

REVIEW

A systematic review of psychotherapeutic interventions for women with metastatic breast cancer: Context matters

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Funding information

Cancer Institute NSW; NHMRC, Grant/Award Number: GNT 1042942

Abstract

Objectives: To summarise the evidence-base of psychological interventions for women with metastatic breast cancer, by mode of delivery (group, individual, or low-intensity interventions). To synthesise data regarding core intervention-elements (eg, intervention duration) and context factors (trial setting, uptake and adherence, and demographic characteristics).

Methods: Four databases were searched (inception-May 2016): MEDLINE (OvidSP), PsycINFO (OvidSP), CINAHL (EBSCO), and SCOPUS; reference lists were examined for additional publications. Grey literature was excluded. Outcome data were extracted for survival, distress, quality of life, coping, sleep, fatigue, and/or pain and summarised through narrative synthesis.

Results: Fifteen randomised clinical trials (RCTs), reported across 23 articles, met inclusion criteria: 7 groups, 4 individuals, and 4 low-intensity interventions. Overall, interventions improved distress (8/13 RCTs), coping (4/5 RCTs), and pain (4/5 RCTs). No evidence of survival benefit was found. For remaining outcomes, evidence was either insufficient, or too mixed to draw conclusions. Group programs had the strongest evidence-base for efficacy; individual and low-intensity therapy had insufficient evidence to form conclusions. Group interventions had longest intervention durations and lowest uptake and adherence; low-intensity interventions had shortest durations and highest uptake and adherence. Disparities in uptake, adherence, and reach were evident, with the demographic profile of participants polarised to young, Caucasian, English-speaking, partnered women.

Conclusions: There remains a paucity of psychological interventions for women with metastatic breast cancer. Those that exist have an inconsistent evidence-base across the range of patient-reported outcomes. Further research is needed to evaluate accessible delivery formats that ensure efficacy as well as uptake.

KEYWORDS

cancer, metastatic breast cancer, oncology, psychological interventions, systematic review, treatment modality

1 | INTRODUCTION

Metastatic breast cancer (MBC) is defined as cancer that has spread from the breast to more distant parts of the body, most commonly the bones, lungs, liver, and brain.¹ Despite the advent of new systemic therapies, the median survival remains approximately 2 to 3 years,² although some women experience extended survival periods of over 10 years.

A diagnosis of advanced breast cancer brings significant adjustment challenges for women, as they face a future of physical symptoms, functional limitations, and existential concerns about their ultimate mortality. Clinically elevated distress and impaired health-related quality of life (HRQOL) occur in 35% to 43%³ of women following diagnosis of MBC. Given that psychological factors, particularly depression, have been implicated in reduced survival time,⁴ increased

severity of symptoms,⁵ and reduced treatment adherence,⁶ psychosocial interventions for this population are warranted.

However, contrasting to the 28 randomised controlled trials (RCTs) of psychological interventions for 3940 participants with early-stage breast cancer,⁷ the Cochrane Collaboration found only 10 RCTs had been specifically conducted for 1378 women with MBC.⁸ This MBC Cochrane Review, a meta-analysis published in 2013, included papers published between 1989 and 2010, with 7 of the 10 included RCTs evaluating a group therapy program of either a supportive-expressive (n = 4), or cognitive behavioural (n = 3) framework.⁸ The remaining 3 studies trialled individual face-to-face therapies and did not use either of these therapeutic frameworks. Overall, the Cochrane Review found a short-term psychological benefit of MBC interventions for improving pain and psychological outcomes (distress, helplessness/hopelessness, social functioning, cancer-related distress, and emotional control) and a modest survival benefit at 1 year that was not sustained at 5 years.⁸

Since publication of the Cochrane Review, there has been a surge of interest in the provision of online self-directed psychological interventions targeting illness-related distress.^{9–18} Furthermore, recent survey data indicate that psychosocial needs remain the key concern of women with MBC, with self-care strategies being listed as the most-preferred or second-preferred option for distress management.¹⁹ These Web-based and self-care strategies form part of an emerging modality for receiving psychological interventions, termed “low-intensity.”²⁰ The primary purpose of low-intensity interventions is to increase access to evidence-based psychological therapies, using the minimum level of intervention necessary to create maximum gain and delivering content in a variety of flexible forms, including email, smart devices, telephone, print-based resources, and internet.²⁰ While definitions vary, the key attributes of low-intensity interventions are that they are more accessible, use fewer formal health care professionals’ resources compared with conventional therapies, and may use nonspecialist practitioners to deliver the programs.^{20,21} In contrast, group and individual psychotherapeutic interventions are high intensity, defined as therapist-administered programs, delivered in-person by health professionals with specialist training,⁸ and can be of either brief (less than 6 hours) or extensive duration.²¹ Considering the research and patient preference developments relating to low-intensity interventions, particularly Web-based psychological therapy, it is timely to investigate whether they have penetrated the MBC literature and to update the evidence in light of any new, or previously missed, trials that have been published.

