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TABLE OF CONTENTS

HEADER	1
ABSTRACT	1
PLAIN LANGUAGE SUMMARY	3
SUMMARY OF FINDINGS	5
BACKGROUND	8
OBJECTIVES	8
METHODS	9
Figure 1.	11
Figure 2.	12
RESULTS	14
Figure 3.	15
DISCUSSION	22
AUTHORS' CONCLUSIONS	24
ACKNOWLEDGEMENTS	25
REFERENCES	26
CHARACTERISTICS OF STUDIES	31
DATA AND ANALYSES	74
Analysis 1.1. Comparison 1: Symptoms, Outcome 1: Anxiety	76
Analysis 1.2. Comparison 1: Symptoms, Outcome 2: Depression	76
Analysis 1.3. Comparison 1: Symptoms, Outcome 3: Fatigue	77
Analysis 1.4. Comparison 1: Symptoms, Outcome 4: Emotional distress	77
APPENDICES	77
HISTORY	78
CONTRIBUTIONS OF AUTHORS	78
DECLARATIONS OF INTEREST	79
SOURCES OF SUPPORT	79
DIFFERENCES BETWEEN PROTOCOL AND REVIEW	79
INDEX TERMS	79

[Intervention Review]

Telephone interventions for symptom management in adults with cancer

Emma Ream¹, Amanda Euesden Hughes², Anna Cox¹, Katy Skarparis³, Alison Richardson⁴, Vibe H Pedersen⁵, Theresa Wiseman⁶, Angus Forbes⁵, Andrew Bryant⁷

¹School of Health Sciences, University of Surrey, Guildford, UK. ²Memorial Sloan Kettering Cancer Center, New York, USA. ³Department of Nursing, Midwifery & Health, Northumbria University, Newcastle, UK. ⁴School of Health Sciences, University of Southampton, Southampton, UK. ⁵Florence Nightingale School of Nursing and Midwifery, King's College London, London, UK. ⁶Health Services Research, The Royal Marsden NHS Foundation Trust, London, UK. ⁷Institute of Health & Society, Newcastle University, Newcastle upon Tyne, UK

Contact address: Emma Ream, e.ream@surrey.ac.uk.

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ABSTRACT

Background

People with cancer experience a variety of symptoms as a result of their disease and the therapies involved in its management. Inadequate symptom management has implications for patient outcomes including functioning, psychological well-being, and quality of life (QoL). Attempts to reduce the incidence and severity of cancer symptoms have involved the development and testing of psycho-educational interventions to enhance patients' symptom self-management. With the trend for care to be provided nearer patients' homes, telephone-delivered psycho-educational interventions have evolved to provide support for the management of a range of cancer symptoms. Early indications suggest that these can reduce symptom severity and distress through enhanced symptom self-management.

Objectives

To assess the effectiveness of telephone-delivered interventions for reducing symptoms associated with cancer and its treatment. To determine which symptoms are most responsive to telephone interventions. To determine whether certain configurations (e.g. with/without additional support such as face-to-face, printed or electronic resources) and duration/frequency of intervention calls mediate observed cancer symptom outcome effects.

Search methods

We searched the following databases: the Cochrane Central Register of Controlled Trials (CENTRAL; 2019, Issue 1); MEDLINE via OVID (1946 to January 2019); Embase via OVID (1980 to January 2019); (CINAHL) via Athens (1982 to January 2019); British Nursing Index (1984 to January 2019); and PsycINFO (1989 to January 2019). We searched conference proceedings to identify published abstracts, as well as SIGLE and trial registers for unpublished studies. We searched the reference lists of all included articles for additional relevant studies. Finally, we handsearched the following journals: *Cancer*, *Journal of Clinical Oncology*, *Psycho-oncology*, *Cancer Practice*, *Cancer Nursing*, *Oncology Nursing Forum*, *Journal of Pain and Symptom Management*, and *Palliative Medicine*. We restricted our search to publications published in English.

Selection criteria

We included randomised controlled trials (RCTs) and quasi-RCTs that compared one or more telephone interventions with one other, or with other types of interventions (e.g. a face-to-face intervention) and/or usual care, with the stated aim of addressing any physical or

psychological symptoms of cancer and its treatment, which recruited adults (over 18 years) with a clinical diagnosis of cancer, regardless of tumour type, stage of cancer, type of treatment, and time of recruitment (e.g. before, during, or after treatment).

Data collection and analysis

We used Cochrane methods for trial selection, data extraction and analysis. When possible, anxiety, depressive symptoms, fatigue, emotional distress, pain, uncertainty, sexually-related and lung cancer symptoms as well as secondary outcomes are reported as standardised mean differences (SMDs) with 95% confidence intervals (CIs), and we presented a descriptive synthesis of study findings. We reported on findings according to symptoms addressed and intervention types (e.g. telephone only, telephone combined with other elements). As many studies included small samples, and because baseline scores for study outcomes often varied for intervention and control groups, we used change scores and associated standard deviations. The certainty of the evidence for each outcome was interpreted using the Grading of Recommendations Assessment, Development and Evaluation (GRADE) approach.

Main results

Thirty-two studies were eligible for inclusion; most had moderate risk of bias, often related to blinding. Collectively, researchers recruited 6250 people and studied interventions in people with a variety of cancer types and across the disease trajectory, although many participants had breast cancer or early-stage cancer and/or were starting treatment. Studies measured symptoms of anxiety, depression, emotional distress, uncertainty, fatigue, and pain, as well as sexually-related symptoms and general symptom intensity and/or distress.

Interventions were primarily delivered by nurses ($n = 24$), most of whom ($n = 16$) had a background in oncology, research, or psychiatry. Ten interventions were delivered solely by telephone; the rest combined telephone with additional elements (i.e. face-to-face consultations and digital/online/printed resources). The number of calls delivered ranged from 1 to 18; most interventions provided three or four calls.

Twenty-one studies provided evidence on effectiveness of telephone-delivered interventions and the majority appeared to reduce symptoms of *depression* compared to control. Nine studies contributed quantitative change scores (CSs) and associated standard deviation results (or these could be calculated). Likewise, many telephone interventions appeared effective when compared to control in reducing *anxiety* (16 studies; 5 contributed quantitative CS results); *fatigue* (9 studies; 6 contributed to quantitative CS results); and *emotional distress* (7 studies; 5 contributed quantitative CS results). Due to significant clinical heterogeneity with regards to interventions introduced, study participants recruited, and outcomes measured, meta-analysis was not conducted.

For other symptoms (*uncertainty, pain, sexually-related symptoms, dyspnoea, and general symptom experience*), evidence was limited; similarly meta-analysis was not possible, and results from individual studies were largely conflicting, making conclusions about their management through telephone-delivered interventions difficult to draw. Heterogeneity was considerable across all trials for all outcomes.

Overall, the certainty of evidence was very low for all outcomes in the review. Outcomes were all downgraded due to concerns about overall risk of bias profiles being frequently unclear, uncertainty in effect estimates and due to some inconsistencies in results and general heterogeneity.

Unsubstantiated evidence suggests that telephone interventions in some capacity may have a place in symptom management for adults with cancer. However, in the absence of reliable and homogeneous evidence, caution is needed in interpreting the narrative synthesis. Further, there were no clear patterns across studies regarding which forms of interventions (telephone alone versus augmented with other elements) are most effective. It is impossible to conclude with any certainty which forms of telephone intervention are most effective in managing the range of cancer-related symptoms that people with cancer experience.

Authors' conclusions

Telephone interventions provide a convenient way of supporting self-management of cancer-related symptoms for adults with cancer. These interventions are becoming more important with the shift of care closer to patients' homes, the need for resource/cost containment, and the potential for voluntary sector providers to deliver healthcare interventions. Some evidence supports the use of telephone-delivered interventions for symptom management for adults with cancer; most evidence relates to four commonly experienced symptoms - depression, anxiety, emotional distress, and fatigue. Some telephone-delivered interventions were augmented by combining them with face-to-face meetings and provision of printed or digital materials. Review authors were unable to determine whether telephone alone or in combination with other elements provides optimal reduction in symptoms; it appears most likely that this will vary by symptom. It is noteworthy that, despite the potential for telephone interventions to deliver cost savings, none of the studies reviewed included any form of health economic evaluation.

Further robust and adequately reported trials are needed across all cancer-related symptoms, as the certainty of evidence generated in studies within this review was very low, and reporting was of variable quality. Researchers must strive to reduce variability between studies in the future. Studies in this review are characterised by clinical and methodological diversity; the level of this diversity hindered comparison across studies. At the very least, efforts should be made to standardise outcome measures. Finally, studies were compromised by inclusion of small samples, inadequate concealment of group allocation, lack of observer blinding, and short length of follow-up. Consequently, conclusions related to symptoms most amenable to management by telephone-delivered interventions are tentative.

PLAIN LANGUAGE SUMMARY

Telephone interventions for managing symptoms in adults with cancer

Background

People with cancer experience a variety of symptoms caused by their disease and its treatment. Symptoms can include depression, anxiety, fatigue and pain. These are often managed, day-to-day, by patients or their family members. If symptoms are not well managed, this can lead to other problems, such as difficulties in carrying out everyday tasks, poor sleep and poor quality of life.

Cancer professionals have developed psychological and educational treatments to help people to manage cancer symptoms. These treatments (or interventions) can be delivered by telephone (telephone interventions) in the patients' homes instead of face-to-face in hospital.

What questions does this review aim to answer?

This Cochrane Review aimed to answer the following questions.

1. Are telephone interventions for adults with cancer effective in relieving symptoms of cancer and cancer treatment?
2. Which symptoms are most reduced when telephone interventions are used?
3. What parts of telephone interventions have the most impact in reducing cancer symptoms?

In this review, telephone interventions were interventions given only, or mainly, by telephone. They were given by health professionals. As well as telephone contact, they could include face-to-face contact, or printed, digital or online information, such as, leaflets, computer programs and websites.

How did we answer these questions?

We searched medical databases and journals to find all randomised controlled trials that used a telephone intervention to reduce any cancer symptoms. Randomised controlled trials allocate people randomly to one treatment or another; they provide the most reliable evidence. Studies could compare telephone interventions with another telephone intervention, with another type of intervention (e.g. face-to-face), or with usual care. Participants in these studies were adults with any kind of cancer at any stage.

Results

We included 32 studies with a total of 6250 participants. Most studies (21) were from the USA. Nine studies recruited women with breast cancer, 11 included people with breast, colorectal, lung, or prostate cancer. Fourteen studies included people with early-stage cancer. Nurses provided interventions in 24 studies. Only 10 studies delivered interventions solely by telephone, and 16 studies combined telephone calls with other materials (printed or digital). Studies measured symptoms of depression, anxiety, emotional distress, uncertainty, fatigue, pain, sexual symptoms, and breathlessness. They also measured the effect of all the symptoms together (the general symptom experience).

Most studies compared a telephone intervention with usual care alone or usual care with additional support. Eight studies compared two telephone interventions against each other; some also compared these with usual care.

Because the studies were so different from each other, we could not combine the results into one analysis for each symptom. However, some studies measured changes in symptoms using standardised or similar scales. They recorded participants' scale scores at the beginning of the intervention, during the intervention, and at the end, resulting in a 'change score'. We analysed the results from studies that recorded change scores.

What does evidence from the review tell us?

Twenty-one studies provided evidence on depression compared to usual care or other interventions, but only nine provided change scores. These found that telephone interventions appeared to reduce symptoms of depression. Likewise, telephone interventions appeared effective compared to usual care or other interventions in reducing anxiety (16 studies; 5 contributed change scores); fatigue (9 studies; 6 contributed change scores); and emotional distress (7 studies; 5 contributed change scores).

Evidence for other symptoms was limited, making it difficult to draw conclusions.

Certainty of the evidence

Telephone interventions appear to relieve some symptoms of cancer and cancer treatment, however, the studies were small and very different from each other, so our confidence (certainty) in the evidence is very low. It is unclear whether telephone interventions alone, or combined with face-to-face meetings, or printed or audio materials, are most effective in reducing the many symptoms that people with cancer experience.

Conclusions

Telephone interventions are convenient for patients, their families and healthcare workers but the results of our review were not conclusive. Further, rigorous research on this topic would help to answer our review questions.

Search date

This review includes evidence published up to January 2019.

SUMMARY OF FINDINGS

Summary of findings 1. Summary of findings

Telephone interventions compared with control interventions for symptom management in adults with cancer

Patient or population: individuals with any cancer at any stage

Settings: randomised controlled trials

Intervention: telephone interventions with or without additional support

Comparison: control intervention

Outcomes	Risks	Effects of interventions	No. of participants (studies)	Certainty of the evidence (GRADE)
Anxiety	We could not present illustrative absolute effects because a representative control group risk could not be ascertained from the studies nor from any external source. Furthermore, results were reported in narrative form and varied considerably	<p>Effect measures (using change score (CS)) ranged from:</p> <p>SMD -5.1 (95% CI -6.1 to -4.1) for breast cancer to SMD -0.3 (95% CI -0.3 to 0.9) for prostate cancer</p> <p>Other cancer sites including colorectal and lung and trials including participants with mixed cancers</p> <p>The 5 trials reporting data where change scores could be calculated were generally very heterogeneous in terms of demographics, including age and gender (gender specific or mixed cancers), FIGO stage (early or advanced disease), and delivery of interventions and controls. This may have differed in the number of telephone calls and whether additional management components were used in intervention arms, and in controls being sufficiently different to consider any data synthesis methods</p>	<p>277 participants (5 studies)</p> <p>Sample sizes were often small, and baseline outcome values for intervention and control groups largely differed widely in 11 further studies. Therefore displaying only studies that used change scores seemed appropriate, and in future updates of the review, meta-analytical approaches will be attempted</p>	<p>⊕○○○</p> <p>Very low^{a,b,c}</p>
Depression		<p>Effect measures (CS) ranged from:</p> <p>SMD -2.2 (95% CI -2.7 to -1.7) for colorectal cancer to SMD 0.3 (95% CI 0.04 to 0.5) for mixed cancers</p> <p>Other cancer sites including breast, lung, and prostate cancer. There was scope in the breast and mixed cancer subgroups to potentially pool results, but even within these more restrictive analyses, there was considerable heterogeneity, imprecision, and inconsistency across trials. Therefore results were reported by single trials, and results were presented narratively</p> <p>The 9 trials reporting data where change scores could be calculated were generally very heterogeneous in terms of demographics, including age and gender (gender specific</p>	<p>1059 participants (9 studies)</p> <p>Sample sizes were often small, and baseline outcome values for intervention and control groups largely differed widely in 12 further studies. Therefore displaying only studies that used change scores seemed appropriate, and in future updates of the review, attempts at meta-analysis will be made</p>	<p>⊕○○○</p> <p>Very low^{a,b,c}</p>

	<p>ic or mixed cancers), FIGO stage (early or advanced disease), and delivery of interventions and controls. This may have differed in the number of telephone calls and whether additional management components were used in intervention arms, and in controls being sufficiently different to consider any data synthesis methods</p>	
<p>Fatigue</p>	<p>Effect measures (CS) ranged from:</p> <p>SMD -0.9 (95% CI -1.5 to -0.3) for breast cancer to SMD 0.0 (95% CI -0.2 to 0.2) for mixed cancers</p> <p>Another cancer site, including prostate cancer. There was scope in the mixed cancer subgroup to potentially pool results, but even within these more restrictive analyses, there were sufficient clinical differences between trials to justify not using this approach. Therefore results were reported by single trials and are presented narratively</p> <p>The 6 trials reporting data where change scores could be calculated were generally very heterogeneous in terms of demographics, including age and gender (gender specific or mixed cancers), FIGO stage (early or advanced disease), and delivery of interventions and controls</p>	<p>895 participants (6 studies)</p> <p>⊕○○○ Very low^{a,b,c}</p> <p>Sample sizes were often small, and baseline outcome values for intervention and control groups largely differed widely in 3 further studies. Therefore displaying only studies that used change scores seemed appropriate, and in future updates of the review, attempts at meta-analysis will be made</p>
<p>Emotional distress</p>	<p>SMDs (CS) in each individual trial all indicated uncertainty as to whether telephone interventions or control interventions were best for minimising emotional distress (all estimates were imprecise)</p> <p>Cancer sites included breast, prostate, and mixed cancers. There was scope in the breast cancer subgroup to potentially pool results, but there were sufficient clinical differences between the 2 trials in terms of including participants at different stages and ages to justify not using this approach. Therefore results were reported by single trials and are presented narratively</p> <p>The 5 trials reporting data where change scores could be calculated were generally very heterogeneous in terms of demographics, including age and gender (gender specific or mixed cancers), FIGO stage (early or advanced disease), and delivery of interventions and controls. This may have differed in the number of telephone calls and whether additional management components were used in intervention arms, and in controls being sufficiently different to consider any data synthesis methods</p>	<p>968 participants (5 studies)</p> <p>⊕○○○ Very low^{a,b,c}</p> <p>Sample sizes were often small, and baseline outcome values for intervention and control groups largely differed widely in 2 further studies. Therefore displaying only studies that used change scores seemed appropriate, and in future updates of the review, attempts at meta-analysis will be made</p>

Other outcomes included uncertainty, pain, sexually related symptoms, dyspnoea, and general symptoms. Data for any of these outcomes were not pooled due to considerable heterogeneity across all aspects. Magnitudes of effect were not reported

Studies for each outcome ranged from 2 to 6 (10 were included in the wider 'general symptoms' outcome)

⊕○○○

Very low^{a,b,c}

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: confidence interval; CS: change score; FIGO: International Federation of Gynecology and Obstetrics; SMD: standardised mean difference.

GRADE Working Group grades of evidence.

High certainty: further research is very unlikely to change our confidence in the estimate of effect.

Moderate certainty: further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low certainty: further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low certainty: we are very uncertain about the estimate.

^aDowngraded by one level due to concerns about overall risk of bias being unclear or high.

^bDowngraded by one level due to concerns about precision.

^cDowngraded by one level due to inconsistencies in results and general heterogeneity.

BACKGROUND

Description of the condition

People with cancer often experience a variety of symptoms as a result of their disease and its treatment (Harrington 2010; Kim 2012; Van Lancker 2014). As much cancer treatment is delivered on an ambulatory basis, patients and family members are largely responsible for their day-to-day management (Dodd 2000; McPherson 2014). Inadequate symptom management can result in early discontinuation of, or delays in, treatment (Cleeland 2009), and it has considerable implications for patient outcomes including functioning, psychological well-being, and quality of life (Dodd 2001; Glover 1995; Laugsand 2011).

Symptoms often manifest concurrently and appear related to one another. Symptom clusters - where three or more related symptoms manifest concurrently (Dodd 2001) - have become the subject of much contemporary research, with some evidence suggesting that they may have prognostic capabilities or may influence cancer outcomes (Cheville 2010). It is estimated that 40% of oncology patients experience more than one symptom at any one time (Kim 2009), and that as disease progresses, symptom burden rises. One study of 1000 people with cancer admitted to a palliative care unit determined that people experienced differing numbers of symptoms on admission, ranging between 1 and 29. However, the median number of symptoms that people presented with was 11 (Walsh 2000).

Attempts to reduce the incidence and severity of cancer symptoms have involved the development and testing of psycho-educational interventions to enhance patients' symptom self-management. These interventions may include therapeutic elements such as information exchange, problem-solving, coping skills training, and facilitating expression of emotions and concerns (Barsevick 2002). Although traditionally delivered face-to-face, interventions are increasingly being delivered by telephone (e.g. Freeman 2015), online (e.g. Steel 2016), or by mobile phone (e.g. Kearney 2009). These alternative modes of delivery are convenient for health professionals and patients alike. Integral to these is the delivery of supportive, interactive care provided by health professionals that provides patients with information about symptom management and support and encouragement in adopting effective self-care.

Description of the intervention

This review evaluates the effectiveness of telephone interventions delivered to people with cancer, with the aim of improving symptoms of the disease and/or its treatment. These interventions are typically educational or psychologically based in nature and may entail cognitive-behavioural, motivational, or supportive elements to facilitate patient management of symptoms. They can be delivered to patients alone or in conjunction with informal carers (family or friends). Further, they can be supplemented with face-to-face contact with health professionals and digital/online/printed educational materials.

Such interventions are gaining in popularity as health systems worldwide are challenged fiscally from having to care for increasingly ageing populations with limited available resources and soaring pharmacological and other healthcare costs. Interventions delivered by telephone are feasible and acceptable

to patients and offer health services a cheaper alternative to interventions delivered face-to-face.

How the intervention might work

Telephone interventions for symptom management may vary in terms of the symptom(s) they address, the theoretical frameworks underpinning them, the length of time over which they are delivered, and the training/qualifications of persons providing the telephone contact. However, whatever their make-up, telephone interventions are united in their potential for providing timely information and support to promote behaviour change and/or adherence with prescribed medications and/or recommended self-care, thereby enhancing patient outcomes and quality of life.

Why it is important to do this review

Although historically, information and support in managing symptoms were delivered face-to-face, increasingly this is not the case. The trend is for care to be provided nearer patients' homes, meaning that people with cancer are typically seeing hospital-based staff less often. Thus, there is a greater requirement for information and support in symptom management to be provided by other means, such as by telephone. Telephone interventions have been developed for a range of cancer symptoms (Scura 2004). Early indications suggest that these interventions have benefit, as they:

- reduce symptom severity;
- reduce symptom distress;
- enhance self-management of symptoms; and
- facilitate adaptation to symptoms.

However, evidence published to date has not been subject to rigorous systematic review. Four previous literature reviews have explored allied topics. Cox 2003 and Dickinson 2014 appraised and synthesised literature related to cancer follow-up (by telephone and through use of technology, respectively). Gotay 1998 reviewed outcomes of psychosocial support provided by telephone, and Galway 2012 reviewed psychosocial interventions (but did not focus on their delivery by telephone). Thus, none of these reviews explicitly analysed literature specifically evaluating telephone-delivered interventions for cancer symptoms. Further, the Gotay 1998 and Cox 2003 reviews are very much out-of-date. Thus, there is good justification to undertake a Cochrane systematic review to explore the effectiveness of telephone-delivered interventions for cancer symptoms.

OBJECTIVES

To assess the effectiveness of telephone-delivered interventions for reducing symptoms associated with cancer and its treatment. To determine which symptoms are most responsive to telephone interventions. To determine whether certain configurations (e.g. with/without additional support such as face-to-face, printed or electronic resources) and duration/frequency of intervention calls mediate observed cancer symptom outcome effects.

METHODS

Criteria for considering studies for this review

Types of studies

Randomised control trials (RCTs) and quasi-RCTs that:

- compared a telephone intervention with other types of interventions (e.g. a face-to-face intervention) and/or usual care; or
- compared different models of telephone interventions (i.e. with different content) against each other and/or a third arm comprising usual care.

Types of participants

We included studies evaluating telephone interventions for adult men and women (over 18 years of age) with a clinical diagnosis of cancer, regardless of tumour type, stage of disease, type of treatment, and time of recruitment (e.g. before, during, or after anticancer treatment).

We excluded studies that did not focus on cancer patients, or in which only a portion of the sample consisted of cancer patients.

Types of interventions

We included telephone interventions comprising any number of telephone calls delivered by any health or social care professional to cancer patients, with the stated aim of addressing any physical or psychological symptoms of cancer and its treatment. The interventions were referred to by study author(s) as psychological, psychosocial, psycho-educational, non-pharmacological, or supportive.

We excluded interventions that:

- were not primarily delivered by telephone (e.g. the main form of contact was face-to-face and the patient received a single telephone call to monitor progress), although we did include telephone interventions supported with printed/digital/online materials;
- aimed to improve patients' general well-being or adaptation to cancer including managing fear of recurrence (a common concern following cancer) (i.e. interventions that were not aimed primarily at improving cancer symptoms);
- evaluated triaging or monitoring care or treatment compliance; or
- were not delivered by a health or social care professional, or if details of the background of the person delivering the intervention could not be obtained.

Some interventions within the review incorporated elements other than telephone support. Thus, we categorised them according to whether they comprised solely telephone intervention or included additional supportive elements (e.g. face-to-face consultation, printed materials).

Types of outcome measures

We included data related to symptoms associated with cancer and its treatment, measured by standardised instruments that measured symptoms related to cancer with some evidence of validity and reliability.

Primary outcomes

- Anxiety (measured by validated instruments such as the Hospital Anxiety and Depression Scale (HADS) or the State Trait Anxiety Inventory (STAI))
- Depressive symptoms (measured by validated instruments such as the Hospital Anxiety and Depression Scale (HADS); the Beck Depression Inventory (BDI); or the Centre for Epidemiologic Studies Depression Scale (CES-D))
- Emotional distress (measured by validated instruments such as the Profile of Mood States (POMS))
- Uncertainty from being diagnosed with, and treated for, cancer (as measured by validated instruments such as the Mischel Uncertainty in Illness Scale)
- Fatigue (measured by validated instruments such as the Brief Fatigue Inventory (BFI); the Multi-dimensional Fatigue Inventory (MFI); or the Piper Fatigue Scale)
- Pain (measured by validated instruments such as the Brief Pain Inventory (BPI))
- Nausea/ vomiting (measured by validated instruments such as the Index of Nausea, Vomiting and Retching (INVR))
- Sexually-related symptoms (measured by validated instruments such as the Index of Sexual Satisfaction; the Female Sexual Function Index; or the International Index of Erectile Function)
- Lung cancer symptoms (measured by validated instruments such as the Functional Assessment of Cancer Therapy - Lung Cancer (FACT-L) or the Memorial Symptom Assessment Scale (MSAS) for dyspnoea items)

Secondary outcomes

- Symptom experience
- Symptom distress (as measured by validated instruments such as the General Symptom Distress Scale)

Search methods for identification of studies

We applied no language restriction for this review, so non-English publications were to be translated if necessary. This was not needed.

