

# Phyllis Butow

## List of Publications by Year in descending order

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Version: 2024-02-01

627  
papers

35,052  
citations

3531  
90  
h-index

7348  
152  
g-index

637  
all docs

637  
docs citations

637  
times ranked

26646  
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.  | 2.3  | 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.   | 2.2  | 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.   | 1.6  | 463       |
| 4  | Sharing decisions in cancer care. <i>Social Science and Medicine</i> , 2001, 52, 1865-1878.  | 3.8  | 445       |
| 5  | The dynamics of change: Cancer patients' preferences for information, involvement and support. <i>Annals of Oncology</i> , 1997, 8, 857-863.   | 1.2  | 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. | 1.2  | 437       |
| 7  | Cancer Patient Preferences for Communication of Prognosis in the Metastatic Setting. <i>Journal of Clinical Oncology</i> , 2004, 22, 1721-1730.  | 1.6  | 428       |
| 8  | Communicating prognosis in cancer care: a systematic review of the literature. <i>Annals of Oncology</i> , 2005, 16, 1005-1053.  | 1.2  | 394       |
| 9  | Clinical trials in children. <i>Lancet, The</i> , 2004, 364, 803-811.  | 13.7 | 381       |
| 10 | Truth-telling in discussing prognosis in advanced life-limiting illnesses: a systematic review. <i>Palliative Medicine</i> , 2007, 21, 507-517.  | 3.1  | 357       |
| 11 | Psychological responses of patients receiving a diagnosis of cancer. <i>Annals of Oncology</i> , 2003, 14, 48-56.  | 1.2  | 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.   | 2.2  | 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.   | 1.6  | 318       |
| 14 | When the diagnosis is cancer: Patient communication experiences and preferences. <i>Cancer</i> , 1996, 77, 2630-2637.  | 4.1  | 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.   | 2.2  | 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.  | 1.6  | 296       |
| 17 | A systematic review on communicating with patients about evidence. <i>Journal of Evaluation in Clinical Practice</i> , 2006, 12, 13-23.  | 1.8  | 282       |
| 18 | Oncologists' reactions to cancer patients' verbal cues. <i>Psycho-Oncology</i> , 2002, 11, 47-58.  | 2.3  | 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.                             | 2.3 | 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.   | 3.7 | 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.  | 5.0 | 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. | 1.7 | 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.                           | 7.1 | 249       |
| 24 | Breast cancer survivors'™ supportive care needs 2-10 years after diagnosis. <i>Supportive Care in Cancer</i> , 2007, 15, 515-523.   | 2.2 | 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.  | 4.1 | 239       |
| 26 | Effective health communication "a key factor in fighting the COVID-19 pandemic. <i>Patient Education and Counseling</i> , 2020, 103, 873-876.   | 2.2 | 239       |
| 27 | How to recognize and manage psychological distress in cancer patients. <i>European Journal of Cancer Care</i> , 2005, 14, 7-15.   | 1.5 | 229       |
| 28 | Hospital-based interventions: a systematic review of staff-reported barriers and facilitators to implementation processes. <i>Implementation Science</i> , 2018, 13, 36.  | 6.9 | 228       |
| 29 | Randomized Clinical Trials in Oncology: Understanding and Attitudes Predict Willingness to Participate. <i>Journal of Clinical Oncology</i> , 2001, 19, 3554-3561.  | 1.6 | 221       |
| 30 | Long-term survival from gynecologic cancer: Psychosocial outcomes, supportive care needs and positive outcomes. <i>Gynecologic Oncology</i> , 2007, 104, 381-389.   | 1.4 | 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.   | 2.2 | 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.                  | 1.2 | 219       |
| 33 | Review of Adherence-Related Issues in Adolescents and Young Adults With Cancer. <i>Journal of Clinical Oncology</i> , 2010, 28, 4800-4809.  | 1.6 | 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.  | 2.2 | 207       |
| 35 | Patient participation in the cancer consultation: Evaluation of a question prompt sheet. <i>Annals of Oncology</i> , 1994, 5, 199-204.  | 1.2 | 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.                           | 2.2 | 189       |