Given the variety of formats now available for disseminating psychological interventions, it is important to evaluate how the core elements of an intervention (modality of delivery and their differing treatment durations) impact on outcomes. While the prior Cochrane Review summarised the number of group and individual interventions included, they did not compare outcomes by delivery modality.⁸ Furthermore, it is also becoming increasingly recognised that research data must be considered within the context they were obtained. That is, to identify (1) in what setting, (2) under what conditions, and (3) for which specific section of the population the evidence was derived, to then understand the implications for

subsequent implementation. These context and intervention-elements data have not previously been summarised, but have important implications for reach and uptake.

This current review therefore aimed to (1) update efficacy findings from a previous Cochrane Review of psychological interventions for women with MBC, to include any missed RCTs, or RCTs published in the subsequent 5 years, (2) summarise outcomes by treatment modality, and (3) synthesise the following intervention-duration and trial context data: rates of uptake (defined as the number of consenting participants as a proportion of approached eligible individuals, eg, Brebach et al²²), adherence (the amount of an intervention an individual engages with or completes²³), the geographic and medical settings of trials conducted, and the demographic profile of included participants.

2 | METHOD

2.1 | Methodological framework

The preferred reporting items for systematic reviews and meta-analyses (PRISMA) statement was used,²⁴ and a review protocol was developed (Data S1).

2.2 | Data sources

Two methods of identifying primary studies were used. First, one author (LB) conducted 4 electronic bibliographic database searches (from each database’s inception to May 2016): MEDLINE, PsycInfo, CINAHL, and SCOPUS. A list of keywords and MeSH terms was generated to identify studies, falling within 3 key search strategies (for full details, see Data S1—review protocol):

1. *Terms related to metastatic breast cancer:* “metasta*,” “secondary,” “advanced,” “breast cancer”;
2. *Terms related to psychological intervention:* “psycho*,” “intervention,” “psychotherapy,” “group support*,” “self-help,” “internet*,” “tele-counsel*”;
3. *Terms related to outcomes:* “survival,” “quality of life,” “psychosocial.”

This initial search identified 1151 citations (740 after deduplication). The reference lists of the included primary studies and previous systematic reviews were then screened to identify further articles. The full electronic search strategy used for one database (PsycINFO) is included in Data S2. Data S3 summarises the search and retrieval process used.

2.3 | Study selection

The title and abstract of each citation were examined against the following prespecified inclusion and exclusion criteria:

1. The article was published, or in-press, in a peer-reviewed journal between database inception and May 25, 2016. Grey literature was excluded.

2. The intervention targeted adult women with a diagnosis of metastatic or recurrent breast cancer.
3. Studies including women with MBC as part of a larger sample of patients with other cancer types or stages were eligible for inclusion if data could be extracted specific to women with MBC. There were no restrictions on the minimum MBC sample size required.
4. The intervention was psychotherapeutic in nature (ie, provided psychological strategies, of any therapeutic framework, to manage illness-related challenges). Interventions providing information/education without a therapeutic component were excluded.
5. Survival, distress, coping, quality of life, or somatic symptoms was an outcome.
6. The study was a randomised controlled trial, quasi-randomised trial, or feasibility RCT.
7. The article was published in English.

Initial assessment of eligibility was performed by EK based on titles and abstracts. Database searches were then replicated by one other author (LB or BK). There was 97.6% agreement ($\kappa = 0.767$) on the selected review articles. Disagreement in selection was resolved through discussion between authors. Full texts of citations meeting initial inclusion criteria were obtained. All full texts were independently examined by 2 authors (LB and EK) to confirm inclusion/exclusion, with 100% agreement obtained.

2.4 | Data extraction

Data were extracted independently by 2 members of the authorship group using a standardised coding sheet, based on the Cochrane library's recommendations for data extraction.²⁵ Where discrepancies occurred in the data extracted by the initial reviewing authors, a third author (LB/EK) reviewed the data, and consensus was reached through discussion. Interventions were coded as either "Group," "Individual," or "Low-Intensity" during extraction and synthesis. Group and individual interventions were stated as such in papers and were delivered in person. To be classified as "Low-Intensity," interventions had to meet 4 criteria drawn from definitions of low-intensity therapies^{20,21}: (1) administered in a non-face-to-face setting, (2) delivered by non-mental health practitioners, (3) be more accessible, and (4) be briefer for the practitioner compared to therapist-delivered treatments.

