Integrating culturally informed approaches into physiotherapy assessment and treatment of chronic pain: a pilot randomised controlled trial

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ABSTRACT

Objective To evaluate patient engagement with, and the feasibility of, a novel, culturally adapted physiotherapy pain management approach.

Design A participant-blinded and assessor-blinded pilot randomised controlled trial.

Setting Outpatient physiotherapy departments at two public hospitals and one district pain clinic.

Participants Adults (n=48) with chronic musculoskeletal pain (daily pain >3 months), who self-identified as Mandaeans, Assyrians or Vietnamese, were randomised to one of two physiotherapy treatment conditions.

Interventions 24 participants underwent combined group and individualised treatment described as ‘culturally adapted physiotherapy’, while 24 underwent evidence-informed ‘usual physiotherapy care’. Both treatment arms consisted of up to 10 sessions over a 3-month period.

Outcome measures Patient engagement was measured via participant attendance, adherence and satisfaction data. Secondary outcomes included clinical measures of pain severity, interference and suffering, physical function and negative emotional state.

Results 96% of participants undergoing culturally adapted physiotherapy completed treatment, compared with 58% of the usual physiotherapy group. For the culturally adapted group attendance (87%±18%) and adherence (68%±32%) were higher relative to usual care (68%±32% and 55%±43%). Satisfaction was similar for the culturally adapted (82.7%±13.4%) and usual care (79.3%±17.3) groups. For secondary outcomes, a significant between-group effect for pain-related suffering in favour of the culturally adapted group was observed with a medium effect size (partial $\eta^2$ of 0.086, mean 3.56, 95% CI 0.11 to 7), while results for pain severity, interference, physical function and negative emotional state were similar.

Conclusions Aligning treatment with the beliefs and values of culturally and linguistically diverse communities enhances patient engagement with physiotherapy. These results support the feasibility of a larger, multisite trial to determine if improved engagement with culturally adapted physiotherapy translates to improved clinical outcomes.

Trial registration number ACTRN12616000857404; Pre-results.

Strengths and limitations of this study

- This was a randomised, assessor-blinded and participant-blinded controlled trial.
- It provides evidence of feasibility of culturally adapted physiotherapy approaches for pain management as explored with three culturally and linguistically diverse communities.
- Observed recruitment rates, follow-up rates and preliminary data can inform a future fully powered randomised controlled trial.
- As a pilot study, analyses of clinical outcomes are exploratory.

INTRODUCTION

Patient engagement is paramount for the delivery of efficient and effective healthcare, reflecting a patients’ relationship with the health encounter, such that they participate (attends and adheres) and recognise value in their treatment (satisfaction and treatment completion). Research that has evaluated interventions and models of care to enhance patient engagement has provided evidence of success. Whether this is true for culturally and linguistically diverse (CALD) communities remains uncertain. This is problematic because healthcare must be responsive to the comparatively poorer health status observed in many CALD communities. Further, strategies promoting engagement tailored to the needs of CALD communities is vital, particularly given that many countries around the world are now culturally plural societies.

Culturally adapted approaches have been suggested to be an effective strategy to enhance patient engagement and reduce health disparities in CALD communities. Such approaches speak to more equitable health outcomes for diverse cultures by minimising the risk of a model that results in more favourable outcomes for the dominant, hegemonic culture. Systematic reviews...
and meta-analyses support the use of culturally adapted treatment for mental health conditions, chronic disease management, cancer screening and health promotion.\(^4\)\(^5\)\(^6\)\(^9\) For example, meta-analyses of mental health interventions demonstrated small to large pooled effect sizes in favour of culturally adapted treatments, compared with usual care.\(^5\)\(^6\)\(^9\) Despite evidence supporting the use of culturally adaptive approaches, research is still lacking for many prominent, debilitating conditions, including chronic pain.\(^10\) As such, suboptimal health outcomes continue to be observed in patients from CALD communities with chronic pain.