| #  | ARTICLE   | IF   | CITATIONS |
|----|---|------|-----------|
| 37 | Do patients want doctors to talk about spirituality? A systematic literature review. Patient Education and Counseling, 2015, 98, 1320-1328.   | 2.2  | 185       |
| 38 | Survivorship care plans in cancer: a systematic review of care plan outcomes. British Journal of Cancer, 2014, 111, 1899-1908.  | 6.4  | 183       |
| 39 | Effect of medical Qigong on cognitive function, quality of life, and a biomarker of inflammation in cancer patients: a randomized controlled trial. Supportive Care in Cancer, 2012, 20, 1235-1242.                 | 2.2  | 179       |
| 40 | Impact of Medical Qigong on quality of life, fatigue, mood and inflammation in cancer patients: a randomized controlled trial. Annals of Oncology, 2010, 21, 608-614.   | 1.2  | 176       |
| 41 | A Research Agenda for Communication Between Health Care Professionals and Patients Living With Serious Illness. JAMA Internal Medicine, 2017, 177, 1361.  | 5.1  | 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. Supportive Care in Cancer, 2012, 20, 2651-2659. | 2.2  | 168       |
| 43 | Psychological outcomes and risk perception after genetic testing and counselling in breast cancer: a systematic review. Medical Journal of Australia, 2003, 178, 77-81.   | 1.7  | 166       |
| 44 | Advance care planning for cancer patients: a systematic review of perceptions and experiences of patients, families, and healthcare providers. Psycho-Oncology, 2016, 25, 362-386.                                  | 2.3  | 166       |
| 45 | Fear of cancer recurrence: a theoretical review and novel cognitive processing formulation. Journal of Cancer Survivorship, 2016, 10, 663-673.  | 2.9  | 166       |
| 46 | Misunderstanding in cancer patients: Why shoot the messenger?. Annals of Oncology, 1999, 10, 39-46.   | 1.2  | 164       |
| 47 | The characteristics and effectiveness of Question Prompt List interventions in oncology: a systematic review of the literature. Psycho-Oncology, 2015, 24, 245-252.   | 2.3  | 163       |
| 48 | Psychological morbidity and quality of life of ethnic minority patients with cancer: a systematic review and meta-analysis. Lancet Oncology, The, 2011, 12, 1240-1248.  | 10.7 | 162       |
| 49 | Computer-based interaction analysis of the cancer consultation. British Journal of Cancer, 1995, 71, 1115-1121.   | 6.4  | 157       |
| 50 | Parents' attitudes to children's participation in randomized controlled trials. Journal of Pediatrics, 2003, 142, 554-559.  | 1.8  | 155       |
| 51 | Clinical pathway for the screening, assessment and management of anxiety and depression in adult cancer patients: Australian guidelines. Psycho-Oncology, 2015, 24, 987-1001.                                       | 2.3  | 148       |
| 52 | Randomized Trial of ConquerFear: A Novel, Theoretically Based Psychosocial Intervention for Fear of Cancer Recurrence. Journal of Clinical Oncology, 2017, 35, 4066-4077.   | 1.6  | 148       |
| 53 | Communication styles in the cancer consultation: preferences for a patient-centred approach. Psycho-Oncology, 2000, 9, 147-156.   | 2.3  | 147       |
| 54 | Can a "question prompt list" empower cancer patients to ask relevant questions?. Cancer, 2008, 113, 225-237.  | 4.1  | 147       |