2.5 | Quality assessment

The quality of each paper was assessed by 2 authors independently: the *research design* quality was assessed using the 5 criteria for empirically supported psychotherapies²⁶; and the quality of *reporting* was secondly assessed at the study level using 6 criteria from the Cochrane Collaboration's "risk of bias" tool.²⁵

2.6 | Data synthesis

A narrative synthesis of results was used due to the substantial heterogeneity in study design, population, type of intervention, outcomes,

and timing of assessments. The first author synthesised results, with findings verified by the authorship group.

3 | RESULTS

3.1 | Flow of studies through the review

From the initial pool of 740 screened titles and abstracts, 47 citations met inclusion criteria for full review. Two potential studies were identified from reference lists of the initial full-texts screen. Of these 49 total citations, 26 were subsequently excluded during full-text analysis, because MBC-specific patient data could not be extracted from larger samples of mixed cancer types ($n = 14$); the study was not an RCT ($n = 6$); the intervention was not psychotherapeutic ($n = 4$); or the publication did not evaluate efficacy ($n = 2$). Therefore, 23 articles (from 15 RCTs) were included.²⁷⁻⁴⁹ Five RCTs not summarised by the previous Cochrane review were identified and included in this review; 4 were published prior to the previous Cochrane Review update.^{28,37,41,48} Of note, only one new RCT was published in the past 5 years.⁴³

3.2 | Overview of included studies

Data S4 summarises the characteristics of the 15 RCTs reported across 23 articles, with studies listed by treatment-delivery modality. A total of 1638 women participated in the RCTs; 7 interventions (50%) were delivered in a group format^{31-33,38,40,41,45}; 4 in individual therapy^{27,28,44,48}; and 4 as "low-intensity" (telephone counselling, $n = 2$; expressive writing, $n = 2$).^{37,39,42,43} None of the included studies trialled online/Web-based therapy. Most ($n = 13$) interventions fell into 5 therapeutic frameworks: (1) supportive expressive group therapy (SEGT; $n = 4$)^{31,38,40,45}; (2) cognitive behaviour therapy (CBT; $n = 5$)^{28,33,37,44,48}; (3) SEGT/CBT combined ($n = 1$)³²; (4) expressive writing ($n = 2$)^{42,43}; (5) hypnosis⁴¹; with the remaining 2 RCTs trialling telephone counselling,³⁹ and a nurse-lead counselling program.²⁷ Program duration ranged from 80 minutes (expressive writing), to 78 hours (SEGT); treatment providers ranged from none (expressive writing) to 2 highly skilled cotherapists (CBT, SEGT); and the setting was predominantly urban hospitals or clinics ($n = 11/15$).

3.3 | Methodological quality

Data S5 summarises the methodological quality of the included RCTs. Overall, the methodological quality was moderate. While no RCT met all 11 combined criteria, 8 (53%) RCTs met at least 8 of the 11 combined criteria.^{31-33,38-40,43,48} Most of these higher quality studies were group-delivered^{31-33,38,40}; 1 individual therapy⁴⁸ and 2 low-intensity^{39,43} studies met this quality criteria threshold.

Looking at the criteria separately, 6 trials (40%) met all the *research design* criteria.^{31,33,38-40,43} All trials used acceptable control conditions: 8 RCTs used less methodologically rigorous comparators (usual care, $n = 5$)^{27,28,33,39,45}; waitlist control, $n = 3$)^{41,44,48}; while 7 had stronger control conditions (attention-control, $n = 5$)^{31,38,40,42,43}; active-treatment control, $n = 2$)^{32,37}. Most studies ($n = 11$) had sample sizes larger than 25 participants per group; however, only 6 trials reported their power calculations and were sufficiently powered to detect moderate

effects. All except one study⁴⁴ specified inclusion criteria, and only 2 studies did not clearly report or did not use reliable/well-validated measures.^{37,44}

No RCT met all 6 reporting criteria for risk of bias.²⁵ Twelve studies (80%) gave sufficient information about how the allocation sequence was generated, and 9 of these gave sufficient information about whether allocation was adequately concealed. Only 2 studies specifically met the criteria for blinded assessment by research personnel (no trial reported on participant blinding).^{43,48} Strategies to manage incomplete outcome data were reported in 7 (47%) RCTs, while 9 trials were free of selective reporting of outcomes. Finally, only 5 RCTs (33%) were clearly free of other bias, with the most studies being either unclear in their reporting ($n = 7$) or not meeting this criterion ($n = 3$).