Chronic pain disorders contribute to considerable societal burden and personal suffering.\(^11\) Many physiotherapy interventions for chronic pain, particularly exercise-based approaches, are safe and effective.\(^12\)\(^13\) Current evidence-based recommendations suggest that exercise, when combined with cognitive–behavioural and psychosocial treatments, reduces pain, improves quality of life and reduces long-term disability.\(^12\)\(^14\) However, the efficacy of these approaches has been established in populations speaking the same language, with few studies including CALD and migrant communities.\(^10\) The limited research inclusive of CALD communities suggests limited efficacy for pain, quality of life and psychological health outcomes.\(^10\) Such uncertainty supports investigation of sociocultural factors that could influence implementation of pain management approaches within CALD communities.\(^15\)

Successful management of chronic pain requires a strong therapeutic alliance and patient acceptance of, and engagement with, treatment concepts.\(^16\)\(^17\) Unfortunately, engagement with activity-based treatments is often suboptimal in CALD communities, evidenced by lower attendance, reduced acceptance and premature drop-out from treatment.\(^10\)\(^18\) Discordant expectations, low patient–provider alliance, cultural–spiritual factors and communication problems have been cited as contributors to suboptimal engagement for CALD communities.\(^19\)\(^20\) This is perhaps not surprising in the context of intercultural encounters where there is evidence of healthcare provider ethnocentrism, implicit and explicit bias towards patients from CALD backgrounds.\(^21\)\(^22\)\(^23\) Since engagement with treatment underpins improved patient outcomes,\(^24\) it is imperative that strategies are implemented to optimise engagement by CALD populations for costly and debilitating conditions, such as chronic pain.

Thus, the aim of this pilot study was to determine the feasibility, patient engagement and trends of clinical effectiveness of a culturally adapted physiotherapy assessment and treatment approach compared with evidence-informed ‘usual physiotherapy care’. Thus, the research questions for this pilot randomised trial were:

1. Is a 12-week culturally adapted treatment approach superior to ‘usual physiotherapy care’, in terms of patient engagement (adherence, attendance and satisfaction)?

2. Is it feasible to deliver and evaluate culturally adapted physiotherapy assessment and treatment approaches across three CALD communities using a randomised controlled trial (RCT) design?

**METHODS**

**Design**

This was a prospective, multicentre pilot RCT with concealed allocation, and participant and assessor blinding, using a patient sample with chronic pain drawn from three CALD communities (Mandaean, Assyrian and Vietnamese). The trial was conducted across two hospital-based physiotherapy departments and one district pain clinic, between July 2016 and June 2017. A study protocol with eligibility criteria and intervention descriptions was published previously.\(^22\) The study was registered with the Australian and New Zealand Clinical Trials Registry (ACTRN12616000857404).

**Participants and recruitment**

This pilot RCT was the culmination of 3 years of engagement with local Assyrian, Mandaean and Vietnamese communities, facilitated by the multicultural health unit in South-West Sydney Local Health District. Bilingual community educators and multicultural health workers informed the development of the intervention in earlier qualitative phases\(^15\) and guided processes in this RCT, ensuring the research team were cognisant of the communities needs and vulnerabilities.

Following consultation with multicultural representatives, it was evident that a broad recruitment strategy was required to be inclusive. This included: (1) recognising the complexity of chronic pain in each community by not excluding participants based on pain location (such as only including low back pain) or psychological comorbidity; (2) considering patients from multiple countries of birth (Iraq, Iran, Syria, Turkey, Jordan and Vietnam) and anyone speaking Arabic, Assyrian or Vietnamese as potentially eligible, especially where data on ethnocultural identification were not available. Ethnocultural identification was then established according to self-identification by the participant at the screening assessment.

A total of 94 participants were assessed for eligibility by a physiotherapist not involved in the delivery of interventions and who was bilingual or used the services of an accredited health language interpreter. While a multicultural community representative was not present during recruitment, participants were offered the opportunity to consult community representatives and family members before consenting to participation. This resulted in 48 participants randomised into the study. Inclusion criteria were: adult (≥18 years), non-specific musculoskeletal pain, daily pain of greater than 3 months’ duration, self-identification as a member of the Mandaean, Assyrian or Vietnamese ethnocultural communities, and ability to provide written informed consent in their own language or English. Exclusion criteria were: specific diagnoses...
necessitating other treatment (i.e., complex regional pain syndrome), surgery within the last 3 months and assistance for mobility other than a walking stick, to ensure safety during a group or home-based exercise programme.

