <i>European Journal of Cancer Care</i> , <b>2011</b> , 20, 220-7	2.4	90
517	The use of unproven methods of treatment by cancer patients. Frequency, expectations and cost. <i>Supportive Care in Cancer</i> , <b>1998</b> , 6, 337-47	3.9	87
516	Attitudes and information needs of Chinese migrant cancer patients and their relatives. <i>Australian and New Zealand Journal of Medicine</i> , <b>1999</b> , 29, 207-13		87
515	Family involvement in cancer treatment decision-making: A qualitative study of patient, family, and clinician attitudes and experiences. <i>Patient Education and Counseling</i> , <b>2016</b> , 99, 1146-1155	3.1	85
514	A matter of trust--patient's views on decision-making in colorectal cancer. <i>Health Expectations</i> , <b>2004</b> , 7, 104-14	3.7	84
513	Evaluation of a novel individualised communication-skills training intervention to improve doctors' confidence and skills in end-of-life communication. <i>Palliative Medicine</i> , <b>2013</b> , 27, 236-43	5.5	80
512	Conquer fear: protocol of a randomised controlled trial of a psychological intervention to reduce fear of cancer recurrence. <i>BMC Cancer</i> , <b>2013</b> , 13, 201	4.8	79
511	Seeking informed consent to cancer clinical trials: describing current practice. <i>Social Science and Medicine</i> , <b>2004</b> , 58, 2445-57	5.1	79
510	A systematic review of decision aids for patients making a decision about treatment for early breast cancer. <i>Breast</i> , <b>2016</b> , 26, 31-45	3.6	78

509	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> , <b>2014</b> , 132, 690-7	4.9	78
508	Handling missing items in the Hospital Anxiety and Depression Scale (HADS): a simulation study. <i>BMC Research Notes</i> , <b>2016</b> , 9, 479	2.3	77
507	Immigrants' perceptions of the quality of their cancer care: an Australian comparative study, identifying potentially modifiable factors. <i>Annals of Oncology</i> , <b>2014</b> , 25, 1643-9	10.3	76
506	Genetic testing for melanoma risk: a prospective cohort study of uptake and outcomes among Australian families. <i>Genetics in Medicine</i> , <b>2009</b> , 11, 265-78	8.1	75
505	Physician-identified factors affecting patient participation in reaching treatment decisions. <i>Journal of Clinical Oncology</i> , <b>2008</b> , 26, 1724-31	2.2	75
504	Discussing adjuvant cancer therapy. <i>Journal of Clinical Oncology</i> , <b>2001</b> , 19, 1768-78	2.2	75
503	A comparison of nutritional management with stress management in the treatment of bulimia nervosa. <i>British Journal of Psychiatry</i> , <b>1991</b> , 159, 250-61	5.4	75
502	Discussing end-of-life issues with terminally ill cancer patients and their carers: a qualitative study. <i>Supportive Care in Cancer</i> , <b>2005</b> , 13, 589-99	3.9	74
501	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> , <b>2016</b> , 34, 4405-4414	2.2	72
500	Family perspectives on deceased organ donation: thematic synthesis of qualitative studies. <i>American Journal of Transplantation</i> , <b>2014</b> , 14, 923-35	8.7	70
499	The impact of communication on adherence in pain management. <i>Pain</i> , <b>2013</b> , 154 Suppl 1, S101-S107	8	70
498	Long-term outcomes of genetic counseling in women at increased risk of developing hereditary breast cancer. <i>Patient Education and Counseling</i> , <b>2001</b> , 44, 215-25	3.1	70
497	Advance care planning for adults with CKD: a systematic integrative review. <i>American Journal of Kidney Diseases</i> , <b>2014</b> , 63, 761-70	7.4	69
496	From inside the bubble: migrants' perceptions of communication with the cancer team. <i>Supportive Care in Cancer</i> , <b>2010</b> , 19, 281-90	3.9	69
495	Psychosocial predictors of survival: metastatic breast cancer. <i>Annals of Oncology</i> , <b>2000</b> , 11, 469-74	10.3	69
494	Family information needs at childhood cancer treatment completion. <i>Pediatric Blood and Cancer</i> , <b>2012</b> , 58, 621-6	3	68
493	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> , <b>2009</b> , 77, 103-8	3.1	68
492	Communicating risk information: the influence of graphical display format on quantitative information perception-Accuracy, comprehension and preferences. <i>Patient Education and Counseling</i> , <b>2007</b> , 69, 121-8	3.1	68