3.4 | Trial outcomes

There was marked heterogeneity in outcomes, measures used, and timing of assessments in included trials. Only 7 RCTs specified a primary outcome (group: $n = 4$, individual: $n = 1$, low-intensity: $n = 2$), specifically, survival ($n = 3$), mood disturbance ($n = 2$), quality of life ($n = 1$), and symptom severity response time ($n = 1$). To increase clarity of findings, this review consolidated outcomes into the following domains: (1) survival; (2) distress (incorporating depression, anxiety, stress, cancer-related distress, and intrusive thoughts); (3) coping (incorporating mental adjustment domains such as helplessness, denial, and repression); (4) quality of life; and the commonly reported cancer symptoms of (5) pain (intensity, suffering, sensation, duration, and perceived ability to control); (6) fatigue; and (7) sleeping difficulties. Given the marked heterogeneity of number and timing of assessments, and the analyses conducted, comparing the short- versus long-term benefits of interventions was not possible. This review therefore adopted an inclusive criterion coding an intervention as a positive trial if it yielded a favourable impact on an outcome for at least one of the follow-up assessments. Table 1 summarises the evidence-base per outcome domain, by treatment-delivery modality for the included trials, with an overall (pooled) summary of the evidence-base. This review did not synthesise outcomes analysed in fewer than 2 trials: these included

social support, self-efficacy, and self-esteem; these outcomes were instead reported per individual study in Data S4.

3.4.1 | Survival

Six RCTs investigated survival as an outcome, all delivered as group programs^{32,34,38,40,46,47}; only the first published study⁴⁷ found a statistically significant survival benefit resulting from group participation. Of note, none of the 3 studies that were specifically powered for survival as a primary outcome obtained significant effects.^{32,38,40}

3.4.2 | Distress

Pooling across intervention types, "distress" was the most frequently assessed outcome, investigated in 13 trials^{28,31-33,37-43,45,48}; 8 RCTs demonstrated efficacy for at least 1 distress outcome,^{31,33,37,38,40,41,45,48} of which 7 tested established therapeutic frameworks (ie, SEG, CBT). Breaking results down by treatment modality, Table 1 demonstrates that 6 of the 7 group therapy trials examining distress were positive (86%), including the single trial that specified distress as a primary outcome³¹; one of the 2 individual therapy trials was positive (50%), while 1 low-intensity trial obtained significant effects (25%), with the remaining 3 low-intensity RCTs, including one that specified distress as a primary outcome,³⁹ failing to find effects. Therefore, while the evidence-base for psychological therapy (pooled across treatment modalities) for distress was supported, only group-delivered programs consistently demonstrated positive results.

3.4.3 | Coping

Five trials examined coping domains as an outcome.^{31,32,40,44,45} Four were group-delivered; and all 4 obtained positive effects, therefore reaching criteria for being considered empirically supported. However, findings were significant only for certain domains, helplessness/hopelessness,^{32,40} restraint,³¹ and a "maladaptive coping" subscale,⁴⁵ but not for other commonly reported coping styles including denial, defensive-repressive coping, anxious preoccupation, cognitive avoidance, fatalism, or fighting spirit.^{31,32,40,45} The fifth trial was for individually administered therapy,⁴⁴ where coping was not statistically significantly impacted by the intervention. No trial to date has examined the impact of low-intensity interventions on coping outcomes.

TABLE 1 Outcome summary: number of positive vs negative trials for trial-outcomes, by treatment-delivery modality

Outcome	Number of RCTs	Group			Individual			Low Intensity			RCTs Pooled
		Positive	Negative	Conclusion	Positive	Negative	Conclusion	Positive	Negative	Conclusion	Conclusion
Survival	6	1	5	No support	0	0	n/a	0	0	n/a	No support
Distress	13	6	1	Supported	1	1	Unclear	1	3	No support	Supported
Coping	5	4 ^a	0	Supported	0	1	Unclear	0	0	n/a	Supported
Quality of life	6	1 ^b	2	Unclear	0	3	No support	0	0	n/a	No support
Pain	5	3	0	Supported	1	0	Unclear	0	1	Unclear	Supported
Fatigue	5	1	1	Unclear	0	1	Unclear	0	2	No support	No support
Sleep	4	0	0	n/a	0	1	Unclear	2 ^c	1	Support	Unclear

Abbreviation: RCT, randomised controlled trial.