Sixteen participants from each community were allocated randomly to the experimental or control group after baseline assessment (Figure 1). Group allocation was determined by a computer-generated sequence with a 1:1 allocation ratio, with each ethnocultural community randomised separately. An independent person prepared sealed opaque envelopes containing the intervention arm, labelled with a participant number according to their entrance sequences. These envelopes were managed securely by a central administrative officer responsible for randomising participants and arranging relevant appointments once a participant had been consented. Participants were blind to treatment allocation and were told the trial was comparing two physiotherapy approaches for chronic pain and it was unknown which was more effective. Thus, participants were unaware that they were receiving culturally adapted treatment approaches for the experimental groups. The success of blinding was assessed at the 3-month reassessment with the question; ‘Do you think your physiotherapist has been trained in culturally responsive treatments for chronic pain?’.

Figure 1 Flow diagram of the study following consolidated standards of reporting trial guidelines.
Intervention
Participants from intervention and control groups attended for a maximum of 10 sessions of physiotherapy over a 3-month treatment period. A maximum of ‘10’ sessions was selected to enable the treating physiotherapist to tailor interventions to the individual needs of participants, and was consistent with the average number of physiotherapy sessions reported in clinical trials for the management of chronic pain. All participants were given a home exercise programme designed by their physiotherapist, and they were provided with translated log-books to facilitate recording of exercise adherence. A professional health interpreter was available for all treatment sessions (group and individual), if required, in accordance with best practice.

Culturally adapted physiotherapy assessment and treatment
Participants received a combination of group and individual physiotherapy sessions, adapted to reflect the ethnocultural beliefs and values of the community to which the participant identified. Three ethnocultural-specific group programmes were designed by the research team, informed by qualitative research involving each community and guided by two adaptation frameworks. Sessions were delivered once per week for 6 weeks, included a combination of education and exercise, and were conducted in groups of eight participants from the same ethnocultural community. Sessions were run by a physiotherapist at a local community facility, and facilitated by a bilingual educator in the language of participants. In addition, group sessions were supplemented by up to four individual sessions tailored to the participant according to the culturally informed initial assessment to ensure consistency with the dose of the control group. Components of the cultural adaptation for each ethnocultural community have been previously published and a summary is presented in online supplementary appendix 1.

Evidence-informed ‘usual physiotherapy care’
Participants allocated to this condition attended physiotherapy in the outpatient department where they were referred, for treatment informed by evidence-based recommendations for chronic pain. All treating physiotherapists underwent a training session to familiarise them with evidence-based management of chronic pain. Treatment adherence to these guidelines was monitored by review of therapist treatment logs. Treating physiotherapists used their clinical judgement to guide the specifics of treatment according to principles of patient-centred care. Following the initial assessment, physiotherapists worked with patients to select the treatment mode (individual or group based), frequency and dose (to a maximum of 10 sessions) tailored to the patient’s needs and goals, consistent with best available evidence. It is of note that a substantial proportion of research examining the impact of interventions on chronic pain had excluded patients from CALD backgrounds.

Outcomes
Trained assessors, not involved in the recruitment or treatment of participants and unaware of group assignment, performed assessments according to standardised instructions at baseline (month 0) and (3-month reassessment). Success of assessor blinding was determined with the question; ‘Did you know to which treatment arm the participant belonged?’ If an assessor responded ‘yes’, they were asked to nominate ‘to which group’.

Data to assess feasibility were collected throughout the trial period regarding recruitment rates, treatment withdrawals, therapist fidelity to evidence-based guidelines, success of participant and assessor blinding, and trial drop-outs. Primary outcome measures were: measures of patient engagement, defined by attendance, and adherence to, and satisfaction with treatment. Attendance was measured as the proportion of sessions attended, relative to the number of sessions scheduled. Adherence was calculated as a percentage of the average number of home exercise sessions completed each week, relative to the number of sessions prescribed, determined from participant log-books or self-report (where the participant was unable or did not complete the log-book). Patient satisfaction with treatment was evaluated using the Client Satisfaction Questionnaire (CSQ-8), which evaluates satisfaction with treatment generally, and was selected because it has been validated in Arabic and Vietnamese.