Electronic searches

We searched the following databases.

- Cochrane Central Register of Controlled Trials (CENTRAL; 2019, Issue 1), in the Cochrane Library ([Appendix 1](#)).
- MEDLINE via OVID (1946 to January 2019) ([Appendix 2](#)).
- Embase via OVID (1980 to January 2019) ([Appendix 3](#)).
- Cumulative Index to Nursing and Allied Health Literature (CINAHL) via Athens (1982 to January 2019).
- British Nursing Index (1984 to January 2019).
- PsycINFO (1989 to January 2019).

The search strategies are provided in the appendices.

Searching other resources

We searched conference proceedings to identify published abstracts, along with SIGLE (System for Information on Grey Literature in Europe) and trial registers for unpublished studies. We searched the reference lists of all included articles to identify

additional relevant studies. Finally, we handsearched the following journals from 2007 to 2019.

- *Cancer.*
- *Journal of Clinical Oncology.*
- *Psycho-oncology.*
- *Cancer Practice.*
- *Cancer Nursing.*
- *Oncology Nursing Forum.*
- *Journal of Pain and Symptom Management.*
- *Palliative Medicine.*

We found no additional studies, We found the studies identified for this review from the databases listed above.

Data collection and analysis

Selection of studies

Two review authors (from the pool of AEH, VHP, AC, KC, and ER) independently assessed the potential relevance of all titles and abstracts identified through the literature searches. We retrieved in full text studies identified by either review author as potentially relevant. Two review authors (of AEH, VHP, AC, and ER) independently assessed each of these studies against the review inclusion and exclusion criteria. A third review author resolved disagreements. Studies that appeared eligible for inclusion but were subsequently judged to not meet the selection criteria were detailed in the [Characteristics of excluded studies](#) table, including the specific reason(s) for exclusion (e.g. intervention not delivered by health or social care professionals).

Data extraction and management

We used standardised data extraction forms to extract all available data. Two independent review authors extracted data from each included study. We checked the forms against each other, and when we noted discrepancies, we referred to the original papers. We addressed unresolved discrepancies through discussion and consensus, involving the entire review team when necessary. We contacted study authors to obtain missing data. We extracted and reported on the following data.

- Geographic location.
- Sample demography (age, gender, tumour type, disease stage, treatment).
- Number of participants (including those lost to follow-up).
- Details of randomisation and allocation concealment.

- Aim of the intervention.
- Details of the intervention (number and frequency of telephone calls; duration of calls; health or social care professional(s) delivering intervention; incorporation of additional elements (face-to-face contacts, printed/digital/online materials, email contact)).
- Details of control/usual care.
- Primary and secondary outcome measures.
- Time points at which outcomes were collected and reported (frequency, length of follow-up).
- Reported statistics used to assess validity of results.
- Quality assurance processes used to ensure uniformity of intervention delivery (e.g. if intervention providers were trained and/or supervised; if a protocol was used; if an integrity check was described).

When possible, all data extracted were those relevant to an intention-to-treat analysis in which participants were analysed in the groups to which they were assigned. When study authors reported on the same piece of research in a series of publications, we considered the main study as the one that depicted the study design in detail and reported on primary outcomes of the study.

We managed data using Review Manager 5.3 ([RevMan 2014](#)).

Assessment of risk of bias in included studies

All review authors independently assessed and reported potential bias for each trial using the data extraction form. A third review author (ER) resolved any conflicts. We used the following criteria from the *Cochrane Handbook for Systematic Reviews of Interventions* as a guide for assessment.

- Random sequence generation.
- Allocation concealment.
- Blinding of participant, providers, outcome assessors, data analysts.
- Completeness of outcome data; adequate if less than 20% of patients were lost to follow-up and reasons for loss to follow-up were similar in both treatment arms.
- Selective reporting/intention-to-treat analysis.
- Other potential sources of bias.

We incorporated results of the assessment into the review through systematic narrative description and commentary about each of these domains. Further, we constructed a risk of bias graph ([Figure 1](#)) and a risk of bias summary ([Figure 2](#)).

Figure 1. Risk of bias graph: review authors' judgements about each risk of bias item presented as percentages across all included studies.

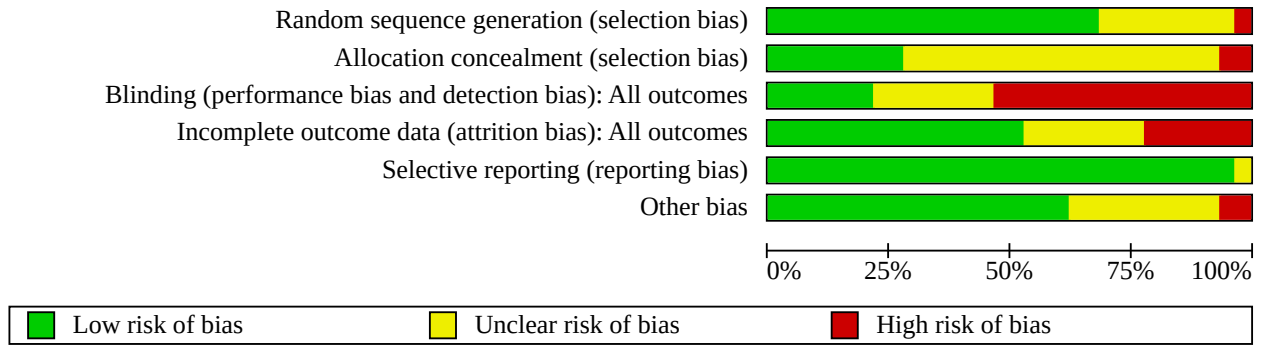


Figure 2. Risk of bias summary: review authors' judgements about each risk of bias item for each included study.

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding (performance bias and detection bias): All outcomes	Incomplete outcome data (attrition bias): All outcomes	Selective reporting (reporting bias)	Other bias
Allard 2007	+	?	-	?	+	?
Allen 2002	?	?	-	?	+	-
Badger 2005	-	?	-	?	+	+
Badger 2007	?	?	-	+	+	+
Badger 2013a	?	?	-	-	+	+
Badger 2013b	?	?	-	+	+	?
Badr 2015	?	?	-	+	+	?
Bailey 2004	+	?	-	+	+	+
Barsevick 2004	?	?	-	?	+	+
Barsevick 2010	+	+	-	+	+	+
Chambers 2014	+	+	?	-	+	?
Chambers 2015	+	+	-	+	+	+
Dong 2018	+	+	?	+	+	?
Downe-Wamboldt 2007	+	+	-	+	+	+
Girgis 2009	+	?	-	+	+	+
Kronke 2010	+	?	+	-	+	+
Livingston 2010	+	?	-	+	?	+
Mishel 2002	?	?	?	?	+	+
Mishel 2005	+	?	-	+	+	?
Molassiotis 2009	+	+	+	+	+	+
Mosher 2016	+	?	+	?	+	+
Porter 2011	+	+	+	?	+	?
Rawl 2002	?	-	+	-	+	+
Ream 2015	+	?	+	+	+	?
Reese 2014	?	?	-	+	+	?
Reese 2018	+	-	?	+	+	+
Shannon 2007	+	?	-	+	+	+

Figure 2. (Continued)

Reese 2018	+	-	?	+	+	+
Sherwood 2005	+	?	+	-	+	+
Sikorskii 2007	+	?	?	?	+	+
Thomas 2012	+	+	-	-	+	+
Traeger 2015	+	?	?	+	+	-
Watson 2017	+	?	?	-	+	?
Yates 2005	+	+	?	+	+	+

Measures of treatment effect

We processed data in accordance with guidance provided by the *Cochrane Handbook for Systematic Reviews of Interventions*. We analysed all outcomes in the review (anxiety, depressive symptoms, emotional distress, fatigue, pain, uncertainty, sexually-related issues) as continuous variables, reflecting how they were presented by study authors. As some studies had small samples, and because baseline scores for study outcomes tended to vary between intervention and control, we determined to input change scores and their associated standard deviations into the analyses. We extracted these statistics (when reported) and analysed them alongside (1) baseline, endpoint, and follow-up mean scores and associated standard deviations of outcomes of interest; (2) P values; and (3) numbers of patients who provided data at each assessment point to estimate the standardised mean difference (SMD) of change scores between treatment arms and its standard error.

Dealing with missing data

We did not impute missing outcome data for any of the outcomes other than to calculate missing standard deviations of change scores, as few authors reported these. We imputed these values using the approach of [Follmann 1992](#) and [Abrams 2005](#), as detailed in Section 16.1.3.2 of the *Cochrane Handbook for Systematic Reviews of Interventions*. This entailed calculating the correlation coefficient from one study in the symptom group that reported study outcomes in detail - including the standard deviation of the change score - and then using the summary statistics to determine standard deviation of change from baseline across other studies. Two study authors provided sufficient detail to enable calculation of standard deviations of change scores for symptoms of anxiety, depression, emotional distress, and fatigue ([Downe-Wamboldt 2007](#); [Ream 2015](#)).

Assessment of heterogeneity

We assessed and found considerable heterogeneity in included studies in terms of (1) interventions introduced; (2) types of participants; and (3) outcomes measured. All interventions were delivered primarily by telephone, but some also incorporated face-to-face elements and/or printed, digital, or online materials. Interventions varied in length and frequency and were provided at different times in the cancer journey from treatment to survivorship. There was some standardisation with regards to outcomes measured.

We did not assess methodological and statistical heterogeneity due to considerable clinical heterogeneity across trials. Most studies compared a telephone intervention with usual care, five compared

two different interventions with usual care ([Badger 2007](#); [Dong 2018](#); [Girgis 2009](#); [Livingston 2010](#); [Thomas 2012](#)), and six compared two interventions without a usual care arm ([Badger 2013a](#); [Badger 2013b](#); [Chambers 2014](#); [Reese 2018](#); [Sikorskii 2007](#); [Watson 2017](#)).

For future updates, we plan to assess statistical heterogeneity between study outcomes by visually inspecting forest plots and by calculating the I² statistic (estimation of the percentage of heterogeneity between trials that cannot be ascribed to sampling variation ([Higgins 2003](#))), and when possible, by conducting subgroup analyses (see later). If we find evidence of substantial heterogeneity, we will investigate and report the possible reasons for this.

Data synthesis

We did not perform meta-analyses due to considerable heterogeneity.

For future updates of the review, we will do the following.

- We will use random-effects models with inverse variance weighting for all meta-analyses ([DerSimonian 1986](#)).
- We will calculate SMDs in outcomes between telephone and control groups (rather than mean differences, if appropriate) to take account of the different scales used across studies to measure different symptom outcomes.
- When we are unable to obtain required data to incorporate studies into meta-analyses, or when we identify insufficient studies related to management of a particular symptom, we will continue to report study findings in a narrative fashion.

Subgroup analysis and investigation of heterogeneity

For future updates, we will present subgroups on forest plots and will aim to determine whether there is a difference in outcomes according to whether telephone interventions are provided on their own or in conjunction with other elements (e.g. printed materials, face-to-face meetings).

Sensitivity analysis

We found an insufficient number of studies (and no meta-analyses were conducted) to allow review authors to undertake sensitivity analysis to determine the effect of including/excluding studies with high risk of bias (e.g. as a result of inadequate concealment of allocation).

RESULTS

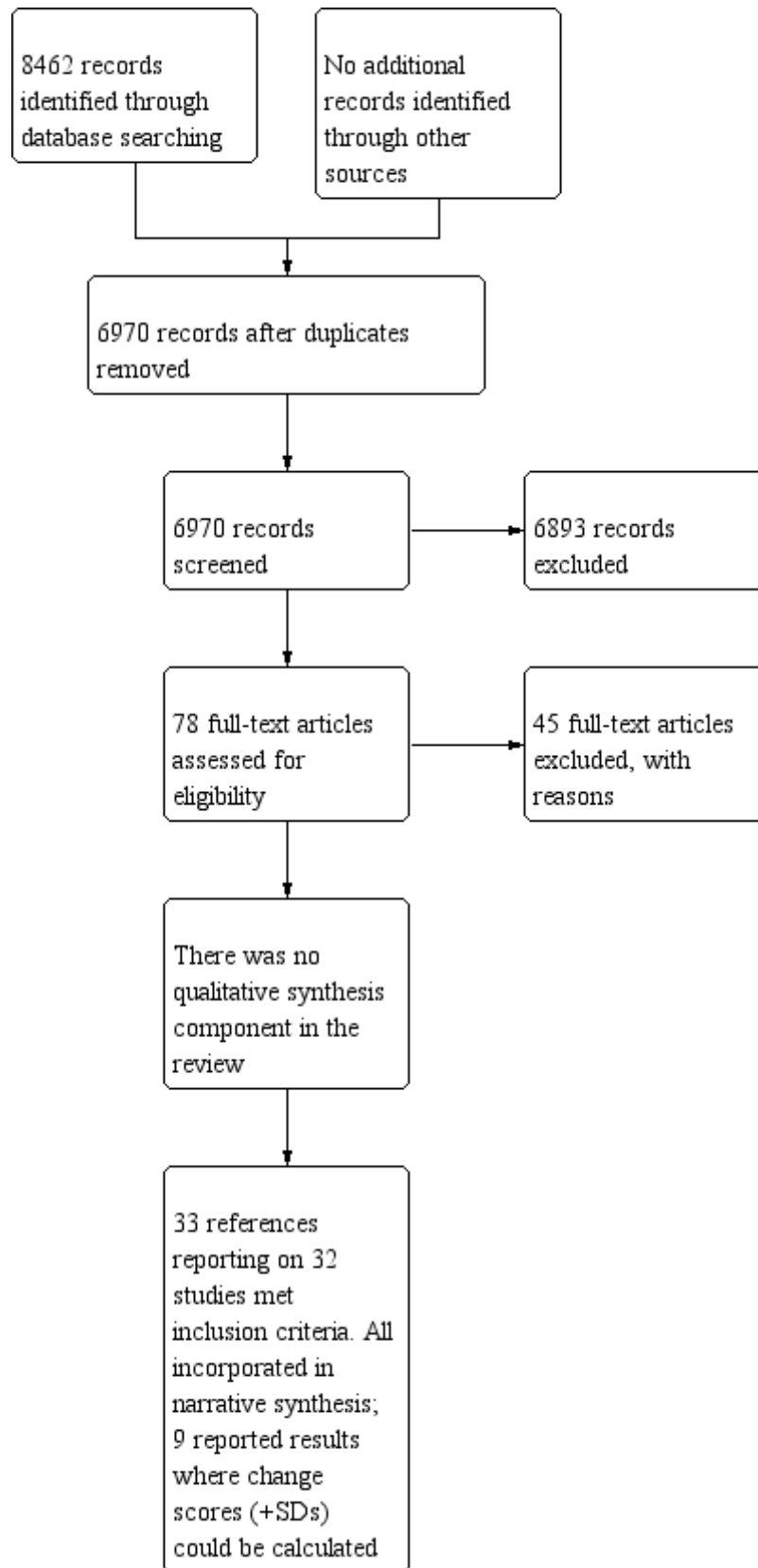
Description of studies

See [Characteristics of included studies](#) and [Characteristics of excluded studies](#).

Results of the search

After duplicates and clearly irrelevant articles were eliminated, the electronic and manual search (to January 2019) yielded 78 studies that were potentially eligible for inclusion. After assessing the full text of studies against the inclusion criteria, we excluded 45 studies, leaving 33 studies for inclusion in the review. [Gil 2006](#) was nested in the [Mishel 2005](#) study because it reported on long-term outcomes of this study. The PRISMA flow chart is presented in [Figure 3](#).

Figure 3. Study flow diagram.



Included studies

Study design

We included 31 randomised controlled trials (RCTs); the final study employed a repeated-measures experimental design but did not use random principles for assignment to intervention and control groups (Badger 2005). Ten studies appeared adequately powered (i.e. study authors had determined the sample size required to show effect and had managed to recruit to this target) (Allard 2007; Chambers 2014; Girgis 2009; Kroenke 2010; Molassiotis 2009; Mosher 2016; Sherwood 2005; Sikorskii 2007; Traeger 2015; Yates 2005). Three further studies calculated sample sizes required, but due to under-recruitment (Livingston 2010; Watson 2017), or higher than predicted attrition (Thomas 2012), these studies were underpowered. Most studies did not specify required sample size to attain adequate power. Five were pilot studies with small sample sizes (Badger 2005; Badr 2015; Bailey 2004; Reese 2014; Reese 2018).

Most studies compared a telephone-delivered intervention with usual care. However, of the included studies, five compared an intervention delivered via telephone with an attentional, rather than usual care, control (Badger 2005; Badger 2007; Barsevick 2004; Barsevick 2010; Mosher 2016). A further study provided a wait-list control whereby participants in the control group received the intervention on completion of study assessments (Reese 2014). Four studies compared a telephone-delivered intervention against augmented usual care, which incorporated additional elements including passive referral to a help line; education/support; nutrition information; or extra information about symptoms reported by patients given to the treating oncologist (Barsevick 2004; Kroenke 2010; Mosher 2016; Porter 2011). Six studies compared two/three interventions but incorporated no control group (Badger 2013a; Badger 2013b; Chambers 2014; Reese 2018; Sikorskii 2007; Watson 2017). One was an equivalence trial (Watson 2017). Eight studies included three study arms that incorporated telephone-delivered intervention(s) with alternative intervention with/without a control group (usual care or attentional control) (Badger 2007; Badger 2013a; Chambers 2015; Dong 2018; Girgis 2009; Livingston 2010; Mishel 2002; Thomas 2012).

Contact with study authors

We contacted five study authors to clarify issues around provision of the intervention; we subsequently excluded two articles from the review, as data obtained from the study author made it clear that these studies were ineligible. We included three studies once the nature of the interventions and who delivered them were clarified (Badger 2013a; Reese 2014; Reese 2018). Further, we contacted one study author with regards to the Mishel 2002 study, to query numbers of participants in each study arm.

Sample size

Sample sizes ranged from 23 in Reese 2014 to 575 in Mishel 2005, with a total of 6250 cancer patients recruited across the 32 studies.

Setting

Twenty-one studies were conducted in the USA (Allen 2002; Badger 2005; Badger 2007; Badger 2013a; Badger 2013b; Badr 2015; Bailey 2004; Barsevick 2004; Barsevick 2010; Kroenke 2010; Mishel 2002; Mishel 2005; Mosher 2016; Porter 2011; Rawl 2002; Reese 2014; Reese 2018; Sherwood 2005; Sikorskii 2007; Thomas 2012; Traeger

2015). Four were conducted in Australia (Chambers 2014; Chambers 2015; Girgis 2009; Yates 2005), two in Canada (Allard 2007; Downe-Wamboldt 2007), three in the UK (Molassiotis 2009; Ream 2015; Watson 2017), and one in China (Dong 2018), and one recruited across two countries - Australia and Canada (Livingston 2010).

Participants

Nine studies addressed breast cancer exclusively (Allard 2007; Allen 2002; Badger 2005; Badger 2007; Badger 2013a; Badger 2013b; Mishel 2005; Reese 2018; Yates 2005), three solely recruited men with prostate cancer (Bailey 2004; Chambers 2015; Mishel 2002), three recruited only lung cancer patients (Badr 2015; Mosher 2016; Porter 2011), and two recruited only people diagnosed with colorectal cancer (Dong 2018; Reese 2014).

Eleven studies included heterogeneous samples of cancer patients, most commonly with a combination of breast, colorectal, lung, and prostate cancer (Barsevick 2010; Chambers 2014; Downe-Wamboldt 2007; Girgis 2009; Kroenke 2010; Livingston 2010; Molassiotis 2009; Rawl 2002; Ream 2015; Traeger 2015; Watson 2017); four studies did not specify the type of cancer diagnosis patients had received (Barsevick 2004; Sherwood 2005; Sikorskii 2007; Thomas 2012).

In addition to including people with cancer, 11 studies addressed partners or carers of cancer patients; five included partners of women with breast cancer (Badger 2005; Badger 2007; Badger 2013a; Badger 2013b; Reese 2018), one included partners of men with prostate cancer (Chambers 2015), one included partners of patients treated for colorectal cancer (Reese 2014), one included partners of people diagnosed with lung cancer (Badr 2015), two included carers of people diagnosed with lung cancer (Mosher 2016; Porter 2011), and one included carers of people with a range of cancers (Chambers 2014). Outcomes for the partners/carers reported in these studies are not included in this review.

Most studies (n = 14) recruited patients with early-stage cancer. Remaining studies recruited patients with early/locally advanced cancer (n = 2) (Badger 2005; Reese 2018), or people with advanced cancer (n = 4) (Badr 2015; Girgis 2009; Molassiotis 2009; Sherwood 2005), or they did not (n = 11) specify the disease stage for eligible patients (Badger 2005; Barsevick 2010; Chambers 2014; Dong 2018; Kroenke 2010; Mosher 2016; Sikorskii 2007; Ream 2015; Reese 2014; Thomas 2012; Watson 2017). One study specifically targeted cancer survivors who were between five and nine years post treatment (Mishel 2005).

Most studies recruited consecutive patients irrespective of symptom intensity; few (n = 6) recruited people whose symptoms had attained a threshold level (Chambers 2014; Dong 2018; Kroenke 2010; Mosher 2016; Ream 2015; Reese 2018).

Symptoms

Interventions introduced across studies aimed to reduce a variety of symptoms caused by cancer and its treatment; psychological and emotional symptoms were frequently assessed. Sixteen studies measured effects of telephone-delivered interventions on anxiety (Badger 2007; Badger 2013b; Badr 2015; Bailey 2004; Dong 2018; Girgis 2009; Livingston 2010; Molassiotis 2009; Mosher 2016; Porter 2011; Rawl 2002; Ream 2015; Reese 2018; Traeger 2015; Watson 2017; Yates 2005). Depressive symptoms was an outcome measured in 21 studies (Badger 2005; Badger 2007; Badger 2013a;

Badger 2013b; Badr 2015; Bailey 2004; Barsevick 2010; Dong 2018; Downe-Wamboldt 2007; Girgis 2009; Kroenke 2010; Livingston 2010; Molassiotis 2009; Mosher 2016; Porter 2011; Rawl 2002; Ream 2015; Reese 2018; Traeger 2015; Watson 2017; Yates 2005). Seven studies focused more broadly on emotional distress (Allard 2007; Allen 2002; Bailey 2004; Chambers 2014; Downe-Wamboldt 2007; Livingston 2010; Mishel 2005). Finally, three interventions aimed to alleviate uncertainty resulting from diagnosis of, and treatment for, cancer (Bailey 2004; Mishel 2002; Mishel 2005).

With regards to symptoms with a physical element, nine studies measured the impact of interventions on fatigue (Badger 2005; Badger 2013b; Bailey 2004; Barsevick 2004; Barsevick 2010; Molassiotis 2009; Mosher 2016; Ream 2015; Yates 2005), six on cancer-related pain (Barsevick 2010; Kroenke 2010; Molassiotis 2009; Mosher 2016; Porter 2011; Thomas 2012), three on sexually-related symptoms (Chambers 2015; Reese 2014; Reese 2018), and two on dyspnoea (Mosher 2016; Porter 2011).

Ten studies measured general symptom experience and reported overall symptom intensity and/or symptom distress (Badger 2013a; Badger 2013b; Barsevick 2010; Kroenke 2010; Mishel 2002; Molassiotis 2009; Porter 2011; Sherwood 2005; Sikorskii 2007; Traeger 2015).

Many symptoms were incorporated as secondary outcomes across studies. For example, a study evaluating a telephone-delivered intervention for fatigue may also have incorporated anxiety and/or depressive symptoms as secondary outcomes.

Intervention format

A summary of intervention characteristics is provided in the table titled [Characteristics of included studies](#). All interventions were delivered primarily by telephone to participants in their homes. However, only ten were delivered solely by telephone (Allard 2007; Badger 2005; Badger 2007; Badger 2013b; Bailey 2004; Dong 2018; Downe-Wamboldt 2007; Livingston 2010; Reese 2014; Traeger 2015).

Sixteen of the remaining study interventions were delivered by telephone in combination with printed materials and/or online/digital materials (Badger 2013a; Badr 2015; Barsevick 2004; Barsevick 2010; Chambers 2014; Chambers 2015; Girgis 2009; Mishel 2002; Mishel 2005; Mosher 2016; Porter 2011; Ream 2015; Reese 2018; Sikorskii 2007; Thomas 2012; Watson 2017).