^aSupported for helplessness, restraint and "maladaptive-coping" subscale; not for denial or defensive-repressive.

^bOnly for social functioning.

^cFor one of these studies, benefits found only among those recently diagnosed.⁴²

3.4.4 | Quality of life

Six RCTs evaluated quality of life: 3 were group programs^{32,38,40}; 3 were individual therapy.^{27,44,48} Only one group therapy trial reported statistically significant effects,⁴⁰ in the specific QOL domain of social functioning, with the remaining 5 RCTs failing to find an effect, including one that specified QOL as a primary outcome.⁴⁴ Psychological interventions for quality of life are therefore currently unsupported, both within delivery modalities and overall.

3.4.5 | Pain

Pain was evaluated in 5 RCTs: 3 group trials,^{29,30,49} 1 individual,²⁸ and 1 low-intensity.³⁷ When pooling across the RCTs, 4 of the 5 psychological interventions trialled had an impact on pain intensity, sensation, suffering, or perceived control over pain.^{28-30,49} Only the low-intensity intervention did not have an impact on pain.³⁷ Overall, the use of psychological interventions, delivered in a face-to-face format (either group or individually), is currently supported; however, the lack of trials for individual and low-intensity modalities preclude definitive recommendations being formed.

3.4.6 | Fatigue

Fatigue was evaluated in 5 RCTs: 2 group interventions,^{29,45} 1 individual therapy,⁴⁸ and 2 low-intensity interventions.^{37,43} Only 1 of the 5 trials, a group intervention,⁴⁵ demonstrated efficacy. Based on the current evidence, there is insufficient data to draw conclusions regarding the use of psychological intervention to treat cancer-related fatigue in MBC.

3.4.7 | Sleeping difficulties

Four trials examined sleep: 1 individual intervention⁴⁸ and 3 low-intensity.^{37,42,43} Two of the 3 low-intensity RCTs found evidence of

benefit^{37,42}; however, for one of these trials, the benefits only applied to those “recently” diagnosed with MBC, defined as within the past 20 months.⁴² The remaining 2 trials failed to obtain significant results for improving sleep. The impact of psychological interventions, both overall or by treatment modality, on sleep-related difficulties therefore remains unclear based on the limited data currently available.

3.5 | Trial context

Table 2 summarises intervention duration and contextual factors associated with trial participation. As can be seen, the overall number of participants and average sample size per trial differed between treatment modalities: Group and low-intensity intervention trials had comparably large sample sizes, with an average sample size per trial of 128 and 129.5 women, respectively, in contrast to the average trial sample size of 54 women for RCTs of individually administered treatments.

3.6 | Participant characteristics

As Table 2 indicates, participant characteristics were not reported uniformly. Of the available data, most studies included women who were young (mean age = 53.08 y), less than 2 years postdiagnosis of MBC at study entry (pooled time since diagnosis = 1.96 y), partnered/married (68.4% of included study participants), English speaking (86.7% of studies), and Caucasian (89.8% of included studies). While 62.2% of participants overall were tertiary educated, only 31.9% of participants were employed. While participant characteristics in general did not differ between treatment modalities, a notable exception was education level: based on the 8 studies reporting education, only 47.3% of group participants were tertiary educated, compared with 58.5% and 80.9% of individual and low-intensity participants, respectively. Group participants were nearly 5 years younger than those in individual therapy,

TABLE 2 Trial context factors and intervention duration, by treatment delivery modality