Secondary outcomes included core measures recommended by the Initiative on Methods, Measurement and Pain Assessment in Clinical Trials. This included measures for pain severity and interference (Brief Pain Inventory: BPI), pain-related suffering (Pictorial Representation of Illness and Self Measure: PRISM), physical function (6 min walk test: 6MWT, and 1 min sit to stand test: STS test) and severity of symptoms for Depression, Anxiety and Stress (DASS-21). The reliability and validity of these measures, including for Arabic and Vietnamese translations, have been reported previously and were documented in the trial protocol.

Patient involvement
The research questions were developed following qualitative enquiry into the experience of chronic pain among CALD communities. Specifically, challenges raised by participants accessing and participating in pain management in South-West Sydney were incorporated in the study design. As such, participant engagement was considered a primary outcome measure. While patients were not involved in the recruitment and conduct of the study, all participants were given the opportunity to attend a feedback session following trial completion, held in local community venues.

Sample size and statistical analysis
A total sample of 48 participants was deemed appropriate to allow the piloting of a novel culturally adapted programme with three communities (eight participants per programme), while ensuring equal numbers in both
treatment arms (24 culturally adapted and 24 usual care) and allowing for the detection of medium to large effects (effect sizes of 0.5–0.8), should they exist.36 37

Descriptive statistics were used to report the characteristics of participants, including means and SDs for continuous variables, and frequencies and proportions (%) for categorical variables. Primary outcome measures (attendance, adherence and satisfaction) were evaluated using descriptive statistics and Mann-Whitney U tests, because data were not normally distributed and transformations did not achieve normality. Effect sizes for non-parametric tests were reported using r and interpreted as large (0.5), medium (0.3) and small (0.1).38

Exploratory examination for group differences in secondary outcome measures was undertaken using a repeated-measures analysis of variance (ANOVA), with the treatment condition (usual care/culturally adapted intervention) as the between-group factor and time of assessment (preintervention or reassessment) as the repeated, within-group factor. One-way repeated measures ANOVAs compared within-group main effects at each time point. Effect sizes were classified as small, medium or large (partial $\eta^2$ 0.01, 0.06, 0.14, respectively).39 If the assumptions of ANOVA were violated, data were transformed to achieve a normal distribution before repeating the ANOVA. Intention-to treat analyses were performed for all participants and missing data were addressed by carrying the last data point forward.40 Analyses were performed using the SPSS, V.24.

RESULTS
Feasibility and treatment characteristics
Forty-eight participants, 16 from each ethnocultural community, were randomised within 4 months (figure 1). For the culturally adapted treatment arm, all group sessions were delivered by the physiotherapist who developed the culturally adapted treatment protocols, according to the session manual and verified by review of the therapist log-book. On average, three individual sessions were recommended to supplement the six group sessions (range 1–4). One participant discontinued treatment prematurely, citing illness. For the usual care arm, 14 participants completed the treatment they were allocated. Ten participants withdrew from treatment citing reasons that included illness (n=1), treatment not helping (n=4), lack of time (n=1) and changed mind/sought care elsewhere (n=4). Treating physiotherapists in the usual care arm used both group and individual modes of delivery for 8/24 participants, while individual therapy alone was recommended for 16 participants. Fidelity was evaluated from logbooks completed by each therapist as the percentage of core treatment components included. The components included pain education, goal setting, activity pacing, active coping strategies, flare-up management and a tailored home exercise programme. For the 14 participants who completed treatment, there was 100% therapist fidelity to six core treatment components while for the other participants, an average of four of the six core components were included prior to drop-out, with flare up management and active coping strategies the most commonly omitted elements. Therapist fidelity to evidence-based principles was confirmed for all participants, except for the two participants who withdrew following their initial assessment.

Blinded reassessment data were available for 45 participants, with 3 participants (usual care group) withdrawing from the trial and declining final assessment for similar reasons: ‘treatment has not helped me’, ‘treatment has not done anything to help my leg pain at all’ and ‘treatment has been a waste of time’. As such, the last data point for each was carried forward for all outcomes except satisfaction, for which an initial data point was not available. Success rates for assessor blinding was 91%, while 44% of participants correctly answered the blinding question regarding their therapists’ cultural responsiveness. No participant experienced an adverse event due to participation in the trial.

Demographic and baseline symptom characteristics of participants are displayed in table 1. There were no significant differences between the groups for baseline characteristics.