Two studies combined telephone calls with face-to-face sessions (Molassiotis 2009; Sherwood 2005), and three combined telephone calls with both face-to-face sessions and digital/printed materials (Allen 2002; Rawl 2002; Yates 2005). One final study evaluated an intervention that incorporated automated symptom monitoring and prescribing recommendations (regarding depressive symptoms and fatigue) made to participants by oncologists, in addition to telephone calls (Kroenke 2010).

Health professionals delivering interventions

Most interventions (n = 24) were delivered by nurses (Allard 2007; Allen 2002; Badger 2005; Badger 2007; Bailey 2004; Barsevick 2004; Barsevick 2010; Chambers 2014; Chambers 2015; Downe-Wamboldt 2007; Girgis 2009; Kroenke 2010; Livingston 2010; Mishel 2002; Mishel 2005; Molassiotis 2009; Porter 2011; Rawl 2002; Ream

2015; Sherwood 2005; Sikorskii 2007; Thomas 2012; Traeger 2015; Yates 2005), including:

- oncology nurses (Barsevick 2010; Chambers 2014; Chambers 2015; Girgis 2009; Livingston 2010; Rawl 2002; Ream 2015; Sherwood 2005; Sikorskii 2007; Thomas 2012; Traeger 2015; Yates 2005);
- research nurses (Allen 2002; Barsevick 2004); and
- psychiatric nurses (Badger 2005; Badger 2007).

Eight studies did not specify the specialty or training of nurses who delivered the interventions (Allard 2007; Bailey 2004; Downe-Wamboldt 2007; Kroenke 2010; Mishel 2002; Mishel 2005; Molassiotis 2009; Porter 2011). In nine studies, the nurses had received training in other skills necessary for intervention delivery, including counselling (Badger 2005; Badger 2007; Barsevick 2004; Downe-Wamboldt 2007; Girgis 2009; Livingston 2010; Ream 2015), communication (Chambers 2015), and education (Kroenke 2010).

In nine studies, interventions were delivered by professionals other than nurses. These included psychologists (Chambers 2014; Dong 2018; Reese 2014; Reese 2018; Watson 2017), social workers (Badger 2013a; Mosher 2016), a mental health counsellor (Badr 2015), and master's prepared social workers and para-professionals/psychologists/counsellors (Badger 2013b).

Theoretical basis of interventions

Twenty-one studies used theoretical models to inform the telephone intervention: Self-Regulation Theory (Allard 2007; Rawl 2002; Ream 2015), Interpersonal Therapy (Badger 2005; Badger 2007; Badger 2013a), the Stress Process Model (Badger 2013b), the Common Sense Model of Illness (Barsevick 2004; Barsevick 2010), the Transtheoretical Model (Thomas 2012), Self-Determination Theory (Badr 2015), Cognitive-Behavioural Theory/Coping Skills Training (Downe-Wamboldt 2007; Porter 2011; Sherwood 2005), the PRECEDE Model of Health Behaviour (Yates 2005), Social Cognitive Theory (Mosher 2016), the Theory of Uncertainty in Illness (Bailey 2004; Mishel 2002; Mishel 2005), and the Cognitive-Behavioural and Sex Therapy Theory (Reese 2014; Reese 2018).

The theoretical basis informing the design of the interventions was unclear - or unreported - in 11 studies (Allen 2002; Chambers 2014; Chambers 2015; Dong 2018; Girgis 2009; Kroenke 2010; Livingston 2010; Molassiotis 2009; Sikorskii 2007; Traeger 2015; Watson 2017). Although Allen 2002 did not specify the theoretical basis for the intervention provided, the intervention model for this study was based around problem-solving and made use of motivational techniques in its delivery. Finally, Sikorskii 2007 employed a multi-dimensional interactive approach, which drew on various strategies around coping, re-framing, providing education, and eliciting support for adapting to, or overcoming, cancer-related symptoms.

Number of calls/duration/timing

Most (n = 29) of the interventions provided a standardised number of telephone calls to participants; these ranged from one call in Chambers 2014 to 18 calls delivered weekly over 18 weeks in Molassiotis 2009. In the remaining three studies, the number of calls provided varied by need (Downe-Wamboldt 2007; Watson 2017), or the numbers of calls and the intervals between them were unclear (Girgis 2009).

Although most ($n = 17$) intervention calls were delivered weekly, three interventions were delivered every other week (Allen 2002; Rawl 2002; Thomas 2012). One intervention delivered three calls over three successive cycles of chemotherapy (Ream 2015), one provided four calls over two successive cycles of chemotherapy (Traeger 2015), and seven others provided a series of calls with increasing time intervals between them (Chambers 2015; Girgis 2009; Kroenke 2010; Livingston 2010; Porter 2011; Sherwood 2005; Sikorskii 2007). In one study, calls were made over a three-month period at participants' convenience (Downe-Wamboldt 2007), and in another, up to eight calls were again scheduled over an approximate 12-week period (Watson 2017). Finally, Chambers 2014 made five calls but did not specify the timing of these calls.

Most frequently, the intervention comprised three or four calls (Allen 2002; Badger 2013a; Barsevick 2004; Barsevick 2010; Girgis 2009; Livingston 2010; Mishel 2005; Mosher 2016; Ream 2015; Reese 2014; Reese 2018; Sherwood 2005; Thomas 2012; Traeger 2015; Watson 2017). In three studies, participants received more than 10 calls: Porter 2011 delivered 14; Molassiotis 2009 provided 18; and although Kroenke 2010 planned to deliver four calls, automated symptom reports with high symptom scores triggered extra calls - participants received on average 11 calls.

Twenty-two studies reported the duration of calls provided (Badger 2005; Badger 2007; Badger 2013a; Badger 2013b; Badr 2015; Bailey 2004; Barsevick 2004; Chambers 2014; Chambers 2015; Dong 2018; Kroenke 2010; Livingston 2010; Molassiotis 2009; Mosher 2016; Porter 2011; Rawl 2002; Ream 2015; Reese 2014; Reese 2018; Thomas 2012; Traeger 2015; Yates 2005); duration ranged on average from 10 minutes in Yates 2005 to 70 minutes in Reese 2018.

Recording and documentation of calls

Only 15 studies recorded telephone calls in some way for quality assurance (Badger 2005; Badger 2007; Badger 2013b; Barsevick 2004; Barsevick 2010; Chambers 2014; Chambers 2015; Livingston 2010; Mosher 2016; Porter 2011; Rawl 2002; Ream 2015; Reese 2018; Sherwood 2005; Traeger 2015). Downe-Wamboldt 2007 assessed fidelity by reviewing the interventionist's notes to determine goals recorded and degree of problem-solving achieved. Watson 2017 addressed fidelity by observing some of the intervention sessions.

Telephone calls were additionally documented in some way as part of the intervention (not explicitly for quality purposes): two used patient-completed worksheets completed after calls (Allen 2002; Badger 2007); four obtained feedback surveys/documentation completed on a computer or via a touchpad telephone (Girgis 2009; Kroenke 2010; Livingston 2010; Sikorskii 2007); one reviewed and assigned homework during intervention calls (Badr 2015); another required the nurse delivering the intervention to keep a diary recording participants' engagement with calls (Ream 2015); and two used patient diaries completed between telephone calls to help tailor the intervention (Barsevick 2004; Barsevick 2010).

Excluded studies

In the [Characteristics of excluded studies](#) table, we list the 45 studies excluded after assessment of full text (according to the criteria specified at [Types of interventions](#)) and specify the reasons for exclusion.

We excluded studies primarily because:

- they did not address management of symptoms ($n = 21$); or
- interventions were not delivered primarily by telephone ($n = 12$).

Risk of bias in included studies

Assessments of risk of bias and methodological certainty are provided in the [Characteristics of included studies](#) table and the Risk of bias graph (Figure 1).

Here we summarise risk of bias.

- Adequate sequence generation: fulfilled in 22 studies (Allard 2007; Bailey 2004; Barsevick 2010; Chambers 2014; Chambers 2015; Dong 2018; Downe-Wamboldt 2007; Girgis 2009; Kroenke 2010; Livingston 2010; Mishel 2005; Molassiotis 2009; Mosher 2016; Porter 2011; Rawl 2002; Ream 2015; Reese 2018; Sikorskii 2007; Thomas 2012; Traeger 2015; Watson 2017; Yates 2005).
- Allocation concealment: fulfilled in 10 studies (Barsevick 2010; Chambers 2014; Chambers 2015; Dong 2018; Downe-Wamboldt 2007; Molassiotis 2009; Porter 2011; Thomas 2012; Traeger 2015; Yates 2005).
- Blinding: fulfilled in six studies (Kroenke 2010; Molassiotis 2009; Porter 2011; Rawl 2002; Ream 2015; Sherwood 2005).
- Incomplete outcome data assessed: fulfilled in 17 studies (Badger 2007; Badger 2013b; Badr 2015; Bailey 2004; Barsevick 2010; Chambers 2015; Dong 2018; Downe-Wamboldt 2007; Girgis 2009; Livingston 2010; Mishel 2005; Molassiotis 2009; Ream 2015; Reese 2014; Reese 2018; Traeger 2015; Yates 2005).
- Free of selective reporting: fulfilled in 30 studies (Allard 2007; Allen 2002; Badger 2005; Badger 2007; Badger 2013a; Badger 2013b; Badr 2015; Bailey 2004; Barsevick 2004; Barsevick 2010; Chambers 2014; Chambers 2015; Dong 2018; Downe-Wamboldt 2007; Girgis 2009; Kroenke 2010; Mishel 2002; Mishel 2005; Molassiotis 2009; Mosher 2016; Porter 2011; Rawl 2002; Ream 2015; Reese 2014; Reese 2018; Sherwood 2005; Sikorskii 2007; Thomas 2012; Traeger 2015; Yates 2005).
- Other bias: detected in 12 studies (Allard 2007; Allen 2002; Badger 2013b; Badr 2015; Chambers 2014; Dong 2018; Mishel 2005; Porter 2011; Ream 2015; Reese 2014; Traeger 2015; Yates 2005).

Effects of interventions

See: [Summary of findings 1](#) Summary of findings

1. Findings by symptom

1.1 Anxiety; telephone intervention versus control

Sixteen studies measured effects of their interventions on anxiety (Badger 2007; Badger 2013b; Badr 2015; Bailey 2004; Dong 2018; Girgis 2009; Livingston 2010; Molassiotis 2009; Mosher 2016; Porter 2011; Rawl 2002; Ream 2015; Reese 2018; Traeger 2015; Watson 2017; Yates 2005). These studies used several different validated scales.

- Hospital Anxiety and Depression Scale (Girgis 2009; Livingston 2010; Molassiotis 2009; Ream 2015; Watson 2017; Yates 2005).
- State version of the State-Trait Anxiety Inventory (STAI) (Badger 2013b; Rawl 2002).
- Trait Anxiety version of the State-Trait Anxiety Inventory (STAI) (Porter 2011).

- Anxiety subscale of the Profile of Mood States - Short Form (Bailey 2004).
- Hamilton Anxiety Scale (HAMA) (Chinese version) (Dong 2018).
- Self-Rating Anxiety Scale (SAS) (Chinese version) (Dong 2018).
- Investigator-designed instrument composed of the Positive and Negative Affect Schedule (PANAS), the Short Form-12 (SF-12) Scale, and the Index of Clinical Stress (ICS) (Badger 2007).
- 6-item Patient-Reported Outcomes Measurement Information System short form anxiety measure (Badr 2015).
- 2-item Generalised Anxiety Disorder (GAD-2) Scale (Traeger 2015); 7-item Generalised Anxiety Disorder (GAD-7) Scale (Mosher 2016; Reese 2018).

In nine studies, anxiety was a primary outcome (Badger 2007; Badger 2013b; Badr 2015; Girgis 2009; Livingston 2010; Mosher 2016; Porter 2011; Rawl 2002; Watson 2017). Two of these (Badger 2007; Badr 2015), plus three studies in which anxiety was a secondary outcome (Bailey 2004; Dong 2018; Ream 2015), provided data from 277 participants that reported change scores and associated standard deviations - or these could be calculated. Dong 2018 measured anxiety outcomes with two measures (HAMA and SAS); only those related to SAS are displayed on the forest plot. This decision was based on the allied Self-Rating Depression Scale, which was used to power the study. Three individual studies appeared to suggest that use of a telephone had a significant impact on symptom management of anxiety in participants with various types of cancer (Badger 2007; Badr 2015; Dong 2018), but another two trials found no evidence of a difference (Bailey 2004; Ream 2015). We did not pool the overall effect estimates due to considerable heterogeneity, but we have depicted results of individual trials in the forest plot in [Analysis 1.1](#).

It is interesting to note that Dong 2018, a three-arm trial incorporating a telephone support arm in addition to telephone-delivered reminiscence therapy and usual care control, determined that generic telephone support generated similar improvements in anxiety as telephone-based reminiscence therapy.

Of the trials that did not report a magnitude of effect explicitly nor address baseline imbalance using change scores (or these and their associated standard deviations could not be calculated) - and in which anxiety was a primary outcome - three reported some effect on anxiety. Badger 2013b tested two different forms of telephone-delivered intervention without usual care control. Although the study determined that interventions were associated with statistically significant decreases in anxiety, without a control group for comparison it is difficult to conclude that these improvements were generated by the intervention rather than by an alternative factor (including passage of time). Watson 2017 also had no control group, but this was an equivalence trial - researchers were seeking to determine equivalence between standard and telephone-delivered cognitive-behavioural therapy (CBT) in people with high psychological needs; conventional CBT has established efficacy. Watson 2017 determined that with regards to anxiety, telephone-delivered and face-to-face CBT approaches were equally effective. Finally, Rawl 2002, in the trial report, noted improvements in anxiety nearing statistical significance ($P = 0.09$). The other four studies did not detect any significant intervention effect (Girgis 2009; Livingston 2010; Mosher 2016; Porter 2011).

1.2 Depressive symptoms; telephone intervention versus control

Twenty-one studies measured effects of interventions on symptoms of depression (Badger 2005; Badger 2007; Badger 2013a; Badger 2013b; Badr 2015; Bailey 2004; Barsevick 2010; Dong 2018; Downe-Wamboldt 2007; Girgis 2009; Kroenke 2010; Livingston 2010; Molassiotis 2009; Mosher 2016; Porter 2011; Rawl 2002; Ream 2015; Reese 2018; Traeger 2015; Watson 2017; Yates 2005).

The following validated measurement scales were used to measure depressive symptoms.

- Centre for Epidemiologic Studies Depression Scale (CES-D) (Badger 2005; Badger 2007; Badger 2013a; Badger 2013b; Downe-Wamboldt 2007; Rawl 2002).
- Hospital Anxiety and Depression Scale (Girgis 2009; Livingston 2010; Molassiotis 2009; Ream 2015; Watson 2017; Yates 2005).
- Profile of Mood States - Short Form (POMS-SF) (Bailey 2004; Barsevick 2010).
- Hamilton Depression Scale (HAMD) (Chinese version) (Dong 2018).
- Self-Rating Depression Scale (SDS) (Chinese version) (Dong 2018).
- Depression severity subscale of the 36-item Short Form Health Survey (SF-36) (Kroenke 2010).
- Short Form-12 (SF-12) (Barsevick 2010).
- Hopkins Symptom Checklist (HSCL-20) (Kroenke 2010).
- Patient Health Questionnaire (PHQ-9) (Kroenke 2010; Mosher 2016; Reese 2018).
- 2-item Patient Health Health Questionnaire (PHQ-2) (Traeger 2015).
- Beck Depression Inventory (BDI) (Porter 2011).
- 6-item Patient-Reported Outcomes Measurement Information System short-form depression measure (Badr 2015).

In all but six studies (Bailey 2004; Barsevick 2010; Molassiotis 2009; Ream 2015; Traeger 2015; Yates 2005), depressive symptoms was the primary outcome. Nine studies, including 1059 participants, reported quantitative results such as change scores - or data enabling these and their associated standard deviations to be calculated (Badger 2005; Badger 2007; Badr 2015; Bailey 2004; Barsevick 2010; Dong 2018; Downe-Wamboldt 2007; Kroenke 2010; Ream 2015) ([Analysis 1.2](#)). Dong 2018 measured depressive symptom outcomes with two measures (HAMD and SDS); only those related to SDS were included in the forest plot. This decision was based on use of the SDS to power the study. Three of these studies had measured depression as a secondary outcome (Bailey 2004; Barsevick 2010; Ream 2015). Four individual studies appeared to suggest that use of a telephone had significant impact on symptom management of depression among participants with various types of cancer (Badger 2007; Badr 2015; Dong 2018; Kroenke 2010); Downe-Wamboldt 2007 was of borderline significance. Many trials were small and underpowered and did not report magnitude of effect. We did not pool the overall effect estimates due to considerable heterogeneity, but we have depicted the results of individual trials in the forest plot in [Analysis 1.2](#).

Of trials that did not report a magnitude of effect explicitly nor address baseline imbalance using change scores (or these and their associated standard deviations could not be calculated) - and that had symptoms of depression as a primary outcome -

four reported some improvement in the intervention arm. [Rawl 2002](#) measured the impact of a telephone-delivered intervention on depressive symptoms in patients with newly diagnosed cancer. Researchers found that patients who received the intervention had significantly fewer depressive symptoms ($P = 0.05$) midway through the intervention when compared to those in the usual care group, although this was not maintained. One month post intervention, the difference was no longer statistically significant ($P = 0.07$). [Porter 2011](#) detected reduced symptoms of depression in lung cancer patients following provision of both of their two telephone-delivered interventions (coping skills training (CST) versus education/support) ($B = -5.55$, standard error = 0.28, $P = 0.05$). As with the intervention's effects on anxiety, levels of depressive symptoms dropped more for patients with stage I cancer given the education/support intervention, and more for patients with stage II-III cancer given the CST intervention ($B = -2.38$, standard error = 2.86, $P = 0.006$). [Badger 2013a](#) and [Badger 2013b](#) reported statistically significant decreases in depressive symptoms over time associated with delivery of two forms of telephone-delivered intervention. However, neither study incorporated a usual care control for comparison. Finally, [Watson 2017](#), an equivalence trial comparing face-to-face and telephone-delivered CBT, determined that the telephone-delivered version generated a statistically significant reduction in depressive symptoms equivalent to that seen with CBT delivered in-person.

Three studies whose primary outcome was reduced depressive symptoms found no significant reductions in these generated by the interventions ([Girgis 2009](#); [Livingston 2010](#); [Mosher 2016](#)).

1.3 Fatigue; telephone intervention versus control

Nine studies reported on effects of interventions on fatigue. Six evaluated interventions delivered specifically to reduce fatigue ([Badger 2005](#); [Barsevick 2004](#); [Barsevick 2010](#); [Mosher 2016](#); [Ream 2015](#); [Yates 2005](#)). Another three studies measured fatigue as a secondary outcome arising from the interventions ([Badger 2013b](#); [Bailey 2004](#); [Molassiotis 2009](#)).

Fatigue was measured using the following measurement tools.

- Multi-Dimensional Fatigue Inventory (MFI) ([Badger 2005](#); [Badger 2013b](#)).
- Brief Fatigue Inventory (BFI) ([Ream 2015](#)).
- Fatigue Distress Scale ([Ream 2015](#)).
- Schwartz Cancer Fatigue Scale ([Barsevick 2004](#)).
- General Fatigue Scale (GFS) ([Barsevick 2004](#); [Barsevick 2010](#)).
- Profile of Mood States (POMS), fatigue subscale ([Barsevick 2004](#); [Barsevick 2010](#)).
- National Cancer Institute Common Toxicity Criteria (NCI-CTC) - single item ([Molassiotis 2009](#)).
- Fatigue Symptom Inventory ([Mosher 2016](#)).

Six studies reported quantitative results including change scores - or data enabling these and their associated standard deviations to be calculated ([Badger 2005](#); [Bailey 2004](#); [Barsevick 2004](#); [Barsevick 2010](#); [Ream 2015](#); [Yates 2005](#)). All but [Bailey 2004](#) incorporated fatigue as a primary outcome. Collectively, trial authors had attained data on 895 participants. Three of the individual studies appeared to suggest that use of the telephone had a significant impact on symptom management of fatigue for participants with various types of cancer ([Badger 2005](#); [Barsevick 2004](#); [Yates](#)

[2005](#)). Although the intervention evaluated by [Ream 2015](#) in a feasibility trial did not significantly reduce overall (global) fatigue, study authors reported that it did generate significant reductions in distress caused by the symptom ($P < 0.05$). Other studies found no evidence of any differences between arms ([Bailey 2004](#); [Barsevick 2010](#)). We did not pool the overall effect estimates due to considerable heterogeneity, but we have depicted the results of individual trials in the forest plot in [Analysis 1.3](#).

[Molassiotis 2009](#) reported significant reductions in fatigue generated by a telephone-delivered home care intervention delivered weekly over 18 weeks to people given a course of oral capecitabine. However, differences in comparison with the usual care control declined over time. Greatest improvement in fatigue was found from cycle 0 to 2 ($P = 0.005$); by the end of treatment, these benefits were no longer statistically significant ($P = 0.93$). [Badger 2013a](#), which evaluated two forms of telephone-delivered health education and counselling (without usual care control), reported statistically significant reductions in fatigue ($P < 0.05$) through telephone-delivered interventions. However, the pilot study [Mosher 2016](#) did not detect any statistically significant benefit of a telephone-delivered intervention aimed at reducing a broad range of symptoms associated with lung cancer (one of which was fatigue).

1.4 Emotional distress; telephone intervention versus control

Seven studies investigated effects of interventions on emotional distress ([Allard 2007](#); [Allen 2002](#); [Bailey 2004](#); [Chambers 2014](#); [Downe-Wamboldt 2007](#); [Livingston 2010](#); [Mishel 2005](#)).

Definitions of emotional distress and tools used to measure it varied across studies. The following measurement tools were used.

- Profile of Mood States (POMS) ([Allard 2007](#); [Bailey 2004](#); [Mishel 2005](#)).
- Brief Symptom Inventory-18 (BSI-18) ([Chambers 2014](#)).
- Medical Outcomes Study 36-item Short Form (SF-36), Mental Health Index subscale ([Allen 2002](#)).
- Derogatis Psychosocial Adjustment to Illness Scale - Self-Report (PAIS-SR) ([Downe-Wamboldt 2007](#)).
- Rosebaum's Self-Control Schedule (SCS) ([Bailey 2004](#)).
- Investigator-developed scale originally developed for breast cancer patients ([Livingston 2010](#)).

Five studies including 968 participants reported quantitative results including change scores - or data enabling these and their associated standard deviations to be calculated ([Allard 2007](#); [Allen 2002](#); [Bailey 2004](#); [Downe-Wamboldt 2007](#); [Mishel 2005](#)) ([Analysis 1.4](#)). None of these studies reached significance. We did not pool the overall effect estimates due to considerable heterogeneity, but we have depicted the results of individual trials in the forest plot in [Analysis 1.4](#).

[Livingston 2010](#) evaluated effects of two telephone interventions on emotional distress but provided insufficient data to enable calculation of change scores (and standard deviations). Study authors report that although both interventions appeared to reduce cancer-specific distress, neither generated statistically significant improvements when compared with the control. [Chambers 2014](#) determined that both study arms (single-session oncology nurse-delivered telephone intervention and five-session psychologist-delivered telephone intervention) were associated

with decreased emotional distress. However, once again, lack of a 'no treatment' control group renders it impossible to conclude whether this resulted from the interventions over other factors such as passage of time.

1.5 Uncertainty; telephone intervention versus control

Three papers reported the impact of interventions on feelings of uncertainty in relation to a patient's illness (Bailey 2004; Mishel 2002; Mishel 2005). All three based their interventions on the same theoretical framework - Mishel's Uncertainty Theory - and focused on reducing uncertainty through strategies such as cognitive re-framing of threatening events and problem-solving strategies. These papers did not report change scores, or it was impossible to calculate these alongside the standard deviation of the change score, and meta-analysis was not possible.

Uncertainty was measured using the following instruments.

- Mishel's Uncertainty in Illness Scale (Mishel 2002).
- Self-Control Scale - Problem-Solving and Cognitive Re-framing subscales (Bailey 2004; Mishel 2002; Mishel 2005).
- Growth Through Uncertainty Scale (GTUS) (Bailey 2004).
- Confusion subscale of the Profile of Mood States - Short Form (POMS-SF) (Bailey 2004).

Bailey 2004 delivered a watchful waiting intervention by telephone to men with prostate cancer. Researchers detected no statistically significant differences between group overall scores for uncertainty management. However, the intervention group displayed a significant improvement on the 'New view of life' subscale compared to the usual care group ($P = 0.02$). Mishel 2002 also addressed uncertainty in men with prostate cancer; these investigators delivered two telephone-delivered interventions to men following surgery for their disease. The interventions differed only in that one group had supplementary delivery of it to a close family member. Study authors reported significant improvements in uncertainty management ($F[16,438] = 1.96$; $P = 0.01$), cognitive re-framing ($F[4456] = 3.81$; $P = 0.005$), and problem-solving ($F[4456] = 2.40$; $P = 0.049$) in both intervention groups when compared with the control group immediately following intervention delivery, but these differences were not maintained over time.