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|----|---|-----|-----------|
| 55 | Do personal stories make patient decision aids more effective? A critical review of theory and evidence. BMC Medical Informatics and Decision Making, 2013, 13, S9.   | 3.0 | 147       |
| 56 | When and How to Initiate Discussion About Prognosis and End-of-Life Issues with Terminally Ill Patients. Journal of Pain and Symptom Management, 2005, 30, 132-144.   | 1.2 | 146       |
| 57 | Psychological Responses and Coping Strategies Among Patients With Malignant Melanoma. Archives of Dermatology, 2009, 145, 1415-27.  | 1.4 | 146       |
| 58 | Parental adjustment to the completion of their child's cancer treatment. Pediatric Blood and Cancer, 2011, 56, 524-531.   | 1.5 | 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. British Journal of Cancer, 2012, 106, 1053-1061.   | 6.4 | 144       |
| 60 | Does Age Really Matter? Recall of Information Presented to Newly Referred Patients With Cancer. Journal of Clinical Oncology, 2008, 26, 5450-5457.  | 1.6 | 143       |
| 61 | Symptom Clusters in Patients With Advanced Cancer: A Systematic Review of Observational Studies. Journal of Pain and Symptom Management, 2014, 48, 411-450.   | 1.2 | 142       |
| 62 | Cancer Consultation Preparation Package: Changing Patients but Not Physicians Is Not Enough. Journal of Clinical Oncology, 2004, 22, 4401-4409.   | 1.6 | 140       |
| 63 | Doctors discussing religion and spirituality: A systematic literature review. Palliative Medicine, 2016, 30, 327-337.   | 3.1 | 140       |
| 64 | Analysis of cancer risk and BRCA1 and BRCA2 mutation prevalence in the kConFab familial breast cancer resource. Breast Cancer Research, 2006, 8, R12.   | 5.0 | 135       |
| 65 | Epidemiological evidence for a relationship between life events, coping style, and personality factors in the development of breast cancer. Journal of Psychosomatic Research, 2000, 49, 169-181.   | 2.6 | 133       |
| 66 | Increasing oncologists' skills in eliciting and responding to emotional cues: evaluation of a communication skills training program. Psycho-Oncology, 2008, 17, 209-218.  | 2.3 | 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. Supportive Care in Cancer, 2010, 18, 1241-1262. | 2.2 | 131       |
| 68 | Asking questions can help: development and preliminary evaluation of a question prompt list for palliative care patients. British Journal of Cancer, 2003, 89, 2069-2077.   | 6.4 | 130       |
| 69 | The needs of terminally ill cancer patients versus those of caregivers for information regarding prognosis and end-of-life issues. Cancer, 2005, 103, 1957-1964.  | 4.1 | 130       |
| 70 | Discrepant Perceptions About End-of-Life Communication: A Systematic Review. Journal of Pain and Symptom Management, 2007, 34, 190-200.   | 1.2 | 130       |
| 71 | Meeting the Decision-Making Preferences of Patients With Breast Cancer in Oncology Consultations: Impact on Decision-Related Outcomes. Journal of Clinical Oncology, 2012, 30, 857-862.   | 1.6 | 128       |
| 72 | Psychosocial Predictors of Survival in Metastatic Melanoma. Journal of Clinical Oncology, 1999, 17, 2256-2256.  | 1.6 | 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.  | 2.2 | 121       |
| 74 | Handling missing items in the Hospital Anxiety and Depression Scale (HADS): a simulation study. <i>BMC Research Notes</i> , 2016, 9, 479.  | 1.4 | 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.   | 2.8 | 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.   | 3.8 | 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.        | 2.2 | 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. | 1.5 | 117       |
| 79 | Promoting fruit and vegetable consumption. Testing an intervention based on the theory of planned behaviour. <i>Appetite</i> , 2012, 58, 997-1004.   | 3.7 | 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.                         | 1.6 | 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.      | 2.2 | 111       |
| 82 | What is important in cancer care coordination? A qualitative investigation. <i>European Journal of Cancer Care</i> , 2011, 20, 220-227.  | 1.5 | 109       |
| 83 | Information booklets about cancer:. <i>Patient Education and Counseling</i> , 1998, 33, 129-141.   | 2.2 | 108       |
| 84 | Lay constructions of decision-making in cancer. <i>Psycho-Oncology</i> , 2002, 11, 295-306.  | 2.3 | 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.     | 3.1 | 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.  | 2.2 | 104       |
| 87 | The use of unproven methods of treatment by cancer patients. <i>Supportive Care in Cancer</i> , 1998, 6, 337-347.  | 2.2 | 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 patient'™s perspective. <i>Annals of Oncology</i> , 2001, 12, 365-371.   | 1.2 | 100       |
| 90 | Family Perspectives on Deceased Organ Donation: Thematic Synthesis of Qualitative Studies. <i>American Journal of Transplantation</i> , 2014, 14, 923-935.   | 4.7 | 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.   | 2.3 | 98        |
| 92  | A matter of trust - patient's views on decision-making in colorectal cancer. <i>Health Expectations</i> , 2004, 7, 104-114.  | 2.6 | 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.  | 2.6 | 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.  | 1.4 | 92        |
| 95  | The impact of communication on adherence in pain management. <i>Pain</i> , 2013, 154, S101-S107.   | 4.2 | 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. | 1.6 | 91        |
| 97  | Seeking informed consent to cancer clinical trials. <i>Social Science and Medicine</i> , 2004, 58, 2445-2457.  | 3.8 | 89        |
| 98  | Advance Care Planning for Adults With CKD: A Systematic Integrative Review. <i>American Journal of Kidney Diseases</i> , 2014, 63, 761-770.  | 1.9 | 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.   | 4.0 | 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.  | 2.8 | 84        |
| 101 | Fear of cancer recurrence and death anxiety. <i>Psycho-Oncology</i> , 2018, 27, 2559-2565.   | 2.3 | 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.   | 2.9 | 84        |
| 103 | Discussing Adjuvant Cancer Therapy. <i>Journal of Clinical Oncology</i> , 2001, 19, 1768-1778.   | 1.6 | 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.   | 2.4 | 83        |
| 105 | From inside the bubble: migrants' perceptions of communication with the cancer team. <i>Supportive Care in Cancer</i> , 2011, 19, 281-290.   | 2.2 | 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.  | 1.7 | 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.   | 2.2 | 81        |
| 108 | Physician-Identified Factors Affecting Patient Participation in Reaching Treatment Decisions. <i>Journal of Clinical Oncology</i> , 2008, 26, 1724-1731.   | 1.6 | 81        |