Primary outcomes
Attendance
Overall mean (±SD) attendance at physiotherapy was 8.0±3.1 visits. The culturally adapted treatment group attended a higher number of scheduled sessions compared with ‘usual physiotherapy care’ (mean difference=4.0 sessions, 95% CI 2.6 to 5.3). There was an 87% (±18) attendance rate in the culturally adapted programme, compared with 68% (±32) in the usual care group with a medium between group effect size (U=170, $r=0.36$).

Home exercise adherence
Home exercise adherence data were available for all participants in the culturally adapted programme (n=24) and 22 participants from the usual care group. Data were absent for two participants who dropped out after their initial visit. Overall, adherence varied from 0% to 100%. The average number of home exercises prescribed was similar for the culturally adapted (n=7, range 2–10) and usual care group (n=6, range 3–11). Overall, the culturally adapted group had a higher adherence rate (88%±15%) relative to usual physiotherapy care (55%±43%), consistent with a moderate between group effect size (U=145, $r=0.39$).

Satisfaction
Satisfaction data were available for all participants who attended the 3-month blinded assessment (n=45). Overall, 93% of participants were satisfied with treatment. Overall, 71% were highly satisfied, evaluated by a score
of greater than 50% and 75%, respectively, for the CSQ-8. Satisfaction between the two groups did not differ. Mean CSQ-8 scores for the culturally adapted and usual physiotherapy care groups were 82.7 (±13.4) and 79.3 (±17.3).

### Secondary outcomes

Culturally adapted treatment resulted in greater improvements in pain-related suffering than ‘usual physiotherapy care’, with a medium effect size observed (partial $\eta^2$ 0.086) (table 2). A small effect size was observed for between-group difference in favour of the culturally adapted group for BPI pain interference (partial $\eta^2$ 0.02) and 6MWT (partial $\eta^2$ 0.053), while no effect was observed for BPI pain severity, STS test or the DASS-21 (table 2).

### Sample size estimates

With respect to feasibility for a larger trial based on trial data, for power of 80%, alpha of 5% and a drop-out rate of 20%, a sample size of 124 in each group would be required to detect a clinically significant difference of 50m for walking distance\(^{42}\) for the intervention group, based on the SD observed in our study of 128m. This sample size would also be sufficient to identify between-group differences for the BPI Severity (2.2-point difference, SD 2.51) and interference subscales (2.2-point difference, SD 2.55), the PRISM suffering score (3.3cm difference, SD 8.46) and the DASS total score (13-point difference, SD 31.88). A sample size of 300 would also allow for clinically important between-group differences to be detected for the 1 min STS test (3 repetition difference, SD 8.46).

### DISCUSSION

The culturally adapted programme was designed to target-specific language, cultural and access barriers faced by CALD communities that participate in pain management treatments. Results from this pilot study suggest there is an advantage in favour of a culturally adapted physiotherapy programme relative to usual physiotherapy care for addressing barriers to optimal patient engagement. The culturally adapted programmes were well received by all three communities, demonstrated by significantly higher patient engagement (attendance, completion of treatment and adherence) compared with the usual care group. While specific conclusions regarding the efficacy of treatment for clinical outcomes cannot be made, the moderate to small effect sizes observed for the secondary outcomes of pain-related suffering, pain interference and physical function warrant further investigation. Recent systematic reviews of multidisciplinary and exercise-based treatments for chronic pain have revealed pooled effect sizes that are small for function and disability, while pain and psychological health were associated with small effect sizes or no effect, depending on whether care was interdisciplinary or single disciplinary.\(^{43-45}\) In the context of such evidence, the results of this trial support further research into cultural adaptation to maximise the effect on pain and psychological outcomes.

Attendance and treatment retention is an important aspect of patient engagement essential to ensure positive outcomes from cognitive–behavioural and exercise treatments for chronic pain are realised.\(^{16, 17}\) Despite this, drop-out from pain management programmes has been reported to be as high as 40%,\(^{46}\) while for exercise-based physiotherapy, drop-out rates of 30%–40% are common.\(^{47, 48}\) In the current study, drop-out rates in