Later, Mishel 2005 evaluated an uncertainty intervention in long-term breast cancer survivors and found a statistically significant difference for cognitive re-framing (proxy for uncertainty management) ($P = 0.01$). Outcomes were more pronounced in African American women ($P = 0.03$) than in White women. Late outcomes (20 months) of the intervention were reported in the subsequent publication of Mishel 2005 by Gil 2006; these results confirmed that benefits generated by the intervention were maintained over time ($F[1479] = 3.94$; $P < 0.05$, $d = 0.06$, $n_2 = 0.008$). Further, women in the intervention group reported decreased illness uncertainty, whereas there was no change for women in the control group from baseline to 20 months (Wilk's lambda $F(1479) = 4.85$; $P < 0.03$, $d = 0.09$, $n_2 = 0.010$).

1.6 Pain; telephone intervention versus control

Six studies examined interventions to relieve cancer-related pain (Barsevick 2010; Kroenke 2010; Molassiotis 2009; Mosher 2016; Porter 2011; Thomas 2012). These trials did not report change scores, or it was impossible to calculate these alongside the

standard deviation of the change score, and meta-analysis was not possible. Researchers measured outcomes using the following tools.

- Brief Pain Inventory (BPI) (Barsevick 2010; Kroenke 2010; Mosher 2016; Porter 2011; Thomas 2012).
- Cancer Institute Common Toxicity Criteria (NCI-CTC) pain item (Molassiotis 2009).
- Bodily Pain Scale from the Medical Outcomes Study 36-item Short Form (SF-36) (Kroenke 2010).
- Barriers Questionnaire (BQ) (Thomas 2012).

Three of these studies reported statistically significant reductions in pain. Kroenke 2010 measured pain at four study points following telephone intervention (1, 3, 6, and 12 months). Across all points, the intervention group reported significantly greater improvements than the usual care group for pain severity ($P < 0.0001$), interference in functioning caused by pain ($P < 0.0001$), and bodily pain ($P = 0.004$). Thomas 2012 tested effectiveness of two interventions compared to usual care in decreasing intensity of cancer pain by lowering patients' attitudinal barriers to pain management. Trial authors found no differences between groups at the end of the study in terms of barriers to pain management, average pain intensity scores ($F = 2.58$; $P = 0.08$), pain relief ($F = 2.63$; $P = 0.07$), or overall body pain ($F = 2.817$; $P = 0.062$). However, post hoc contrasts demonstrated that the coaching group had significantly lower mean pain interference scores compared to the education and usual care groups ($F = 4.53$; $P = 0.03$ and $P = 0.02$, respectively) at the end of the study. Molassiotis compared an 18-week home care intervention for people with breast or colorectal cancer given oral chemotherapy. Measures were taken weekly, and across all, the intervention group reported significantly less pain when compared with the control group (ranging between $P < 0.0005$ and $P < 0.001$).

The remaining three studies did not generate statistically significant results (Barsevick 2010; Mosher 2016; Porter 2011).

1.7 Sexually-related symptoms; telephone intervention versus control

Three studies focused on the impact of the intervention on sexually-related symptoms including sexual function and satisfaction (Chambers 2015; Reese 2014; Reese 2018). These trials did not report change scores, or it was impossible to calculate these alongside the standard deviation of the change score, and meta-analysis was not possible.

These researchers measured outcomes using the following.

- International Index of Erectile Function (Chambers 2015; Reese 2014; Reese 2018).
- Female Sexual Function Index (Reese 2014; Reese 2018).
- Female Sexual Distress Scale (Reese 2018).
- PROMIS SexFS v2. Global Sexual Satisfaction Scale (Reese 2018).
- Sexual needs subscale of the Supportive Care Need Survey (Chambers 2015).
- Psychological Impact of Erectile Dysfunction - Sexual Experience (Chambers 2015).
- Index of Sexual Satisfaction (Reese 2014).

Reese 2014 evaluated a telephone intervention intended to improve physical intimacy and sexual concerns for couples in whom one was diagnosed with colorectal cancer. Study authors reported a large effect size related to sexual functioning associated with the intervention for female (0.85) participants and a moderate effect size for males (0.58). However, they observed no effects of the intervention on sexual distress. They followed this study by evaluating an adapted version of the intervention in breast cancer survivors (adapted to address specific needs of breast cancer survivors and their intimate partners). Similar to their findings from research with people with colorectal cancer (Reese 2014), Reese 2018 reported medium to large positive effects on all sexual outcomes measured.

Conversely, Chambers 2015 found no evidence of differences between intervention and control groups for any of the measures above. However, these researchers did report statistically significant differences at 12 months for use of erectile dysfunction medication. Results demonstrated that participants in the intervention group were 3.14 times more likely to use medical treatment for erectile dysfunction than those in the control group ($P = 0.0008$).

1.8 Dyspnoea; telephone intervention versus control

Only two studies addressed this symptom (Mosher 2016; Porter 2011), even though people with lung cancer were the target population for a number of interventions. Once more, these investigators did not report change scores, or it was impossible to calculate these alongside the standard deviation of the change score, and meta-analysis was not possible.

These studies measured outcomes using the following.

- Four items on the Memorial Symptom Assessment Scale (MSAS) related to breathlessness (Mosher 2016).
- Functional Assessment of Cancer Therapy - Lung Cancer subscale (Porter 2011).

Dyspnoea was solely a primary outcome in Mosher 2016. However, this trial did not report any statistically significant benefit regarding breathlessness arising from a generic telephone-delivered symptom management intervention - involving carers - when compared with telephone education/support. Conversely, comparison between a telephone-delivered carer-assisted coping skills programme and telephone-delivered education/support conducted by Porter 2011 did generate statistically significant improvements ($B = 0.76$, standard error = 0.21, $P = 0.0003$) in lung cancer-specific symptoms. This subscale included shortness of breath, coughing, weight loss, and loss of appetite. Neither of these studies included a usual care control, making it impossible to posit firm conclusions.

1.9 General symptom experience; telephone intervention versus control

General symptom intensity and/or symptom distress scores were study outcomes in 10 of the studies incorporated in the review (Badger 2013a; Badger 2013b; Barsevick 2010; Kroenke 2010; Mishel 2002; Molassiotis 2009; Porter 2011; Sherwood 2005; Sikorskii 2007; Traeger 2015). The following tools were used to measure this.

- General Symptom Distress Scale (Badger 2013a; Badger 2013b).
- Side Effect Checklist (SCL) (Barsevick 2010).

- Symptom Distress Scale (Mishel 2002).
- National Cancer Institute Common Toxicity Criteria (NCI-CTC) (Molassiotis 2009).
- Functional Assessment of Cancer Therapy - Lung Cancer (FACT-L) lung cancer-specific subscale (Porter 2011).
- Memorial Symptom Assessment Scale - Short Form (MSAS-SF) (Traeger 2015).
- Various investigator-developed symptom severity scales (Kroenke 2010; Sherwood 2005; Sikorskii 2007).

Again, these studies did not report change scores, or it was impossible to calculate these alongside the standard deviation of the change score, and meta-analysis was not possible. Of the 10 studies, only two reported positive effects of telephone interventions on general symptom experience (Badger 2013b; Kroenke 2010). Kroenke 2010 measured study participants' physical symptom burden and calculated an overall symptom score. Study authors determined that the telephone intervention for pain and depression that was introduced generated statistically significant reductions in general symptom severity by the end of the study when compared with the control ($M -1.0$, 95% confidence interval (CI) -2.7 to -0.7 ; $P = 0.014$). The other study that reported statistically significant improvements in general symptom experience evaluated two different telephone interventions for Latina women with breast cancer (Badger 2013b). Badger 2013b determined that both interventions (telephone interpersonal counselling (TIP-C) and telephone health education (THE)) generated significant improvements in general symptom distress, with participants in the THE group reporting slightly greater improvements ($F[1.60, 55.83] = 17.13$; $P < 0.001$) than TIP-C participants ($F[2.66] = 7.63$; $P < 0.01$).

None of the eight remaining studies generated statistically significant results. However, although Molassiotis 2009 detected no significant improvement - when introducing telephone-delivered symptom-focused interventions for people with colorectal or breast cancer receiving oral chemotherapy - in a composite symptom score derived from nine symptoms (oral mucositis, hand-foot syndrome, diarrhoea, constipation, nausea, vomiting, pain, fatigue, and insomnia), they did find improvements across most individual symptoms over time. During treatment cycles 1 and 2, the intervention group reported significant improvements across all symptoms ($P < 0.0005$ to 0.005), except for hand-foot syndrome ($P = 0.08$) and vomiting ($P = 0.062$). These improvements were maintained until the sixth cycle for all symptoms except fatigue.

DISCUSSION

Summary of main results

Telephone-delivered interventions are growing in popularity within cancer care, arguably reflecting the trend for health care to be delivered in day care settings with limited capacity for delivering interventions face-to-face. This review incorporated data from 6250 people with cancer across 32 studies. Most of the interventions incorporated had been developed for - or tested ability to reduce - psychological symptoms, notably depressive symptoms ($n = 21$), anxiety ($n = 16$), and emotional distress ($n = 7$). The predominance of telephone-delivered interventions for psychological symptoms is unsurprising; telephone counselling has been shown to be effective in reducing psychological symptoms including depression and anxiety in patient populations other than those with diagnosed

cancer (Reese 2002), in people specifically with breast cancer (Chen 2018), and more recently among carers (Lins 2014). Not all studies had theoretical frameworks underpinning the intervention that was delivered and evaluated; those that did have such a framework used one from a diverse array of 11 differing frameworks.

Twenty-one of the 32 studies reported a significant effect ($P < 0.05$) in favour of the intervention condition generated by the telephone-delivered intervention (Allard 2007; Allen 2002; Badger 2007; Badger 2013a; Badger 2013b; Badr 2015; Bailey 2004; Barsevick 2004; Barsevick 2010; Chambers 2015; Dong 2018; Girgis 2009; Kroenke 2010; Mishel 2002; Mishel 2005; Molassiotis 2009; Rawl 2002; Porter 2011; Thomas 2012; Watson 2017; Yates 2005). However, meta-analyses were not possible for any of the outcomes due to considerable heterogeneity. Findings provided in the review suggest that telephone-delivered interventions may be effective in addressing some common symptoms associated with cancer, but this would need to be established in future updates of the review, when more evidence is available for each cancer type. At present, no firm conclusions can be drawn. It is encouraging, however, that findings from the review appear to be largely consistent with those from the Lins 2014 and Chen 2018 systematic reviews. The only difference is related to anxiety; neither Chen 2018 nor Lins 2014 reported positive outcomes for anxiety. Reasons for this are unclear, although populations eligible for inclusion in these reviews differed.

Further, this review has demonstrated that telephone-delivered interventions are being developed for managing a range of physical as well as psychological cancer-related symptoms. Symptoms including pain, dyspnoea, and sexually-related symptoms, in addition to fatigue, have been subject to investigation. Sexually-related symptoms are a relatively recent addition to the range of symptoms being addressed in this area of research. This mirrors research more generally into sexually-related symptoms in cancer care. They have been orphan symptoms - comparatively neglected - when compared with other cancer-related symptoms (White 2011). Further, this review identified that, with regards to telephone-delivered interventions, dyspnoea could also be viewed as an orphan symptom. This may reflect perceived challenges of dealing with a complex and often acute symptom like dyspnoea by telephone. It may be that brief psycho-educational interventions that are delivered primarily by telephone may be unsuitable for this group of cancer patients. However, further research is needed before such conclusions can be drawn.

For a number of symptoms - uncertainty, pain, dyspnoea, and sexually-related symptoms - data from the studies incorporated are inconsistent, making conclusions regarding their management through telephone-delivered interventions difficult to draw; around half failed to generate statistically significant findings.

Most interventions were delivered over a relatively short period, typically ranging from two to eight weeks; most frequently, interventions were provided weekly. However, lack of data prevented comparison of effectiveness by length or frequency of the intervention (dose). Likewise, evidence was insufficient to enable comparison by stage of disease. Some preliminary evidence from Porter 2011 suggests that different forms of interventions may be more/less effective according to stage of disease. This theory needs further investigation. Differential responses to interventions are important to determine. Although likely to be more cost-effective than interventions delivered face-to-face (cost-

effectiveness of these interventions has not been reported on), those delivered by telephone remain relatively costly in terms of health professional time. It is clearly optimal to target interventions towards people who would benefit most, to enable effective resource use.

Telephone-delivered interventions may provide greater benefit for particular people. Arguably, people with high symptom intensity could benefit the most, as they would also have high possibility for symptom reduction. However, as few studies specifically targeted people with moderate or greater symptoms, it is not possible to draw conclusions about the relative efficacy of interventions according to symptom burden. Further, the telephone can provide a degree of anonymity that may be helpful with certain symptoms (e.g. sexually-related ones), and no requirement for face-to-face contact is likely to be of particular importance to people whose immunity is compromised through disease and/or treatment. However, data are insufficient to enable these hypotheses to be confirmed or refuted. It would appear that telephone-delivered interventions may also generate practical benefit, for example, being particularly pertinent for people living at a distance from treatment centres, or for those with fatigue for whom the effort required to attend delivery of an intervention in person may be prohibitive.

Many people with cancer report feelings of fatigue and the negative effect this can have on their motivation for symptom self-management (Ahlberg 2003; Ream 2015). Further, health professionals may find it more difficult to motivate patients in their patient group to adopt behaviour change when delivering interventions remotely (e.g. by telephone) rather than in person. This would suggest that techniques such as motivational interviewing (MI) could offer important benefit in telephone-delivered interventions for symptom self-management - in particular, when patients may be experiencing fatigue. MI was used to generate behaviour change in two studies within the review (Thomas 2012; Ream 2015). Two further trial authors referred to techniques within their interventions aimed at enhancing motivation to change behaviour (Allen 2002; Yates 2005). Incorporation of goal-setting and monitoring used by MI, as well as interventions based around it, provides impetus for behaviour change.

All interventions were delivered by health and social care professionals, most by nurses. Most papers ($n = 22$) referred to training needed by those delivering interventions - notably in counselling skills, communication, and education. This is not attained without cost. Unfortunately, the papers in this review did not feature any form of cost evaluation; thus it is impossible to determine the cost implications of interventions such as these. This is an important topic that needs consideration in future research.

Overall completeness and applicability of evidence

This is the first review to appraise and synthesise findings from studies evaluating telephone-delivered interventions for management of cancer-related symptoms. This review indicates that such interventions may be effective in ameliorating cancer-related symptoms - most notably, depressive symptoms, fatigue, and emotional distress. Some symptoms (uncertainty, pain, dyspnoea, sexually-related symptoms) have been subject to relatively less research; robust conclusions cannot be drawn from available evidence.

This review does have some limitations. First, it is limited by the narrative nature of the synthesis and the relatively small number of studies addressing certain symptoms, such as sexually-related symptoms and dyspnoea, which rendered study conclusions difficult to determine. Further, it was impossible to determine whether better outcomes were achieved if the telephone was used alongside other elements (e.g. face-to-face contact, printed/digital/online materials). Although some studies that were incorporated into the review reported findings that failed to attain statistical significance, the review is likely to have been subject to a degree of publication bias. Finally, studies incorporated into the review were largely conducted in the United States, Australia, and the United Kingdom - thus their findings may not translate well to other continents and nations. The findings presented need to be viewed in light of these limitations.

Further, symptoms are multi-dimensional. This review focused on severity of symptoms; it did not address the distress generated by symptoms. Several studies in this review measured and reported on symptom distress, in addition to intensity. A symptom may be of low intensity but still may give rise to considerable distress. This may be important clinically - arguably symptoms should be prioritised according to how distressing they are. The authors of this review are unable to draw conclusions about the effectiveness of telephone interventions with regards to symptom distress.

This review did incorporate psychological symptoms including uncertainty. However, it did not include fear of recurrence, as this is a concern associated with cancer that can give rise to symptoms like anxiety and uncertainty (rather than being a symptom per se). A recent study has evaluated an intervention for fear of recurrence delivered by telephone (Dieng 2016). We did not include this paper in the review.

Quality of the evidence

We used the GRADE approach to interpret findings based on the certainty of evidence generated across the included studies. Most of the 32 trials that met our inclusion criteria were at moderate risk of bias. All but one were randomised controlled trials, so the evidence generated by them should be interpreted as highest certainty available, despite blinding not being achieved across almost all of these trials; lack of blinding can give rise to performance or detection bias. However, it is acknowledged that in research of this nature, blinding of participants would be possible only if a form of sham intervention was delivered to those in the control group. This in itself introduces difficulties, as some of the effects of a telephone-delivered intervention are likely to reflect characteristics of the telephone conversation (that would be experienced by those in the sham intervention control). This could render effects of the telephone intervention difficult to detect. Alternatively, an intervention could be tested against an active control for which the control group receives an intervention of known positive effect.

Review authors noted heterogeneity in results generated by the interventions studied (i.e. inconsistency within the body of evidence). This is likely to reflect differences between the interventions themselves. Although all were predominantly delivered by telephone, they varied in length, in content, and in some cases in incorporation of additional elements such as face-to-face contact or additional supportive media. Further, the cancer populations studied varied, and this appeared to introduce

additional levels of heterogeneity. Therefore, meta-analyses were not performed.

Overall, the certainty of evidence was very low for all outcomes in the review, as we are very uncertain about the estimates. We downgraded all outcomes due to concerns about overall risk of bias profiles being unclear or high, imprecision, inconsistency in results, and general heterogeneity.

Potential biases in the review process

We performed a comprehensive search, including a thorough search of the grey literature, and two review authors independently sifted all studies and extracted study data. We restricted the review to randomised controlled trials (RCT) and one quasi-RCT. By searching a wide range of databases and grey literature, we attempted to ensure that we did not overlook any relevant evidence.

A threat to the validity of the review is likely to be publication bias; studies that did not find the treatment to be effective may not have been published. We did not report any meta-analyses due to considerable clinical and methodological heterogeneity. Meta-analysis may have led to misleading results that lacked any kind of generalisability.

Agreements and disagreements with other studies or reviews

The results of this review are largely consistent with those published previously by Chen 2018. That review evaluated tele-health interventions for quality of life and psychological outcomes in people with breast cancer. Those review authors similarly reported that compared with usual care, telephone-delivered interventions led to statistically significantly less depressive symptoms, distress, and perceived stress. Consistency across reviews such as these does provide some confidence in the veracity of findings. However, unlike the present review, Chen 2018 reported that anxiety did not differ between tele-health and usual care control. As with this review, the findings of Chen 2018 have to be considered with a degree of caution owing to between-study heterogeneity.

AUTHORS' CONCLUSIONS

Implications for practice

This review found some evidence supporting use of telephone-delivered interventions for managing cancer-related symptoms - most evidence is related to managing anxiety, depressive symptoms, emotional distress, or fatigue. This would suggest that telephone interventions should be considered as one component for managing these cancer-related symptoms. Arguably, these interventions would not need to be provided within statutory services; they could be provided by voluntary sector providers, who frequently provide patients support and information via health professional-delivered telephone help lines. We found limited evidence regarding potential management of some symptoms by telephone-delivered intervention, notably regarding sexually-related symptoms and pain.

Interventions evaluated in this review varied considerably in terms of (1) the number of calls provided; (2) the length and timing of calls; (3) the content of calls; and (4) provision or not of

additional supportive material. It appears that for some symptoms (i.e. depressive symptoms), telephone-only interventions may be indicated. However, given the small, mostly biased studies - and lack of meta-analysis - this conclusion is tentative. For other symptoms, arguably those for which greater behaviour change may be required (e.g. fatigue), it may be the case that additional supportive materials are required to optimise outcomes. It has not been possible however through this review to conclude how many calls, delivered over what period, of particular duration and content, are required to generate effect. It appears that it may be beneficial for those delivering interventions to be trained in motivational interviewing skills.

It is noteworthy that most studies included within the review did not target samples with high baseline symptom burden. Arguably, it would be wise to do so, as there could be greater possibility for symptom reduction in such patient groups and greater associated cost savings.

Healthcare services are facing an unprecedented level of austerity that is likely to persist given ageing communities and finite health expenditure. There is a requirement for care to shift closer to home, and for cost containment to be applied. Telephone-delivered interventions could make an important contribution to both requirements. Further research is needed to determine clearly which symptoms are most amenable to be managed through telephone-delivered interventions, which patients should be targeted to achieve clinically important and sustained benefit, and which interventions are most cost-effective.

There is risk of bias in the studies reported on in this review (most studies were at risk of some bias). When considered alongside other factors including between-study heterogeneity and imprecise estimates of effect, the overall certainty of evidence is best considered very low, as we are very uncertain about the estimates. A lot of the heterogeneity was due to different cancer sites. Thus, the review findings need to be interpreted with a degree of caution. Consistency between this review and that of [Chen 2018](#) is however encouraging and can lend support to the conclusions drawn.

Implications for research

Further work is necessary to determine:

- which cancer-related symptoms are amenable to management by telephone-delivered intervention; particular attention needs to be directed towards symptoms that have been subject to little research (e.g. sexually-related symptoms, dyspnoea);
- what augmentation of telephone-delivered interventions (e.g. with face-to-face meetings or additional resources) works, and in what particular circumstances;

- cost implications of delivering interventions by telephone to manage cancer-related symptoms;
- consensus on the most appropriate measures to be used to measure outcomes and minimal clinically important differences for each one;
- consensus on standardising reporting of the intervention theory underpinning the intervention and its components;
- longevity of effects generated by telephone-delivered interventions for cancer-related symptoms;
- effectiveness of telephone-delivered interventions in reducing fear of recurrence (a phenomenon that is increasingly being written about in the literature but is not deemed a symptom and thus was excluded from this review); and
- effectiveness of telephone-delivered interventions for enhancing quality of life for people with cancer.

Future research is needed to address methodological limitations identified in the included studies. Notably, future randomised controlled trials need to have better methodological conduct and design to minimise risk of bias and to provide more extensive reporting of pertinent outcomes. Further, they need sufficient statistical power and length of follow-up to generate much needed definitive evidence concerning efficacy of telephone-delivered interventions for management of cancer symptoms across diverse patient groups.