Outcomes	Group (n = 7)	Individual (n = 4)	Low Intensity (n = 4)	Pooled (n = 15)
Trial characteristics				
Sample size (SS) average SS per trial	N = 900 (range, 37-235) M = 128	N = 217 (range, 24-205) M = 54.3	N = 518 (range, 47-308) M = 129.5	N = 1638 (range, 24-308) M = 109
Participant characteristics				
Age, years (n RCTs)	51.60 (N = 6)	56.29 (N = 3)	54.73 (N = 4)	53.08 (N = 13)
Years since MBC diagnosis (n RCTs)	2.50 (N = 4)	0.75 (N = 3)	2.64 (N = 3)	1.96 (N = 10)
English language (n RCTs)	100% (N = 7)	50% (N = 4)	100% (N = 4)	86.7% (N = 13)
Caucasian ethnicity (n RCTs)	85.5% (N = 2)	98.0% (N = 2)	86.0% (N = 4)	89.8% (N = 8)
Married (n RCTs)	67.5% (N = 6)	66.3% (N = 3)	71.3% (N = 3)	68.4% (N = 12)
Tertiary educated (n RCTs)	47.3% (N = 2)	58.5% (N = 3)	80.9% (N = 3)	62.2% (N = 8)
Employed (n RCTs)	36% (N = 2)	31.3% (N = 2)	28.5% (N = 2)	31.9% (N = 6)
Intervention characteristics				
Uptake (n RCTs; range)	41.9% (N = 6; 19%-79%)	60.5% (N = 2; 60%-61%)	79.1% (N = 2; 69%-89%)	53.02 (N = 10; 19%-89%)
Adherence (n RCTs; range)	66.1% (N = 7; 43%-82%)	62.6% (N = 4; 40%-100%)	85.9% (N = 4; 80%-96%)	69.3% (N = 15; 40%-100%)
Intervention average duration (range)	60 h (16-78)	6.83 h (2-16.5) ^a	3.79 h (1.33-6.5)	32.6 h (1.33-78)

Abbreviation: RCT, randomised controlled trial.

^aOne study did not report intervention duration, beyond stating sessions were unlimited (range not reported). Therefore, calculation based on N = 3 trials with available data. Attrition calculated from treatment or posttreatment attrition rates, where reported, it does not include follow-up assessment attrition.

and 3 years younger than low-intensity intervention participants, while women in individual therapy were more recently diagnosed than the other 2 modalities.

3.6.1 | Intervention characteristics

As Table 2 summarises, group interventions had the highest participation-burden/commitment, as measured by the maximum number of hours of treatment a participant might receive. Group interventions had an average intervention duration of 60 hours (ranging from 16 to 78 h). This was notably higher than in individually delivered (6.83 h), and low-intensity (3.79 h) interventions, and nearly double the pooled interventions' average duration of 32.6 hours.

Only 10 trials^{27,32,33,38,40-43,45,48} summarised uptake data, while all 15 trials reported on adherence. As the number of intervention-hours decreased, the uptake rate increased: uptake was highest at 71.9% amongst low-intensity interventions (range, 69%-89%) and lowest at 41.9% for group delivery formats (range, 19%-79%), with individually administered therapy falling in between (60.5%; range 60%-61%). As Table 2 demonstrates, a similar pattern was observed for adherence, with the exception that individual therapy had lower adherence than group programs.

4 | DISCUSSION

Since 1980, only 15 trials (published across 23 articles) using rigorous randomised controlled designs have examined psychotherapeutic interventions for MBC. Only one new study has been published in this area in the last 5 years, and no studies of Web-based interventions have been published. This was contrary to expectations, given that research and service provision for women with MBC has increasingly been recognised as a priority area,¹⁹ combined with the interest in Web-based interventions for cancer.⁹⁻¹⁶

No evidence of a survival benefit was found in the present review; while this contrasts with the 1-year survival benefit found in the earlier Cochrane Review,⁸ the present review examined overall survival, rather than separating 1-year versus 5-year survival rates, and was therefore consistent with the Cochrane Review finding of no survival benefit at 5 years. This finding was unsurprising, given that only one additional RCT was identified that examined this relationship.⁴⁴ This study had a very small sample size ($n = 43$), did not have survival as its primary aim, and was underpowered for such an analysis; furthermore, the final assessment period was only 6 months after study entry, therefore too short to examine survival advantage adequately. While the mechanism for increasing survival is plausible, given the postulated relationship between depression and reduced survival^{4,50}; findings indicating stress may lead to cancer progression⁵¹; that social support/marital status can have as powerful an impact on survival as chemotherapy⁵²; and that depression reduces adherence to medical treatments,⁵³ to date none of the interventions trialled appear to have yielded large enough effects to impact on these mechanisms. A trial examining whether treating depression among those most vulnerable—such as socially isolated individuals—can improve survival would be beneficial. While promising, such a study would be difficult to power

and conduct, given the large sample size and very long-term follow-up required.

Consistent with earlier reviews, this review did obtain evidence to support the use of psychological therapies overall (pooling across delivery modality) for distress, select coping domains (helplessness/hopelessness and restraint-coping), and pain. This positive impact on distress is particularly notable, given that for all included studies, except one,⁴⁸ elevated distress was *not* an inclusion criteria, therefore floor effects may have been operating. For the remaining psychosocial outcomes, namely, quality of life, fatigue, and sleep, the evidence was either insufficient, or too mixed to draw conclusions, indicating that further research on the use of psychological therapies to improve these outcomes in women with MBC is warranted.