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Participant baseline demographic and symptom characteristics</th>
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<tbody>
<tr>
<td></td>
<td>Culturally adapted (n=24)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>55 (10.0)</td>
</tr>
<tr>
<td>Sex, (n) male:female</td>
<td>5:19</td>
</tr>
<tr>
<td>Length of time in Australia, years</td>
<td>15.5 (12.9)</td>
</tr>
<tr>
<td>Migration circumstances</td>
<td></td>
</tr>
<tr>
<td>Voluntary migrant, n (%)</td>
<td>6 (25)</td>
</tr>
<tr>
<td>Refugee, n (%)</td>
<td>18 (75)</td>
</tr>
<tr>
<td>Marital status—married, n (%)</td>
<td>16 (67)</td>
</tr>
<tr>
<td>Level of education</td>
<td></td>
</tr>
<tr>
<td>No school or primary, n (%)</td>
<td>9 (38)</td>
</tr>
<tr>
<td>Secondary, n (%)</td>
<td>13 (54)</td>
</tr>
<tr>
<td>Tertiary, n (%)</td>
<td>2 (8)</td>
</tr>
<tr>
<td>Duration of pain (years)</td>
<td>10.0 (7.9)</td>
</tr>
<tr>
<td>Work status</td>
<td></td>
</tr>
<tr>
<td>Full-time or part-time work, n (%)</td>
<td>1 (4)</td>
</tr>
<tr>
<td>Unemployed due to pain, n (%)</td>
<td>18 (75)</td>
</tr>
<tr>
<td>Retired, n (%)</td>
<td>2 (8)</td>
</tr>
<tr>
<td>Other, n (%)</td>
<td>3 (13)</td>
</tr>
<tr>
<td>Receiving pension or benefit, n (%)</td>
<td>23 (96)</td>
</tr>
<tr>
<td>Mean classes of pain medication/5</td>
<td>2.08 (0.78)</td>
</tr>
<tr>
<td>BPI (pain severity)/10</td>
<td>7.3 (1.8)</td>
</tr>
<tr>
<td>BPI (pain interference)/10</td>
<td>7.7 (1.6)</td>
</tr>
<tr>
<td>DASS subscores/42</td>
<td></td>
</tr>
<tr>
<td>Depression</td>
<td>27.6 (12.2)</td>
</tr>
<tr>
<td>Anxiety</td>
<td>23.9 (12.4)</td>
</tr>
<tr>
<td>Stress</td>
<td>26.8 (11.4)</td>
</tr>
<tr>
<td>Pain suffering (PRISM)/27</td>
<td>3.4 (5.0)</td>
</tr>
<tr>
<td>6MWT</td>
<td>266.8 (142.3)</td>
</tr>
<tr>
<td>1 min STS test</td>
<td>9.6 (6.5)</td>
</tr>
</tbody>
</table>

Data are presented as mean (±SD) unless otherwise indicated. *Classes included simple analgesics, compound analgesics, anti-inflammatory, anticonvulsant and opioids. 1 min STS test, 1 min sit to stand test; BPI, Brief Pain Inventory; DASS, Depression Anxiety and Stress Scale; 6MWT, 6 min walk test; PRISM, Pictorial Representation of Illness and Self Measure Separation.
the ‘usual physiotherapy care’ group (42%) were consistent with rates observed in the literature, while for the ‘culturally adapted’ group, drop-out was less (4%). Further, attendance at scheduled sessions was higher in the ‘culturally adapted’ group, and participants were willing to attend for a greater number of sessions. In combination, such findings suggest that attention to social and ethnocultural dimensions unique to CALD migrant communities successfully engaged participants. For the culturally adapted group, a combination of both surface-level (language, food, music, group interaction and setting) and deep-level (reframing content to align with explanatory models of pain and ethnocultural values) adaptations were included to enhance the cultural relevance of programme content and facilitate patient engagement. While programmes were conducted in a similar geographical location (ie, suburb) to the usual care group in the hospital outpatient service, the use of a community venue was an important technique for balancing power differentials in therapeutic relationships and reducing access barriers, thereby contributing to engagement outcomes. As such broad multidimensional adaptations should be considered in future research.