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* Indicates the major publication for the study

CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Allard 2007

Study characteristics

Methods	<p><u>Setting</u>: 5 regional centres in different geographic areas in Quebec, Canada</p> <p><u>Recruitment</u>: eligible women were over the age of 18, were French speaking, and were diagnosed with primary breast cancer or a suspected lesion. They were scheduled to undergo day surgery and had no hearing impairment and no access to a phone at home. Staff nurses identified eligible patients who were given a pamphlet describing the study and subsequently, if willing to participate, were telephoned by a member of the research team</p> <p><u>Randomisation</u>: randomised clinical block trial</p>
Participants	117 women with breast cancer aged 18 years or older
Interventions	<p><u>Intervention</u>: based on self-regulation theory, which emphasises concrete objective information and regulation of emotional and functional processes to achieve optimal coping. Comprising 2 telephone sessions delivered weekly for 2 weeks during the immediate postoperative period. In each session, par-</p>

Allard 2007 (Continued)

Participants were asked to identify and describe symptoms in concrete terms. Actions taken by women to manage each symptom were rated according to their effectiveness from 1 to 5 (1 = not effective, 5 = very effective). Effective actions were encouraged; ineffective actions were explored and women encouraged to find other, potentially helpful, actions. The interventionist suggested new or additional self-care strategies and acknowledged feelings expressed by women during the call

Control group: usual care consisting of perioperative teaching and a follow-up phone call within 24 hours of discharge from the hospital

Interventionist: the intervention was delivered by the principal investigator, who is a trained nurse

Outcomes	<ul style="list-style-type: none"> Psychological symptoms - emotional distress Impact on quality of life/functioning - functional status <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> Profile of Mood States Short Form (POMS-37) (α0.75-0.94) Symptom Impact Profile (SIP); subscales 'recreation and pastimes' and 'home management'
Notes	Some participants may have already started adjuvant hormonal therapies when recruited for this study

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	The research assistant randomly allocated the women with or without axillary node dissection to the experimental or control group at each site using a table of random numbers
Allocation concealment (selection bias)	Unclear risk	The method of concealment is not described; however, the research assistant allocated participants using a table of random numbers, and the research assistant did not perform the intervention. Detection bias may exist, as it is unclear if the intervener or the research assistant collected the data
Blinding (performance bias and detection bias) All outcomes	High risk	Owing to the nature of the intervention, study participants were not blinded. It is unclear who collected the data and by what method
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Attrition rate is reported. Reasons for attrition are not reported
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	Unclear risk	The process of screening was reliant on staff nurses identifying participants at the doctor's office. It is unclear if all eligible participants were approached; some bias may exist, as these staff nurses may have excluded or not approached participants for many reasons

Allen 2002
Study characteristics

Methods	<u>Setting</u> : a network of sites (hospitals and private oncology clinics) in Rhode Island, Massachusetts, Pennsylvania, New Jersey, USA
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Allen 2002 (Continued)

Recruitment: eligible women were aged 50 or younger, had no prior history of breast carcinoma, had stage I-IIIa tumours, and were beginning their first course of chemotherapy treatment. Medical staff provided information about the study to eligible women. Women interested in participating were sent a letter describing the study, followed by a telephone call several days later to answer questions and invite participation. Women who agreed to participate were asked to complete a baseline telephone interview and were mailed questionnaires. They were also asked to nominate a primary support person (PSP) to participate as a partner in the intervention. Nominated partners were then contacted by research staff to invite their participation and to complete similar surveys

Randomisation: RCT. Women were stratified by recruitment site location and PSP status (spouse/significant other, other family member or friend, no participating PSP)

Participants	164 women with breast cancer younger than 50 years
Interventions	<p>Intervention: based on motivational technique, using an investigator-developed “home care training model”. Nurse delivered to patient and other supportive person if nominated. The intervention consisted of 5 interactive components: problem orientation, problem definition, generation of alternatives, decision-making, and solutions implementation. Patients were given a printed manual with chapters that paralleled the intervention components. The intervention was conducted as 1 × 2-hour individual face-to-face training session held in the participant's home, followed by 4 telephone contacts 2 weeks apart, and a final 2-hour individual face-to-face training session held in the participant's home, where the nurse took a less active role while letting the participant and the PSP take over. In each session, worksheets were completed and interventionist feedback was provided. Participants could instigate telephone contact to discuss additional questions</p> <p>Control group: usual care (not described)</p> <p>Interventionist: oncology research nurse</p>
Outcomes	<ul style="list-style-type: none"> • Psychological symptoms - emotional distress • Impact on quality of life/functioning - quality of life, rehabilitation needs, unmet needs for assistance <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> • Medical Outcomes Study 36-item Short Form General Health Survey (SF-36) - MHI-5 subscale • Cancer Rehabilitation Evaluation System (CARES) • Investigator-generated list of met/unmet needs for personal care, meal preparation, housekeeping, shopping, transportation, and child care in the past month • Impacts of Events Scale (IES) • Social Problem-Solving Inventory - Revised (SPSI-R) ($\alpha = 0.89$ to 0.93)

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Generation of random sequence is not described
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	Owing to the nature of the intervention, study participants were not blinded. It is unclear who collected the data and by what method
Incomplete outcome data (attrition bias)	Unclear risk	Attrition rate is reported. Reasons for attrition are not reported

Allen 2002 (Continued)

All outcomes

Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	High risk	Due to unavoidable delays in completing all 6 sessions for some participants receiving the intervention, women in the experimental group completed both of their follow-up assessments approximately 1 month later than those in the control arm

Badger 2005
Study characteristics

Methods	<p><u>Setting</u>: academic cancer centres and urban private oncology offices in Arizona, USA</p> <p><u>Recruitment</u>: eligible women had a diagnosis of stage I-III breast cancer, were receiving adjuvant treatment, were able to speak English and talk on the telephone, were married, and had a partner, who were also willing to take part in the intervention and were employed at the time of the study. Participants were recruited from a local cancer centre, oncologists' offices, and support groups, and through self-referral after reading brochures displayed in the various settings</p> <p><u>Randomisation</u>: pilot quasi-randomised controlled trial (experimental with repeated measures)</p>	
Participants	48 women in their mid-50s with breast cancer	
Interventions	<p><u>Intervention</u>: telephone interpersonal counselling (TIP-C). Based on theories of interpersonal therapy and cancer education. Participants received 6 weekly telephone calls × 30 minutes (approximately) while they were undergoing treatment for breast cancer. Sessions focused on cancer education, interpersonal role disputes, social support, awareness, management of depressive symptoms, and role transitions. Partners of participants received 3 telephone-delivered TIP-C sessions (weeks 1, 3, and 5) during the same 6-week period as the women. These sessions also focused on issues such as cancer education, role disputes, role transitions, and social support</p> <p><u>Control group</u>: participants in the usual care arm received a resource list about cancer and brief, focused telephone calls (6 for women and 3 for their partners × 5 to 10 minutes) to inquire about general well-being and to answer general questions (no counselling)</p> <p><u>Interventionist</u>: nurse counsellors (master's prepared clinical nurse specialists in psychiatric/mental health nursing who had additional oncology training)</p>	
Outcomes	<ul style="list-style-type: none"> Psychological symptoms - depression Fatigue Impact on quality of life/functioning - stress, positive and negative effects <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> Center for Epidemiologic Studies Depression Scale (CES-D) Multi-dimensional Fatigue Inventory (MFI) 	
Notes		
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	High risk	Pilot study, quasi-randomised sample

Badger 2005 (Continued)

Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	It is unclear if study participants were blinded. Interveners were not blinded. It is unclear who collected the data and by what method
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Attrition is not reported
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Badger 2007
Study characteristics

Methods	<p><u>Setting</u>: Arizona, USA (not specified)</p> <p><u>Recruitment</u>: eligible women had a diagnosis of stage I-III breast cancer, were receiving adjuvant treatment, were able to speak English and talk on the telephone, had no physical or psychological disabilities, and had a partner who was available and willing to take part in the intervention. Participants were recruited from a local cancer centre, oncologists' offices, and support groups, and through self-referral after reading brochures displayed in the various settings</p> <p><u>Randomisation</u>: 3-arm RCT</p>
Participants	96 women with breast cancer and their supportive partners
Interventions	<p><u>Intervention</u>: participants and their partners were randomised to receive 1 of 3 programmes: (1) telephone interpersonal counselling (TIP-C); (2) self-managed exercise (SE); or (3) attention control (AC). In the TIP-C arm, participants received 6 × weekly calls, which consisted of cancer education, social support, and awareness and management of depressive and anxiety symptoms. Partners received 3 × calls (every other week of the intervention) to discuss their well-being and relationship. Participants in the SE arm also received 6 × weekly calls. The calls encouraged participants to exercise and recorded information about progress and intensity of exercise. The SE partners also received calls every other week to encourage exercise and to track progress</p> <p><u>Control group</u>: the AC group received printed information about breast cancer and 6 brief weekly calls. Their partners received 3 biweekly calls during the same period. The calls averaged 7 minutes in duration. If these participants reported any problems, they were directed to the primary physician. No counselling or encouragement to exercise was offered</p> <p><u>Interventionist</u>: psychiatric nurse counsellors</p>
Outcomes	<ul style="list-style-type: none"> Psychological symptoms - depression, anxiety <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> Center for Epidemiologic Studies Depression Scale (CES-D) ($\alpha \geq 0.85$) Investigator-designed survey devised from parts of the Positive and Negative Affect Schedule (PANAS), 1 item from the Short Form-12 (SF-12) Scale, and 3 items from the Index of Clinical Stress Scale (ICS) (in this study, $\alpha = 0.91, 0.89, 0.88$ for breast cancer participants; and $0.89, 0.88, 0.91$ for partners at T1, T2, and T3, respectively)

Badger 2007 (Continued)

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The generation of random sequence is not described
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	It is not clear whether participants were blinded to group. Interveners were not blinded. It is unclear who collected the data and by what method
Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Badger 2013a
Study characteristics

Methods	<p><u>Setting</u>: Arizona, USA (not specified)</p> <p><u>Recruitment</u>: eligible women had a diagnosis of breast cancer, were currently receiving treatment, were at least 21 years of age, had access to a telephone, and had a supportive partner (SP) who was available and willing to take part in the intervention. SPs were not restricted to spouses. Participants were recruited from a local cancer centre, oncologists' offices, and support groups, and through self-referral</p>
Participants	52 women with breast cancer and their supportive partners
Interventions	<p><u>Interventions</u>: women and their supportive partners were randomly assigned to 1 of 3 interventions: (1) telephone health education (THE), (2) telephone interpersonal counselling (TIC), or (3) videophone interpersonal counselling (VIC). All participants received 8 weekly telephone calls of 30 minutes in their preferred language (English or Spanish), and their SP received 4 biweekly sessions. Participants in THE received leaflets on breast cancer terminology, treatments, side effect management, nutrition, physical activity, and resources. These materials were reviewed with participants and SPs over the telephone. Participants in the TIC and VIC arms received counselling via video or telephone, which addressed mood and affect management, emotional expression, interpersonal communication, relationships with family and providers, social support, and referral to support service as required</p> <p><u>Interventionist</u>: an information specialist provided THE; TIC and VIC were provided by a social worker</p>
Outcomes	<ul style="list-style-type: none"> • Depression • Symptom distress • Social well-being • Spiritual well-being

Badger 2013a (Continued)

Methods for assessing outcomes:

- Center for Epidemiologic Studies Depression Scale (CES-D)
- General Symptom Distress Scale (GSDS)
- Social Well-Being Scale
- Spiritual Well-Being Scale

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The generation of random sequence is not described
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	It is unclear if participants were blinded to group. Interveners were not blinded
Incomplete outcome data (attrition bias) All outcomes	High risk	Over 20% loss to follow-up; significantly higher attrition in the THE group compared to the TIC or VIC group
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Badger 2013b
Study characteristics

Methods	<p><u>Setting</u>: Arizona, USA (not specified)</p> <p><u>Recruitment</u>: eligible women had a diagnosis of stage I-III breast cancer, were receiving adjuvant treatment, spoke Spanish or English, had no physical or psychological disabilities, had access to a telephone, and had a supportive partner (SP) who was available and willing to take part in the intervention. Eligible SPs were 21 years of age or older, spoke Spanish or English, had no physical or psychological disabilities, and had access to a telephone. Participants were recruited from a local cancer centre, oncologists' offices, and support groups, and through self-referral</p> <p><u>Randomisation</u>: RCT (3-wave repeated measure with a between-subjects factor)</p>
Participants	90 Latina women with breast cancer and their supportive partners
Interventions	<p><u>Intervention</u>: women and their supportive partners were randomly assigned to 1 of 2 interventions: (1) telephone interpersonal counselling (TIP-C), or (2) telephone health education (THE). The TIP-C intervention addressed mood and affect management, emotional expression, interpersonal communication and relationships, social support, and cancer information. Participants received 8 weekly phone calls × 30 minutes (approximately) in their preferred language (Spanish or English), and their SPs received 4 sessions every other week. Interventions were tailored to the cultural values and beliefs of participants</p>

Badger 2013b (Continued)

Control group: participants in the THE group also received 8 weekly phone calls in their preferred language (Spanish or English), and their SPs received 4 biweekly sessions. The THE focused on normal breast health and breast cancer, routine tests for diagnosis and prevention and associated terminology, treatment, side effects of treatment and strategies to combat these side effects, lifestyle interventions such as nutrition and physical activity, and referrals and resources

Interventionist: bilingual, bicultural master's-prepared social workers

Outcomes	<ul style="list-style-type: none"> • Psychological symptoms - depression, anxiety • Fatigue • Physical symptoms - symptom distress • Impact on quality of life/functioning - social well-being, spiritual well-being, psychosocial resources of cancer knowledge, and social support <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> • Center for Epidemiologic Studies Depression Scale (CES-D) ($\alpha \geq 0.86$ for participants and SPs at all 3 time points) • Multi-dimensional Fatigue Inventory ($\alpha \geq 0.82$ for participants and SPs) • General Symptom Distress Scale ($\alpha \geq 0.66$ for participants and SPs) • State version of the State-Trait Anxiety Inventory ($\alpha \geq 0.86$ for participants and SPs) • Quality of Life Breast Cancer Instrument; 8-item social well-being subscale ($\alpha \geq 0.65$ for participants and SPs) and 8-item spiritual well-being subscale ($\alpha \geq 0.55$ for participants and $\alpha \geq 0.77$ for SPs) • Investigator-designed cancer knowledge instrument • Perceived social support - family ($\alpha \geq 0.89$ for participants and SPs)
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Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The generation of random sequence is not described
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	It is not clear whether participants were blinded to group. Interveners were not blinded. It is unclear who collected the data and by what method
Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	Unclear risk	Participants received \$20 for completing assessments

Badr 2015
Study characteristics

Methods Setting: Mount Sinai Hospital, New York, USA

Telephone interventions for symptom management in adults with cancer (Review)

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Badr 2015 (Continued)

Recruitment: eligible participants had advanced lung cancer (LC), were within 1 month of treatment initiation, were spending more than 50% of their time out of bed each day, and had a partner (or other close family member) whom they identified as their primary carer. Participants and carers had to be at least 18 years old and able to read and understand English, and had to provide informed consent

Randomisation: 2-arm RCT

Participants	39 lung cancer participants and their partners
Interventions	<p>Participants and their partners were randomised to receive usual medical care or a psychosocial intervention</p> <p>Psychosocial intervention: participants and carers in the intervention group received a manual, covering the topics of self-care, stress and coping, symptom management, effective communication, problem-solving, and maintaining and enhancing relationships. In addition, they participated in 6 × weekly 60-minute telephone counselling sessions</p> <p>Usual medical care: primary palliative care was provided by the participant's medical oncologist, including basic management of pain and other symptoms, including depression and anxiety, with referral to outpatient supportive oncology practice if required</p> <p>Interventionist: trained interventionist with a master's degree in mental health counselling</p>
Outcomes	<ul style="list-style-type: none"> • Psychological symptoms - depression, anxiety • Caregiver burden • Autonomy – participant and carer • Competence – participant and carer • Relatedness – participant and carer <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> • 6-item Patient-Reported Outcomes Measurement Information System (PROMIS) short form depression measure • 6-item PROMIS short form anxiety measure • 12-item short form of the Zarit Burden Interview • Pierce measure of carers' autonomous motivation for tending to participant needs • Treatment Self-Regulation Questionnaire to assess participant autonomy for engaging in self-care • Measure of competence based on the work of Lorig • 4-item relatedness measure

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The generation of random sequence is not described
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	Owing to the nature of the intervention, study participants and interveners were not blinded. It is unclear who collected the data and by what method
Incomplete outcome data (attrition bias)	Low risk	Loss to follow-up < 20%; similar reasons between groups

Badr 2015 (Continued)

All outcomes

Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	Unclear risk	Participants received \$20 gift cards on completing surveys

Bailey 2004
Study characteristics

Methods	<p><u>Setting</u>: North Carolina, USA (not specified).</p> <p><u>Recruitment</u>: eligible men were diagnosed with stage B1, B2, or C1 prostate cancer and had been in watchful waiting from 1 to 124 months. Participants were recruited from the urology practices of 3 physicians at 1 hospital. The sample was a convenience sample, which was then randomised to intervention and control groups</p> <p><u>Randomisation</u>: pilot RCT</p>	
Participants	41 men with prostate cancer	
Interventions	<p><u>Intervention</u>: based on Mishel's Reconceptualised Uncertainty in Illness Theory. Participants in the intervention group received telephone calls from the interventionist weekly × 5 weeks. During each call, participants' problems were identified and the nature of their uncertainty was assessed, then the intervention was delivered. The intervention consisted of 4 elements: (1) re-framing uncertainty through encouraging probabilistic thinking, discouraging negative perspectives and providing information, (2) helping incorporate uncertainty into life structure through emphasising empowering activities, promoting self-care, and encouraging assertive communication with healthcare providers about symptom concerns, (3) supporting participants' belief in future treatment, and (4) encouraging self-monitoring and vigilance</p> <p><u>Control group</u>: usual care (not described in detail). Had access to naturally occurring sources of support</p> <p><u>Interventionist</u>: male nurse</p>	
Outcomes	<ul style="list-style-type: none"> • Psychological symptoms - expression, anxiety, anger, vigour, confusion • Uncertainty - new view of life, acceptance of the situation, continual uncertainty, negative consequences • Fatigue • Impact on quality of life/functioning - quality of life <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> • Growth Through Uncertainty Scale (GTUS) ($\alpha = 0.94$ in this study) • Profile of Mood States - Short Form (POMS-SF) ($\alpha = 0.94$ in this study) • Quality of Life Cantril's Ladder • Rosebaum's Self-Control Schedule (SCS) ($\alpha = 0.81$ in this study) 	
Notes		
Risk of bias		
Bias	Authors' judgement	Support for judgement

Bailey 2004 (Continued)

Random sequence generation (selection bias)	Low risk	A table of random numbers was used to assign men to either an experimental group that received the intervention or a control group that received usual care
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	Nurse data collectors did not deliver the intervention, but owing to the nature of the intervention, study participants were not blinded
Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Barsevick 2004
Study characteristics

Methods	<p><u>Setting</u>: a university health science centre in Utah and a comprehensive cancer centre in Philadelphia, USA</p> <p><u>Recruitment</u>: eligible participants spoke English; were beginning curative or local control treatment for breast, lung, colorectal, advanced prostate, gynaecological, or testicular cancer; and planned to receive at least 3 cycles of chemotherapy, 6 weeks of radiotherapy, or concurrent chemotherapy and radiotherapy. Prior treatment other than surgery had to be completed at least 1 month previously. Participants were excluded if they were having stem cell transplantation, interleukins, interferons, or tumour necrosis factor; had chronic fatigue syndrome; were enrolled in another study involving a psycho-educational intervention; had overt evidence of psychiatric disorder; or had initiated treatment for anaemia or depression in the past 3 weeks</p> <p><u>Randomisation</u>: RCT</p>
Participants	396 men and women with various cancer diagnoses
Interventions	<p><u>Intervention</u>: the Telephone Energy Conservation and Activity Management (ECAM) intervention was based on the Common Sense Model, which guided participants through 3 stages of information-processing to help them manage cancer-related fatigue. The intervention consisted of 3 weekly telephone sessions commencing at the beginning of chemotherapy treatment or at week 3 to 5 during radiation therapy. Session 1 consisted of coping skills training. Participants kept a journal and prioritised their usual activities. During session 2, participants were assisted to develop a care plan to minimise the interference of fatigue. In session 3, participants appraised and revised the plan. Sessions 1 and 2 each lasted 30 minutes, and session 3 lasted 15 minutes</p> <p><u>Control group</u>: attention control consisting of information about nutrition and a healthy diet. Participants kept a dietary record for 24 hours</p> <p><u>Interventionist</u>: research nurse</p>
Outcomes	<ul style="list-style-type: none"> Psychological symptoms - psychosocial well-being Fatigue

Barsevick 2004 (Continued)

- Impact on quality of life/functioning - physical functioning

Methods for assessing outcomes:

- Short Form of the Profile of Mood States Fatigue Scale (POMS-SF) ($\alpha = 0.89$ for this sample)
- Schwartz Cancer Scale (SCFS) - Physical Fatigue subscale ($\alpha = 0.97$)
- General Fatigue Scale (GFS) ($\alpha = 0.95$)
- Functional Performance Inventory (FPI) ($\alpha = 0.91$ for this sample)

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The generation of random sequence is not described
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	It is unclear whether participants were blinded to group. Interveners were not blinded
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Loss to follow-up and between-group differences regarding reasons for it are unclear
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Barsevick 2010
Study characteristics

Methods	<p>Setting: 2 university health centres, a community cancer centre, and a comprehensive cancer centre in Philadelphia, USA</p> <p>Recruitment: eligible patients were 18 years of age or older and were beginning a new chemotherapy regimen with at least 2 drugs administered intravenously in a cyclical manner for breast, lung, colorectal, prostate, gynaecological, bladder, or testicular cancer or lymphoma. Prior treatment other than surgery had to be completed at least 1 month previously, and the individual could receive concurrent radiotherapy. Participants had to be able to read and write English. Individuals were excluded if they were having marrow or stem cell transplantation, interleukins, interferons, or tumour necrosis factor; had chronic fatigue syndrome; were being treated for diagnosed sleep disorder; were enrolled in another study involving a psycho-educational intervention; had a communication impairment; had overt evidence of psychiatric disorder; or had initiated treatment for anaemia or depression in the past 3 weeks. Potential participants were approached by telephone or in the clinic, and the study was explained</p> <p>Randomisation: RCT. Participants were stratified by diagnosis (breast cancer vs non-breast cancer) at each site and then were randomly assigned to intervention or control group</p>
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Barsevick 2010 (Continued)

Participants	292 men and women aged 18 years or older with various cancers	
Interventions	<p><u>Intervention:</u> The EASE intervention is based on the Common Sense Model and includes providing information about fatigue and sleep disturbance, coping skills training, and appraisal of used coping strategies. Participants received 3 telephone sessions during the second, third, and fourth weeks after the first chemotherapy treatment. Participants also received a handbook with information about symptoms and examples of energy conservation and sleep management strategies. Between sessions 1 and 2, participants completed a daily diary of their symptoms and sleep patterns, as well as a priority list of usual activities</p> <p><u>Control group:</u> attention control consisting of information about nutrition and a healthy diet. Participants kept a dietary record for 24 hours in preparation for session 2. The 3 control sessions were equivalent to the intervention session in terms of the amount of time spent with the individual</p> <p><u>Interventionist:</u> research nurse</p>	
Outcomes	<ul style="list-style-type: none"> • Psychological symptoms - depression • Fatigue • Physical symptoms - pain, sleep disturbance • Impact on quality of life/functioning - functional status <p>Method of assessing outcome measures:</p> <ul style="list-style-type: none"> • General Fatigue Scale (GFS) ($\alpha = 0.92$ for this sample) • Profile of Mood States (POMS) - fatigue subscale ($\alpha = 0.94$ for this sample) • Pittsburgh Sleep Quality Index (PSQI) ($\alpha = 0.75$ for this sample) • Octagonal Basic Motion Logger Actigraph (measures sleep-wake and activity-rest patterns) • Morin Sleep Diary ($\alpha = 0.83$ to 0.99 for this sample) • Brief Pain Inventory (BPI) ($\alpha = 0.92$ for this sample) • Profile of Mood States (POMS) - depression subscale ($\alpha = 0.90$ for this sample) • Side Effect Checklist (SCL) ($\alpha = 0.87$ for this sample) • Brief Pain Inventory (BPI) - adapted to apply to symptoms rather than to pain only • SF-12 • Eastern Cooperative Oncology Group Performance Status 	
Notes		
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random assignments were generated by the statistician
Allocation concealment (selection bias)	Low risk	Random assignments were generated by the statistician and placed in sealed envelopes that were numbered and selected sequentially for each stratification group
Blinding (performance bias and detection bias) All outcomes	High risk	It is unclear whether participants were blinded to group. Interveners were not blinded. It is unclear who collected the data and by what method
Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups

Barsevick 2010 (Continued)

Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Chambers 2014
Study characteristics

Methods	<p><u>Setting</u>: community-based; 2 large and well-established state-based Cancer Helplines in Australia</p> <p><u>Recruitment</u>: eligible participants were adult patients and carers who called cancer information and support Cancer Helplines in 2 Australian states (Queensland and New South Wales). Patients and carers were independent callers and, therefore, were not a dyad. Study inclusion criteria included (a) having a score of 4 or greater on the Distress Thermometer, (b) being able to read and speak English, and (c) having no previous history of head injury and/or dementia. Individuals under current psychiatric care and those who presented with grief or bereavement were excluded. The Cancer Helpline operator offered callers who met selection criteria entry into the study at the time of the call</p> <p><u>Randomisation</u>: randomised trial. Participants were randomised following completion of baseline measures. Randomisation was stratified by participant and carer and state</p>	
Participants	354 participants with various cancer diagnoses and 336 carers	
Interventions	<p><u>Intervention</u>: participants and carers randomised between (a) single nurse-delivered telephone self-management session and (b) 5 telephone CBT sessions delivered by a psychologist</p> <p>Both groups were provided a self-management resource kit comprising:</p> <ul style="list-style-type: none"> • printed self-help advice about stress management; problem-solving; healthy lifestyle; strategies for mobilising personal and community support networks; and • digital media about relaxation exercises 	
Outcomes	<ul style="list-style-type: none"> • Psychological distress • Cancer-specific distress • Perceived positive life changes <p>Method of assessing outcome measures:</p> <ul style="list-style-type: none"> • Brief Symptom Inventory-18 (BSI-18) • Impact of Events Scale (IES) • Posttraumatic Growth Inventory (PTGI) 	

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation was stratified by participant and carer and state (Queensland vs New South Wales) and occurred in blocks of 10, with each condition randomly generated 5 times within each block to ensure an unpredictable allocation sequence with equal numbers of participants in each group at the completion of each block

Chambers 2014 (Continued)

Allocation concealment (selection bias)	Low risk	Unpredictable allocation sequence was undertaken by the project manager and was concealed from investigators
Blinding (performance bias and detection bias) All outcomes	Unclear risk	Project staff tracking assessments were blinded to treatment allocation
Incomplete outcome data (attrition bias) All outcomes	High risk	Loss to follow-up over 20%; loss to follow-up considerably higher in the 5-session psychologist-delivered CBT sessions intervention group
Selective reporting (reporting bias)	Low risk	Balanced reporting for all measures
Other bias	Unclear risk	93% completed the single-intervention session with the nurse; only 53% completed all 5 sessions in the psychologist arm