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|-----|--|------|-----------|
| 109 | Family information needs at childhood cancer treatment completion. <i>Pediatric Blood and Cancer</i> , 2012, 58, 621-626.  | 1.5  | 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.   | 1.2  | 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.   | 5.6  | 81        |
| 112 | Communicating risk information: The influence of graphical display format on quantitative information perceptionâ€”Accuracy, comprehension and preferences. <i>Patient Education and Counseling</i> , 2007, 69, 121-128.   | 2.2  | 80        |
| 113 | Psychosocial predictors of survival: Metastatic breast cancer. <i>Annals of Oncology</i> , 2000, 11, 469-474.  | 1.2  | 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.  | 2.2  | 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.   | 11.4 | 79        |
| 116 | A bridge between cultures: interpretersâ€™ perspectives of consultations with migrant oncology patients. <i>Supportive Care in Cancer</i> , 2012, 20, 235-244.   | 2.2  | 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.   | 2.2  | 76        |
| 118 | The role of psychosocial factors in the development of breast carcinoma: Part II. <i>Cancer</i> , 2001, 91, 686-697.   | 4.1  | 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.  | 1.2  | 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.  | 3.1  | 75        |
| 121 | Can consumers learn to ask three questions to improve shared decision making? A feasibility study of the <sup>ASK</sup> (AskShareKnow) Patientâ€”Clinician Communication Model<sup>Â®</sup> intervention in a primary healthâ€”care setting. <i>Health Expectations</i> , 2016, 19, 1160-1168. | 2.6  | 75        |
| 122 | Exploring the screening capacity of the Fear of Cancer Recurrence Inventoryâ€”Short Form for clinical levels of fear of cancer recurrence. <i>Psycho-Oncology</i> , 2018, 27, 492-499.   | 2.3  | 75        |
| 123 | On the receiving end IV: Validation of quality of life indicators. <i>Annals of Oncology</i> , 1991, 2, 597-603.   | 1.2  | 74        |
| 124 | Pediatricians' attitudes toward randomized controlled trials involving children. <i>Journal of Pediatrics</i> , 2002, 141, 798-803.  | 1.8  | 74        |
| 125 | Survivorship care after breast cancer treatment â€” Experiences and preferences of Australian women. <i>Breast</i> , 2011, 20, 271-277.  | 2.2  | 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.  | 1.6  | 74        |



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|-----|--|------|-----------|
| 127 | Management of common clinical problems experienced by survivors of cancer. <i>Lancet</i> , The, 2022, 399, 1537-1550.  | 13.7 | 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.  | 1.2  | 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.  | 2.3  | 73        |
| 130 | Responding to the active and passive patient: flexibility is the key. <i>Health Expectations</i> , 2002, 5, 236-245.   | 2.6  | 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.  | 2.3  | 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.   | 2.2  | 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.  | 2.6  | 71        |
| 134 | Implementing patient question-prompt lists into routine cancer care. <i>Patient Education and Counseling</i> , 2012, 86, 252-258.  | 2.2  | 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.   | 3.1  | 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. | 6.4  | 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.   | 1.4  | 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.  | 4.1  | 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.   | 3.8  | 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.  | 2.2  | 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.                    | 2.3  | 68        |
| 142 | Attentional bias and metacognitions in cancer survivors with high fear of cancer recurrence. <i>Psycho-Oncology</i> , 2015, 24, 416-423.   | 2.3  | 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.   | 1.2  | 68        |
| 144 | Attitudes to randomized clinical trials amongst outpatients attending a medical oncology clinic. <i>Health Expectations</i> , 1999, 2, 33-43.  | 2.6  | 67        |

| #   | ARTICLE   | IF   | CITATIONS |
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