Treatment adherence is an aspect of patient engagement that has been positively related to patient outcomes in rehabilitation programmes. Nevertheless, adherence to exercise interventions for chronic pain conditions is suboptimal. For example, adherence rates for osteoarthritis exercise programmes can be as low as 50%, and varies between 64% and 71%, respectively, for neck pain and low back pain. For the current study, there was wide variation in adherence rates for the ‘usual care’ group with a mean of 55% (±43%), while for the ‘culturally adapted’ group, adherence was significantly higher and less variable (88%±15%). Low adherence rates in the ‘usual care’ group could have been due to suboptimal communication, patient–provider interactions, and failure to adequately tailor interventions to the sociocultural needs of the individual patient. A systematic review cited the association between anxiety and depression, highly prominent symptoms in our sample, and low adherence to physiotherapy. However, since both treatment arms experienced similar symptoms, this association alone, does not account for the different adherence rates observed. Similarly, the low adherence rate for the ‘usual care’ group could not be ascribed to language barriers, since both groups had similar access to professional interpreting services and translated exercise diaries. Instead, the current findings emphasise a potential role for physiotherapists to optimise the intercultural therapeutic interaction by attending to a patient’s beliefs and values, and aligning treatment components accordingly.

Baseline outcome data from the three CALD communities highlighted participants’ severe pain and psychological symptoms. Participants had higher mean pain duration, and average pain severity scores, than those observed in cohorts attending multidisciplinary pain clinics. Similarly, average scores for depression, anxiety and stress according to the DASS-21 were all in the ‘severe’ range, and higher than mean scores observed in a large Australian pain clinic cohort. Potentially, such observations were not surprising given 71% of our sample identified as refugees. However, in the context of severe depression, the efficacy of rehabilitation programmes for chronic pain programmes is known to be reduced. As such, the physiotherapy approaches employed in our study might be insufficient to induce meaningful changes in pain and psychosocial functioning. While the individualised design of both treatment arms allowed for the involvement of other specialities, such as psychology, participants did not pursue this recommendation in 85% of cases. Such low uptake, in combination with high pain engagement outcomes.

Table 2: Between-group comparison

<table>
<thead>
<tr>
<th>Outcome</th>
<th>ANOVA time x group</th>
<th>Between-group comparison of change scores culture–usual mean (95% CI)</th>
<th>F (1,46) value</th>
<th>Partial η²</th>
</tr>
</thead>
<tbody>
<tr>
<td>BPI pain severity</td>
<td>−0.14 (−1.25 to 0.97)*</td>
<td>0.063</td>
<td>0.01</td>
<td></td>
</tr>
<tr>
<td>BPI pain interference†</td>
<td>−0.57 (−1.73 to 0.60)*</td>
<td>0.962</td>
<td>0.020</td>
<td></td>
</tr>
<tr>
<td>Pain-self separation†</td>
<td>3.56 (0.11 to 7.0)</td>
<td>4.322</td>
<td>0.086</td>
<td></td>
</tr>
<tr>
<td>6MWT (m)</td>
<td>28.44 (−7.40 to 64.28)</td>
<td>2.551</td>
<td>0.053</td>
<td></td>
</tr>
<tr>
<td>STS test (reps)</td>
<td>1.13 (−2.44 to 4.69)</td>
<td>0.405</td>
<td>0.009</td>
<td></td>
</tr>
<tr>
<td>DASS depression</td>
<td>−2.67 (−9.03 to 3.69)*</td>
<td>0.712</td>
<td>0.015</td>
<td></td>
</tr>
<tr>
<td>DASS anxiety</td>
<td>−2.0 (−8.28 to 4.28)*</td>
<td>0.411</td>
<td>0.009</td>
<td></td>
</tr>
<tr>
<td>DASS stress</td>
<td>0.58 (−4.80 to 5.97)*</td>
<td>0.048</td>
<td>0.001</td>
<td></td>
</tr>
</tbody>
</table>

*Minus score in favour of experimental group.
†Transformed data.
ANOVA, analysis of variance; BPI, Brief Pain Inventory; DASS, Depression, Anxiety and Stress Scale; 6MWT, 6 min walk test; STS, sit to stand test.
and psychological symptom scores, emphasises a need for
treatment adaptations to engage other disciplines and
align comprehensive multidisciplinary approaches with
the beliefs, values and unique needs of diverse ethno-
cultural communities. However, the maintained high
adherence and attendance data for the culturally adapted
group in the presence of high pain scores and psycholog-
ical symptoms was a positive finding.