Chambers 2015
Study characteristics

Methods	<p><u>Setting</u>: private clinics, public/private hospitals, and public service announcements in Queensland, Australia</p> <p><u>Recruitment</u>: eligible participants were men who had been scheduled for/had undergone radical prostatectomy within the last 12 months and their female partners. Participants were required to be able to read and write in English, without any other concurrent cancer, and with no history of head injury, dementia, or psychiatric illness</p> <p><u>Randomisation</u>: randomised in blocks of 12, with each condition randomly generated 4 times within each block</p>
Participants	189 men with prostate cancer and their female partners
Interventions	<p><u>Intervention</u>: men and their partners were randomly assigned to 1 of 3 groups: (a) couples-based peer-delivered telephone support, (b) couples-based nurse-delivered telephone counselling, or (c) usual care. All participants in the intervention arms received at least 6 sessions of telephone support/counselling by nurse counsellors or peer support volunteers. Couples recruited pre-surgery received an additional 2 calls before undergoing surgery. Both intervention arms included skills training in couple communication and conjoint coping, in addition to a DVD with tip sheets. The nurse counselling intervention included educating about prostate cancer, menopause, and sexuality; providing behavioural homework on increasing expression of affection and non-demanding sexual touch; challenging negative beliefs about prostate cancer, ageing, and sexuality; and choosing and integrating a medical treatment for erectile treatment into the relationship. The peer support intervention included psycho-education about diagnosis; common experiences with surgery and recovery; managing side effects; improving communication; maintaining intimacy; sexual problems; and managing erectile dysfunction. Men and their partners were required to attend all sessions. Participants randomised to usual care received standard medical management and published participant education materials</p> <p><u>Interventionist</u>: 2 prostate cancer nurse counsellors</p>
Outcomes	<ul style="list-style-type: none"> • Uptake of medical treatment for erectile dysfunction • Sexual function/satisfaction • Sexuality support needs • Masculine self-esteem • Marital satisfaction

Chambers 2015 (Continued)

Methods for assessing outcomes:

- Utilisation of erectile dysfunctions scale
- International Index of Erectile Function
- Sexuality needs subscale of the Supportive Care Needs Survey
- Psychological Impact of Erectile Dysfunction – Sexual Experience
- Masculine Self-Esteem Scale
- Revised Dyadic Adjustment Scale
- Miller Social Intimacy Scale

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation occurred in blocks of 12, with each condition randomly generated four times with each block to ensure an unpredictable allocation sequence
Allocation concealment (selection bias)	Low risk	Sequence was undertaken by the project manager and concealed from investigators
Blinding (performance bias and detection bias) All outcomes	High risk	Owing to the nature of the intervention, participants and interveners were not blinded
Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Dong 2018
Study characteristics

Methods	<p><u>Setting</u>: China</p> <p><u>Recruitment</u>: All participants had colorectal cancer diagnosis confirmed by pathology and/or cytology before enrolment. Inclusion criteria were (1) ≥ 18 years of age; (2) Self-Rating Depression Scale score of 50 to 70; (3) willing to comply with study protocols; (4) expected survival time over 6 months; (5) awareness of personal disease status; (6) understanding of the study and being voluntarily recruited; (5) no history of other malignant tumours or personal psychiatric conditions, family history of psychosis, or special medication history of psychoactive drugs; and (6) not treated with antidepressant drugs</p> <p><u>Randomisation</u>: randomised by computer-generated random numbers to 1 of 3 arms - 2 treatment arms, 1 control arm</p>
Participants	135 colorectal cancer participants

Dong 2018 (Continued)

Interventions	<p>Before recruitment, all participated in a 90-minute physiological and psychological education lecture based on the SIGN guidelines on colorectal cancer</p> <p><u>Control</u>: participants in the control group received usual care</p> <p><u>Interventions</u>: participants in intervention arms received 6 weekly telephone sessions of 20 to 40 minutes. The content of sessions for the 2 intervention arms is given below:</p> <p><i>Telephone Support (TS)</i>: each week, participants were asked questions related to changes in condition, new symptoms, fear of recurrence, treatment and side effects, genetic risk, self-care (diet, support groups, finances), and family concerns</p> <p><i>Telephone-Based Reminiscence Therapy (TBR)</i>: Week 1: the psychologist guided the participant to reminisce about people who had a positive influence on his/her life. Week 2: the participant reminisced about happy times in the past. Week 3: the participant talked about his or her past achievements and the significance of these achievements. Week 4: the participant recalled the important turning points in his/her life and the influence of each. Week 5: the participant talked about his/her struggles with cancer and their positive significance. Week 6: the participant talked about his/her hopes for the future</p> <p><u>Interventionist</u>: psychologist</p>
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Outcomes	<ul style="list-style-type: none"> • Depression • Anxiety • Subjective well-being • Perceived social support <p>Method of assessing outcome measures:</p> <ul style="list-style-type: none"> • Chinese version of the Self-Rating Depression Scale (SDS) • Hamilton Depression Scale (HAMD-24) • Chinese version of the Self-Rating Anxiety Scale (SAS) • Hamilton Anxiety Scale (HAMA) • Memorial University of Newfoundland Scale of Happiness (MUNSH) • Perceived Social Support Scale (PSSS)
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Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated random numbers (EPIDAT 3.1)
Allocation concealment (selection bias)	Low risk	Random numbers were provided in numbered opaque envelopes by an external staff member, ensuring that assessors were blinded to treatment group assignment
Blinding (performance bias and detection bias) All outcomes	Unclear risk	No details
Incomplete outcome data (attrition bias) All outcomes	Low risk	No loss to follow-up
Selective reporting (reporting bias)	Low risk	Balanced reporting for all measures

Dong 2018 (Continued)

Other bias	Unclear risk	Did not collect pathological grading and staging data for colorectal cancer, and previous studies showed that the severity of depression in colorectal cancer participants is related to pathological grade
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Downe-Wamboldt 2007
Study characteristics

Methods	<u>Setting:</u> academic cancer centre, Canada <u>Recruitment:</u> eligible participants were 50 years or older, had surgery within the past 3 months, were receiving care from the Academic Clinic's Oncology Program, were living within 200 km of the academic clinic, had access to a telephone, were aware of their diagnosis, understood English, and were able to provide informed consent. Women with breast cancer stage I or II who had a lumpectomy or a mastectomy were eligible, as were men with prostate cancer stage T1 or T2M0 who had had a radical prostatectomy, and individuals with small cell lung cancer stage I, II, or IIIA. Patients receiving chemotherapy were excluded. Participants were recruited from the offices of private surgical oncologists and urologists and site-specific clinics (breast, lung, and prostate) at an academic cancer centre clinic <u>Randomisation:</u> RCT	
Participants	175 men or women aged 50 years or older with breast, lung, or prostate cancer	
Interventions	<u>Intervention:</u> individualised telephone problem-solving counselling based on cognitive-behavioural principles. A series of counselling sessions were offered at participants' convenience over 3 months. Collaborative in nature; the participant and the interventionist worked together to define participant concerns, identify solutions, and evaluate consequences of these solutions <u>Control group:</u> usual care, including diagnostic and follow-up care at 3-month intervals <u>Interventionist:</u> nurse counsellor	
Outcomes	<ul style="list-style-type: none"> Psychological symptoms - depression, psychological adjustment Impact on quality of life/functioning - service use Methods for assessing outcomes: <ul style="list-style-type: none"> Centre for Epidemiologic Studies Depression Scale (CES-D) ($\alpha = 0.89$ in this study) Derogatis Psychosocial Adjustment to Illness Scale - self-report (PAIS-SR) ($\alpha = 0.82$ in this study) Expenditures for Health and Social Service Utilisation Questionnaire Jalowiec Coping Scale (JCS) ($\alpha = 0.88$ and 0.94 for total scores; $\alpha = 0.70$ for 7 of the subscales in this study) 	
Notes		

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Participants were randomised to groups using concealed random computer-generated numbers that randomly blocked after every second and fourth subject to ensure equivalent numbers of subjects in both arms of the trial
Allocation concealment (selection bias)	Low risk	Participants were randomised to groups using concealed random computer-generated numbers that randomly blocked after every second and fourth subject to ensure equivalent numbers of subjects in both arms of the trial

Downe-Wamboldt 2007 (Continued)

Blinding (performance bias and detection bias) All outcomes	High risk	Owing to the nature of the intervention, participants and personnel were not blinded to the intervention
Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting of outcomes
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Girgis 2009
Study characteristics

Methods	<p><u>Setting</u>: New South Wales, Australia</p> <p><u>Recruitment</u>: eligible participants were 18 years of age or older, had non-localised breast or colorectal cancer, were diagnosed within the last 6 months, resided in New South Wales province at diagnosis, were able to complete study measures in English, and were physically and mentally capable of study participation. Potential participants were identified by the New South Wales Central Cancer Registry, and eligibility was confirmed by participants' treating physicians. Eligible individuals were sent a letter from the CCR seeking consent to receive study information from the research team. If a signed consent form was returned, the participant was contacted by telephone to complete baseline measures</p> <p><u>Randomisation</u>: parallel factorial RCT</p>
Participants	356 men and women aged 18 years or older with non-localised breast or colorectal cancer
Interventions	<p><u>Intervention</u>: based on the notion of providing feedback for patient-reported outcomes (PROs) to clinicians to improve supportive care for cancer patients. Participants completed 3 computer-assisted telephone interviews at baseline and at 3 and 6 months. Patient-reported outcomes collected during these interviews were summarised onto a feedback sheet by a specially developed computer programme, which identified issues of concern (e.g. severe symptoms). In the TCW arm, feedback sheets were forwarded to telephone care workers (TCWs), who then telephoned participants to discuss reported issues of concern and used a modified version of the Cancer Helpline database to refer participants to appropriate resources/services consistent with recommended feedback sheet strategies. TCWs also followed up with participants at 6-week intervals to assess coping. In the oncologist/GP arm, 2 hard copies of feedback sheets were mailed to participants' nominated oncologists and GPs for discussion at their next appointments. Clinicians were asked to keep 1 feedback sheet for their records and to return the second, which indicated which issues of concern were discussed and whether any actions were taken</p> <p><u>Control group</u>: usual care (not described)</p> <p><u>Interventionist</u>: oncology nurses with telephone counselling training</p>
Outcomes	<ul style="list-style-type: none"> • Psychological symptoms - anxiety, depression • Fatigue • Physical symptoms - nausea, vomiting • Impact on quality of life/functioning - quality of life, perceived needs, communication with health care professionals <p>Methods for assessing outcomes:</p>

Girgis 2009 (Continued)

- Hospital Anxiety and Depression Scale
- European Organisation for Research and Treatment of Cancer (EORTC) Quality of Life Questionnaire, version 3
- Supportive Needs Survey - Short Form
- Needs Assessment for Advanced Cancer Patient Questionnaire

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random assignment to one of the three study groups, by using a computer-generated algorithm, occurred at completion of the baseline CATI and included stratification by sex and tumour type
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	Owing to the nature of the intervention, participants and personnel were not blinded
Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting on outcomes
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Kroenke 2010
Study characteristics

Methods	<p><u>Setting</u>: Indiana, USA</p> <p><u>Recruitment</u>: eligible participants had a diagnosis of cancer and were screened positive for either pain or depression with scores of 6 or higher for cancer-related pain, or 10 or higher for depression, as measured using 2 validated measurement instruments. Participants were recruited from 16 urban and rural oncology practices. Participants presenting for oncology clinic visits underwent an eligibility interview, and eligible participants willing to participate provided audiotaped oral informed consent (followed up by posted written consent forms) and completed a baseline interview</p> <p><u>Randomisation</u>: RCT; block-randomised in block sizes of 4, 8, and 12 and stratified by symptom (pain vs depression)</p>
Participants	405 men and women suffering from cancer-related depression or pain
Interventions	<p><u>Intervention</u>: consisted of centralised tele-care management coupled with automated home-based symptom monitoring by interactive voice recording or Internet. Participants received a baseline call and 3 follow-up calls (1, 4, and 12 weeks) during the first 3 months of treatment, as well as triggered telephone calls when automated monitoring indicated inadequate symptom improvement, non-adherence to medication, adverse effects, suicidal ideation, or a participant request to be contacted. The automated symptom monitoring survey was administered twice a week for the first 3 weeks, then</p>

Telephone interventions for symptom management in adults with cancer (Review)

Kroenke 2010 (Continued)

weekly during weeks 4 through 11, twice a month during months 3 through 6, and once a month during months 7 through 12

Control group: usual care. Participants in this arm were informed of their depressive and pain symptoms, and their screening results were provided to the oncologist. No further attempts were made by study personnel to influence depression or pain management unless a psychiatric emergency arose (e.g. suicidal ideation was detected on baseline or follow-up outcome assessment)

Interventionist: nurse care manager trained in assessing symptom response and medication adherence, in providing pain and depression-specific education, and in making treatment adjustments according to evidence-based guidelines

Outcomes

- Psychological symptoms - depression
- Physical symptoms - pain, physical symptom burden
- Impact on quality of life/functioning - quality of life, disability, self-reported health care use

Methods for assessing outcomes:

- Patient Health Questionnaire (PHQ-9)
- Brief Pain Inventory (BPI) Interference Scale
- Hopkins Symptom Checklist (HSCL-20)
- Short Form Health Survey (SF-36) Mental Health Inventory and Bodily Pain Scale
- Generalised Anxiety Disorder Scale
- Sheehan Disability Scale
- Short Form Health Survey (SF-12)

Notes
Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation was computer-generated in randomly varying block sizes of 4, 8, and 12 and was stratified by symptom type (pain only, depression only, or both pain and depression)
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	Low risk	Outcome assessments were conducted by a research assistant blinded to study group
Incomplete outcome data (attrition bias) All outcomes	High risk	Loss to follow-up 34%
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting on outcomes
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Livingston 2010
Study characteristics
Telephone interventions for symptom management in adults with cancer (Review)

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Livingston 2010 (Continued)

Methods	<p>Setting: Victoria, Australia</p> <p>Recruitment: eligible participants were men recently diagnosed with colorectal or prostate cancer stage I-III, who spoke sufficient English, had no psychiatric illness, and had a prognosis longer than 52 weeks. Participants were recruited at the end of the consultation, at which time they received their diagnosis. Specialists followed a standardised script and introduced participants to the concept of the Cancer Helpline and stated that they would arrange for the Helpline to contact them, or (if in the usual care group) recommended that participants contact the Helpline themselves. The specialist provided a referral slip detailing the number of outcalls the participants would receive, or the Helpline toll-free number to call themselves. The specialist then advised participants that a study was being conducted about the needs of men with cancer, gave them an information package about the study, and asked them to consider taking part</p> <p>Randomisation: RCT. Block randomisation: medical specialists were randomised to 1 of 3 trial arms and then enrolled 15 consecutive participants to that arm, 15 participants to the next arm, and so forth. Specialists were blinded to which arm they were referring to</p>	
Participants	571 men with colorectal or prostate cancer	
Interventions	<p>Intervention: consisted of 2 different interventions: (1) active referral - 4 outcalls, in which men received 4 telephone calls from the Cancer Helpline within 1 week of diagnosis, and at 6 weeks, 3 months, and 6 months post diagnosis, and (2) active referral - 1 outcall, in which men received 1 telephone call within 1 week of diagnosis. In both arms, participants were called and offered to discuss 10 different topics: the cancer diagnosis; treatment/management issues; what to expect from surgery; management of side effects; communication with the specialist; partner/family issues; psychological/emotional and communication concerns; understanding cancer language; diet and nutrition; other support services; and availability of printed resources. If participants did not mention a topic, the cancer nurse raised the topic. All topics discussed and information requested were documented</p> <p>Control group: passive referral arm; designed to resemble usual care. Participants were referred to the Cancer Helpline, with contact at their initiative</p> <p>Interventionist: Cancer Helpline cancer nurses with postgraduate counselling qualifications and minimum 5 years clinical oncology experience</p>	
Outcomes	<ul style="list-style-type: none"> Psychological symptoms - cancer-specific distress, anxiety, depression <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> Hospital Anxiety and Depression Scale (HADS) ($\alpha = 0.78$ to 0.86 for anxiety, $\alpha = 0.78$ to 0.81 for depression at all time points in the current study) Control Preference Scale 	
Notes		
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Participants were block randomised (15 at a time) to each of the 3 study arms by specialist clinicians via computer-generated random numbers produced by the project co-ordinator under supervision of the investigators
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	Owing to the nature of the intervention, participants and personnel were not blinded

Livingston 2010 (Continued)

Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups
Selective reporting (reporting bias)	Unclear risk	Significant differences between active referral - 4 outcalls and passive referral arms were presented. Non-significant differences between the active referral - 1 outcall and active referral - 4 outcalls arms were mentioned but were not presented. However, lack of effect of the active referral intervention is discussed
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Mishel 2002
Study characteristics

Methods	<p><u>Setting</u>: central and eastern North Carolina, USA</p> <p><u>Recruitment</u>: eligible patients were African American and Caucasian men who were diagnosed with localised prostate cancer and were within 2 weeks of catheter removal post surgery and/or < 3 weeks of starting radiation therapy, had no major cognitive impairment, had no concurrent other cancer, had access to a telephone, had a family member willing to participate in the study, and planned to reside in their current community for 12 months. Participants were recruited from 9 treatment facilities. Members of the research team introduced the study to participants during a clinic visit, and interested participants were subsequently mailed an information package and were called to determine willingness to participate</p> <p><u>Randomisation</u>: RCT using 3 × 2 block randomisation</p>
Participants	239 men with prostate cancer
Interventions	<p><u>Intervention</u>: based on the Theory of Uncertainty in Illness, which assumes that uncertainty occurs when participants lack the information or knowledge needed to fully understand their illness and treatments. Through cognitive re-framing, participants can learn to cope better with uncertainty. The intervention consisted of 8 × weekly phone calls, which were delivered in different ways in the 2 intervention arms: (1) direct, in which only the participant received the intervention, and (2) supplemented, with supplemented delivery to a close family member. An assessment was made by telephone to identify the participant's cancer-related concerns. In the supplemented arm, the spouse or designated family support person also received a weekly phone call for 8 weeks from a nurse who conducted a similar assessment of family members' concerns about the participant using a list similar to that used for the participants. Based on identified concerns, an appropriate intervention was then selected from a standardised list, which included validating and reinforcing views and behaviours, providing information, activating resources, teaching symptom management strategies, structuring expectations, applying problem-solving, and teaching assertion techniques for communicating with health care providers. These interventions were applied to concerns about diagnosis, treatment, response to treatment, living with cancer, caring for oneself, and social/lifestyle issues. Printed materials, audiotapes, and videotapes for managing specific problems were mailed to participants after each weekly call. Any materials that were sent to the participant were reviewed at the next phone call</p> <p><u>Control group</u>: usual care, which included printed general health information not related to prostate cancer or the side effects of treatment</p> <p><u>Interventionist</u>: trained nurses matched with the participant and the family member by ethnicity and gender</p>
Outcomes	<ul style="list-style-type: none"> • Psychological symptoms - symptom distress • Uncertainty - uncertainty and uncertainty management were measured

Telephone interventions for symptom management in adults with cancer (Review)

Mishel 2002 (Continued)

Methods for assessing outcomes:

- Mini-Mental State Examination (MMSE) ($\alpha = 0.85$ to 0.98 in various studies)
- Mishel's Uncertainty in Illness Scale, shortened version ($\alpha = 0.82$ to 0.88 in this sample)
- Self-Control Scale - Problem-Solving and Cognitive Re-framing subscales ($\alpha = 0.86$ to 0.90 in this sample)
- Cancer Knowledge Scale (KR-21) ($\alpha = 0.64$ to 0.73 in this sample)
- Symptom Distress Scale ($\alpha = 0.68$ to 0.74 in this sample)

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The generation of random sequence is not described
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	Unclear risk	"Data collectors who were not involved in the delivery of the intervention collected all data". It is unclear if participants were blinded to group
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Low loss to follow-up, but reasons for this are not described
Selective reporting (reporting bias)	Low risk	Results of all measures are reported, including non-significant findings. However, the paper focuses slightly more on significant differences found at T2 than on non-significant differences at T3
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Mishel 2005
Study characteristics

Methods	<p><u>Setting</u>: cancer centres and hospitals in North Carolina, USA</p> <p><u>Recruitment</u>: eligible participants were African American and Caucasian women who were 5 to 9 years post treatment for breast cancer, had no cognitive impairment, had no concurrent treatment for another cancer, were recurrence free, could be on tamoxifen or similar agents, had access to a telephone, and planned to reside in their current community for 2 years following entry into the study. Participants were identified using the tumour registries from 13 cancer centres and hospitals. Introductory letters were sent to these women to ask permission to release their name and contact details to the research team, who then mailed letters about the study to interested women and later called them to determine willingness to participate. Additional African American women were recruited from the community via announcements in local media and community volunteers</p> <p><u>Randomisation</u>: RCT, block-randomised by ethnicity</p>
Participants	509 African American and Caucasian breast cancer survivors

Mishel 2005 (Continued)

Interventions

Intervention: based on the Theory of Uncertainty in Illness, which assumes that uncertainty occurs when participants lack the information or knowledge needed to fully understand their illness and treatments. Through cognitive re-framing, participants can learn to cope better with uncertainty. The intervention had 2 main components: (1) cognitive strategies delivered via audiotapes to teach active emotion-focused coping responses to threats of recurrence; and (2) behavioural strategies packaged in a self-help manual designed to provide management skills, information and resources on long-term treatment side effects, and cancer resources. Using a standardised protocol, interventionists guided women through the intervention over the course of 4 weekly telephone calls. The women practised 1 of the 4 cognitive coping skills each week during the calls: (1) relaxation, (2) pleasant imagery, (3) calming self-talk, and (4) distraction. The women were instructed to use these skills when confronting a trigger of fear of recurrence, or when anticipating such a trigger. During the third and fourth telephone calls, women were guided in use of the manual, which contained behavioural strategies in the form of information about long-term treatment side effects. During each telephone session, women were encouraged to practise the skills by listening to audiotapes and using the manual until the next call

Control group: usual care. No attempt was made to limit exposure to naturally occurring learning contexts such as media coverage of cancer issues, public health programmes, or cancer support groups

Interventionist: nurses (specific training not described)

Outcomes

- Psychological symptoms - psychological distress
- Uncertainty - uncertainty and uncertainty management were measured

Methods for assessing outcomes:

- Self-Control Scale - Problem-Solving and Cognitive Re-framing subscales ($\alpha = 0.84$ to 0.86 in this sample)
- Cancer Survivor Knowledge Scale (KR-20) ($\alpha = 0.58$ in this sample)
- Participant/Provider Communication Scale ($\alpha = 0.76$ to 0.78 in this sample)
- Social Support Short Form Questionnaire Satisfaction subscale ($\alpha = 0.90$ to 0.89 in this sample)
- Cognitive Coping Strategies Questionnaire (CSQ), modified version ($\alpha = 0.71$ to 0.89 in this sample)
- Total Information Received and Helpfulness of Information Questionnaire (investigator developed) (on various subscales, $\alpha = 0.82$ to 0.94 in this sample)
- Profile of Mood State - Short Form (POMS-SF) (total scale: $\alpha = 0.94$ to 0.95 in this sample; subscales: $\alpha = 0.80$ to 0.90 in this sample)

Notes
Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	The SAS program Proc Plan was used to construct a randomisation plan
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	Owing to the nature of the intervention, participants and personnel were not blinded
Incomplete outcome data (attrition bias) All outcomes	Low risk	Low loss to follow-up, but reasons for this are not described
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting on outcomes

Mishel 2005 (Continued)

Other bias	Unclear risk	At each data collection point, each woman received a "gasoline card worth \$20"
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Molassiotis 2009
Study characteristics

Methods	<p><u>Setting</u>: Manchester, UK</p> <p><u>Recruitment</u>: eligible participants were 18 years of age or older, were diagnosed with breast or colorectal cancer with a life expectancy longer than 6 months and starting oral capecitabine, were able to self-care and communicate in English, lived within 35 miles of the hospital, and had no past experience in home care nursing programmes. Participants were recruited</p> <p><u>Randomisation</u>: RCT</p>	
Participants	164 men and women aged 18 years or older with colorectal or breast cancer	
Interventions	<p><u>Intervention</u>: no theoretical basis; developed on best evidence for symptom management from the nursing literature. Participants received a home care nursing (HCN) programme for 18 weeks. The HCN programme consisted of symptom assessment, participant education, and/or treatment of symptoms. The nurse conducted 1 standard home visit (1 to 1.5 hours) during the first week of capecitabine treatment, in which chemotherapy and its adverse effects were discussed, questions participants had were sought and answered, and support was given. Subsequent home visits were offered when participants experienced multiple grade 3 toxicities or had difficulty coping with chemotherapy. All home care participants also received 1 monitoring phone call per week during all cycles (i.e. a minimum of 18 phone contacts during the intervention). These lasted between 10 and 25 minutes and included toxicity assessment followed by discussion of possible strategies to deal with reported symptoms. Participants also had access to a 24-hour, on-call specialist nursing service</p> <p><u>Control group</u>: usual care consisting of information about oral capecitabine and its adverse effects provided by clinicians and accompanied by printed information. Relevant medications to relieve symptoms were prescribed as needed. Participants were given the hospital's 24-hour emergency hotline phone number</p> <p><u>Interventionist</u>: nurses with training and experience in home care and in cancer care who received additional training relevant to the intervention</p>	
Outcomes	<ul style="list-style-type: none"> • Psychological symptoms - anxiety, depression • Fatigue • Physical symptoms - symptom toxicity including oral mucositis, hand-foot syndrome, diarrhoea, constipation, nausea, vomiting, pain, insomnia • Impact on quality of life/functioning - quality of life, health services utilisation <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> • National Cancer Institute Common Toxicity Criteria (NCI-CTC) • Hospital Anxiety and Depression Scale • European Organisation for Research and Treatment of Cancer Quality of Life Questionnaire C30 (EORTC-QLQ-C30) 	
Notes		
Risk of bias		
Bias	Authors' judgement	Support for judgement