When summarising outcomes by treatment modality, this study indicates that the evidence-base for psychological therapies has been largely generated by group therapy studies, which were the most intensive (ie, weekly for up to a year). Individual interventions did not meet criteria for a supported intervention for any outcome; however, as most outcomes were only investigated in single studies, further research in this area is required to clarify findings. Only tentative support was found for low-intensity interventions regarding one outcome, sleep; no support was found for fatigue and distress; and an unclear evidence-base was noted for pain. Therefore, at this stage, both individual and low-intensity interventions have insufficient evidence to form conclusions regarding efficacy or to make recommendations for their use, and it remains premature to directly compare group interventions with other modalities of support. While groups were most efficacious, the program efficacy must be balanced against cost-effectiveness considerations that have not been explicitly summarised.

While it was outside the scope of the present review to comprehensively summarise outcomes by specific therapeutic frameworks, it was noted that 7 of the 8 positive studies for distress used either SEGT or CBT.^{31,33,37,38,40,45,48} Whilst the study combining CBT with SEGT reported only one significant result, improved helplessness/hopelessness,³² it should be noted that this RCT used an active treatment (self-help CBT) as the comparator. While it is possible that this self-help (low-intensity) CBT comparator provided equivalent benefit to more intensive psychological treatments, this was not tested, as the trial was not a non-inferiority study; head-to-head trials comparing efficacy across treatment modalities could be usefully examined in future research. While low-intensity interventions did not have a sufficient evidence-base to recommend dissemination from the current review, this may be due to the fact that 3 of the 4 included low-intensity RCTs did not test established therapeutic frameworks. Indeed, the only low-intensity intervention testing an established framework (CBT) obtained more significant results than the other 3 low-intensity interventions that trialled other therapeutic frameworks.³⁷

While group programs clearly had the strongest empirical base for their use, it is notable that no new group-based RCTs have emerged since the last Cochrane Review on this topic. A number of reasons likely exist for this, including reduced research funding, general debate over the efficacy and acceptability of psychological interventions for cancer patients, and hence the value of further research on such interventions (eg, Lepore and Coyne⁵⁴ and Manne and Andrykowski⁵⁵). However, it is important to consider that the burden of participation

and high-cost/resource use are significant barriers to uptake of group interventions. Indeed, this study highlighted that groups required the highest levels of commitment, with an average of 60 hours of participation, nearly 10 times the commitment required for individual therapy and 15 times greater than low-intensity interventions. Overall, as intervention duration increased, uptake and adherence decreased. Disparities between those who took up interventions and those who declined were also discernible from this review: across the included studies that reported demographic characteristics, there was a general pattern for young, married/partnered, Caucasian, English-speaking women to be over-represented. Eligibility criteria for most studies also required women to live within close proximity of the Cancer Centre where the studies were being conducted (for face-to-face interventions), therefore reducing the likelihood of rural/regional women participating. Thus, given the high level of commitment required for group therapy, restrictions in uptake on rural/regional women, low adherence rates for individual therapies and higher adherence rates for low-intensity therapies found in this review, one way in which to address the need for accessible low-burden interventions may be to explore low-intensity interventions *based on established therapeutic frameworks*, such as SEG, CBT, or acceptance and commitment therapy (ACT). Moreover, given the success of Web-based interventions as a specific example of low-intensity intervention in treating distress related to a range of chronic physical health conditions,¹⁷ online interventions based on therapeutic frameworks may be the most obvious choice in balancing the need for efficacy with uptake and adherence in interventions available to women with MBC. This could also offer the benefit of providing online social support, as many online programs incorporate forums and can be run in group formats.

4.1 | Methodological quality

None of the studies met all quality criteria, suggesting need for improvements in the methodological quality of research design and reporting. This does not imply that the studies were of poor quality; rather, that reporting was not transparent and many criteria could not be determined from the available information. It should be noted that over the 35-year timeframe that these studies were conducted, the reporting criteria required by journals have markedly changed, with methods developed to improve the quality of *reporting* of trials such as the CONSORT statement and checklist.⁵⁶ The substantial heterogeneity of research design, analytic strategy, comparator groups, methodological quality, sample sizes, and outcome measures used further increases the challenges arising when evaluating efficacy. For example, only 7 of the 15 included RCTs specified a primary outcome; even when stated, these primary outcomes varied markedly across survival and psychosocial outcomes so it was not possible to summarise efficacy by primary outcomes. Methods for improving the quality of *design* could therefore include standardising the range of outcome measures used, specifying and powering to a primary outcome, and using treatment-as-usual or appropriate placebo-controls rather than waitlist or active-treatment comparators. As prior reviews have noted, waitlist designs are problematic as control-participants anticipate future therapeutic input and are therefore less likely to seek constructive action compared with other control-group types.⁵⁷ While active-comparators