491	Comparison of group and individual cognitive-behavioral therapy for patients with bulimia nervosa. <i>International Journal of Eating Disorders</i> , <b>2003</b> , 33, 241-54; discussion 255-6	6.3	68
490	Discussing life expectancy with terminally ill cancer patients and their carers: a qualitative study. <i>Supportive Care in Cancer</i> , <b>2005</b> , 13, 733-42	3.9	68
489	The role of psychosocial factors in the development of breast carcinoma: Part II. <i>Cancer</i> , <b>2001</b> , 91, 686-694	6.7	67
488	Pediatricians' attitudes toward randomized controlled trials involving children. <i>Journal of Pediatrics</i> , <b>2002</b> , 141, 798-803	3.6	67
487	Psychological intervention targeting distress for cancer patients: a meta-analytic study investigating uptake and adherence. <i>Psycho-Oncology</i> , <b>2016</b> , 25, 882-90	3.9	67
486	Survivorship care after breast cancer treatment--experiences and preferences of Australian women. <i>Breast</i> , <b>2011</b> , 20, 271-7	3.6	66
485	Responding to the active and passive patient: flexibility is the key. <i>Health Expectations</i> , <b>2002</b> , 5, 236-45	3.7	65
484	Attitudes toward prophylactic oophorectomy and screening utilization in women at increased risk of developing hereditary breast/ovarian cancer. <i>Gynecologic Oncology</i> , <b>1999</b> , 75, 122-9	4.9	65
483	On the receiving end. IV: Validation of quality of life indicators. <i>Annals of Oncology</i> , <b>1991</b> , 2, 597-603	10.3	65
482	Prevalence and predictors of anxiety and depression in women with invasive ovarian cancer and their caregivers. <i>Medical Journal of Australia</i> , <b>2010</b> , 193, S52-7	4	64
481	Therapeutic life review in palliative care: a systematic review of quantitative evaluations. <i>Journal of Pain and Symptom Management</i> , <b>2015</b> , 49, 747-61	4.8	62
480	A bridge between cultures: interpreters' perspectives of consultations with migrant oncology patients. <i>Supportive Care in Cancer</i> , <b>2012</b> , 20, 235-44	3.9	61
479	Towards better communication in cancer care: a framework for developing evidence-based interventions. <i>Patient Education and Counseling</i> , <b>2004</b> , 55, 32-9	3.1	61
478	Attitudes to randomized clinical trials amongst out-patients attending a medical oncology clinic. <i>Health Expectations</i> , <b>1999</b> , 2, 33-43	3.7	60
477	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> , <b>2008</b> , 36, 459-72	6	59
476	Analyzing the process and content of genetic counseling in familial breast cancer consultations. <i>Journal of Genetic Counseling</i> , <b>2004</b> , 13, 403-24	2.5	59
475	Improving doctors' letters. <i>Medical Journal of Australia</i> , <b>2002</b> , 177, 516-20	4	59
474	A qualitative analysis of reasons for leaving, or not attending, a cancer support group. <i>Social Work in Health Care</i> , <b>2008</b> , 47, 14-29	2	58

473	Consultation audio tapes: an underused cancer patient information aid and clinical research tool. <i>Lancet Oncology, The</i> , <b>2002</b> , 3, 431-7	21.7	58
472	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> , <b>2014</b> , 111, 1-7	8.7	57
471	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> , <b>2011</b> , 1, 87-94	2.2	56
470	Current approaches to managing fear of cancer recurrence; a descriptive survey of psychosocial and clinical health professionals. <i>Psycho-Oncology</i> , <b>2014</b> , 23, 390-6	3.9	55
469	Implementing patient question-prompt lists into routine cancer care. <i>Patient Education and Counseling</i> , <b>2012</b> , 86, 252-8	3.1	55
468	Patient perspectives regarding communication about prognosis and end-of-life issues: how can it be optimised?. <i>Patient Education and Counseling</i> , <b>2013</b> , 90, 307-14	3.1	55
467	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> , <b>2012</b> , 12, 339	4.8	55
466	Sharing vs. caring—the relative impact of sharing decisions versus managing emotions on patient outcomes. <i>Patient Education and Counseling</i> , <b>2011</b> , 82, 233-9	3.1	55
465	When the treatment goal is not cure: are patients informed adequately?. <i>Supportive Care in Cancer</i> , <b>2002</b> , 10, 314-21	3.9	55
464	Adapting the nominal group technique for priority setting of evidence-practice gaps in implementation science. <i>BMC Medical Research Methodology</i> , <b>2016</b> , 16, 110	4.7	55
463	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> , <b>2011</b> , 42, 710-20	4.8	54
462	Interpretation in consultations with immigrant patients with cancer: how accurate is it?. <i>Journal of Clinical Oncology</i> , <b>2011</b> , 29, 2801-7	2.2	54
461	Health professional and consumer views on involving breast cancer patients in the multidisciplinary discussion of their disease and treatment plan. <i>Cancer</i> , <b>2007</b> , 110, 1937-44	6.4	54
460	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> , <b>2016</b> , 51, 88-98	4.8	53
459	Anxiety in the context of cancer: A systematic review and development of an integrated model. <i>Clinical Psychology Review</i> , <b>2017</b> , 56, 40-54	10.8	53
458	Looking different, feeling different: women's reactions to risk-reducing breast and ovarian surgery. <i>Familial Cancer</i> , <b>2012</b> , 11, 215-24	3	53
457	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> , <b>2013</b> , 22, 2797-806	3.9	53
456	Women's preferences for discussion of prognosis in early breast cancer. <i>Health Expectations</i> , <b>2001</b> , 4, 48-57	3.7	53

455	Grappling with cultural differences; communication between oncologists and immigrant cancer patients with and without interpreters. <i>Patient Education and Counseling</i> , <b>2011</b> , 84, 398-405	3.1	52
454	Measuring cancer care coordination: development and validation of a questionnaire for patients. <i>BMC Cancer</i> , <b>2011</b> , 11, 298	4.8	52
453	The motherhood choice: themes arising in the decision-making process for women with multiple sclerosis. <i>Multiple Sclerosis Journal</i> , <b>2008</b> , 14, 701-4	5	52
452	Informing breast cancer patients about clinical trials: a randomized clinical trial of an educational booklet. <i>Annals of Oncology</i> , <b>2002</b> , 13, 1414-23	10.3	52
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155	Development and pilot testing of a Decision Aid (DA) for women with early-stage breast cancer considering contralateral prophylactic mastectomy. <i>Breast</i> , <b>2018</b> , 40, 156-164	3.6	8
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