Molassiotis 2009 (Continued)

Random sequence generation (selection bias)	Low risk	Random assignment was carried out by an independent statistician through a computer-generated program
Allocation concealment (selection bias)	Low risk	No clinician or researcher could anticipate or direct allocation
Blinding (performance bias and detection bias) All outcomes	Low risk	Interventionist and participants not blinded, but outcome assessor blinded
Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting on outcomes
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Mosher 2016
Study characteristics

Methods	<p><u>Setting</u>: Indiana, USA</p> <p><u>Recruitment</u>: eligible participants were diagnosed with small cell or non-small cell lung cancer; had at least 1 symptom of moderate severity, defined by validated cutoff points for depressive symptoms, anxiety, pain, vitality, or breathlessness; were fluent in English; and had a consenting carer</p> <p><u>Randomisation</u>: RCT; randomisation by a person not involved with study delivery using an SAS procedure</p>
Participants	106 dyads comprising a man/woman with symptomatic lung cancer and his/her carer
Interventions	<p><u>Intervention</u>: based on Social-Cognitive Theory. Participants took part in four 45-minute telephone symptom management (TSM) sessions. Both dyad members participated simultaneously through speaker phone. Received instruction in symptom management strategies. Mailed handouts detailing major points discussed during sessions, along with home practice assignments and digital media with instructions for relaxation exercises. Primary goal of intervention was to teach participants and carers evidence-based cognitive-behavioural and emotion-focused strategies for managing symptoms. All sessions had dual focus on participant and carer concerns</p> <p><u>Control group</u>: four 45-minute telephone sessions delivered to dyads by speaker phone aimed at directing participants to resources for practical health information and psychosocial services</p> <p><u>Interventionist</u>: clinical social workers trained by a PhD psychologist</p>
Outcomes	<ul style="list-style-type: none"> • Depression • Anxiety • Physical symptoms (pain, fatigue, breathlessness) • Self-efficacy • Social constraints • Caregiver burden <p>Methods for assessing outcomes:</p>

Mosher 2016 (Continued)

- 8-item Patient Health Questionnaire (PHQ-8)
- 7-item generalized anxiety disorder scale (GAD-7)
- Brief Pain Inventory (BPI)
- Fatigue Symptom Inventory
- 4 items from the Memorial Symptom Assessment Scale (MSAS) to measure frequency and severity of breathlessness and associated distress

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Generated by an individual independent from the study. Stratified by participant gender and performance status
Allocation concealment (selection bias)	Unclear risk	No details
Blinding (performance bias and detection bias) All outcomes	Low risk	Research assistants blind to study condition conducted all assignments by telephone
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Loss to follow-up over 20%; reasons similar between groups
Selective reporting (reporting bias)	Low risk	Balanced reporting for all measures
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Porter 2011
Study characteristics

Methods	<p><u>Setting</u>: North Carolina, USA</p> <p><u>Recruitment</u>: eligible participants had a diagnosis of non-small cell lung cancer stage I-III or limited-stage small cell lung cancer, had no other cancers in the past 5 years, were able to read and speak English, and had a carer who was also willing to participate. Participants were recruited from a university hospital oncology programme and from community oncology clinics</p> <p><u>Randomisation</u>: RCT</p>
Participants	233 lung cancer patients and their carers
Interventions	<p><u>Intervention</u>: coping skills training based on cognitive-behavioural therapy principles. Participants and their carers received either (1) carer-assisted coping skills training, or (2) education/support involving the carer. Both arms consisted of 14 × 45-minute telephone-based sessions delivered to the participant and the carer simultaneously. Sessions were conducted over an 8-month period tapered from weekly (sessions 1 to 3) to biweekly (sessions 4 to 10) to monthly (sessions 11 to 14). Telephone sessions were supplemented with printed and digital/audiovisual materials with instructions for progressive muscle relaxation. Sessions 1 to 7 focused on specific coping skills; sessions 8 to 12 focused on smoking cessation, relaxation and imagery exercises, and application of coping skills to particular challenges faced</p>

Porter 2011 (Continued)

by the participant and/or carer; session 11 focused specifically on the carer, encouraging the carer to explore his/her own resources and sources of stress and discussing how to use coping skills to ease the burden of care giving; sessions 13 and 14 focused on maintenance strategies including a review of coping skills learned and how to maintain regular practice of skills to prevent and cope with possible setbacks

Control group: participants and carers in the education/support arm received information about lung cancer and its treatment via a presentation and discussion format. Sessions were supplemented with handouts summarising major points and listing additional resources (e.g. websites, books) that participants could access if desired. Participants in this condition did not receive any training in coping skills

Interventionist: both treatment conditions were delivered by registered nurses trained by PhD-level psychologists and medical oncologists

Outcomes

- Psychological symptoms - psychological distress
- Fatigue
- Physical symptoms - pain
- Impact on quality of life/functioning - quality of life, self-efficacy for symptom management

Methods for assessing outcomes:

- Brief Pain Inventory (BPI)
- Beck Depression Inventory (BDI) ($\alpha = 0.86$ in this study)
- State Trait Anxiety Inventory (STAI) ($\alpha = 0.92$ in this study)
- Functional Assessment of Cancer Therapy - Lung Cancer (FACT-L) (in this study, $\alpha = 0.70$ to 0.86 on various subscales)
- Investigator-developed instrument to measure self-efficacy based on the Chronic Pain Self-Efficacy Scale ($\alpha = 0.95$ in this study)

Notes
Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation assignments were generated by an individual not involved in the study via a random-numbers table
Allocation concealment (selection bias)	Low risk	Assignments were concealed in envelopes that were not opened until participants had completed their pre-treatment evaluation
Blinding (performance bias and detection bias) All outcomes	Low risk	Research assistants conducting assessments were blind to treatment condition
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	High loss to follow-up; reasons for this not described in detail
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting
Other bias	Unclear risk	Lack of a no treatment control group

Rawl 2002
Study characteristics

Methods	<p><u>Setting</u>: urban, tertiary cancer centre and community-based cancer centre in a medium-sized midwestern city, USA</p> <p><u>Recruitment</u>: eligible participants were aged 18 years or older; were newly diagnosed with breast, colorectal, or lung cancer; were undergoing chemotherapy; had identified carers; and spoke English. Participants were recruited by a trained recruiter, who explained the study</p> <p><u>Randomisation</u>: RCT</p>
Participants	109 men and women with breast, colon, or lung cancer and their partners
Interventions	<p><u>Intervention</u>: based on self-regulation theory, which suggests that concrete, objective information in combination with appropriate emotional support can help minimise disruption of a participant's usual activities resulting in decreased psychological distress. A computer programme guided the clinical intervention, which consisted of 5 face-to-face visits and 4 telephone calls delivered alternately every second week over 18 weeks. During the initial visit, the nurse took a brief history, assessed the participant's symptoms, and developed a plan of care tailored to the participant. In each following visit/call, nurses assessed participants' physical, mental, and resource needs and their symptoms, and tailored interventions to participants using the computer programme. Effectiveness of interventions used to cope with these needs was assessed, and nurses offered emotional and counselling support to participants</p> <p><u>Control group</u>: usual care, which consisted of verbal information about what to expect from chemotherapy and symptoms that should be reported to the doctor. Participants received any education normally delivered during chemotherapy but no attention outside of medical visits</p> <p><u>Interventionist</u>: master's prepared oncology nurse specialist trained specifically in the intervention protocol and in use of the computer-based nursing system</p>
Outcomes	<ul style="list-style-type: none"> Psychological symptoms - psychological functioning, anxiety, depression <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> Medical Outcomes Study 36 Short Form (SF-36) State-Trait Anxiety Inventory (STAI) Center for Epidemiological Studies Depression-20 Scale (CESD-20)

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Group assignment was generated by computer
Allocation concealment (selection bias)	High risk	Not stated
Blinding (performance bias and detection bias) All outcomes	Low risk	Interviewers who collected outcome data were blind to respondents' group assignments
Incomplete outcome data (attrition bias) All outcomes	High risk	Attrition over 20%, and the intervention had twice the level of attrition as the control. Nine withdrew from the study due to the time-consuming nature of the intervention

Rawl 2002 (Continued)

Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Ream 2015
Study characteristics

Methods	<p><u>Setting</u>: UK</p> <p><u>Recruitment</u>: patients were eligible if they were receiving their first course of intravenous (IV) chemotherapy for treatment of breast or colorectal cancer or a lymphoma, were aged 18 years or over, had experienced at least moderate fatigue during previous treatment cycles, and had sufficient proficiency in English to complete the study. Participants were recruited when they attended their third cycle of chemotherapy</p> <p><u>Randomisation</u>: RCT via a table of random numbers</p>
Participants	44 participants with breast or colorectal cancer were recruited to the trial; 37 participants completed the trial
Interventions	<p><u>Intervention</u>: the Beating Fatigue by Telephone intervention comprised education on fatigue; assessment and monitoring of fatigue; coaching in self-care; and provision of emotional support. Participants in this arm were provided with a Coping with Fatigue booklet (Macmillan Cancer Support) and an investigator-designed handbook and fatigue diary. Telephone calls were scripted and specifically addressed fatigue through motivational interviewing techniques. Participants received 3 telephone calls</p> <p><u>Control</u>: people randomised to the control group were provided usual care: basic screening for fatigue and limited self-care advice without onward referral</p> <p><u>Interventionist</u>: a cancer nurse with experience of working on a Cancer Helpline</p>
Outcomes	<ul style="list-style-type: none"> • Primary outcomes: fatigue intensity and distress • Secondary outcomes: fatigue self-efficacy and psychological well-being <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> • Brief Fatigue Inventory • Fatigue Distress Scale • Hospital Anxiety and Distress scale • Scale based on Fuchs' brief health-specific self-efficacy scale
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Participants were randomised by simple randomisation via a table of random numbers between intervention and usual care
Allocation concealment (selection bias)	Unclear risk	Not stated

Ream 2015 (Continued)

Blinding (performance bias and detection bias) All outcomes	Low risk	Researchers collecting outcome measures were blind to the intervention. Health professionals providing care were blind to the intervention
Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting
Other bias	Unclear risk	The intervention group had higher scores at baseline compared to the control group in all variables under investigation - given small sample size, this may be important

Reese 2014
Study characteristics

Methods	<p><u>Setting</u>: Baltimore, USA (not specified)</p> <p><u>Recruitment</u>: eligible patients had undergone surgery/other treatment for colorectal cancer and had answered yes to having sexual concerns. In addition, they were required to be over 21 years old with a partner of at least 1 year, with sufficient reading/writing skills in English. Partners were required to be over 21 with sufficient English to participate</p> <p><u>Randomisation</u>: RCT stratified by participant gender and current ostomy use</p>
Participants	18 participants treated for colorectal cancer and their partners
Interventions	<p><u>Intervention</u>: participants and their partners were randomly assigned to the intimacy enhancement (IE) intervention or a wait-list control group (WL). The IE intervention consisted of 4 50-minute (weekly) telephone sessions, which included techniques from sex therapy and couple/marital therapy such as sensual touching exercises, improving sexual communication, identifying/challenging overly negative or rigid sexually related cognitions, and intimacy-building activities</p> <p><u>Control</u>: after both assessments had been completed in the waiting-list condition, the couple was able to participate in the IE intervention</p> <p><u>Interventionist</u>: previous paper outlining this intervention says, "all sessions of the intervention were delivered by the first author, J.B.R., a clinical psychologist"</p>
Outcomes	<ul style="list-style-type: none"> • Sexual distress • Sexual communication • Intimacy • Sexual function • Medical impact on sexual function • Self-efficacy <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> • Index of Sexual Satisfaction • 13-item Dyadic Sexual Communication Scale • Miller Social Intimacy Scale • Female Sexual Function Index (FSFI) and International Index of Erectile Functioning (IIEF) • Medical Impact subscale of Sexual Function Questionnaire

Reese 2014 (Continued)

- 3 self-efficacy questions rated on a 10 to 100 scale

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Randomisation was stratified by participant gender and ostomy use
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	High risk	Owing to the nature of the intervention, participants and interveners were not blinded
Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting
Other bias	Unclear risk	Couples were given \$25 if they completed both assessments

Reese 2018
Study characteristics

Methods	<p><u>Setting</u>: USA</p> <p><u>Recruitment</u>: breast cancer survivors were identified through providers' schedules and the institutional tumour registry. Women were eligible if they (a) were age ≥ 21 years, (b) were in a stable relationship (i.e. living with a romantic partner for ≥ 6 months) that could involve sexual activity, (c) had completed active treatment 6 months to 5 years ago for non-recurrent stage I–III breast cancer (current use of endocrine therapy was acceptable), and (d) scored ≥ 3 on the Patient Care Monitor sexual concerns screening item. Women were excluded if they had a past history of any cancer other than non-melanoma skin cancer, an ECOG performance score > 2 (or were judged as being too ill to participate), or overt cognitive dysfunction or psychiatric disturbance; were pregnant; or were currently engaged in couple or marital therapy; or if either they or their partner had hearing impairment or were not able to speak English. No exclusions were made based on sexual orientation</p> <p><u>Randomisation</u>: 2-arm RCT</p>
Participants	29 women with breast cancer and their partners (20 IE, 9 LHT)
Interventions	<p>Both arms consisted of four 60 to 75 minute weekly sessions delivered by telephone to both members of the couple by the interventionist and included session handouts for both survivors and partners sent in advance of the scheduled session</p> <p><u>Intimacy Enhancement (IE) arm</u>: 4 sessions of education and skills training to address survivors' sexual concerns and to improve intimacy. Techniques include sensual touching exercises adapted to incorporate specific instructions for how to deal with thoughts and feelings about breast touching during these exercises, communication skills with regard to sex and intimacy, identifying and challenging overly negative or inflexible sexually related cognitions, and broadening the repertoire of both sexual</p>

Reese 2018 (Continued)

and non-sexual intimacy building activities. Each of the 4 sessions includes a detailed agenda and topic, 1 or more skills practice sessions, and assignment of home practice exercises

Living Healthy Together (LHT) arm: education and support across a range of topics: education on finding support and the breast cancer experience, what it means to “live healthy together” (session 1), stress and stress management (session 2), fatigue and sleep (session 3), and diet and nutrition (session 4). Each session consisted of interactive discussion and activities including self-assessment and discussions both with the interventionist and between members of the couple in relation to the session topic

Interventionist: trained health care professionals (social work or counselling/psychology)

Outcomes

- Satisfaction with therapeutic services
- Programme evaluation
- Sexual function
- Sexual satisfaction
- Survivors’ sexual distress
- Self-efficacy for coping with sexual concerns
- Perceived quality of communication about sex
- Emotional intimacy
- Relationship quality
- Cancer-related distress
- Survivors’ body image distress
- Survivors’ psychological distress

Method of assessing outcome measures:

- Client Satisfaction Questionnaire (CSQ-8)
- Brief programme evaluation survey
- Female Sexual Function Index (FSFI)
- International Index of Erectile Functioning (IIEF)
- PROMIS SexFS version 2 Global Sexual Satisfaction Scale
- Female Sexual Distress Scale - Revised (FSDS-R)
- 3-item survey of self-efficacy for coping with sexual concerns
- Dyadic Sexual Communication Scale (DSCS)
- Emotional Intimacy subscale of the Personal Assessment of Intimacy in Relationships (PAIR)
- Dyadic Adjustment Scale (DAS-7)
- Impact of Event Scale - Revised Body Image Scale (BIS) Patient Health Questionnaire (PHQ-9) 7-item measure of Generalized Anxiety Disorder (GAD-7)

Notes
Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Study biostatistician generated the randomisation sequence (2:1 ratio)
Allocation concealment (selection bias)	High risk	Project manager assigned participants to interventions
Blinding (performance bias and detection bias) All outcomes	Unclear risk	No details
Incomplete outcome data (attrition bias)	Low risk	Study retention 97%

Reese 2018 (Continued)

All outcomes

Selective reporting (reporting bias)	Low risk	Balanced reporting for all measures
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Sherwood 2005
Study characteristics

Methods	<p><u>Setting</u>: Michigan, Indiana, and Ohio, USA</p> <p><u>Recruitment</u>: eligible participants were aged 21 years or older; were newly diagnosed with any solid tumour or non-Hodgkin's lymphoma stage III, IV, or recurrent; were undergoing chemotherapy; had no cognitive disabilities; had access to a telephone; and were able to read and speak English. Participants were recruited from 6 urban cancer centres by trained recruiters, who explained the study and obtained written consent</p> <p><u>Randomisation</u>: RCT</p>	
Participants	124 men and women aged 21 years or older with various cancers	
Interventions	<p><u>Intervention</u>: based on cognitive-behavioural theory, which states that an individual's perception of a situation affects his/her behaviour and beliefs regarding ability to control it. Participants' ability to manage symptoms can therefore be improved by changing their perception of them. The intervention consisted of 5 contacts spread over 8 weeks. The first and last contacts were face-to-face and were used mainly to establish rapport and to facilitate closure. The second, third, and fourth contacts were delivered by telephone. During each contact, nurses assessed symptoms and participants rated the severity of symptoms. Participants selected symptoms they would like to focus on, and nurses tailored a list of interventions. Nurses helped participants re-frame attitudes and beliefs and proposed cognitive-behavioural strategies. Participants then agreed to implement the interventions. Participants were responsible for choosing and implementing the strategies</p> <p><u>Control group</u>: usual care (not described)</p> <p><u>Interventionist</u>: nurse with experience in oncology trained in the study protocol</p>	
Outcomes	<ul style="list-style-type: none"> Psychological symptoms - depression Fatigue Physical symptoms - pain, nausea, vomiting, insomnia, dyspnoea, weakness, anorexia, xerostomia, fever, constipation, mouth sores <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> Symptom severity was measured by asking participants to rate the severity of each symptom on a scale of 0 to 10, and then summing severity rates for each symptom between sessions Center for Epidemiological Studies Depression-20 Scale (CESD-20) ($\alpha = 0.89$) 	
Notes		
Risk of bias		
Bias	Authors' judgement	Support for judgement

Sherwood 2005 (Continued)

Random sequence generation (selection bias)	Low risk	Following baseline data collection, a stratified randomisation schema was used to randomly assign participants from each recruitment site
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	Low risk	Data collection interviewers were not nurses and were not aware of which arm of the study participants were in
Incomplete outcome data (attrition bias) All outcomes	High risk	Attrition over 20%; reasons for this not reported
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Sikorskii 2007
Study characteristics

Methods	<p><u>Setting</u>: Michigan and Indiana, USA</p> <p><u>Recruitment</u>: eligible patients were 21 years or older, had a diagnosis of a solid tumour cancer or non-Hodgkin's lymphoma, were undergoing chemotherapy, were able to speak and read English, and had access to a touch-tone telephone. Participants were recruited from 2 comprehensive cancer centres, 2 community cancer oncology programme, and 6 hospital-affiliated community oncology centres by nurses from the clinical trial offices at these sites</p> <p><u>Randomisation</u>: RCT; participants were randomised according to recruitment location and site of cancer</p>
Participants	435 men and women with various cancers
Interventions	<p><u>Intervention</u>: no specified theoretical basis but incorporates elements of coping, re-framing, education, and eliciting support for adapting to or overcoming problems such as managing cancer symptoms. Consisted of (1) an automated telephone symptom management (ATSM) intervention delivered via an automated system, or (2) nurse-assisted symptom management (NASM). Participants in both arms received a total of 6 telephone calls over 8 weeks; weekly calls during the first 4 weeks; then 2 biweekly calls. For participants assigned to the NASM group, nurses delivered up to 4 strategies for each symptom, supplemented with references to a symptom management guide. At each subsequent contact, assigned strategies were evaluated. If a strategy was not tried, or was tried but was found not helpful, participants were counselled as to how they might fit strategy into their daily activities, or they were offered different strategies. Successful strategies were reinforced and continued. In the ATSM arm, a pre-recorded female voice queried participants regarding severity of the 17 symptoms. To rate severity, participants pressed the appropriate numbers on their telephone keypads. For symptoms rated at 4 or higher, participants were directed to the section of the symptom management guide that informed them about strategies to manage each symptom</p> <p><u>Control group</u>: no usual care group</p> <p><u>Interventionist</u>: experienced cancer nurses; specific training not described</p>
Outcomes	<ul style="list-style-type: none"> Psychological symptoms - distress

Sikorskii 2007 (Continued)

- Fatigue
- Physical symptoms - pain, dyspnoea, distress, nausea, fever, difficulty remembering, lack of appetite, dry mouth, vomiting, numbness and tingling, diarrhoea, cough, constipation, weakness, and alopecia

Methods for assessing outcomes

- Symptom severity was scored by participants on a scale of 0 to 10, and severity rates for each symptom were summed between sessions

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation was conducted via a computer minimisation programme
Allocation concealment (selection bias)	Unclear risk	Not stated
Blinding (performance bias and detection bias) All outcomes	Unclear risk	Not stated
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Loss to follow-up 20%; reasons not detailed
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Thomas 2012
Study characteristics

Methods	<p><u>Setting</u>: California and New Jersey, USA</p> <p><u>Recruitment</u>: eligible participants were 18 years of age or older, were able to read and understand English, had access to a telephone, had a life expectancy longer than 6 months, had cancer-related pain with an average pain intensity score of 2 or higher (on a 0 to 10 scale), had no cognitive or psychiatric condition, had no substance abuse problem, had no severe pain unrelated to their cancer, or resided in a setting where they could not self-administer pain medication (e.g. nursing home). Potential participants were identified by clinical staff and were recruited from 6 outpatient oncology clinics (4 Veterans Affairs facilities, 1 county hospital, and 1 community-based practice) by a member of the research team</p> <p><u>Randomisation</u>: RCT; participants were stratified by pain intensity and treatment via permuted blocks with variable sizes</p>
Participants	318 men and women aged 18 years or older with various types of cancer-related pain
Interventions	<p><u>Intervention</u>: based on change theory, specifically the Transtheoretical Model, in which behavioural change is a function of a person's state of readiness or motivation to modify a particular behaviour. Uses principles of coaching and motivational interviewing to modify participants' attitudes towards pain</p>

Thomas 2012 (Continued)

management. Participants received either (1) an educational intervention or (2) a coaching intervention. Both groups viewed a video on overcoming attitudinal barriers and received a pamphlet on managing cancer pain. In addition, all participants received 4 × 30 minute biweekly telephone calls conducted over 6 weeks. In the coaching group, the calls explored participants' beliefs about pain, use of pain medication, non-pharmacological pain management strategies, and communication about pain management. In the educational and usual care groups, the calls were for attention control (content not described)

Control group: usual care. Participants viewed a video on cancer produced by the American Cancer Society

Interventionist: advanced practice oncology nurse with expertise in cancer pain management trained in the Transtheoretical Model and in motivational interviewing by a cognitive-behavioural psychologist, and in the specific coaching protocol used in the study. Research associates were trained in providing attention control telephone calls for the control group

- Outcomes
- Psychological symptoms - attitudinal barriers to pain management
 - Physical symptoms - pain
 - Impact on quality of life/functioning - functional status, quality of life

Methods for assessing outcomes:

- Barrier Questionnaire (BQ)
- Brief Pain Inventory (BPI)
- MOS 36-item Short Form Health Survey (SF-36)
- Functional Assessment of Cancer Therapy Scale (FACT-G)

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Participants were randomised via permuted blocks with variable sizes
Allocation concealment (selection bias)	Low risk	Participants and clinicians were blinded to participants' group assignments
Blinding (performance bias and detection bias) All outcomes	High risk	Owing to the nature of the intervention, participants and personnel were not blinded
Incomplete outcome data (attrition bias) All outcomes	High risk	Loss to follow-up over 20%; no detail provided on reasons for loss to follow-up between study groups
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

Traeger 2015
Study characteristics
Telephone interventions for symptom management in adults with cancer (Review)

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Traeger 2015 (Continued)