are methodologically strong, they are generally only adopted in settings where strong evidence-based gold-standard treatments exist and non-inferiority designs are appropriate. Given the overall paucity of psychological intervention trials in MBC, it is premature to be testing non-inferiority, and efforts should instead be directed to building the evidence-base for psychological treatments. Two of the included studies used an active comparator.^{32,37} Finally, ensuring that sample sizes are sufficient to meet power requirements for statistical analysis would greatly benefit this area, as only 40% of included RCTs were adequately powered.

4.2 | Limitations and future directions

The current review has several limitations. First, the effect sizes of outcomes were not summarised, as these data were not routinely or consistently reported across trials. Second, studies often included multiple follow-up assessment points; however, these points differed immensely between studies, or assessments were aggregated into an average/composite follow-up score making it difficult to summarise outcomes by time-periods. While all available data was extracted, this review elected to summarise the overall impact on psychosocial domains (that is, whether the intervention led to a significant group difference at any of the assessments), rather than differentiating the short- versus long-term impact, or changes over time within outcomes. However, for many of the included studies, psychosocial effects were short-term/not sustained or required a lengthy course of psychological treatment to obtain effects (eg, a full year of SEG to obtain statistically significant group differences). Third, in addition to the changes in trial reporting requirements over the 35-year review period, the oncology landscape for treating metastatic breast cancer (including average survival time, side effects, and resulting quality of life) has also changed significantly, with the advent of new systemic treatments.² Therefore, any conclusions made when comparing psychological interventions delivered across this broad time-frame should be made with caution. Likewise, the discipline of psycho-oncology—particularly our understanding of the mechanisms through which psychological interventions lead to change—has also matured over that time-period. While SEG and CBT remain the gold-standard treatments, more emphasis in recent times has been placed on the role of perseverative thinking (worry and rumination), attentional biases, and metacognitive processes as maintenance factors for psychological morbidity.⁵⁸ None of the included studies have examined interventions that specifically target these factors, such as the “third-wave” psychological intervention strategies (metacognitive or mindfulness-based therapies), despite a recent Breast Cancer Gap Analysis identifying this as a priority for research.⁵⁹ Given the promising evidence emerging for these new intervention types in non-metastatic populations⁶⁰ and in metastatic/advanced disease populations that are not breast cancer-specific,^{61–63} this is a useful avenue for future research. Finally, it should be noted that grey literature was excluded from this review, as there is a high discordance in the abstract-to-publication rates and quality; grey literature often only reports preliminary results that are unrepresentative of final observations once all data is collected and analysed,⁶⁴ with results changing in up to 42% of the abstract-to-publication

comparisons.⁶⁵ Given this, only final fully published (and data-extractable) peer-reviewed articles were eligible for inclusion.

4.3 | Summary

In summary, this review confirms the paucity of research addressing the unmet psychological needs of women with MBC and highlights the challenge of balancing efficacious treatments against accessibility, reduced burden, and increased acceptability. There is a lack of research evidence on efficacy of interventions in women of diverse ethnic background, socioeconomic status, and broader age ranges. It is paramount to ensure interventions reach, and meet the needs of, a broad range of women with MBC, while simultaneously providing evidence-based benefits. While psychological therapies had consistent support for their use in improving distress, aspects of coping, and pain, when separated by treatment modality, this evidence is only applicable to group-delivered programs at this stage; the evidence-base is insufficient for individual therapies and low-intensity interventions. More research is therefore required on how to deliver evidence-based interventions in an accessible low-burden format.

ACKNOWLEDGEMENTS

Dr Beatty is funded from an NHMRC project grant (GNT 1042942). Prof Girgis is funded from a Cancer Institute NSW grant. The authors would like to thank Dr Addie Wootten for contributing to the design of this systematic review.

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How to cite this article: Beatty L, Kemp E, Butow P, et al. A systematic review of psychotherapeutic interventions for women with metastatic breast cancer: Context matters. *Psycho-Oncology*. 2017;0:1-9. <https://doi.org/10.1002/pon.4445>