Feasibility

Previous research involving CALD communities has
identified significant challenges in engagement and retent-
ion in clinical research. Williams et al. enrolled and ran-
domised 78 participants from three CALD backgrounds (Greek, Italian and Vietnamese) living
with chronic disease to a medication self-management
programme and found less than half completed the
post-treatment reassessment (3 months). Similarly, Sver-
issen et al. found a 35% drop-out rate among CALD
communities in Australia enrolled to a chronic disease
self-management programme. Despite this, our expe-
rience supports research inclusive of, and specifically
targeted towards, CALD communities. Our high recruit-
ment rates, short recruitment time, absence of adverse
outcomes and low trial drop-out rate of 6%, support
the feasibility of implementing randomised controlled
research trial designs within CALD communities. Specific
attention should be given towards involvement of bilin-
gual support workers, professional translation and inter-
preting services, and engagement of ethnocultural
community members in trial design and implementa-
tion, to optimise the prospects of the success of our pilot
study. Further, while cost-effectiveness was not a specific
outcome, there were no substantial cost disadvantages of
delivering culturally adapted treatment. Both treatment
arms were delivered by public health outpatient services.
While the cost of hire of community venues was greater
($A1595), this cost was offset by delivering 67% of cultur-
ally adapted treatment in groups. Similarly, there were no
cost disadvantages of engaging a bilingual support worker
in lieu of a health language interpreter, both of which are
funded by different sectors of the public health service.
This provides further support for feasibility. Finally,
sample size estimates using our pilot data inform the
feasibility of a fully powered RCT to evaluate the clinical
effectiveness of culturally adapted approaches, with the
potential to maintain participant engagement.

Study limitations

While the ‘culturally adapted’ programme was success-
fully piloted across the three ethnocultural communi-
ties, it is important to note the study’s limitations. First,
participant adherence data relied on self-report. A log
book was developed to facilitate recording of adher-
ence, but many participants (15/48) had difficulty
completing and/or did not complete the log-book. As
such self-report during sessions was used, and therefore
data could have been compromised by recall error, or
desire to please the treatment provider. This is a chal-
gen for researchers working with CALD communities
who have linguistic limitations, with a need for reli-
able, valid measures for recording patient adherence to
address such issues. Second, some participants with low
education and literacy levels (33% of the sample had
either no or primary-level schooling) were challenged
by the log-book and scale outcome measures, potentially
compromising results. However, the challenge of literacy
was similar for both groups and is unlikely to explain
any between-group differences because all participants
were provided with assistance from the bilingual blinded
assessor to interpret and complete outcome measures.
Third, 44% of participants were potentially unblinded,
based on their responses to the participant blinded
question. However, since the difference between the
two treatment arms (‘culturally adapted’ versus ‘usual
physiotherapy’) and study hypothesis was not disclosed
to participants, it is unlikely that this substantially influ-
enced their treatment outcomes. Fourth, since there was
no follow-up beyond treatment conclusion, we cannot
report the sustainability of treatment gains. Thus, there
is a need for long-term outcomes. Finally, current results
only relate to the three ethnocultural communities of
interest and are not generalisable to broader CALD
communities within Australia or internationally. Non-
etheless, improved engagement by all three communities
highlights that treatment approaches can be effectively
adapted to suit individual communities, using a struc-
tured adaptation framework.

A final consideration is the healthcare context within
which this study was conducted. Australia is a multi-
cultural society and healthcare providers, including
participating physiotherapists, comprise a multitude of
ethnocultural, religious and professional identities, that
influence their provision of healthcare and the intercul-
tural relationship. Future studies may wish to consider the assessment of healthcare
provider cultural responsiveness are factors
that may have influenced treatment outcomes. As such, cultural concordance and
healthcare provider cultural responsiveness are factors
to be delineated between adaption elements and ther-
pist characteristics. Culture is a highly complex construct
and it must be considered that the culture of healthcare
providers, along with the health system itself, will influ-
ence treatment outcomes.

CONCLUSIONS

To meet the needs of multicultural populations, inter-
ventions should be tailored to the individual, social and
ethnocultural factors that influence health. Novel inter-
ventions, such as the culturally adapted physiotherapy
approaches documented in this study, are likely to be crit-
ical for the development of effective pain management
approaches that fully engage CALD patients with chronic
pain.
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Patient consent Not required.

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