Methods	<p>Setting: Massachusetts, USA</p> <p>Recruitment: eligible participants were 18 years or older; had a diagnosis of stage I-III breast, lung, or colorectal cancer and were scheduled to start chemotherapy; and were able to respond to questionnaires in English. Participants were recruited by a study co-ordinator during their first chemotherapy visit. They were recruited consecutively until target enrolment was reached</p> <p>Randomisation: by a clinical trials office (1:1 allocation sequence) with stratification by tumour type</p>
Participants	120 participants initiating chemotherapy for non-metastatic disease (60 breast cancer; 30 colorectal cancer; 30 lung cancer)
Interventions	<p>Intervention: provocative telephone-based nursing guidance and support during first 2 chemotherapy cycles. Interventionists had a brief outline to structure calls but were encouraged to complete them according to clinical judgement to reflect participant-centred practice. Two calls were provided during the first week after first chemotherapy administration, and 2 in the first week after the second cycle. Calls lasted around 15 minutes</p>
Outcomes	<ul style="list-style-type: none"> • Number of symptoms • Symptom distress • Depression • Anxiety <p>Methods for assessing outcomes:</p> <ul style="list-style-type: none"> • Memorial Symptom Assessment Scale-Short Form (MSAS-SF) • Patient Health Questionnaire-4 (PHQ-4)
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Dana Faber/Harvard Cancer Center Quality Assurance Officer for clinical trials randomly assigned participants
Allocation concealment (selection bias)	Unclear risk	For control participants, the nurse practitioner was not notified of study enrolment or group assignment
Blinding (performance bias and detection bias) All outcomes	Unclear risk	No details
Incomplete outcome data (attrition bias) All outcomes	Low risk	Study retention rate 97.5%
Selective reporting (reporting bias)	Low risk	Balanced reporting of all measures
Other bias	High risk	If the electronic health record showed a documented in-person oncology visit during the intervention delivery window, the in-person visit was counted as a completed call

Watson 2017

Study characteristics

Methods	<p><u>Setting</u>: UK</p> <p><u>Recruitment</u>: a consecutive series of participants referred to the Royal Marsden Hospital's Psychological Care Service over an 18-month period by clinical staff</p> <p><u>Randomisation</u>: prospective randomised equivalence trial</p>
Participants	<p>118 cancer participants greater than 8 weeks post diagnosis with a minimum prognosis greater than 3 months who were considered to have high psychological needs were randomised. Sixty participants were randomised to telephone-CBT (T-CBT), and 58 to treatment as usual-CBT (TAU-CBT) (43 and 35 provided complete analysable data)</p>
Interventions	<p>Prospective randomised equivalence trial comparing TAU-CBT with T-CBT. A no-treatment control group was not used, given prior data indicating the efficacy of standard care CBT for cancer participants</p> <p><u>Telephone CBT</u></p> <p>Participants telephoned at pre-arranged times and sessions scheduled. Up to 8 sessions offered therapy over an approximate 12-week period. Core therapy components include:</p> <ul style="list-style-type: none"> • establishing a collaborative therapeutic relationship between participant and therapist; • focusing sessions through agenda setting; • using a Socratic questioning-guided discovery technique; • teaching problem-focused coping; • using “homework” as a didactic method to advance coping efficacy; • scheduling activities to provide positive behavioural structure and pre-selected goals in everyday life; • using relaxation to assist in management of worry; • teaching participants to use distraction/thought-stopping to limit negative mood; • teaching monitoring/re-scripting/challenging of unhelpful negative automatic thoughts; and • using graded goal-setting, which is central; ventilation of concerns is encouraged. <p>Also provided a patient workbook and a digital relaxation resource</p> <p><u>CBT Face-to-Face Treatment As Usual (TAU-CBT)</u></p> <p>Not described</p> <p><u>Interventionist</u>: level 3/4 mental health professionals/psychologists</p>
Outcomes	<ul style="list-style-type: none"> • Anxiety and depression • Adjustment to cancer • Cancer concerns • Participant satisfaction <p>Method of assessing outcome measures:</p> <ul style="list-style-type: none"> • Hospital Anxiety and Depression Scale • Mental Adjustment to Cancer - helpless/hopeless subscale only • 14-item checklist of cancer concerns • Study-specific service questionnaire
Notes	

Risk of bias

Watson 2017 (Continued)

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation by independent statistician, stratified by therapist
Allocation concealment (selection bias)	Unclear risk	No details
Blinding (performance bias and detection bias) All outcomes	Unclear risk	No details
Incomplete outcome data (attrition bias) All outcomes	High risk	Attrition over 20%; loss to follow-up higher in TAU-CBT
Selective reporting (reporting bias)	Low risk	Balanced reporting for all measures
Other bias	Unclear risk	Referred based on clinician judgement of high psychological need – not screened

Yates 2005
Study characteristics

Methods	<p><u>Setting</u>: Melbourne and Brisbane, Australia</p> <p><u>Recruitment</u>: eligible patients were aged 18 years or older, had stage I or II breast cancer, were commencing adjuvant chemotherapy, and had an Eastern Cooperative Oncology Group performance rating of 1 or 2; their haemoglobin level was at least 11.6 g/mL. Participants were recruited from 5 outpatient clinics at 3 major metropolitan hospitals</p> <p><u>Randomisation</u>: RCT</p>
Participants	109 women aged 18 years or older with breast cancer
Interventions	<p><u>Intervention</u>: psycho-educational intervention aiming to improve participants' knowledge and skills to enable them to perform self-care behaviours designed to minimise fatigue. Consisted of 3 sessions tailored to participants' individual needs and circumstances. The first session consisted of a 20-minute face-to-face meeting in the clinic at the participant's second chemotherapy course, and focused on techniques of information-giving, problem-solving, rehearsal, and reinforcement. The second and third sessions were delivered by telephone, were conducted a week apart, and lasted 10 minutes each. In these sessions, participants' fatigue management strategies were reviewed. Printed information supplemented the face-to-face/telephone interactions</p> <p><u>Control group</u>: participants received general cancer education in the form of verbal and printed information about cancer in sessions equivalent in number and timing to the sessions provided to the intervention group</p> <p><u>Interventionist</u>: oncology nurses trained in the research programme and in the fatigue management or control intervention</p>
Outcomes	<ul style="list-style-type: none"> Psychological symptoms - anxiety, depression, self-efficacy with coping with cancer Fatigue - use of fatigue-management behaviours, confidence with managing fatigue, fatigue intensity, and impact Impact on quality of life/functioning - quality of life

Yates 2005 (Continued)

Methods for assessing outcomes:

- Investigator-designed questionnaire based on elements from the Revised Piper Fatigue Scale (RPFS)
- Functional Assessment of Cancer Therapy - Fatigue (FACT-F)
- 30-item European Organisation for Research and Treatment of Cancer Quality of Life Questionnaire C30 (version 3)

Notes

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Participants were randomly assigned to intervention or control conditions through a central telephone system via computer-generated random numbers
Allocation concealment (selection bias)	Low risk	Group allocation was concealed from research assistants involved in recruitment and in baseline and follow-up assessments
Blinding (performance bias and detection bias) All outcomes	Unclear risk	Not stated
Incomplete outcome data (attrition bias) All outcomes	Low risk	Loss to follow-up < 20%; similar reasons between groups
Selective reporting (reporting bias)	Low risk	There appears to be no selective reporting
Other bias	Low risk	The trial appears to be free of other problems that could put it at high risk of bias

CBT: cognitive-behavioural therapy.
 ECOG: Eastern Cooperative Oncology Group.
 GP: general practitioner.
 RCT: randomised controlled trial.
 SIGN: Scottish Intercollegiate Guidelines Network.
 TAU: treatment as usual.

Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion
Aranda 2006	Not primarily telephone intervention
Ashing-Giwa 2008	Not focused on symptom management
Bakitas 2009	Not focused on symptom management
Beaver 2006	Not an RCT
Beaver 2009	Not focused on symptom management
Beney 2002	Not focused on symptom management

Study	Reason for exclusion
Bohnenkamp 2004	Not primarily telephone intervention
Budin 2008	Not focused on symptom management
Campbell 2007	Not focused on symptom management
Cheville 2013	Not focused on symptom management
Coleman 2005	Not focused on symptom management
Collie 2006	Not primarily telephone intervention
Craddock 1999	Not focused on symptom management
Dalton 2004	Not primarily telephone intervention
De Wit 1997	Not primarily telephone intervention
DuHamel 2015	Not primarily telephone intervention
Given 2008	Subanalysis of included article (Sikorskii 2007)
Haddad 2003	Not focused on symptom management
Hagiopan 1990	Not focused on symptom management
Halbert 2004	Not exclusively cancer patients
Hanks 2002	Not exclusively cancer patients
Harrison 2011	Not focused on symptom management
Hawkes 2009	Not an RCT
Hawkins 2010	Not focused on symptom management
Hayes 2012	Not focused on symptom management
Kearney 2009	Not primarily telephone intervention
Kimman 2010	Not focused on symptom management
Koller 2013	Not primarily telephone intervention
Kornblinth 2006	Not delivered by an HCP
Marcus 2010	Not delivered by an HCP
McCorkle 2009	Not primarily telephone intervention
Meneses 2009	Not primarily telephone intervention
Nguyen 2018	Not primarily telephone intervention
Park 2012	Not focused on symptom management

Study	Reason for exclusion
Rustoen 2014	Not focused on symptom management
Salonen 2009	Not focused on symptom management
Sandgren 2000	Not focused on symptom management
Sandgren 2003	Not focused on symptom management
Sandgren 2007	Not focused on symptom management
Santacroce 2010	Not adults
Sikorskii 2009	Duplicate of included article (Sikorskii 2007)
Sikorskii 2009b	Duplicate of included article (Sikorskii 2007)
Spoelstra 2013	Not delivered by an HCP
Williams 2004	Not primarily telephone intervention
Winger 2018	Secondary analysis of included article (Mosher 2016)

HCP: health care practitioner.
 RCT: randomised controlled trial.

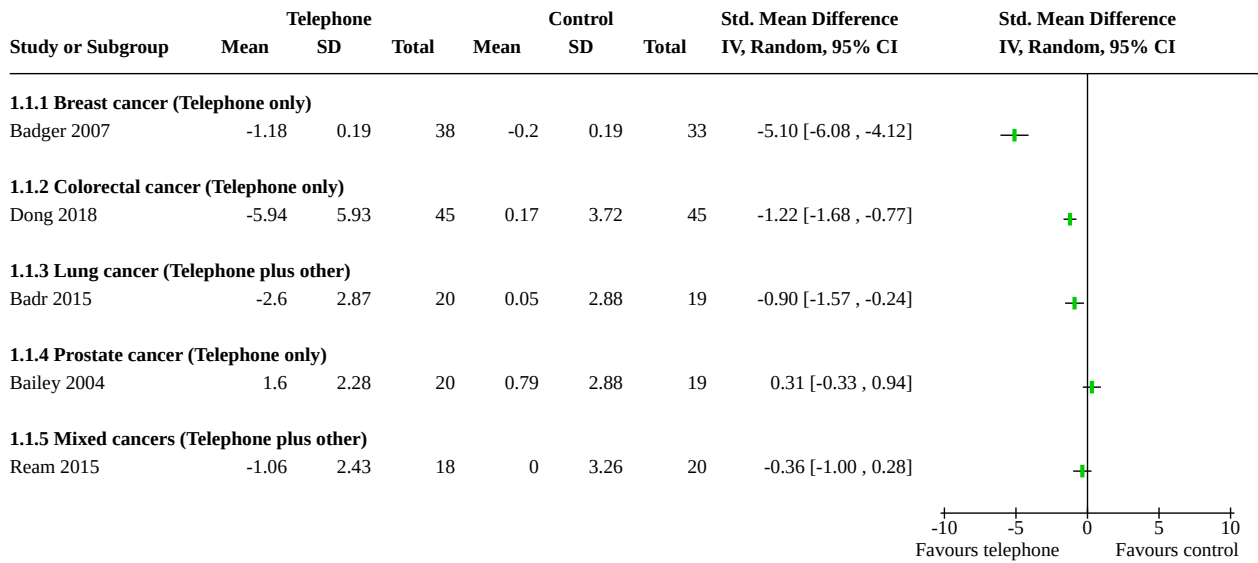
DATA AND ANALYSES

Comparison 1. Symptoms

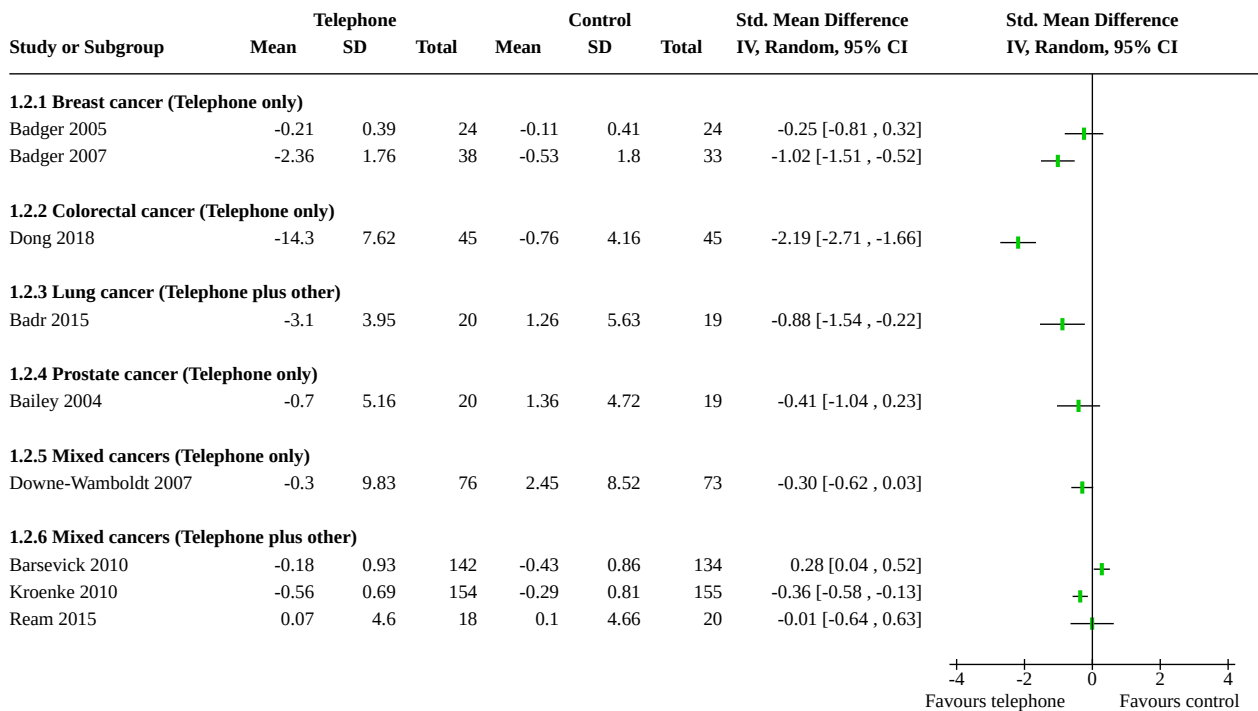
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1.1 Anxiety	5		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.1.1 Breast cancer (Telephone only)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.1.2 Colorectal cancer (Telephone only)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.1.3 Lung cancer (Telephone plus other)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.1.4 Prostate cancer (Telephone only)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.1.5 Mixed cancers (Telephone plus other)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.2 Depression	9		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1.2.1 Breast cancer (Telephone only)	2		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.2.2 Colorectal cancer (Telephone only)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.2.3 Lung cancer (Telephone plus other)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.2.4 Prostate cancer (Telephone only)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.2.5 Mixed cancers (Telephone only)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.2.6 Mixed cancers (Telephone plus other)	3		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.3 Fatigue	6		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.3.1 Breast cancer (Telephone only)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.3.2 Breast cancer (Telephone plus other)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.3.3 Prostate cancer (Telephone only)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.3.4 Mixed cancers (Telephone plus other)	3		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.4 Emotional distress	5		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.4.1 Breast cancer (Telephone only)	2		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.4.2 Breast cancer (Telephone plus other)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.4.3 Prostate cancer (Telephone only)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected
1.4.4 Mixed cancers (Telephone only)	1		Std. Mean Difference (IV, Random, 95% CI)	Totals not selected

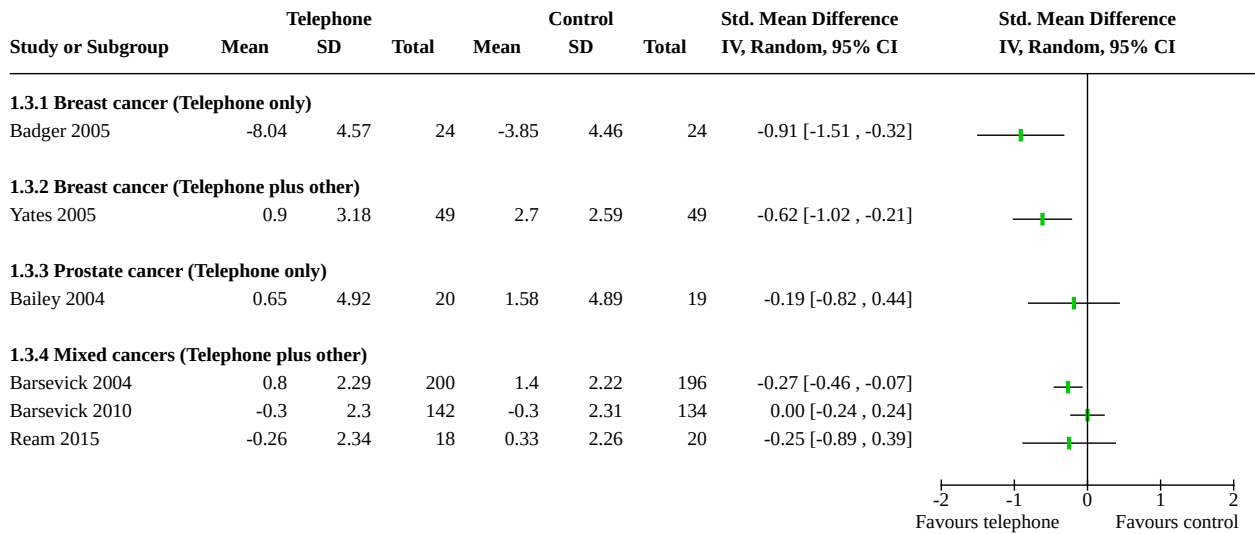
Analysis 1.1. Comparison 1: Symptoms, Outcome 1: Anxiety



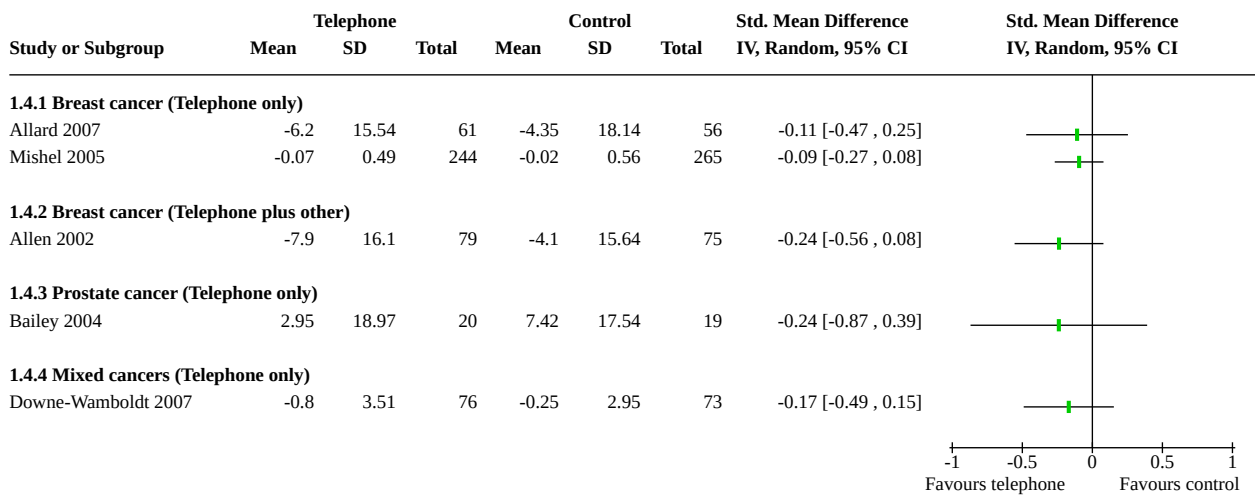
Analysis 1.2. Comparison 1: Symptoms, Outcome 2: Depression



Analysis 1.3. Comparison 1: Symptoms, Outcome 3: Fatigue



Analysis 1.4. Comparison 1: Symptoms, Outcome 4: Emotional distress



APPENDICES

Appendix 1. CENTRAL search strategy

- #1 MeSH descriptor Neoplasms explode all trees
- #2 neoplasm* or cancer* or carcinoma* or tumour* or adenocarcinoma* or leukemia* or leukaemi* or lymphoma* or tumor* or malignan* or myeloma*
- #3 MeSH descriptor Radiotherapy explode all trees
- #4 radiotherap* or radiation or radiochemotherap* or chemoradi* or chemotherap*
- #5 (#1 OR #2 OR #3 OR #4)
- #6 MeSH descriptor Telemedicine explode all trees
- #7 telemedicine or (tele next medicine)
- #8 teleconsultation or (tele next consultation)
- #9 telephone*
- #10 phone*
- #11 cellphone*

#12 remote* near/5 consultation*

#13 (#6 OR #7 OR #8 OR #9 OR #10 OR #11 OR #12)

#14 (#5 AND #13)

Appendix 2. MEDLINE search strategy

1 exp Neoplasms/

2 (neoplasm* or cancer* or carcinoma* or tumour* or adenocarcinoma* or leukemia* or leukaemi* or lymphoma* or tumor* or malignan* or myeloma*).mp.

3 exp Radiotherapy/

4 (radiotherap* or radiation or radiochemotherap* or chemoradi* or chemotherap*).mp.

5 1 or 2 or 3 or 4

6 exp Telemedicine/

7 (telemedicine or (tele adj medicine)).mp.

8 (teleconsultation or (tele adj consultation)).mp.

9 (remote* adj5 consultation*).mp.

10 telephone*.mp.

11 phone*.mp.

12 cellphone*.mp.

13 8 or 6 or 11 or 7 or 10 or 9 or 12

14 "randomized controlled trial".pt.

15 "controlled clinical trial".pt.

16 randomized.ab.

17 randomly.ab.

18 trial.ab.

19 groups.ab.

20 18 or 19 or 16 or 17 or 15 or 14

21 13 and 20 and 5

Appendix 3. Embase search strategy

1 exp Neoplasm/

2 (neoplasm* or cancer* or carcinoma* or tumour* or adenocarcinoma* or leukemia* or leukaemi* or lymphoma* or tumor* or malignan* or myeloma*).mp.

3 exp Radiotherapy/

4 (radiotherap* or radiation or radiochemotherap* or chemoradi* or chemotherap*).mp

5 1 or 2 or 3 or 4

6 exp Telemedicine/

7 (telemedicine or (tele adj medicine)).mp.

8 (teleconsultation or (tele adj consultation)).mp.

9 (remote* adj5 consultation*).mp.

10 telephone*.mp.

11 phone*.mp.

12 cellphone*.mp.

13 8 or 6 or 11 or 7 or 10 or 9 or 12

14 exp Controlled Clinical Trial/

15 randomized.ab.

16 randomly.ab.

17 trial.ab.

18 groups.ab.

19 18 or 16 or 17 or 15 or 14

20 19 and 13 and 5

HISTORY

Protocol first published: Issue 1, 2009

Review first published: Issue 6, 2020

CONTRIBUTIONS OF AUTHORS

ER: background, objectives, and review development; data extraction and analysis; manuscript writing. AR: background, objectives, and review development; data extraction; manuscript review. AH, VHP, AC, KC: literature search; organising/co-ordinating of data extraction; data extraction and analysis; manuscript writing. TW, AF: data extraction; manuscript review.

DECLARATIONS OF INTEREST

Emma Ream: principal investigator of a trial that assessed the effectiveness of a telephone intervention for cancer patients experiencing fatigue.

Amanda Amanda Euesden Hughes: none known.

Anna Cox: none known.

Katy Skarparis: none known.

Alison Richardson: none known.

Vibe H Pedersen: none known.

Theresa Wiseman: none known.

Angus Forbes: none known.

Andrew Bryant: none known.

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External sources

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- Cochrane Gynaecological, Neuro-oncology, and Orphan Cancers, UK

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

The protocol stated that the review would incorporate quality of life as a secondary outcome. However, quality of life resulting from a telephone-delivered intervention for symptom management proved an oft measured outcome that warrants a review in its own right. Consequently, these data have not been reported here. Rather, general symptom experience is presented as the secondary outcome.

INDEX TERMS

Medical Subject Headings (MeSH)

Anxiety [etiology] [therapy]; Cancer Pain [therapy]; Depression [etiology] [therapy]; Dyspnea [etiology] [therapy]; Fatigue [etiology] [therapy]; Neoplasms [*complications] [psychology] [therapy]; Randomized Controlled Trials as Topic; Self Care; Sexual Dysfunction, Physiological [etiology] [therapy]; Stress, Psychological [etiology] [therapy]; Symptom Assessment; Telemedicine [*methods]; *Telephone [statistics & numerical data]; Time Factors; Uncertainty

MeSH check words

Adult; Female; Humans; Male